THE CONSERVATION OF ANTIQUITIES AND WORKS OF ART
DECORATED IVORY PLAQUE FROM NIMRÜD (9TH CENT. B.C.)

After treatment

For conditions before treatment see Plate 20
THE CONSERVATION OF ANTIQUITIES AND WORKS OF ART

Treatment, Repair, and Restoration

BY

H. J. PLENDERLEITH

KEEPER OF THE RESEARCH LABORATORY
BRITISH MUSEUM

LONDON
OXFORD UNIVERSITY PRESS
NEW YORK TORONTO
My previous books, *The Preservation of Antiquities* and *The Conservation of Prints, Drawings, and Manuscripts* (now out of print), were written at the request of the Museums Association, and it gives me great pleasure to be associated with them once again in the present work, which has been accepted as their official textbook on the conservation of museum objects.

H. J. P.
TO MY WIFE
ELIZABETH
PREFACE

This book is concerned with the broad field relating to the restoration of antiquities and works of art, and with their subsequent conservation. It is intended as a handbook for the collector, the archaeologist, and the museum curator, and as a workshop guide for the technician.

As collectors know only too well, the acquisition of objects is but the first step towards their incorporation in the collection. In order to be able to appreciate and study the objects, it is usually necessary to clean, restore, and repair them, and always necessary to maintain a suitable environment which will ensure their stability whether in storage or on exhibition. In the following chapters simple instructions are given for cleaning and preservation, and the collector with a practical turn of mind who desires to carry out for himself the methods described can do so without any special technical training.

The archaeologist will find interest in the methods that science has to offer for the restoration of antiquities fresh from excavation and for revealing evidence of value to him in his researches. Many examples are given throughout the work, showing how unsuspected facts are brought to light during the normal course of laboratory investigation and treatment.

The requirements of the museum curator, who is not always in a position to call in the museum scientist, have been particularly considered, and it is hoped that this publication, which deals with the numerous causes of deterioration in museum objects and their treatment, will enable him to detect and arrest decay in its early stages, and also to carry out the simple cleaning operations that so often add interest and value to the material in a collection.

The subject-matter is necessarily very varied, based as it is upon the day-to-day problems that are presented in the Research Laboratory of the British Museum—problems relating to books, prints,
drawings, manuscripts, textiles, coins, objets d’art, ethnographical specimens, and antiquities of all kinds. From this wide range of material examples have been selected that illustrate common types of deterioration, and an attempt has been made to recommend from among the various methods of treatment available those that have proved most effective and are at the same time relatively easy to apply. In order to preserve the balance, however, detailed descriptions are included of some of the major tasks of restoration that have been carried out. This has been done partly for the sake of interest, and partly to emphasize that it is impossible to prescribe for all contingencies; each specimen that is submitted for treatment presents its own individual problems, and standard methods of treatment may have to be adapted or new methods devised before a satisfactory restoration can be achieved. Whichever method of treatment is chosen, it should be applied so as to yield results that lie between the extremes of over- and under-cleaning, the aim being to realize the golden mean which will satisfy at the same time the requirements of science, art, and archaeology.

The special problems of the picture gallery have not been overlooked. In common with the museum, the picture gallery is vitally concerned with the stability of materials and methods of conservation, but the restoration of easel paintings is a highly specialized undertaking, and while instructions are given for carrying out some of the simpler studio processes, it is not the intention of the author to encourage the amateur to attempt intricate operations on valuable material. Such work is for the professional artist technician—one who has practical experience based upon a knowledge of the methods used by the Old Masters in the different schools of painting. It is important nevertheless that the collector or curator of paintings should himself be familiar with all aspects of picture conservation. He will then be able to discuss his problems in a knowledgeable way with the restorer, and take a personal interest in any treatment that may be required.

All the processes described herein have been tested, most of them at first hand, by the author. Many are standard methods that have been
handed down through several generations, but some are offered for the first time. While the methods recommended are all based upon scientific investigation, the book is not written for the scientist. On the contrary, a conscious effort has been made to write for the non-scientist who has the responsibility of caring for art treasures. By presenting the material in this way it is hoped that the work will be of service to a wide range of readers not only in the museum world, but also in the home, where, indeed, many of our greatest treasures are still to be found.

I am greatly indebted to the members of my staff who have put their specialized knowledge at my service. In particular, my thanks are due to Dr. A. E. Werner for reading the manuscript and making many valuable suggestions; to Mr. R. M. Organ (metals); to Miss Mavis Bimson (stone, ceramics, and glass); to Miss Sylvia Schwegge for her help in collecting and collating the material; and to Mr. L. H. Bell for his help with photography.

H. J. P.

1956
CONTENTS

PREFACE vii
LIST OF ILLUSTRATIONS xiii
ACKNOWLEDGEMENTS xv
INTRODUCTION: THE INFLUENCE OF ENVIRONMENT 1

PART I. ORGANIC MATERIALS 19
I. ANIMAL SKIN AND SKIN PRODUCTS
II. PAPYRUS, PARCHMENT, AND PAPER 41
III. PRINTS, DRAWINGS, AND MANUSCRIPTS 68
IV. TEXTILES 93
V. WOOD 116
VI. BONE AND IVORY 144
VII. BASEL PAINTINGS 157

PART II. METALS 185
VIII. GENERAL INTRODUCTION 207
IX. GOLD AND ELECTRUM
X. SILVER 212
XI. COPPER AND ITS ALLOYS 232
XII. LEAD, TIN, AND PEWTER 258
XIII. IRON AND STEEL 271
CONTENTS

PART III. SILICEOUS AND RELATED MATERIALS

XIV. STONE 294
XV. CERAMICS 326
XVI. GLASS 333

APPENDIXES

I. STRENGTH OF SOLUTIONS 343
II. DANGEROUS CHEMICALS 345
III. PREPARATION AND USE OF FLOUR PASTE 347
IV. PEARSON'S SQUARE FOR SIMPLIFYING DILUTION CALCULATIONS 349
V. ACIDITY, NEUTRALITY, ALKALINITY 350
VI. THERMOMETRIC EQUIVALENTS 352
VII. METHODS OF RECORDING RELATIVE HUMIDITY 353
VIII. THE PEROXIDE TEST FOR VEGETABLE-TANNED LEATHER 355
IX. POTASSIUM LACTATE SOLUTION 356
X. BRITISH MUSEUM LEATHER DRESSING 357
XI. SILICA GEL 358
XII. MICROCRYSTALLINE POLISHING WAXES 359
XIII. METHOD OF ADAPTING MAINS ELECTRIC SUPPLY FOR ELECTROLYTIC REDUCTION 360

INDEX 361
LIST OF ILLUSTRATIONS

PLATES

Decorated Ivory Plaque from Nimrud (9th cent. B.C.)  frontispiece
After Treatment

1. Portrait Vase from Peru (Mochica Culture)  facing page 20
2. Section through a Piece of Ox-Hide  21
3. Leather Bindings and their Treatment with Leather Dressing  21
4. Decay of Leather Bindings by Chemical Action  28
5. Reaction of Vegetable-Tanned Leathers to an Accelerated Ageing Test  28
6. Egyptian Mathematical Leather Roll (c. 1700 B.C.)  29
7. Egyptian Papyri, before and after Unrolling  48
8. Method of Mounting Birch-Bark Manuscripts  49
9. Manuscripts recovered from a Buddhist Figure  64
10. Drawings by Leonardo da Vinci  65
11. Stages in the Removal of a Stain from a Printed Book  80
12. Water-colour Drawing by Guardi  81
13. Fragment of Chasuble from Lincoln Cathedral (13th cent.)  96
14. Maniple of St. Cuthbert from Durham Cathedral (10th cent.)  97
15. Pyx Cloth from Hessett Church, Suffolk (15th or early 16th cent.)  112
16. Apparatus for Fumigating Objects with Hydrogen Cyanide  113
17. Apparatus for Impregnating Objects with Molten Wax  128
18. Specimens of Waterlogged Wood, Treated and Untreated, after Drying  129
19. Sword and Scabbard from Stanwick (Iron Age)  144
20. Decorated Ivory Plaque from Nimrud (9th cent. B.C.) Before Treatment  145
21. Diagram of the Cross-Section of a Panel Painting  160
22. Silver Lamp from Ur of the Chaldees (c. 3000 B.C.)  161
23. Gilt-Bronze Ornament from Sutton Hoo Shield (7th cent.)  176
LIST OF ILLUSTRATIONS

24. Gold Head of a Bull from Ur of the Chaldees (c. 3000 B.C.) 177
25. Section through a corroded Silver Bead 192
26. Section through a corroded Bronze 192
27. Silver Cup from Enkomi, Cyprus (c. 1400 B.C.) 193
28. Silver-Gilt Buckle (Frankish, 7th cent.) 193
29. Silver-Mounted Glass Beaker (1st cent. A.D.) 208
30. Hoard of English Silver Pennies (Edward I and Edward II) 209
31. The Emesa Helmet from Damascus (1st cent. A.D.) 224
32. Hair-like Cracks on Speculum Mirror (Han Period) 225
33. Oyster-shell Pitting on Speculum Mirror (Han Period) 225
34. Egyptian Bronze Group of Isis and Horus (Ptolemaic) 240
35. Bronze Bust of Ethiopian Queen (7th–6th cent. B.C.) 241
36. Bronze Shield-Boss from Anglesey (Early Iron Age) 258
37. Fragmentary Bronze Bowl mounted on Perspex Model 259
38. Fragmentary Bronze Bowl mounted on Perspex Rings 259
39. British Museum Research Laboratory: Part of Metal Conservation Section 270
40. Stages in the Disintegration of Leaden Objects 271
41. Pewter Medal of the Capture of the Bastille 288
42. Iron Steelyard with Leaden Weights (Romano-British, c. 3rd cent. A.D.) 289
43. Sword-Hilt from Sutton Hoo (7th cent.) 304
44. Shield-Boss from Sutton Hoo (7th cent.) 305
45. Helmet from Sutton Hoo (7th cent.) 320
46. Fragment of Iron Sword with Inlaid Ornament (Roman, c. 200 A.D.) 321
47. Iron T’suba inlaid with Silver (Japanese) 324
48. Iron Buckle inlaid with Silver (Frankish, 7th cent.) 325
49. Corroded Iron Purse-Mount: Decoration revealed by X-Radiography (Merovingian Period) 325
50. Limestone: Egyptian Sculptor’s Trial Piece (Ptolemaic) 332
51. Carved Stone Figure from Mexico (probably Huaxtec Culture) 333
52. Marble Head of Mithras, from Walbrook, London. 336
53. Marble Sculpture from the West Frieze of the Parthenon 337
LIST OF ILLUSTRATIONS

54. Cuneiform Tablets of Clay 352
55. Apparatus for Mounting Prints 353

FIGURES

1. Hygrometric Chart 6
2. Synchronous Variations in the Temperature and Relative Humidity of the Atmosphere in Different Locations 7
3. Causes of Damage to Museum Objects 15
4. Making Mounts for Prints 90
5. The Vacuum Desiccator 152
6. Equipment for Electrolytic Reduction 194
7. Use of Needle in Mechanical Cleaning 202
8. Section of Corroded Bronze Spear 237
9. Air-tight Case kept Dry by Silica Gel 245
10. Use of Ion Exchange Resins for the Treatment of Leaden Objects 263
11. Sand-blast Apparatus for Field Use 322

APPENDIX XIII. Adapting Mains Electric Supply for Electrolytic Reduction 360

ACKNOWLEDGEMENTS

Pl. 10 is reproduced by gracious permission of Her Majesty the Queen. Acknowledgements are due to the Illustrated London News for contributing the frontispiece; The Times for permission to reproduce Pl. 52B; the Laboratoire Central des Musées de Belgique for Pl. 17; the British Leather Manufacturers’ Research Association for Pl. 2; Nickeloid Photography for Pl. 16; B. S. Cron for Pl. 48; Walter Hege for Pls. 53A and B; for permission to reproduce all other illustrations the author is indebted to the Trustees of the British Museum. The drawings were contributed by Mr. C. O. Waterhouse, M.B.E., of the British Museum.
INTRODUCTION

THE INFLUENCE OF ENVIRONMENT

The condition of an antiquity or a work of art depends on the material of which it is composed and on the environment it has experienced during its life history. Its ultimate survival will depend, not only on the application of any treatment necessary for its preservation, but on its being provided with an environment in which it will remain stable. Environment plays a major role in conservation, and for this reason any study of the innumerable aspects of the conservation of museum objects must be preceded by a general consideration of the effects of environmental change—in particular, change of temperature and relative humidity.¹

The problem relates both to the effect of change of environment on objects immediately after excavation, and to the question of creating a museum climate suitable for the miscellaneous collections housed in museums and art galleries.

HUMIDITY AND TEMPERATURE CHANGES ON EXCAVATION

Before archaeologists were aware of the effect of sudden changes of relative humidity, it was not an unusual experience for them to open up a tomb in which the contents were in perfect condition, only to see them shrink and warp and sometimes even turn to dust when exposed to a change of atmosphere.

¹ Relative Humidity (R.H.): The relative humidity of a sample of air is the ratio of the amount of moisture actually present (m) in any volume, to the amount needed for saturation (M) at the same temperature, i.e. R.H. = (m/M × 100) per cent.

Air is said to be saturated when it is unable to take up any more water in the form of vapour. On cooling saturated air, water is at once deposited in the form of dew, because the air is unable to carry the same amount of water at the lower temperature.
INTRODUCTION

The principle behind this is that when objects are buried for a long time under conditions that are reasonably constant, they tend to attain a state of equilibrium with their surroundings. When excavation takes place, the materials are submitted to an entirely new environment, the equilibrium is upset, and some objects suffer profoundly. When a tomb is opened, the access of cold air increases the relative humidity, even to the extent of causing deposition of moisture, while the entrance of hot dry air causes a reduction in the relative humidity. Such variations give rise to dimensional changes in organic material—due either to the absorption of moisture or to the loss of moisture—and such dimensional changes manifest themselves in swelling or in shrinking and warping.

A similar kind of problem arises when objects that have been excavated from damp soil are dried quickly instead of being allowed to adapt themselves gradually to the new environment above ground. In the case of organic materials, the sudden loss of moisture may cause shrinkage and severe distortion, and in the case of stone and pottery, crumbling. In order to prevent such deterioration, it is essential, on excavation, to keep objects in a cool place out of the sun, where they can give up their moisture slowly.

When objects having a cellular structure, such as wood, bone, and ivory, have become waterlogged, the problem, on excavation, is accentuated. It is essential to keep such objects immersed in water until such time as they can be packed in wet cloths and transported to a laboratory where they are treated by one or other standard process, or, in some cases, merely dried out slowly under controlled conditions so that shrinkage and distortion are reduced to a minimum. If allowed to dry too quickly, the cellular structure may collapse, and in the event of this happening no amount of resoaking in water will restore the original shape. An extreme case is that of damp organic material having an impervious covering that must be preserved, e.g., Chinese lacquer. With objects of this kind, the most careful treatment is not always a guarantee of success as the following example will show. In 1949 four trays made of very thin lacquered wood were excavated at Chang Sha in a waterlogged condition. They were
immediately packed in zinc-lined boxes containing damp wool and sent by air for treatment in the Research Laboratory of the British Museum. There, in a series of closed vessels containing air of successively decreasing relative humidity, they were dried for twelve months. Although all four objects received exactly the same treatment, only three of the trays could be preserved; the fourth contracted, warped, and finally fell to pieces. The reason for this anomalous behaviour may have been due to the nature, quality, or direction of cut of the wood used in making this particular tray; on the other hand, the tray may have had a more chequered history than the others, in that it may have been subjected to periods of intermittent drying while still in the ground or even to partial drying when it was excavated.

Objects buried in salty ground are exposed not only to moisture but also to the action of salts dissolved in the ground water. When damp materials containing soluble salts are excavated, the water evaporates and the salt crystallizes, and in so doing weakens the object and often disrupts the surface (Pl. 1). In this case, provided a source of salt-free water is available, it is generally advisable to wash out the salts before crystallization takes place.

Metallic objects also become contaminated in salty ground. Although they corrode in the ground there may come a time when they attain a state of equilibrium with their surroundings and the process of corrosion is brought, virtually, to a standstill. On being excavated and exposed to a new environment, corrosion may break out afresh as they proceed to adapt themselves to the new conditions.

**HUMIDITY CHANGES IN THE MUSEUM**

While antiquities may suffer deterioration through exposure to the sudden change in temperature and humidity consequent upon excavation, certain works of art may suffer as severely by exposure

---

1 At a temperature of 70° F. the R.H. of the air within a closed space containing saturated aqueous solutions of the following salts is as follows: potassium nitrate 94 per cent., potassium chloride 86 per cent., sodium chloride 76 per cent., sodium nitrite 66 per cent., magnesium nitrate 53 per cent.
to ordinary everyday atmospheric changes. Easel paintings, for example, whether on panel or canvas, are very sensitive to changes in the atmosphere, and in particular to varying relative humidity. When a painted panel is subjected to variations in relative humidity, the wood undergoes sympathetic dimensional changes which impose rhythmical strains on the painting ground (gesso) and the paint layer. As a result of this movement, cracks and blisters may develop, and the paint may flake off. Before the Second World War a technician was employed for eight months in the year to deal with such defects in pictures at the National Gallery where the atmosphere was not conditioned. On the other hand, in the repository which housed the same pictures during the war and where the air was conditioned to 58 per cent. R.H. at 63° F. the work of the technician was reduced to approximately one month during the first year, progressively less in the following years, and finally after five years his visits became a mere formality as there was no longer anything for him to do. The significance of the relative humidity factor in the conservation of pictures was further emphasized by the fact that, when the pictures were returned to London, they behaved in the same way as before the war, the paint flaking off as badly as ever in rooms that were not air-conditioned.¹

Similarly, in a large repository used during the war to house the British Museum and Victoria and Albert Museum collections (antiquities, books, textiles, prints, drawings, manuscripts, furniture, &c.), the air was controlled at the constant figures 60 per cent. R.H. at 60° F., and not a single case of deterioration was recorded in this repository.

No more striking evidence could be brought forward to illustrate the paramount importance of maintaining temperature and atmospheric relative humidity constant at an appropriate level in museums and picture galleries. Where there exists an air-conditioning plant incorporating a dust extractor, as is customary in modern installations, conditions are ideal. Under such circumstances there is no longer any risk of damage by atmospheric pollution; soot particles are absent

and it is safe to remove glasses from frames so that pictures may be enjoyed without the complication of specular reflections.

While for the majority of collections such strict atmospheric control as 60 per cent. R.H. at 60° F. is not essential, there is everything to be gained by fixing an upper and lower limit of relative humidity and temperature and ensuring as far as possible that conditions are maintained within these predetermined limits. This involves keeping records (see Appendix VII) and having the means of controlling heating and ventilation.

A graphic representation of the relationship between temperature and relative humidity is shown in Fig. 1, from which it will be seen that if at 70° F. the reading is 50 per cent. R.H. (curved line)—conditions that might well be recorded by day—it only requires a fall in temperature of 10° F. for the relative humidity to rise to 70 per cent. This humidity is within the danger limit for the storage of material that is subject to staining by damp and mould growths, and if allowed to persist damage would be unavoidable. Incidentally, this diagram may be used in conjunction with the sling hygrometer to give an approximate figure for the relative humidity instead of using tables. Thus, when the dry-bulb thermometer reads 70° F. and the wet bulb reads 60° F., the relative humidity will be seen to lie between the curved lines 50 per cent. and 60 per cent. and may be judged to be about 54 per cent.

It is interesting to compare continuous records of relative humidity and temperature that have been taken out of doors (where the instrument was screened under standardized conditions) with records taken at the same time indoors, and these, in turn, with the continuous recording from an instrument placed within a closed exhibition case. The daily extremes of humidity out-of-doors in London during June—September were found to be approximately 30—90 per cent. R.H., and these were observed to be reduced in a museum gallery to 40—70 per cent. R.H., the mean temperature of the room being 67° F., whereas, within the case, changes in relative humidity were so far reduced as to be almost imperceptible. The comparative results are shown diagrammatically in Fig. 2. The particular exhibition case
MAXIMUM AND MINIMUM TEMPERATURES
AND RELATIVE HUMIDITIES

IN THE OPEN

IN THE GALLERY

IN THE CASE

RH

Fig. 2. Summary of records taken concurrently over a period of four months in London (June-September). The variations of temperature and relative humidity in the open (broad bands) are to be compared with the less extreme conditions in the museum gallery, and variations are still further reduced in an exhibition case where the record of relative humidity is shown to be practically constant.
used for this test had a capacity of 176 cub. ft., and a glass area of 88 sq. ft. The effect of the closed case was to slow down atmospheric changes until conditions were almost static within the case. The function of a glazed case or frame in excluding dirt may be more spectacular, but its usefulness in minimizing atmospheric changes is no less important when it is a question of long-term conservation. This comparative test demonstrates the value of keeping inside a closed case or frame objects that are susceptible to moisture changes.

In a private house, temperature may assume an importance as great as that of relative humidity, because collections are usually kept in small rooms where the air movement tends to be intensified by the opening of doors and windows and by the presence of heating appliances. In such circumstances there is usually a greater variation of temperature than in the larger rooms of a museum, and the use of a glass case or frame for susceptible material has undoubted advantages.

THE MUSEUM CLIMATE

Temperature and relative humidity have been shown to be inter-dependent. In a museum gallery the choice of temperature will be pre-determined to suit the comfort of visitors,¹ say, within the limits of 60–75° F., and for the collections it will be a question of establishing limits of relative humidity that can be considered safe within this range of temperature. The limits are determined by observing how the most susceptible materials are affected by exposure to extreme conditions and to variations of relative humidity.

A. The lower permissible limit of relative humidity

Hygroscopic materials are the most sensitive to over-drying because they normally contain moisture. Timber that is dried until it is in equilibrium with its surroundings is said to be ‘seasoned’ but it still contains some 12–15 per cent. of its weight of moisture, and if

¹ Comfort is of course as much a matter of R.H. as of temperature. In a discussion on the effects of the water-vapour content of the atmosphere (MacIntyre, J., Journal of the Inst. of Heating and Ventilating Engineers, 1937, 4, No. 48, p. 570) it was claimed that for mental activity an R.H. of under 55 per cent. was ideal, while for rest and recuperation something above 55 per cent. and possibly in the region of 70 per cent. was desirable.
dried below this it will warp. Seasoned timber continues to adjust itself in sympathy with changes in the relative humidity of the atmosphere, and the amount of adjustment will depend on the nature and dimensions of the wood, and the variations to which it is exposed. Small variations of relative humidity within the range 55–65 per cent. are only of significance in the case of thin painted panels where the recurrent adjustments may in time cause cleavage between the surface of the wood and the paint layer. In the case of heavier structures of wood, significant effects are only produced if the variations in relative humidity are large and take place quickly.

The materials other than wood that suffer damage if dried beyond a certain point are paper, parchment, leather, and notably the adhesives used in making furniture and in bookbinding. Furniture that is decorated with marquetry or inlays is particularly liable to be damaged through desiccation of the glue, and over-drying has also much to do with the physical deterioration of bookbindings. Bindings soon become dilapidated if they are kept near a radiator or on a mantelpiece, or even if exposed for long periods to direct sunlight.

Taking into consideration the susceptibility of all organic materials to damage by desiccation, the lower safety limit should be fixed at 30 per cent. R.H. This means that we choose to regard 50 per cent. R.H. as the arbitrary limit of dryness within the agreed temperature range. If, on taking observations, the figure is observed to fall, say, to 45 per cent., no great harm would be likely to result, but if such a figure persisted over a period, this might well be dangerous, and it would be necessary to counter the tendency to over-dryness by reducing the heating, or perhaps by increasing ventilation. With books or papers either or both methods may be used, but with painted panels cracking may be caused by sudden draughts of air. In this case it would be safer to increase the relative humidity by cutting down the heating.

B. The upper permissible limit of relative humidity

The greatest danger that can arise from an excessively high humidity is the tendency for moulds to grow on any material that can
provide nutriment, such as glue, leather, paper. The presence of mould growths is a warning that the atmospheric relative humidity is above the limit of safety. When conditions are favourable to mould growth, for example in a library, a grey dusty bloom is observed in the first instance on the darker bindings, and it soon becomes fluffy with a tendency to be organized in circular patches. The words mould and mildew are used indiscriminately in a popular way to describe growths of minute fungi of which there are many different species, though superficially they are all very much alike. They consist of very tiny threads called hyphae. The hyphae felt together forming a mycelium which throws up fruiting bodies containing spores, and these are present in enormous numbers. Fungi thrive under conditions of damp, warmth, and darkness, and the materials upon which they grow may be stained by contact with the mycelium or with coloured matter formed during growth. The growth of moulds can be prevented by keeping the relative humidity of the atmosphere below 68 per cent., and this figure is therefore to be regarded as the absolute danger limit at temperatures between 60–75°F. In actual practice 65 per cent. R.H. is preferable to 68 per cent. for the upper limit of humidity, as this takes account of the fact that papers vary widely in hygroscopic nature according to the sizing, filling, &c.

Closed island cases afford considerable protection from excessive humidity. This was demonstrated by a war-time observation. In a flooded basement containing ethnographical material, mould growths spread apace, whereas in the same basement a closed cupboard containing some of the same class of material was found, when opened, to be free from mould. As we have already seen, a closed case can slow down atmospheric changes, and in this instance although the cupboard was by no means air-tight, the air inside had remained sufficiently dry to preserve the contents. Conversely, it has been noticed that when parchment and paper have been packed on a wet day, or in a damp locality, there may be a subsequent growth of mildew, arising no doubt from the moist air included in the package.

The limits of atmospheric relative humidity are thus defined as
lying between 50 per cent. R.H. and 65 per cent. R.H. at temperatures 60–75° F. and these conditions are not difficult to maintain in temperate climates. Such conditions are satisfactory for museums, libraries, and muniment rooms. Picture galleries are, however, in a special category because of the highly sensitive nature of the collections they contain, and in this case a constant relative humidity of 58 per cent. at about 63° F. is the ideal to be aimed at.

ATMOSPHERIC POLLUTION

It remains, finally, to refer to the problems that arise from various impurities present in the atmosphere. These are regional in character.

In the first place, there are difficulties peculiar to the preservation of museum materials near the sea-side. Sea air, damp with salt spray, may blow well inland, give up its moisture, and become charged with a fine dispersion of minute salt crystals; these in turn may serve as nuclei for the deposition of moisture, resulting in the formation of mists and fogs. In one or other condition, sea air finds its way indoors and deposits salt in minute amounts. The salt is hygroscopic and maintains a local relative humidity which will support the growth of micro-organisms, even in surroundings that are apparently quite dry. This may be a serious matter where there are collections of books and papers and, as it is difficult to prevent such contamination, it would seem advisable in such localities to keep things covered as far as possible, and to store susceptible material in cupboards or glazed cases rather than on open shelves.

In the second place, there is the problem of damage arising from the serious forms of atmospheric pollution in the neighbourhood of towns and industrial areas. This is indeed a menace to collections of all kinds and one that is substantially beyond control, for, while soot and grit may be excluded, there is no ready means of preventing access of sulphurous gases save by the installation of expensive air-conditioning plant. Sulphurous gases are the cause of widespread deterioration. Hydrogen sulphide reacts with all the metals of antiquity (save gold) to form dark-coloured sulphides, and is particularly damaging to paint films containing white lead (p. 80). Sulphur dioxide is even
more damaging as it is eventually converted into sulphuric acid which attacks a wide variety of materials. The corrosive effects are obvious in specimens exposed to the weather—building stones, or metals—but a more subtle and equally damaging form of deterioration due, in the first instance, to sulphur dioxide is to be observed in organic materials which have been exposed to traces of the resulting acid over a long period of time. The rotting of paper (p. 64) and textiles (p. 112) can often be traced to the action of sulphuric acid, and it is this acid, generated perhaps by a local coke stove, that is so often responsible for the powdery and decayed condition of old leather bookbindings (p. 34).

All the sulphur in the atmosphere comes from the burning of fuel—coal, coke, or oil—and as the sulphur content of these materials is known, as well as the annual amount of fuel consumed, it has been possible to calculate, with what is claimed to be a reasonable degree of accuracy, that the annual total of sulphur compounds polluting the atmosphere in the United Kingdom corresponds to the equivalent of 9 million tons of sulphuric acid—a quantity more than five times the annual commercial production of this acid! Such figures serve to emphasize the magnitude of the problem which is almost beyond comprehension, but the tragic effects are to be seen everywhere.

For industrial areas as for sea-side districts the only protection is exclusion, as far as possible, of the agencies of decay and this means using glazed frames and closed cabinets whether the collections be exhibited in a museum or maintained in the more intimate surroundings of the home. In a closed cabinet the sulphuric acid contamination of the air will soon be exhausted by its reaction with the contents, and no further contamination will take place until the case is reopened and a fresh supply of polluted air admitted.

Such are the main features of the environment as they affect antiquities and works of art. The only way to secure absolute protection is by employing methods of hermetic sealing or by using containers

---

1 The Times, London, 20 April 1953.
filled with an inert gas (nitrogen or helium),\(^1\) but such procedures are only for use in exceptional cases and can hardly be applied to museum objects on a large scale. In general, the most the museum curator can do is to recognize the potential dangers inherent in the atmosphere, and to counteract them by establishing the best possible museum climate for his collections. Where antiquities and works of art have already suffered damage, treatment to prevent further deterioration is essential, and it is hoped that in the pages which follow, the curator will find the means to effect this end.

In Fig. 3 (p. 15) the various factors that contribute to deterioration are shown diagrammatically.

SAFEGUARDING EVIDENCE

Under the influence of its environment, an object may acquire certain features that are of significance in making an assessment of its age, authenticity, or provenance. These relate mostly to surface effects such as incrustations or weathering, but in the preliminary examination the structure and technique should also be considered and care taken to select methods of conservation that will not cause the loss of any essential characteristics.

1. **Incrustation.** Before undertaking any wholesale cleaning of an encrusted object it is as well to remember that the incrustation may, itself, be of value. It may be a key to the exact find spot; it may contain pollen grains that are of potential value for dating purposes, or, if the object is a food-vessel, the incrustation may provide interesting information if the contents can be identified. Records should be kept of any details of texture found on muddy incrustations that may be evidence of contact with rush mats, basketry, textiles, &c. In the event of a number of the same type of objects existing,

---

\(^1\) A Merovingian helmet found at Baldenheim and now in the Strasbourg Museum is preserved in a hermetically sealed Perspex case in which air has been replaced by nitrogen (France-Lanord, A., *Musée Historique*, Lorraine, Nancy). The original documents of the Declaration of Independence and of the Constitution have been hermetically sealed in helium-filled glass cases and are now on display in the Library of Congress, Washington, D.C. Helium was chosen because it is comparatively simple to detect leaks (National Bureau of Standards).
examples should be set aside with their surface deposits uncleaned, as a record of the condition of the objects as found—always provided that they can be relied upon to remain stable under museum conditions. Incrustations that are germane to an object and that do not conceal inscriptions or ornament should, as a general rule, only be removed if they are unstable, or constitute a serious disfigurement, and if the object is strong enough to withstand treatment. If an object is in fragments, it is usually easier to tell (from the surface staining) which pieces fit together before, rather than after, cleaning.

2. Structure. Examination of the micro-crystalline structure of a metal can supply information regarding ancient metal-working techniques. It is possible, for example, by a metallographic examination to determine whether a metal object has been cast or wrought or what heat treatment it has been given in antiquity. To heat the object would vitiate the results of any such inquiry, and methods are restricted, for preference, to those that can be applied without the risk of changing the micro-structure. This is not always possible, however, and certainly not in the case of brittle silver objects which can only be toughened by a course of heat treatment.

3. Decoration and technique. Any material that has been fashioned by man is liable to bear decoration applied to or imposed upon its surface. This may be concealed under incrustations and the greatest care is necessary when scraping or chipping or treating with chemicals to avoid removing evidence of decoration, and to avoid marking the object in such a manner as to lead to possible confusion or misinterpretation of evidence. For the same reason preference should be given to the use of modern materials in certain classes of repair work. This is important in repairing old textiles or embroideries where, unless a synthetic fibre is used, it might be impossible to distinguish old threads from new.
CAUSES OF DAMAGE TO MUSEUM OBJECTS

Humidity
- Excessive dryness
  - Embrittlement by desiccation
  - Damage to marquetry
  - Slackening of canvas
  - Rapid changes
    - Movement of hygroscopic materials
    - Warping of wood
    - Flaking of paint
    - Activation of soluble salts

Contaminated Air
- Sulphur dioxide
  - Bleaching
  - Tendering
- Hydrogen sulphide
- Soot
- Dust
  - Staining

Neglect
- Accident
  - Exposure to excessive light, heat, and humidity
  - Careless handling and packing

- Pests
  - Blackening of lead pigments
  - Tarnishing of metals

Excessive wetness
- Damp
- Heat
- Fungi
- Bacteria
- Weakening of adhesives
- Rotting of size
- Staining of paper, vellum, &c.
- Blurring of inks
- Mildewing of leather
- Metallic corrosion encouraged
- Loss of adhesion of illuminations
- Adhesion of loaded papers
- Tightening of canvas
  - Moth
  - Wood Beetles
  - Silverfish
  - White Ants
  - &c.
  - Rats
  - Mice

Fig. 3
PART I

ORGANIC MATERIALS
CHAPTER I

ANIMAL SKIN AND SKIN PRODUCTS

In 1843 Michael Faraday,¹ when lecturing at the Royal Institution in London, exhibited leather-bound volumes belonging to the Athenaeum Club that were in a shocking state of decay. This condition he attributed to the products of combustion of coal gas, and he proceeded to demonstrate that the moisture from a gas flame, condensed on cold metal, contained sulphuric acid. The Athenaeum ventilating pipes were, in fact, thickly coated with green vitriol (iron sulphate), from the action upon iron of the acid fumes evolved on combustion and these same acid fumes were responsible for the decay of the leather bookbindings and upholstery.

Faraday’s observations seem to be the earliest on record concerning the decay of leather by sulphuric acid, and nothing appears to have been done at the time to carry his contribution farther. About the turn of the century a committee was appointed by the Royal Society of Arts to investigate the cause of leather deterioration, again relating to bookbindings: a report was published, and certain recommendations were made, but actually no concentrated attack was made on the many practical aspects of the problem until the British Leather Manufacturers’ Research Association took the matter up in 1920. The Association has made an intensive study of the whole problem of the decay and preservation of skin and skin products, and the results of twenty-five years’ research have been published in Progress in Leather Science.² This comprehensive review has been freely used in compiling the present chapter.

¹ Faraday, Michael, On the Ventilation of Lamp Burners, Royal Institution Lecture, 7 April 1843.
Museum collections contain skin products of all kinds, in all stages of deterioration, from every part of the globe, and often dating back to remote periods. The skins may be plain or ornamented with tooling or with colour; they may be in the raw condition, or dressed by various processes, but whatever their nature it is the task of the museum laboratory to take measures for their preservation. Before considering what may be done to restore and conserve such objects, some attention must be given to the variety of material that is presented, and to the methods that have been adopted through the ages in preparing animal skins.

**Skin in Primitive Cultures**

In the earliest times skin had to serve many purposes for which it has long since been superseded, and we can gain some idea of its wide application by studying our ethnographical collections. The entire skin of an animal was sometimes sewn up and used as a water-carrier, or filled with air and used as a float or buoy; skin was used to cover primitive canoes or coracles; it was used for sails, tents, domestic utensils, hunting gear, harness, and accoutrements of every kind. The internal organs of animals were also utilized. Vases and waterproof clothing were made from the intestines of the cow and walrus, floats from bladders, and utensils from the stomach of the camel. Masks were made by moulding moist skins or bladders over a suitably modelled shape and allowing them to dry in position; the shape of the mould was retained by the tissue, which could then be stabilized by waterproofing with fat or oil.

Untanned skins and skin products are, however, not permanent materials; they are liable to attack by micro-organisms, and are very sensitive to moisture. These weaknesses can be overcome to a certain degree by methods of curing and dressing, and there is no doubt that such methods were employed in primitive times to protect the skins. Even at a time as remote as the last interglacial period¹ there is evidence to suggest that simple methods of skin-dressing (smoking and treatment with oil) may have been practised with the object of preventing

I. PORTRAIT POTTERY VASE, MOCHICA CULTURE, FROM PERU

Flaking due to crystallization of salts on the surface
2. SECTION THROUGH A PIECE OF OX-HIDE (×15) SHOWING GRAIN WITH HAIR FOLLICLES AND FIBRE BUNDLES IN THE CORIUM

(Courtesy of British Leather Manufacturers' Research Association)

3. SIX LEATHER BINDINGS OF 1828-35

(1) Physical Decay—(Left Pair)—Hinges dry and cracked.
(2) Chemical Decay—(Centre Pair)—General deterioration to powdery red surface.
(3) Use of B.M. Leather Dressing—(Right Pair)—Books were in the condition of (1) before treatment.
putrefaction and enhancing flexibility. However, skin does not last long in damp earth, and no actual examples from this age are still extant.

It is otherwise in the dry and sterile tombs of Egypt where many specimens of ancient skin have survived from Predynastic times. These are rigid and brittle as a result of desiccation, and it is, therefore, not an easy matter to determine whether they have been cured or processed. There is ample evidence that skin was exploited in ancient Egypt for utilitarian as well as for decorative purposes. In Tut-ankhamûn’s tomb it was used embedded in gesso to form a resilient backing for gold leaf that was later ornamented with tooling, and it was even used by the Egyptian scribes in the form of a scroll for writing upon (p. 39). This latter use need occasion no surprise, as the flesh side when scraped and cleaned yields a smooth surface of pleasant colour which ‘takes’ ink satisfactorily. In this respect, however, it falls far short of parchment, both as regards its suitability for the purpose and its lasting properties.

Parchment did not appear until the second century B.C., and then not in Egypt but in Pergamon (Asia Minor), from which its name is derived. The material is made from the tough, white corium, or inner layer of animal skin, by a process that relaxes and flattens the capillaries so that they are unable to reabsorb moisture. Though it is easily stained, and deformed by heat and damp, it is not subject to chemical deterioration, and has proved to be extraordinarily durable as a material for fine writing. Parchment is dealt with in detail on pp. 44 seq.

PHYSICAL FEATURES AND MICRO-STRUCTURE

The word ‘skin’ is used as a general term for all classes of material, whether raw, cured, or processed, and skin from the larger animals, such as horse and cow, is known specifically as ‘hide’. ‘Curing’ refers to a first aid or field treatment to prevent putrefaction, and ‘processing’ to any more permanent treatment, including the manufacture of leather.

The science of the preservation of leather begins with the study of
the untanned skin—the raw material before it is processed—and it may contribute to an understanding of the problem if we consider briefly the main physical features and the micro-structure of animal skin.

When an animal is skinned the pelt is seen to take the form of a continuous membrane of fibrous tissue, having a hair side and a flesh side, the hair side being patterned with sweat glands and hair follicles (grain), and the flesh side lined with fat, muscles, and blood-vessels (Pl. 2). Skins from the different animal species exhibit great variations in physical structure, and this is most pronounced as between warm-blooded and cold-blooded animals. Even in the same species, however, where the pattern or grain tends to be constant, variation in micro-structure may be observed, and is caused by such factors as the age of the animal and quality of feeding. Some skins are valued primarily for their surface covering of hair, fur, or wool, some for the skin itself (rather than for the surface covering), whereas others, notably in the arctic regions, serve both purposes, the skin being impervious to the elements, and the fur acting as an insulator and a conserver of heat.

In the micro-structure of animal skin the true skin tissue or corium is composed of a reticulated network of protein fibres, the main constituent being the protein known as collagen—'the glue producer'. (If skin is boiled in water for a sufficient time, it undergoes hydrolysis and is converted into glue.) Recent chemical research has shown that collagen has a chain structure, i.e. it consists essentially of long molecules composed of many atoms attached together like links in a chain, with smaller side chains branching from the main backbone. These complicated molecules of collagen tend to orient themselves in the same direction, to associate in bundles, and to retain water pertinaciously in a loose form of chemical combination.

PROCESSING

In the normal course of events, the skin of a slaughtered animal is speedily a prey to the growth of moulds and bacteria, but the mechanical removal of associated fats, muscles, and blood-vessels eliminates
much nutrient material, and the drying of the skin goes a long way towards rendering the tissue proof against putrefaction. In this condition, however, the skin is of little practical value; when the collagen fibres have been deprived of their water, the skin becomes horny and brittle, and flexibility can only be restored by relaxing the fibrous bundles so that they slide freely over each other again, as they did before the skin was dried. This relaxing of the skin may be accomplished in several ways, notably by some form of prolonged manipulation, or by the incorporation of lubricants, or, usually, by a combination of both methods.

The tramping of skins in tubs has been a traditional method of preparation in some parts of the world for many years. The Eskimos soften skins by chewing, a process which doubtless is enzymatic in action as well as being mechanical. Manipulative processes may, in themselves, be sufficient, but generally improved results are obtained by the use of dressings, such as castor oil, or sulphonated neat's-foot oil. These are often applied in emulsified form, and they penetrate and lubricate the tissue, partially replacing moisture as the skin dries.

Fur skins

An animal skin 'in the hair' (or wool) prepared for the taxidermist may have been softened by oil dressing, or merely worked mechanically to a condition of relative flexibility. Equally suitable as a method of preparing soft-dressed fur skins is the following process. The fresh skin is stretched out fur downwards and pinned to a board; the flesh side is scraped with a knife having a convex blade, until the surface of the skin is approximately uniform, and then the skin is allowed to dry. It is wetted again, and the process repeated. There seems to be a positive advantage in alternate drying and wetting, but as it is important in the preparation of fur skins to prevent putrefaction, the sequence of operations must not be interrupted; a little carbolic acid may be used if required to keep the skin sweet, or alternatively some salt and alum. Finally, the skin may be pulled over a beam-knife or stake. In the hands of an expert this yields a very soft and flexible product.
Alum dressing

The mention of alum suggests a process of skin dressing called tawing, which dates back to remote antiquity. This consists in treating the skin, after preliminary cleaning, with excess of alum, the resulting product being a substance light in colour, and having the properties of leather so long as the alum is present. The action is reversible, however, and if the alum is removed by washing, as is easily possible, the skin is ruined. Water must not be used, therefore, with tawed skins. Tawing is still practised today in the manufacture of wool skins; tawed leather is used for making gloves, and to a limited extent for bookbinding.

Oil dressing

Oil tannage is employed today in the preparation of chamois leather. For this purpose the skin is split on a band-knife machine, the grain-split or surface side of the skin (skiver) being worked up as leather for bookbinding, while the flesh side is drummed with cod oil and thus converted into chamois. In chamois leather the fibres are protected with oxidized oil, and the skin absorbs water readily as there are comparatively wide capillaries between the fibres. Alternatively, the flesh side may be converted into parchment (p. 45).

Tanning

Leather is animal skin which has been dehaired, defatted, neutralized, and rendered non-putrescent by a form of treatment that makes the skin impervious to water, while preserving the natural pliability. This treatment is carried out with a tanning agent which may be of vegetable or mineral origin.

Vegetable-tanned leather. One of the earliest materials used for tanning leather in northern countries was an infusion of oak bark—the Breton word tann is the equivalent of oak. This infusion contains tannins which are chemical substances that have the property of combining with the collagen and other protein fibres of the skin, replacing the loosely bound water, and reinforcing the side chains or cross
links of the protein molecules. The result is a material having qualities greatly superior to those of raw hide, particularly as regards water resistivity, durability, and what is known as ‘feel’—an elusive quality, but one that is perhaps sufficiently self-explanatory.

In ancient Egypt it is probable that tanning was first carried out by the use of acacia pods, rather than oak bark, and Lucas¹ mentions that articles of leather are common in Tausian, Badarian, and Predynastic tombs. The Egyptian tomb paintings provide some interesting illustrations of contemporary methods of leather-working, notably in a mural at Gurnah (seventeenth dynasty), and also in the tomb of Aba (No. 36) in the Asasif at Thebes (twenty-sixth dynasty). Some of the processes seem to differ little from the methods employed at the present day.

Besides oak bark and acacia pods there are many other plant products widely distributed in nature that have the property of stabilizing the protein structure of skin, and giving it the properties of leather. These vegetable tannins may be classified in two chemical groups, the catechol group, and the pyrogallol group, though certain of them—and oak bark is an example—belong to both categories. These chemical groups are of significance and will be referred to later, in studying the permanence of leathers used in bookbinding and upholstery.

Mineral-tanned leather. Alum tawing may be regarded as a form of mineral tanning, but this description is generally reserved for the modern process carried out by treating the skin with a salt of chromium which reinforces the side-chains of the collagen fibres as efficiently as the vegetable tanning agents, to produce leather of a distinctive quality. Chrome leather is practically non-wettable and very durable both physically and also chemically, since it is not so vulnerable to attack by sulphuric acid as vegetable-tanned leather.

If the two types are compared as potential bookbinding leathers, the main difference lies in the fact that vegetable-tanned leathers can be relaxed and manipulated in the damp condition, and moulded to the shape of the book, whereas chrome leathers are intractable and

difficult to relax and to mould permanently to any required shape. Vegetable-tanned leathers take gold leaf and stamped decoration or tooling well; chrome leathers badly, or not at all. Chromes, on the other hand, are pre-eminent in their resistance to attack by moulds and chemicals. One day, perhaps, by some form of combined vegetable and mineral tanning, it may be possible to produce a new kind of bookbinding leather of improved quality and durability, in which the characteristic weaknesses of the chrome and vegetable-tanned leathers are suppressed, and the virtues of both preserved. Meantime, we are faced with the problem of preserving objects of skin and leather that differ widely in stability.

ACTION OF MOISTURE AND FUNGOID GROWTHS ON SKIN PRODUCTS

Moisture. In extreme cases the degradation of leather by moisture may proceed to the limit, and nothing may remain in an excavation to indicate that leather was once present, save, perhaps, indefinite traces in the sand, such as were found at Sutton Hoo between the various pieces of the purse complex. Gold coins were lying loose in the sand beneath the metal ornaments of the purse flap, and these no doubt were originally in a leather bag. Belt fasteners were also found in the sand with only a stain to mark where the leather belt had once been lying. That bacterial action contributes to such degradation is beyond question, and vestiges of leather sometimes remain as recognizable tissue where they have been stained green with the products of corroded copper which have acted as a sterilizing agent. Peaty waters and very salt waters also have a protective effect. Exposure to the prolonged action of a low humidity, as in some Egyptian tombs, has converted skin into a black syrup of bitumastic appearance, which is sometimes found to be still tacky, but sometimes has run into a hard black solid with a superficial resemblance to ebonite. Both the syrup and the solid material are soluble in water. Leather manuscripts have been found that are only converted in part to this black material (see p. 40), and there is then the problem of
ANIMAL SKIN AND SKIN PRODUCTS

salvaging what remains before the writing is obliterated due to staining by the syrupy decomposition products.

Fungus. In ethnographical collections and in libraries, and, indeed, wherever skin and skin products are stored, humidity control is most important, because moulds grow readily on these materials when the relative humidity of the atmosphere is above 68 per cent. at ordinary temperatures. The mould growths cause staining, and erosion may take place as well. Dyed leathers may change colour as a result of fungus attack, and leather bindings are thus particularly prone to damage. The use of fungicides to prevent the growth of mould is merely palliative, and is not as a rule advocated. The real cause of the trouble is the condition of dampness in the collection, which would be deleterious to all organic materials, and it is of first importance to take steps to restore dry conditions. Good ventilation is essential, and will speedily arrest the spread of mould growths. In countries where there is persistent high humidity, however, the use of fungicides is recognized as a necessity.

A suitable fungicide for application to leather is one which does not cause staining or rotting, and is of low volatility. Paranitrophenol, even in moderate concentrations, causes a yellow stain, but it has been proved to be a reliable fungicide for leather when employed in exceedingly dilute aqueous or alcoholic solution, strength 0.35 per cent.; and pentachlorophenol is effective at 0.25 per cent. strength. When these two fungicides are mixed in equal parts by weight, it has been found by the National Bureau of Standards, U.S.A., that they can be incorporated in combined concentrations as low as 0.2 per cent. and still be effective for the preservation of leather against fungoid attack. Such a mixture was an official issue to the American Army in the tropics towards the close of the Second World War, and proved to be highly satisfactory. When it is desired to afford protection over a prolonged period, greater concentrations of fungicide are, in general, necessary, and it is no longer possible to use paranitrophenol on account of its staining properties. Preference may then

be given to derivatives of pentachlorophenol. Thus, the sodium salt known commercially as Santobrite\(^1\) can be dissolved either in alcohol or in water to make a 2 per cent. solution that is useful for impregnation; or, should a spraying technique be preferred, the lauryl derivative may be selected, a convenient preparation being the paraffin solution available commercially as Mystox L.P.\(^2\)

**INSECT ATTACK ON SKIN PRODUCTS**

It is necessary to take special measures to protect skin products from attack by insects, and even under the best conditions skins should be inspected regularly and cleaned from time to time. Furs are especially prone to attack by moth larvae. Leather bindings too are liable to be damaged by moth, as well as by several other insect parasites in northern countries, notably those of the order Coleoptera.

In a museum provided with special equipment, the usual first-aid treatment for an infestation on a large scale is to fumigate with hydrogen cyanide, methyl bromide, or carbon disulphide. These chemicals are all effective, but none confers lasting protection, and it is essential to examine the store and, if necessary, to treat the room also to ensure that reinfection will not immediately take place.

To deal with small outbreaks, spraying is generally adopted, and a good hand-operated spray-gun can be very effective for this purpose. The insecticide should be highly atomized and brought into contact with the tissue. Packages must therefore be opened and folded leather and furs unwrapped, otherwise the labour will be expended in vain. The spray deposits a very fine film of chemicals that retains its insecticidal properties for some considerable time, and, from this point of view, spraying has an advantage over fumigation. Where spraying has to be done regularly on a large scale, a power spray may be regarded as a necessity, and this is generally employed at a pressure of about 30 lb. per sq. in.

An insecticide for the control of moth is usually dissolved in an

\(^1\) Monsanto Chemicals Ltd., Victoria Station House, London, S.W.1.

\(^2\) Catomance Ltd., Welwyn Garden City, Herts.
4. CHEMICAL DECAY OF LEATHER BOOKBINDINGS (ADVANCED STAGE)
(Hinges broken—covers detached—leather scuffed—headbands torn)

5. THE PEROXIDE TEST FOR ASSESSING THE DURABILITY OF VEGETABLE-TANNED LEATHER

Only protected leathers (bottom series and No. 10) have been able to withstand this accelerated ageing test. (See Appendixes VIII and IX)
A. Before unrolling

B. After unrolling

6. EGYPTIAN MATHEMATICAL LEATHER ROLL (c. 1700 B.C.)
odourless paraffin distillate that can be guaranteed not to cause staining, and the spray may act on the parasites either as a stomach poison, or a contact poison, or both. It is a good idea to vary the insecticide occasionally, as insects have been found to build up a resistance to a particular poison. This has been observed in the case of DDT. Though slow in action DDT is a very effective insecticide, particularly when used with pyrethrum, which is immediately lethal to moth, and is said to have a high ‘knock-down’ value. Many effective commercial preparations are available, but if a fair quantity is required it is cheaper in the end and more satisfactory to prepare the solutions oneself than to use preparations of unknown composition.

The following formulae can be recommended.

1. **Pyrethrum/DDT spray solution.** Dissolve 1 oz. of pyrethrum concentrate (Pyefly)\(^1\) and 1 oz. of DDT powder\(^2\) in one pint of odourless distillate.\(^3\)

2. **Lethane spray solution.** Dissolve 1 oz. of deodorized Lethane 384\(^4\) in 1\(\frac{1}{4}\) pints of odourless distillate.

Fur skins used to be prepared as a routine by rubbing with arsenical soap, but this has given place more recently to treatment with boric acid, or with borax, which is claimed to afford just as effective protection against insect attack as the arsenical compound, without the danger to the human subject attendant on the use of arsenic.

Where skin specimens are known to be free from insects, it is a wise precaution to shut them away or isolate them in tissue paper or in Cellophane. This keeps the objects free from dust and may prevent parasites getting at them. A Cellophane hanging wardrobe as used for textiles may sometimes be brought into service for skins, and the Polythene bags provided with zip-fasteners are also very useful. These may be charged with a handful of dichlorobenzene crystals. This also applies in the case of susceptible material kept in drawers,

---

\(^1\) Pyefly obtainable from Messrs. Stafford, Allen & Sons, Ltd., Cowper Street, Finsbury, E.C.2.

\(^2\) DDT obtainable from The British Drug Houses Ltd., Poole, Dorset.

\(^3\) Odourless distillate obtainable from Messrs. Shell Mex & B.P., Shell Mex House, Strand, W.C.2.

\(^4\) Lethane 384 obtainable from Messrs. Lennig & Co., 177 West Street, Erith, Kent.
cases, &c. The insecticide is effective so long as some crystals remain unvolatilized.

Insects in books

Many insects damage books by tunnelling through the bindings or leaves, and, when infestation is general, wholesale fumigation, e.g. by hydrocyanic gas or carbon disulphide, may be necessary in order to eradicate them.

When only a few books are badly infected, these may be exposed for about a month to the vapours from dichlorobenzene crystals (about 2 oz. per cub. ft. of air space). The books should be opened fan-wise and placed, with the crystals, in a small air-tight box, e.g. a biscuit tin sealed with Sellotape. A loose paper lining is inserted to prevent the books coming in contact with the tin or the insecticide.

Often, however, the attack is less severe, and can be controlled by thoroughly cleaning the books and bookcase, wiping over the shelves with a rag moistened with a liquid insecticide, or spraying the insecticide as a fine mist from an atomizer. Isolated attacks may be checked by applying liquid insecticide to the larva holes by means of a fountain-pen filler or water-colour brush. Bindings may be sprayed, but, as some papers are liable to be stained, the paper should not be sprayed. As insecticides do not necessarily kill eggs, the bindings should be inspected later and treatment repeated if required. Either of the spray solutions mentioned above (pyrethrum or Lethane) is satisfactory for this purpose, and they can be relied upon to kill moth and larvae, and to render the foodstuffs distasteful to all the common book pests. If a spray is used, some preliminary practice may be necessary with the atomizer in order to ensure that a mist and not a stream of liquid is delivered from the orifice.

PRESERVATION OF SKIN PRODUCTS OTHER THAN BOOKBINDINGS

For the preservation of skin and leather in the museum collections it is fortunately not essential to be able to identify the method of processing. Raw uncured skin, when damp, putrefies. If kept in a
reasonably dry atmosphere, skins, whether raw or cured, even if they have been flexible, tend in time to revert to the condition described as horny. Leathers are affected to a less degree and vary according to type, but there is generally a cumulative hardening.

It is no uncommon occurrence to find skins neatly folded in parcels where they have become rigid owing to years of neglect. To attempt to unwrap the brittle skin without some form of preliminary treatment would be to court disaster. If cracking has not already taken place, such specimens can usually be recovered in an undamaged condition by going over the skin, and particularly the folds, with a damp sponge, following this with a leather dressing. The moisture helps the dressing to spread easily. It should be given a little time to penetrate, and then it will be possible to work the skin gradually back into its original shape with the fingers. Skin clothing of all kinds should preferably be kept on coat-hangers, where it is less likely to suffer than if wrapped up and put away in drawers.

When skins are damaged by water, there are invariably complications due to the inevitable fluffy growths of moulds that carpet the affected specimens, concealing ornament and disguising colour. A case in point was provided by a skin cloak (carossa) from Angola, which emitted an unbearable odour of putrescence, and was usually hidden under a covering of green mould. Preliminary cleaning was done in the open, using a soft dry brush. This removed the mould and revealed the clammy skin, which was purplish-red in colour, due to the use of the primitive dye-stuff, cam-wood. The cloak was then pinned out in a shallow case for exhibition purposes, and the atmosphere in the case was gradually desiccated by means of dry silica gel (Appendix XI). After changing the silica gel several times, a paper hygrometer that had been placed in the case recorded a figure of 60 per cent. relative humidity, and as this is a perfectly satisfactory figure for the preservation of organic materials susceptible to mould attack, it was unnecessary to treat the cloak with fungicides. The specimen has since given no further trouble.

Leather objects are occasionally found in bogs or damp ground, and these require special treatment. Though greatly weakened and
distorted, the leather may retain a certain flexibility in the wet condition, which it would lose permanently if allowed to dry. In no circumstances should formalin be used on specimens of this kind as it makes the leather rigid. The object should first be photographed and measured, and the surface then washed with water, using a fine brush. It may then be rubbed over with a 2 per cent. solution of carbolic acid in alcohol; the leather should be immersed immediately afterwards in molten vaseline, and kept there for a day or longer at a temperature of about 80–100° C. This improves the appearance, and in favourable circumstances softens the leather still further so that it may be moulded back to shape. But there is usually a fair amount of shrinkage, and as the vaseline leaves a sticky surface, it is generally desirable, having recovered the shape, to replace the vaseline with paraffin wax. This is done by immersion for half an hour in molten wax at 110° C. The leather is then removed, and may be stuffed with soft paper to maintain the shape as it cools. A very little bitumen powder may be added to the wax to darken it, so that it will not show as a light film on the leather.

REPAIRING OLD LEATHER

When leather has to be repaired this is best done with a paste mixture before the application of leather dressing, because, once the tissue has been lubricated with an oil dressing, the paste cannot be made to adhere. When degreasing is necessary prior to repair, this can be accomplished by trichlorethylene, but it should be applied locally to the parts that have to be stuck together, and sparingly, or it may remove tanning material. Leather dressing may be applied, of course, after the repair has been carried out.

When leather is rotten and brittle it is impossible to put new life into it, but even if the surface is cracked or broken it may be possible to strengthen it by sticking a strong material to the back. Such restorations are carried out in the case of flat leathers—screens,

1 Grade No. 753, a cement made from paste and rubber and supplied by National Adhesives Ltd., Slough, Bucks., is ideal. It may require to be thinned with water a little before use. Flour paste (see Appendix III) is only suitable for thin leathers.
leather wall-coverings, and the flat parts of leather upholstery. If the surface of the leather is very dirty, it should first be cleaned by sponging it over with a little size water.\(^1\) The leather is then removed (if necessary) from the object to which it is attached, and laid face downwards on a table covered with glass or linoleum, and if it is wrinkled, an attempt should be made to flatten the skin by damping, allowing time for the moisture to penetrate, and then gradually lowering a piece of plate glass upon it. When the leather is well relaxed, pressure may be applied by placing suitable weights on the glass and the pressure should be retained until the skin is dry. A piece of raw cloth canvas of suitable size and weight is now pasted to the back of the leather, the canvas is covered with newspaper, and pressed overnight. Next day the leather will be found to be considerably strengthened.

This will be all that is required in the case of a wall-covering. A screen panel would be remounted, possibly after applying webbing to the framework to prevent the leather being damaged if the screen were to be carelessly handled. The side of a leather chair is sometimes backed with thin tough millboard prior to remounting the leather.

It does not follow that because an object is of skin or leather it should necessarily be treated with leather dressing. It may be desirable to keep rigid leather stiff, rather than to make it pliable, and caution is required in dealing with tooled leather, as the sharpness of the pattern might be impaired were the leather to be softened. These remarks apply to such things as wooden chests covered with hide, and to accoutrements, scabbards, &c. Circumstances must determine what is the best treatment, whether it is better to consolidate by using some form of thin lacquer or microcrystalline wax (see Appendix XII), and so restore the appearance, or whether to aim at restoring flexibility.

\(^1\) On no account should commercial detergent powders be used for cleaning leather as they are liable to extract the tanning materials and ruin leather and chamois. Castile soap, soft soap, and saddle soap are harmless to leather provided they are used in strict moderation, but they may affect dye-stuffs.
ORGANIC MATERIALS

PRESERVATION OF LEATHER BOOKBINDINGS

Vegetable-tanned leather, the only kind of leather that we have to consider in this section, is subject to two forms of deterioration, physical and chemical, and these should be distinguished, though both may be present together in one and the same binding. Physical deterioration is due to normal wear and tear, and commonly takes the form of a breakdown of the surface, exposing the tough fibrous underlayers of the skin which are not easily scuffed with the fingernail. Chemical deterioration is accompanied by cracking and often by change of colour, and sooner or later it involves a complete breakdown of structure. Leather in this condition may be reduced to a fine powder by scruffing. Chemical decay takes place more quickly than physical decay, and its effects are more profound. Pl. 3 shows a series of leather bindings in which the hinges have suffered decay, partly physical and partly chemical in nature, since the period 1828–35.

Chemical deterioration. When leather volumes are standing together on a book-shelf the sides are protected, but the backs are exposed, as well as the top headbands, and (especially if in an industrial area), the exposed parts absorb sulphur dioxide from the atmosphere. Leather bindings invariably contain traces of iron compounds; these act as a catalyst in the conversion of sulphur dioxide to sulphur trioxide which, in turn, unites with water to form sulphuric acid. The sulphuric acid attacks the leather tissue in presence of oxygen, causing it to become very brittle. The opening of the books cracks the joints at the top, and this is usually the first visible sign of chemical deterioration. It is only the beginning, however; the hinges eventually become so weak that the covers split off, and the back may eventually fall to pieces (see Pl. 4).

In studying the deterioration of leather in a long series of dated bindings, such as the heavy volumes of The Times newspaper, bound in pigskin at the British Museum, certain anomalies can be observed. Some of the original bindings are still in use today after a hundred years' service—they are worn and frayed round the edges, as is only
to be expected, but they show no signs of the powdery chemical decay that occurs on similar volumes later in the series. The younger leathers have decayed, while the older ones have survived. The second example is that of the General Catalogue of Printed Books (‘G.K. 1’), which is in daily use at Bloomsbury. It contains serviceable though badly worn volumes of about fifty years of age in their original bindings of green straight-grained morocco, whereas others in the same series of only about fifteen years of age have had to be repaired, and some of the volumes have actually had to be rebound.

As a result of investigations initiated by the British Leather Manufacturers’ Research Association, and carried out by R. Faraday Innes, it has been found that it is the presence of certain water-soluble substances in the old leathers that has protected them from chemical attack. In the past, leathers used to be coloured by surface-staining, but modern processes involve immersion of the leather in a dye-bath. No doubt colour-matching has been facilitated by the modern methods, but immersion of the leather means that the protective water-soluble substances have been extracted, and the lasting quality of the leather has thereby been impaired. Innes carried his researches a good deal farther, and showed that if the water-soluble substances, ‘non-tans’, have been removed from freshly manufactured leather, it is possible to introduce certain simple salts which will confer protection (Appendix IX). He also devised an accelerated form of test called the ‘Peroxide’ or ‘P.I.R.A.’ test (Appendix VIII), that will indicate whether a given sample of leather is likely to be permanent, i.e. whether the leather will resist deterioration when it is exposed to oxidation in the presence of sulphuric acid.

The results of applying the peroxide test to a series of bookbinding leathers are shown in Pl. 5. The first row contains ‘controls’ set aside for comparative purposes—they all seem to be sound leathers. The second row shows the result of applying the peroxide test to other samples of the same leathers—only No. 10 has survived. The third row gives results similar to the second; in this case the specimens were washed before applying the test, but as the leathers had already

1 British Leather Mfrs’ Res. Assoc., Laboratory Reports, 1933, 12, 228.
been deprived of their ‘non-tans’ during manufacture, the washing made no difference. In the fourth row the leathers were protected, before testing, by potassium lactate in accordance with the method described in the appendix. The photographs show that the treatment recommended makes it possible for the leathers to survive intensive oxidation in the presence of sulphuric acid and it is thus possible to ensure the life of vegetable-tanned leather by lactate treatment.

‘Real leather’ can only be reliable for bookbinding if it is adequately protected by the presence of non-tans or salts. To ascribe any virtue to ‘acid free leather’ is purely illusory, as leather will absorb sulphur dioxide in any case in polluted atmospheres.

It should be added that protective salts are only of value when applied to freshly tanned leather: they have an insignificant protective action on old leather that has already absorbed some acid and is suffering chemical decay.

The problem of protecting leather against chemical deterioration is still receiving attention. There is prospect that it may be solved by the application of reagents to deprive the iron in the leather of its power to catalyse the formation of sulphuric acid. This has been shown to be possible by using a water-soluble oxalate,¹ and also by sequestrating the iron (see p. 278) with potassium pyrophosphate.² The efficacy of these methods can only be determined by the test of time, however, as in this case the peroxyde test is not applicable. This idea of sequestrating the iron is still only of laboratory interest; so far as is known no commercial book-binding leathers are yet available that are claimed to be protected in this way.

A long-term test. In order to test the various factors that might be supposed to have a bearing on the permanence of vegetable-tanned leather, two identical series of bindings have been prepared, one series being kept in the clean atmosphere of the National Library of Wales, Aberystwyth, and the other set in the Department of Printed Books at the British Museum. The bindings have been done in leather from different animals, various tannages are represented (sumac, gambier,

² Innes recommends the use of a 1 per cent. aqueous solution.
oak bark, &c.), and the leathers are either unprotected or else protected in different ways by impregnating with a variety of salts. Some leathers have been degreased, some treated with leather dressings, &c., the aim being to make the series as representative of all the variants as possible.

Each series contains over 600 volumes. These are examined every few years and, although the scheme has only been in operation for twenty years, some interesting information has already been accumulated. Some of the main conclusions may be here enumerated:

1. The volumes at the British Museum show considerably more deterioration than those at Aberystwyth, where the atmosphere is practically free from sulphur dioxide.

2. Salt-free tannages of the pyrogallol type are more resistant than those of the catechol type, and it is beginning to be possible to distinguish which actual tannages are best and which are least durable; thus, sumac and Nigerian acacia pod tannages are the best of the pyrogallol type, and gambier the least durable of the catechol type.

3. Removal of soluble non-tans facilitates decay. Addition of certain salts increases the resistance even of catechol-tanned leather.

4. The peroxide test is a rough guide as to how durable vegetable-tanned leathers are likely to be in polluted atmospheres.

5. The following are not major factors in the resistance of bindings to chemical decay: type of skin, type of cure, presence or absence of grease, method of liming and pickling.

This short account of the scientific work that is being done on vegetable-tanned leather brings to notice three matters of practical importance in the preservation of bookbindings and of leather upholstery. In the first place, these leathers will be inclined to deteriorate most quickly in rooms where the air is polluted by the fumes of combustion of gas, coal, and especially of coke. Secondly, washing the leather may be equally bad, in that it will tend to remove soluble non-tans, and predispose the tissue to attack. Thirdly, leather dress-

---

ings have little effect in protecting leather from chemical decay, however useful they are in the physical sense in lubricating the tissue and thus keeping the leather supple.

Physical deterioration. Physical deterioration is determined in no small degree by bindery practice, and can be controlled to a large extent by the selection of suitable leathers for different sizes and weights of books, as well as by handling. Sharp corners on metal shelving are an obvious cause of fraying. When books are too tightly packed on the shelf the bindings become deformed, and the effort to pull a book out may result in the top edge of the back being torn. The evil practice of building shelving over radiators results in the speedy deterioration of books kept thereon; heat causes leather to harden and glue to become desiccated; and, if the shelving is open, a dust problem is added to the others.

Leather dressings for library use. In the absence of chemical decay the life of a book depends largely on the condition of the hinges. These should not be allowed to become hard and brittle, and this may be prevented by the occasional application of a leather dressing. The most suitable form of leather dressing for library use is considered to be a mixture compounded on a basis of lanolin; the lanolin sinks into the tissue in the course of a few days and lubricates the fibres. The mixture should also contain a little wax; wax does not sink in, but remains on the surface to consolidate any powdery leather, and it may be polished in a day or two with a soft cloth. The dressing should be a liquid so that it can be applied without strain, and it should be used sparingly, as an excess of grease would be likely to stain light leathers and possibly even the pages of the book. For the composition of the British Museum Leather Dressing, see Appendix X. Similar preparations are now available commercially.1 The occasional use of such a leather dressing will prolong the life of a leather binding whether or not it is suffering from chemical decay, but there is no way of stopping chemical decay when this has once set in.

Washing books. When books are in constant use the bindings tend

1 e.g. 'Pliantine'; obtainable from Arthur Rich & Partners, Ltd., Factory A.2, Treforest Trading Estate, Pontypridd, Glam., Wales.
to become very dirty, and it may be desirable to wash them before applying leather dressing. But washing vegetable-tanned leather is only permissible when protective salts are added thereafter. The procedure is as follows. The books are sponged lightly with water containing a little Castile soap. This is removed with a clean damp sponge, and then the books are set up to dry. They are next sponged with potassium lactate (10 per cent.), Appendix IX, and left overnight. The leather dressing is applied next day, and after a suitable time rubbed up with a soft cloth. In cases where books are washed annually, the strength of the lactate may be reduced to 5 per cent. Potassium lactate solution should be freshly prepared and not left lying about exposed to the air, as it tends to go mouldy on keeping.

UNWRAPPING OF DECAYED LEATHER SCROLLS

Two cases involving the salvage of documents of unusual interest, written on leather or skin which had long since decayed, are here cited, because each involves a principle which is, in its way, a novelty.

The first example is a Hieratic leather roll dating from the Hyksos period or at least the beginning of the eighteenth dynasty, that had been found along with the Rhind mathematical papyrus. Of a light buff colour and extremely brittle, this tightly rolled document had been kept in a tin tube since the time of its discovery, as it had been found impossible to relax it for unwrapping. Small pieces had fractured from the edge of the roll, and these could not be softened with water. This was very surprising, particularly as there was no doubt about the identification: it was goatskin, and, on clearing a specimen and examining it under the microscope, hairs were still visible. Experiments with the fragments showed that, of the film-forming media, nitrocellulose was pre-eminent in toughening the fibres, and it was decided to apply this to the roll. Also, nitrocellulose contracts considerably on drying, and it was hoped that this would assist in the unwrapping.

A syrupy solution of celluloid cuttings was accordingly prepared in a solvent consisting of equal volumes of amyl acetate and acetone and this was applied generously to the back of the roll so that it
soaked well into the skin, leaving a good excess on the outside. While plans were actually still being made to control the unwrapping the roll proceeded to unwrap itself, the external convolution spreading in a wide arc as the celluloid dried and contracted. This movement was encouraged by a further application of nitrocellulose to the untreated tissue that had just been exposed. In the course of a couple of hours, the manuscript was flat between plate glasses, the celluloid having been removed after it had done its work, by washing freely with pure solvent. Pls. 6 A and 6 B show the roll before and after unwrapping.

Professor Glanville’s study of this document disclosed that it is a list in duplicate of twenty-six sums in the addition of fractions and in one place at least there is an error in addition. It is supposed that this scroll had been used for teaching purposes and that scholars copied its contents on clay or stone tablets.¹

The second example is that of the Biblical fragments of the first century B.C. that were found in the cave at ‘Ain Feshkha, Palestine, in 1949, accompanying the famous manuscripts known as the Dead Sea scrolls. These fragments were of various materials, leather and parchment, and were in the form of wads of several layers of the membranes, stuck together with the black pitch-like material produced by the decomposition of some of the leather. The problem was to get rid of this black pitch-like material and so separate the membranes. It was found possible to relax the fragments by moisture, but when damp, the black material became viscous and threatened to smear over and obliterate the writing on the more stable fragments. In this emergency it was found that refrigeration could control the stickiness, and while the freezing prevented the tarry matter from engulfing the manuscript, it did not immediately harden the relaxed tissue. It was thus possible, after relaxing and refrigerating, to separate the membranes by the use of a scalpel. After the separation had been accomplished, the membranes were frozen once more, and in this state it was possible to cut away and discard the black residue which was of no value.

CHAPTER II

PAPYRUS, PARCHMENT, AND PAPER

All of the materials that have been worked by man have at some time or other been used for writing upon—metals, stones, clay, wood, ivory, leather—and yet history has been recorded for us, primarily, on three that might be considered to be the most ephemeral—papyrus, parchment, and paper. These three are completely destroyed by fire and water, easily stained, and subject to attack by moulds and insects, and yet, when kept dry, they have shown remarkable powers of survival.

While differing in chemical composition and physical properties, they have this in common, that they are brittle when desiccated, but when moistened they regain their flexibility to such an extent that they may be safely handled, flattened, and mounted in permanent form. The presence of carbon ink does not complicate treatment, and even the more fugitive iron inks, when decomposed in the tissue, leave a tracing of rust that is unaffected by the slight degree of moisture required for relaxation and manipulation. Parchment is a protein substance; papyrus and paper are composed of cellulose, but the techniques of conservation, though differing in the three materials, are sufficiently related to justify their inclusion together within the present chapter.

PAPYRUS

The ‘paper reed’ of Egypt, Cyperus papyrus (Linn.), was grown in the delta of the Nile and exploited from earliest times to make ropes, sails, boats, matting, cloth, and later to make the paper-like material used for writing purposes. Lucas¹ describes how he succeeded in

¹ Lucas, A., op. cit., p. 164.
making papyrus sheets. The process he used is as follows. Longitudinal strips of the pith of the papyrus plant are laid parallel and slightly overlapping, a second series of strips is placed at right angles over the first, and welded to it by beating with a light mallet, the only cementing material being the natural sap, which, in the fresh reed, is sufficiently adhesive to effect a permanent join; it acts at the same time as a sizing material and prevents ink from spreading when writing on the sheet. Pliny refers to flour paste being used to join such sheets together to form a long roll.

Papyrus was used for writings in Egypt from about 3000 B.C. until about the ninth century A.D., when the advent of paper-making supplanted the manufacture of papyrus as an essential industry. While the principal source of documents written on papyrus has been the Egyptian tombs, it is known that a form of papyrus was also cultivated in Sicily, and fragmentary remains of charred papyrus rolls have been recovered from excavations at Herculaneum. In Egypt papyrus writings are not only excavated from tombs, but sometimes taken from the mummies themselves. Sir Flinders Petrie found old papyrus manuscripts used as layers in the construction of the cartonnage of certain mummies of the late Ptolemaic period at Gurob. Some of these writings have been salvaged, but from the nature of the material and its contamination with plaster, it is not surprising that, as documents, they are incomplete and largely indecipherable.

Papyri can always be relied upon to arouse the interest of collectors, especially when they appear for sale in the form of tight rolls, tied and sealed, it may be, with the clay impression of a cartouche, perhaps of an eighteenth-dynasty king. It is not unusual to find, however, on opening such rolls, that the more attractive the parcel the less the interest of the contents. When the roll does contain fragments of documents, and this is by no means always the case, they may be merely the oft-repeated texts of the Book of the Dead, but where they are of literary texts interest is revived, as Egyptologists have at times been successful in discovering the parent document to which the missing fragments could be restored.
Method of unwrapping papyrus rolls

Papyrus is recovered from the tombs of Egypt in the form of tight rolls, the diameter being determined by the length of the document, and the width varying from a few inches to a foot or more in the larger specimens. Such rolls are often parcelled up in strips of linen. The linen may easily be removed, as the flexibility of the textile is unaffected by age, but the papyrus is always very brittle in the dry state, and must be relaxed by moisture before any attempt is made to unroll it, otherwise it would break into many pieces. The relaxing is done by wrapping the roll loosely in several layers of damp white blotting-paper and setting it aside for an hour or so on a sheet of glass. By this time the external convolutions will have become sufficiently limp to be manipulated without cracking, and unrolling can then be commenced.

Pl. 7 A shows a number of papyri in the dry brittle condition as received for unwrapping, and Pl. 7 B shows a stage in the course of the work. The damp blotting-paper has been removed from the relaxed papyrus and is under the bell-jar to the operator’s right; the papyrus placed against a sheet of dry blotting-paper is lying on plate glass, and is gradually being unrolled, a little at a time against the paper. When a short length has been exposed, this is covered with dry blotting-paper, and flattened under glass. After a few inches of the roll have been unwrapped the operation is stopped and the unrolled portion covered with the wet blotting-paper to make a fresh piece flexible. This is repeated as often as may be necessary, until the whole of the papyrus has been unrolled and laid flat.

Once the papyrus is flat it must be dried without delay, by changing the blotting-paper several times in the course of a few hours. As a protection against moulds it is usual to sterilize the document by pressing it for several days in contact with thymol-impregnated blotting-paper (p. 57), after which it may be mounted passe-partout between two sheets of glass.

Complications arise when a papyrus is already broken or where it is incomplete, and it may then be necessary to get the fragments into
register by studying peculiarities of grain, colour, thickness, and, of course, the writing. Cracks are reinforced with narrow strips of gold-beater's skin; for major repair work either flour paste or a good photographic mountant can be used—but it is more usual to fix detached fragments in position with gold-beater's skin. Smaller papyri may be enclosed in silk gauze and bound up in a guard-book, such as is supplied for newspaper cuttings; this is convenient for study as well as for storage purposes.

Birch bark provides the same problems as papyrus. It is as brittle as tinder in the dry condition but it, also, can be relaxed by moisture. Pl. 8 illustrates a guard-book with a birch-bark document before treatment, and a similar document after flattening and mounting. A tightly wrapped wad of writings on birch bark (Pl. 9) was found inside a Buddhist image of the eighteenth century. It was possible to unwrap it, and to recover eleven legible documents from the wad. These documents are copies of sets of Northern Buddhist charms in doggerel Sanskrit, adorations of the Buddha, requests for purification, success, &c., in a series of imperatives fortified by magic syllables.

PARCHMENT

Parchment originated in Pergamon, Asia Minor, in the pre-Christian era. It is a stronger material than either papyrus or paper, and is able to withstand much harder usage, as one would expect, since it is derived from animal skin. Although parchment may be prepared from the skin of many animals, the commonest source is sheepskin, and this has provided the finest grades for the scribe and illuminator. It is, however, almost impossible by examining an old parchment to determine with any degree of certainty from which animal it has been derived. Exceptionally fine qualities are sometimes attributed to the deer, antelope, or smaller animals. In general, the younger the skin, the thinner the parchment made from it, and the less likely is it to be blemished. Calf parchment is known by the manufacturers today as vellum. It is usually harder than sheep parchment, and on this account of more interest to the binder than to the
scribe, but the word vellum has lost its definitive character, and one often hears the finer grades of medieval parchment described as 'uterine vellum', a description which can seldom be justified.

Manufacture and qualities of parchment

The manufacture of parchment involves operations depending for their success on the experience and dexterity of the craftsman. Occasionally one comes across a *tour de force*, an old liturgical manuscript, perhaps, which in fineness and quality is a match for the best that can be made today, even with the modern abrasive and skin-splitting processes. Fineness is not the only criterion of a good parchment—flexibility, perfection of surface, and strength all contribute to the excellence of the material as a ground for fine writing and illumination.

The technique of parchment-making has changed little through the ages, except for the introduction of the splitting process. Parchment was formerly made from the entire skin; in modern practice parchment is made from the flesh side of a split skin. The whole skin is limed, dehaired, and very thoroughly defleshed, and a splitting-machine separates the skiver from the flesh side. The parchment split is then stretched on a frame and scraped down on both sides with a crescent-shaped knife. As it dries the skin tightens considerably, and scraping is continued with a knife of the same shape but having a blade with a burled edge. The parchment is then treated with hot water, scraped again, and, while still wet, rubbed on both sides with a pumice block. It is then dried on the stretcher frame.

The type of finish varies in accordance with requirements. In the case of writing material which must be white and flawless, bleaching and a further degreasing by plastering with lime-wash may be necessary. On the other hand illuminating parchment must be opaque, and the surface is sometimes finished by coating with talc. One of the chief aims of the manufacturer is to suppress the grain, but there is usually no difficulty in deciding by inspection which side of the material was originally the flesh side, and which the grain side; these correspond respectively to the rough side and smooth side of the
sheet. In good-quality parchment the smooth or grain side is usually velvety to the touch, and as it provides a very agreeable surface for writing upon, it is preferred for this purpose to the flesh side. It is delicate, in the sense that it is easily damaged by erasures or staining, whereas the rough side is more robust and tends in the coarser qualities of parchment to be yellowish and horny in character. In a bound manuscript both sides are used for writing and, in this case, like sides face each other, grain to grain, and flesh side to flesh side, so that when the book is open, there is less chance of disparity in the appearance of adjoining sheets.

Parchment may have a transparent quality resulting from less intensive stretching, or transparency may be obtained by chemical means. In 1790 Edwards of Halifax took out a patent for making parchment transparent by treating the flesh side with potash. In the King’s Library at the British Museum there is a sample of Edwards’s ‘transparent vellum’, as it was called, which takes the form of a binding for a ‘Book of Common Prayer’. This transparent vellum is painted on the inner side and the colours show through the transparent membrane to the front. It has kept its appearance surprisingly well, showing no sign of cockling although it has been on exhibition for many years.

Alkalinity of parchment

A feature of parchment of all types is that it is alkaline. Liming of the skins is carried out in the early stages of fellmongering, prior to degreasing, and no subsequent acid treatment is applied to neutralize the alkali. A trace of lime remains, therefore, and this is firmly held by the collagen fibres of which the parchment is composed. This alkalinity confers a certain measure of protection against the action of moulds and micro-organisms, which prefer a substratum of slightly acid character. It may also provide a clue to the chemical stability of the material, as parchment is not affected by acidic atmospheres, and in this sense is a much more durable material than leather. One disadvantage of alkalinity, however, is the tendency to yellowing, so noticeable when parchment is much handled or exposed to dirt and
grease. The main cause is probably the widespread distribution of iron as an ingredient of dust, yielding the coloured hydroxide. This staining action is intensified by damp, and protection from excessive humidity is of great importance in the preservation of parchment.

**Moisture sensitivity**

Parchment is a hygroscopic substance and it will adsorb moisture in any amount. As an extreme example, if parchment is exposed to excess of water for a lengthy period of time, a complete breakdown of structure will take place by the chemical action known as hydrolysis: the proteins are degraded, the organized structures disappear, and a form of gelatine results known as parchment size.

Under normal conditions parchment tends to adsorb and give up moisture in sympathy with the rise and fall in the relative humidity of the atmosphere. A test applied to a representative sample of parchment showed that it contained 10 per cent. of its weight of water when in equilibrium with an atmosphere at 40 per cent. R.H.; when, without changing the temperature, the atmosphere was suddenly raised to 80 per cent. R.H., the parchment adsorbed moisture until, in the course of three days, it attained equilibrium with its new surroundings and the water content was then found to amount to no less than 30 per cent. of its dry weight. This test shows how responsive the material is to changes in relative humidity; it also indicates that there is a time lag. If parchment is exposed to over-dry or moist conditions for a short period no harm will result, but if extreme conditions persist for a time, then deterioration will follow.

When kept in an atmosphere that is too dry, say at 40 per cent. R.H. or less, parchment tends to become rigid. This condition is reversible, however, as flexibility can always be restored by exposing the membrane to moisture, but, meantime, damage may have been caused to inks and colours through desiccation. Certain inks tend to flake from the surface of over-dry parchment and illuminations on gesso are liable to suffer in the same way. On the other hand, the damage resulting from exposure to high humidity is even more severe, especially if the manuscripts are illuminated or decorated with
painted miniatures. The adsorption of moisture results in cockling of the membranes with consequent deformation which may cause loss of paint. Under humid conditions miniatures in colour may actually become off-set against adjacent leaves. When a bound manuscript is exposed for some time in damp surroundings (e.g. in a safe or deed box in a damp basement), the edges of the pages pick up moisture and become cockled. This type of damage is accelerated if the binding is too tight, or if a book has lost its clasp. The book tends to gape along the fore-edge, exposing a larger surface area to damp so that the wrinkling spreads until, in the end, it is no longer possible to close the book. Such distortion cannot be cured by drying or by putting the book in a press. The only satisfactory mode of treatment is to ‘pull the book’—a task requiring special training and experience—and to deal patiently with the sheets one at a time (p. 85).

Although so sensitive to moisture changes, parchment retains its strength and resiliency and is very durable, provided it is not exposed to extreme conditions for long periods. The great codices that have come down to us from the early centuries of the Christian era are often as fresh in appearance as if they had been executed today, and although they may have been cleaned, flattened, and rebound, this in itself is a testimony to the stability of the material.

Parchment documents that are in a hard and shrunken condition, and with individual pages adhering together, are sometimes brought in for treatment. In such cases no attempt must be made to separate the membranes until they have been relaxed. The relaxing process is very similar to that adopted in the case of desiccated papyri (p. 43), except that humidification should be less intense, and it should be given more time to take effect. The horny tissue of parchment takes about twenty times longer to relax than papyrus, and premature efforts to separate the leaves would certainly cause damage. In some cases it may prove to be impossible to restore the sheets to their original size. Even when the membranes are badly shrunken the writing may still be quite readable, and it is better to leave well alone rather than risk destroying the legibility of the writing.

In order to clean a stained vellum binding it is rubbed over quickly
7A. Six rolls of Egyptian papyri wrapped in linen as found

7B. Unrolling papyri. In the foreground fragments awaiting repair
8. BIRCH BARK MANUSCRIPT FROM KASHMIR (18TH CENT.)

with a sponge made damp with size water, and if dried at once no harm will result.

The cleaning and repair of manuscripts is a craft that can only be mastered by experience. The simple operations of relaxing and removing creases, repairing holes, replacing missing corners, &c., should be practised on unimportant material.

For repairs see pp. 85 seq.

Fungoid attack

If parchment has become mouldy, the most convenient and effective method of fumigation is to expose it in a thymol chamber (p. 56). Larger documents may be opened out and reparchelled in contact with paper of the same dimensions that has been previously impregnated with a suitable fungicide (p. 57). This contact method is recommended where continuous protection is required over a period.

Storing parchment

The control of humidity is an essential factor in the storage of parchment. Ideally, the atmospheric humidity should be kept constant at about 55/60 per cent. R.H. at a temperature between 60° and 75° F. This will prevent either embrittlement resulting from conditions that are too dry, or distortion from exposure to conditions that are too moist. Damp heat must be avoided as this causes shrinkage and encourages the growth of moulds.

It is no easy matter to maintain satisfactory conditions day and night in a private house, or in a safe or strong room, and it would have been impossible to suggest a simple remedy to help in this matter, were it not for the fact that humidity conditions can be stabilized by the presence of a bulk supply of hygroscopic material, such as cotton-wool, carpets, curtains, and textiles generally. Shelving in an unheated room without ventilation is no place for parchment; in the case of metal shelving there will be the added danger from condensed moisture. Metal safes and deed boxes are necessary in the interests of security, but are not the best accommodation unless plenty of hygroscopic material surrounds the parchment to minimize the
effect of changes in humidity. Documents should be wrapped in paper, or preferably in textile, and textiles should be chosen with a thought to the subsidiary risk of insect attack. It has been proved that cotton and linen materials are safest.

In storing parchment it should be remembered that parchment and papers are prone to attack by rats and mice, and precautions must be taken accordingly. Small rodents can work havoc with parchment in a very short time, and to leave a storage drawer open, even for one night, may result in damage that is irreparable. Hygiene is the best safeguard, and no foodstuffs should be left about that might attract the pests. If workmen are on the premises they should be asked to exercise care in this respect.

As to fire precautions, there should always be at hand fire-fighting equipment that would not, itself, harm the documents. Carbon tetrachloride equipment (such as is used for electrical fires), extinguishers of the methyl bromide type, and extinguishers that emit a stream of carbon dioxide gas under pressure are all perfectly safe, but the acid type of fire extinguisher is to be avoided. Asbestos blankets should be kept with the fire-fighting appliances.

**PAPER**

The Chinese claim the honour of being the first to discover how to make paper. They teased out silk or vegetable fibres under water, and collected the aqueous suspension on a porous support such as a stretched cloth. As the water drained away, the support was shaken, and this caused the loosened fibres to interlock and so form a thin matted sheet. When dry, such a sheet proved to be remarkably tough and serviceable.

The invention is recorded as having been made at the beginning of the second century A.D., and the earliest examples of paper that have come down to us date to within fifty years of this time. These were discovered by Sir Aurel Stein in the Great Wall of China. Some of the actual papers have been examined by the author and were found to have been made from rags. This is an interesting fact because the fibres of the paper mulberry (Broussonetia papyrifera) were later
preferred in the Orient, as the raw material for paper-making, and it was in the West that paper made from linen and cotton rags was most in favour. Hunter,¹ who has contributed so much to our knowledge of the history of paper-making, points out that in China the soft absorbent papers made from the mulberry fibre commended themselves for brushwork and wood-block printing on one side of the paper only, whereas in Europe the requirement was for a hard paper, made from rags and treated with size, that could be used for quill-pen work on both sides of the sheet. He remarks upon the fact that it took 1,000 years for knowledge of the invention to reach Europe, and its progress, via the Persian Trade Route, can be followed by studying his dates for the beginning of paper-making in each of the countries concerned:

<table>
<thead>
<tr>
<th>Year</th>
<th>Place</th>
<th>Year</th>
<th>Place</th>
</tr>
</thead>
<tbody>
<tr>
<td>105</td>
<td>Lei-Yang</td>
<td>1151</td>
<td>Spain</td>
</tr>
<tr>
<td>600</td>
<td>Korea</td>
<td>1276</td>
<td>Italy</td>
</tr>
<tr>
<td>610–25</td>
<td>Japan</td>
<td>134th cent.</td>
<td>France and Germany</td>
</tr>
<tr>
<td>751</td>
<td>Samarkand</td>
<td>1494</td>
<td>England</td>
</tr>
<tr>
<td>793</td>
<td>Bagdad</td>
<td>1575</td>
<td>Mexico</td>
</tr>
<tr>
<td>10th cent.</td>
<td>Damascus and Cairo</td>
<td>1690</td>
<td>Philadelphia.</td>
</tr>
</tbody>
</table>

Quality and stability of paper

The quality and stability of paper is governed in the first instance by the raw materials from which it is made, and in Western countries many different types of cellulose fibres have been used.² The strongest and most durable papers are hand-made from a mixture of disintegrated linen and cotton rags, the fibres being sized with gelatine, whereas the poorest and least durable grades of paper are those made by machinery from ground wood, pulped and sized with rosin and aluminium resinate. Between these extremes lie many varieties of modern paper. One of the commonest is composed, for the most part, of sulphite pulp, i.e. wood pulp chemically treated in order to

remove lignin and natural resins. This paper is suitable for book printing and the printing of half-tone blocks. Modern newspapers,¹ which are not in the same class as regards permanence, contain a high proportion of crude ground-wood fibre, and only a small quantity of sulphite pulp. It is easy to determine the source of the constituents of paper by teasing out a small fragment in water and examining the fibres under low magnification. They may then be identified by comparing them with published photo-micrographs such as are to be found in a standard reference book.²

An estimate can be made of the permanence of paper by various forms of testing, e.g. by an accelerated ageing test (heating the paper for 72 hours at 100° C.) or by testing the tensile strength or by submitting it to an endurance test in a special machine, which folds it back and forward, and registers the number of folds that it will stand before the sheet cracks. The mechanical tests are often applied after the accelerated ageing test, and of these the endurance folding test is the most useful. By such testing it can be shown that the acidity of paper is of great importance in relation to its permanence. Neutral papers (pH 7 ± 0.3) (see Appendix V) have a high fold-value, while papers that are acidic have a low fold-value in proportion to their degree of acidity. Where papers are very acid and the pH is as low as 4.4, they may actually be too brittle after heating at 100° C. for testing in the standard form of folding-machine.

Acidity may be acquired by paper on ageing, or it may be introduced in the process of manufacture. Alum is used, for example, in rosin-sized papers, and this in itself places them in an inferior category, irrespective of the nature of the fibres from which they are made, because, in solution, alum is an acidic substance. A residue of bleaching agent left in any paper will cause deterioration, and the greatest care has to be taken in the manufacture of paper to dechlorinate the pulp after bleaching. Paper shares with vegetable-tanned leather a susceptibility to attack by sulphur dioxide, and both leather and

paper have been found to survive best in clean atmospheres free from the products of combustion. The mechanism of decay is similar in each case, and Langwell\textsuperscript{1} has shown that sulphuric acid is formed in paper, just as Innes showed it was formed in leather, by the catalytic agency of traces of metals, and he has endeavoured to suppress the action of these metals by the use of complexing agents, or inhibitors. Iron gall inks actually contain sulphuric acid; this causes brittleness and sometimes even puncturing of the paper where there is ink writing. Puncturing may also be caused by the acid liberated during the growth of certain micro-organisms (\textit{Aspergillus}).

Different methods of overcoming acidity and protecting weakened papers have been practised for many years by W. J. Barrow (\textit{vide infra}) who has been concerned not only with the preservation of paper, but with the closely related problem of preserving ink writing.

The exposure of paper to sunlight is another cause of deterioration; this is well illustrated by the behaviour of newsprint which becomes discoloured and brittle even after a few hours’ exposure. The effect is, however, much more general in its application, as has been demonstrated by Kimberley and Scribner,\textsuperscript{2} who found that even the better-class papers were weakened by exposure to bright light, and that weakening could take place even in the absence of any visible yellowing.

\textit{Humidity, Mildew, and Foxing}

The control of humidity is of the greatest importance in places where books and papers are stored, because paper is a hygroscopic material—it absorbs moisture from the atmosphere. Excessive moisture tends to weaken the tissue,\textsuperscript{3} and promotes the growth of micro-


organisms which rot the size and cause staining. How delicate the
hygrometric balance is may be gauged from a calculation that was
made in connexion with an extension of a quadrant book-stack at
the British Museum. One thousand tons of books were calculated to
absorb at least 20,000 lb. of water when the relative humidity of the
atmosphere increased from 57 per cent. to 63 per cent. at 60° F. To
avoid deterioration, books and papers should be stored under con-
ditions that are constant, say 60 per cent. R.H. at 60° F.

Cellulose and gelatine size, as well as the constituents of book-
binder’s paste (starch, flour, or dextrin), are all excellent nutrient
materials for fungi, and when the relative humidity of the atmosphere
is over 70 per cent., the appearance of mould growths on paper is not
long delayed. If there is an outbreak and the trouble is detected and
dealt with at once by drying the affected papers little harm will ensue, but if infection is allowed to continue unchecked, the papers
will be yellowed and stained with coloured spots. Some micro-
organisms live on the size, and others on the cellulose fibres. Where
the sizing material has been attacked, the size is destroyed, and the
affected area becomes highly absorbent, like blotting-paper, and if
tested by wetting, will appear dull and translucent as compared with
the surroundings. When cellulytic micro-organisms are present, the
surface of the paper will be eroded, and the paper may become very
brittle. In both cases there is a tendency for iron salts (iron is a com-
mon impurity in paper) to be accumulated on the damaged areas,
where they form rusty brown spots commonly known as foxing.
But this type of staining may arise from other causes, and is often
due, in part at least, to the coloured products elaborated by organ-
isms in the course of their development.

Methods of dealing with an outbreak of mould growth

When mould begins to grow, the first indication is the appearance

1 McIntyre, J., Journal of the Institution of Heating and Ventilating Engineers, 1937, 4,
No. 48, p. 570.

2 Armitage, F. D., The Cause of Mildew on Books and Methods of Protection, Bull. 8,

of a fine white fluff, not at first easy to discern, but soon forming furry patches of roughly circular shape, and if no action is taken, growth, once started, will be likely to proceed apace. The real source of the trouble is damp, but so long as infected material is in the room, it would be fatal to attempt to deal with the outbreak merely by raising the temperature, as this would stimulate growth and spread infection. A prior requirement is the improvement of ventilation by the use of fans, &c., special attention being paid to any damp spots. Moulds must not be brushed off the books in the room; all mouldy material should first be removed, and then the room can be dried by electric heaters and fans, or by the dual-purpose machine known as a hot-air circulator, which is merely a magnified form of the familiar electric hair-drier.

Infected books and manuscripts can best be dealt with in the open air, or in a large airy room, by brushing and sunning, books being stood on edge, open, to allow free access of air, and documents flattened and suspended on stretched lines.

If the damp condition of the room can be permanently cured, and if the above instructions have been carried out, all the material may safely be returned and no further outbreak should occur, but if there is any doubt about dry conditions being maintained, then sterilization will be necessary.

(A) Fumigation of room. Gaseous formaldehyde may be used for sterilizing a room containing mould-infested papers,¹ provided the room can be effectively sealed for this purpose. The gas is generated by adding aqueous formaldehyde to a substance such as potassium permanganate, with which it reacts exothermically. Details of the procedure are given by Walker,² who recommends adding 1 lb. of aqueous formaldehyde (Formalin) to about 6 ounces of solid permanganate in a large porcelain basin or bucket. This quantity is

¹ Protein materials (parchment, vellum, leather) should, preferably, not be present as these tend to become hardened by the action of the fumigant. Despite this warning there may be cases in which the fumigation in situ by formaldehyde vapour could be justified as this is the most direct way of preventing moulds from spreading.

sufficient for a room of 1,000–1,500 cub. ft. As the gas is evolved briskly, the operator should make a quick exit, and seal the door on the outside. After twenty-four hours or longer, the room should be thoroughly ventilated, and any lingering odours of the fumigant dispersed by sprinkling the floor with ammonia, which converts the residual formaldehyde to hexamethylenetetramine, which is odourless. Alternatively, the sterilization may be carried out using a paraformaldehyde heater, 5 oz. of commercial paraformaldehyde being required for a room of 1,000 cub. ft.

(B) Sterilization of materials. Two methods are available for sterilization—fumigation which is effective but confers no lasting protection, and a method involving the use of papers impregnated with fungicide of low vapour pressure which gives continuous protection over a period.

(i) Fumigation with thymol vapour. A suitable chamber for this purpose may be improvised from any relatively air-tight cupboard, and to be generally useful it should be large enough to accommodate in a horizontal position anything up to an imperial sheet (30 by 22 in.). The material to be sterilized is supported on a framework (stretcher covered with a net or strands of tape or twine), some 2 feet above the bottom of the cupboard. A 40-watt electric lamp is fixed near the bottom of the cupboard, and this emits enough heat to melt the thymol crystals, which are placed some 2 inches above it, in a clockglass or enamel plate, supported on a wire stand. About 1 ounce of thymol is required for sterilizing the contents of a cupboard of 16 cub. ft. capacity.

To use the apparatus, the current is switched on for two hours, then switched off, and the heating should be carried on for periods of about two hours every morning for fourteen days. If much material is being dealt with, the papers should be rearranged in the cupboard each morning before the light is switched on. After each treatment the door of the cupboard should be kept closed for about twenty-four hours.

In stacking the cupboard it is important to arrange for free access of the thymol vapour around the infected material—small manuscripts
may be suspended, rolls set up on edge, books stood on end with their pages open fan-wise, &c. The success of the treatment depends on two factors, the concentration of thymol vapour reaching the fungus, and the drying action which attends the slight rise in temperature from the electric-light bulb.

It has been found by experiment that the method may be safely applied to prints, drawings, manuscripts, pastels, water-colour paintings, books, and also to parchment and vellum. No harm has been found to result from prolonged dosage. It should be noted that oil-paints and varnishes become softened under treatment, and for this reason the method is unsuitable for the sterilization of easel paintings and the like. The thymol chamber itself should of course be free from any internal paintwork.

(ii) Fumigation with formaldehyde vapour. Formalin (40 per cent. formaldehyde) has long been known as a powerful antiseptic and germicide. It may be employed for sterilizing papers, but is not recommended for parchment, vellum, or other protein products, as these are hardened by this reagent. The papers are exposed for about twelve hours over a dish containing formalin (about 2 oz.). The sterilization must be carried out in a sealed air-tight box, and during exposure the temperature inside the box should not be allowed to fall below 65° F. It is important also that the relative humidity should be kept above 60 per cent. as moisture is essential for effective disinfection. After sterilization the papers should be freely exposed to the air for several hours.

(iii) Use of impregnated paper. White blotting-paper may be impregnated by immersing it for a moment in a 10 per cent. solution of thymol in alcohol, after which the excess of solvent is allowed to evaporate, leaving the thymol uniformly dispersed in the sheet. Sheets with a higher but less uniform concentration of thymol are made by scattering a handful of thymol crystals between several layers of absorbent paper and melting the crystals into the tissue by the application of a hot electric iron. Thymolized papers, prepared by

---

1 As formalin vapours are highly irritant, precautions must be taken not to inhale them.
one or other process, are used for sterilizing papyrus after it has been unrolled, or for interleaving mildewed books.

For the permanent protection of large parchment rolls in store, a fungicide should be chosen that is less volatile than thymol and more potent in its action. In this case the quantity of fungicide can be greatly reduced, and the paper used for impregnation may be thin and not greatly absorbent. Protective sheets are prepared by passing a roll of thin paper through a 10 per cent. aqueous solution of the sodium salt of pentachlorophenol (Santobrite). Such paper provides a valuable means of protecting library material and suchlike in tropical countries where the incidence of mould growth is a serious factor in conservation.

_Insect pests_

In common with other organic materials, parchment and paper are attacked by certain insect pests, and there is no doubt that the presence of gelatine and starch increases the number of potential enemies, and that the greatest activity is associated with damp.

McKenny Hughes\(^1\) has described the insect pests of books and papers common in Britain in a communication to the technical section of the British Records Association. He lays emphasis on cleanliness and free circulation of air as being the best deterrents, and suggests that infected manuscripts can be most simply and effectively sterilized by exposure in an air-tight cupboard containing crystals of paradichlorobenzene. The crystals should be present in a quantity of 1 lb. to every 10 cub. ft. of air space; the cupboard should be sealed with gummed paper, and the time of exposure should be for at least a fortnight at a temperature of not less than 70° F.

When attack by insects is a recurrent problem, the vacuum chamber method of fumigation has much to recommend it. This has been applied on an extensive scale for the sterilization of rare books and manuscripts in the Huntington Library, San Marino, California.\(^2\)


\(^2\) Iiams, T. M., _Library Quarterly_, 1932, ii, 4, p. 375.
using ethylene oxide gas, which has been found to be a safe and effective fumigant.

When an attack by insects has been discovered, the accommodation—drawers, cupboard, shelving—should be thoroughly cleaned and dusted with some persistent insecticide such as DDT, or BHC (Gammexane). Chakravorti\(^1\) has warned against exposing the papers themselves to Gammexane smoke, as this brings about a substantial decrease in tensile strength and folding endurance, and tends to cause iron gall ink to fade.

**Resizing and bleaching**

When paper has been weakened by the decay of the sizing material, it loses its characteristic ‘rattle’, becoming limp and absorbent, and in this condition is readily stained. It may be resized (p. 87), but it is often desirable to clean it beforehand by a process of bleaching (pp. 75 seq.).

Bleaching and pressing are operations carried out in the normal course of manufacturing paper, and they are among the commonest procedures in restoring old papers. Solutions of hypochlorite were at one time employed, exclusively, for bleaching pulp in the manufacture of paper, and also for cleaning stained prints in the studio. In 1937, chloramine-T\(^2\) was advocated as a less drastic agent for use in restoration work (p. 77). More recently a method was described by Gettens\(^3\) making use of sodium chlorite, and while this eliminates the risk of oxidation and the consequent weakening of the tissue which is always present when hypochlorites are employed, it requires the use of apparatus which is not normally available in a restorer’s laboratory. It marks such an important advance in the technique of bleaching, however, as to merit every prominence.

Gettens gives details of three methods of application suitable for the treatment of various types of material, and these have all been

---

tested in the British Museum Laboratory and found to be satisfactory. It will suffice to describe the simplest method which is recommended for bleaching prints, engravings, etchings, and drawings in carbon ink or in pencil, when these may safely be immersed in water. This method can be applied, provided an efficient fume cupboard is available in which all operations can be conducted. It is important that the cupboard be well lit so that the progress of the reaction can be observed through glass, and it should be provided with a drain and running water as all operations must be carried out in the closed cupboard.

To prepare the bleaching solution 75 ml. of 40 per cent. formaldehyde (Formalin) are added to a 2 per cent. aqueous solution of sodium chlorite (made by dissolving 60 grams of technical sodium chlorite in 3 litres of water) in an enamel photographic tray of suitable size. The solution is observed to become yellow owing to the formation of chlorine dioxide which is the active bleaching-agent. The stained print is laid on a glass plate to act as a carrier, and the whole immersed in the solution until the stains are removed. This will take at least fifteen minutes, but may run to upwards of an hour, depending on the nature of the stain. The concentration of the bleaching solution is not critical; it may be weakened or strengthened and 10 ml. of a wetting-agent such as Lissapol added if desired.

When bleaching is completed, the print, resting on the glass plate, is removed from the chlorite solution and washed in running water for at least fifteen minutes to remove sodium salts, no intermediate antichlor bath being required. The print should not be lifted from the glass support while still wet, otherwise it is liable to be damaged.

It has been found that this process removes water-stains, fox-marks, and mildew, and does not give the paper a staring white appearance, as is the tendency with processes involving the use of hypochlorite.

**Strengthening and Deacidification**

When papers are damaged or very brittle they may be strengthened by the process known as lamination. Lamination may be carried out either by covering both sides of the paper with fine silk of open
weave (Crepeline), using a starch paste or dextrin adhesive, or by sandwiching the paper between sheets of cellulose acetate, using a hot press, without the application of any adhesive. Laminated papers of various kinds have been tested by Scribner, who found that the sheets were strengthened without impairing the stability of the paper.

The silk method, which requires no special equipment, has been much employed for the repair of valuable books and manuscripts, but it may add considerably to their weight and bulk, and the adhesive has an obscuring effect, limiting subsequent study by photography. The paste fluoresces when the paper is examined by ultraviolet radiation, and this is a great disadvantage. The silk, moreover, is difficult to remove when, in time, it requires to be replaced, and as this may be, on a conservative estimate, every twenty-five years, there are some serious objections to 'siling'. By contrast, the plastic-sheet method has the great advantage of speed in processing, the cellulose acetate film is transparent to all radiations, and it may be readily removed at any time without straining the paper merely by passing it through baths of acetone. There need be little change in the appearance of the paper when laminated according to the cellulose acetate method, and there is the minimum of increase in the weight and bulk.

Any hesitation in recommending the cellulose acetate process for the lamination of valuable documents is concerned with the fact that, while cellulose acetate may be accepted in itself as innocuous and durable, the sheets used for laminating contain a relatively high percentage of plasticizer, and it is by no means certain that this will remain in the film indefinitely, or indeed that one or other of the plasticizers used may not in some as yet undefined way be deleterious to the paper. It is only fair to add that in the Barrow process which

employs no auxiliary adhesive, and which has been tested on a large scale since 1939, there is no evidence that these criticisms have any foundation. It would nevertheless be reassuring if a standard specification were forthcoming for cellulose acetate film suitable for the lamination of valuable documents.

There are two stages in the Barrow process, (1) removal of the acidity of the paper, and (2) lamination. These two operations are complementary and are designed not only to eliminate existing acidity but to protect the paper from acid attack in future. The lamination process confers mechanical strength and may also have the effect of intensifying faded writing. Details are as follows:

1. Removal of acidity. The paper is placed between two smooth flexible grids of copper, so that it can be conveniently manipulated, and immersed for twenty minutes in a saturated solution of lime water (c. 0·15 per cent. strength). The acid in the paper is neutralized and the paper then contains a slight excess of lime. It is transferred to a bath of calcium bicarbonate (0·20 per cent. strength) for a like time. In this bath the excess of lime is converted into calcium carbonate (chalk) which is finely precipitated in the fibres of the paper where it is retained. The actual quantity of chalk is very small, but sufficient in amount to confer protection against any further acid attack to which the paper may be subjected in urban areas.

2. Lamination. After the acidity has been removed and the paper is dry, it is placed between two sheets of cellulose acetate (thickness 0·002 inch, or for delicate work 0·001 inch), and sheets of fine tissue-paper are laid on either side making a five-ply structure. This is placed in the Barrow laminator, the papers and tissue preheated for a few seconds, then hot-pressed at 315–250°F. by passing through a pair of calender rolls from which a single laminated sheet emerges. The tissue-papers seem to disappear in the process of lamination as do the acetate films, and fine writing is made rather more legible owing to the high refractive index of the cellulose acetate with which it is now consolidated. Lamination by this process causes little change in appearance but there is a marked improvement in durability.
The Barrow laminator is a precision instrument consisting of a pair of thermostatically controlled preheating plates, and a pair of calender rolls, between which may be applied a pressure varying from 300 to 2,000 lb. per sq. in. When the papers and films are assembled, a sheet may be laminated in about thirty seconds. Incidentally, the equipment provides a ready means of mounting maps on cloth, and if the map is surfaced with cellulose acetate film (but not with tissue-paper) it becomes impervious to ink stains and grease.

INK

Having considered the principal materials used as grounds for writing, attention must now be given to inks.¹

The oldest writing-ink of any permanence is carbon ink, which, in its primitive form, was no doubt a mixture of fine soot held in suspension in oil, gum, or glue size. Carbon is fast to light, and as it is unaffected by chemical agents, it has lasted well.

The traditional writing-fluid in Ancient Egypt was a carbon ink. It was used on papyrus, wood, cartonnages, and potsherds. When papyrus is exposed to the sun, the natural biscuit colour of the reed is gradually bleached to a pale ivory, and as any writing in carbon ink is unaffected, the net result is that the ink appears to be darker and altogether more intense against the lighter ground. The binding medium of the ink (gum or glue) has probably long since decayed, but the carbon particles are locked permanently in the tissue and remain black. Chinese and Indian inks are carbon inks, and no problem is involved as regards their preservation, except when they happen to have been applied thickly to a non-absorbent ground, so that the ink adheres badly and tends to flake off. In this case the pigment may be fixed by applying thin gelatine size (p. 87). Carbon is the basis of our modern printing inks, and of the black inks used in the etching and engraving processes. Carbon inks are still preferred

for permanent writing today, and, as a class, they are referred to as manuscript inks. Black waterproof drawing inks are usually carbon inks containing a little varnish.

Carbon inks in general are all unaffected by the mild cleaning and bleaching processes carried out in the laboratory.

Iron inks are in a different category. They are compounded from gallotannic acid in presence of iron, and as the tannates are obtainable from a variety of natural sources, the inks vary in quality. Also from early times iron inks have been prepared according to a multitude of household recipes, and this is another reason for the wide variations in substance and permanence. In some cases the black is indistinguishable in appearance from carbon; in other cases the ink may be a rusty brown or yellow, and so faded as to make the writing illegible. It is not safe to assume that because a sample of writing is ancient and dark in colour, it must necessarily be in carbon ink. Iron inks have been identified which date to the second century B.C., and it is reasonable to suppose that they may have been in use before this; their discovery would approximate in time to the discovery of tanning as a means of preserving animal skins from putrefaction.

The acidity of iron-gall ink may be due to tannic acid or sulphuric acid which are present in varying amounts. Writings have been examined in which the acidity has caused the paper to become perforated and reduced to a frail membrane of lace-like appearance. This phenomenon is not confined, however, to iron-gall inks. Any acid ink may destroy the tissue in the same way, and as all sorts of mixtures have been employed as writing-inks, it is not surprising to find occasionally that even a carbon ink has been sufficiently acid to deteriorate the paper.

Iron inks are recognizable by a simple chemical test. When a tiny drop of 1 per cent. acetic acid is placed by capillary tube on dried iron ink, the ink will become locally soluble; the drop is absorbed into filter paper and it will be turned to Prussian blue if treated with a drop of 1 per cent. potassium ferrocyanide. It is seldom, however, that such a test is called for. In bleaching a print which bears a signature, it is safest to assume that the signature is in iron ink and
The tight roll of birch bark (×1½) as removed from the inside of a Buddhist image

Eleven thin birch bark sheets, inscribed in doggerel Sanskrit, recovered from the roll. (×¼)

9. INDIAN BIRCH BARK WRITINGS (18TH CENT.)
10. PAGE OF DRAWINGS BY LEONARDO DA VINCI

Left: Panchromatic photograph. (To the naked eye nothing was visible)
Right: Ultra-violet fluorescence photograph showing the drawings

By gracious permission of Her Majesty the Queen
therefore fugitive, and to protect it before treatment by a thin wash of celluloid (5 per cent.) dissolved in a solvent composed of equal volumes of acetone and amyl acetate. The film of nitrocellulose prevents the action of chemicals on the inks, and may easily be removed after treatment when the drawing is quite dry, by an application of a little of the solvent, the area being pressed afterwards with white blotting-paper.

As carbon is insoluble, the ink made by grinding it in a binding medium cannot be other than a suspension of fine particles, so that when applied to paper the medium will tend to be absorbed, leaving the particles in the interstices near the surface. With iron inks there is a greater tendency for the pigment to spread,¹ and, indeed, it is the flow of such inks that commends them for use with the pen, whereas in the case of carbon inks, the brush is the more suitable instrument for calligraphy.

In studying yellowed inscriptions that seem to be executed in iron ink, one should bear in mind that artists and writers have often been attracted by rusty colours, and at times have chosen to write or draw in materials which are recognized today as being impermanent. Such are the inks made from sepia (cuttle-fish ink), and from bistre (beech-wood soot). In cleaning and restoration work it is safest to assume that all pale rusty ink lines are fugitive. Coloured inks also are all fugitive. The most stable appear to be the reds. These vary in composition depending upon their source: some, such as madder and logwood, are extracted from plants, and others are obtained from insects, for example, kermes from Persia, grain from Poland, and cochineal from Mexico. The so-called Tyrian purple comes from several species of shell-fish. But even the reds are evanescent, and care should be taken not to expose them to any form of treatment that might result in discoloration or loss.

To try to protect any of these fugitive colours by applying a thin wash of nitrocellulose is not recommended, as in some cases celluloid solution might cause the colours to run.

¹ For the effect of the migration of inks, &c., on the permanence of paper see Barrow, W. J., Archivum, 1953, 3, p. 105.
Reading faded writing

Chemical methods of intensifying writing should not be used until the possibilities of obtaining information by modern photographic processes have been exhausted.

When it is desired to read writing that is illegible, it is first examined by different forms of illumination with the help of light filters, in the hope that by this means details will be revealed which are not visible in the ordinary way.

A source of ultra-violet illumination in a darkened room is sometimes particularly valuable in making indistinct writing readable, and at other times the infra-red viewer may give results that are otherwise unobtainable. When an inscription is partially legible under ultra-violet light, the chances are that a photograph taken using this source will be even more rewarding; and likewise, should the infra-red viewer afford any suggestion that the writing is intensified, it will be worth while taking photographs on specially sensitized infra-red plates using a tungsten lamp source. It is quite impossible to predict in advance which method, if any, is likely to be the more effective, and the results are often entirely negative. It may be stated, in general, that the ultra-violet type of examination is more useful than the infra-red, though with dark subjects the latter is sometimes strikingly successful.

Photographic processes are of great value in studying faded or damaged drawings as they provide a means of intensifying faint lines and suppressing stains, in this way making it possible to appreciate the artist’s work in facsimile. Fluorescence photographs obtained by using a filtered ultra-violet source occasionally yield surprising results as in the case of certain pages from a sketch-book of Leonardo da Vinci in the Royal Collection at Windsor Castle, one of which is illustrated in Pls. 10A and B. An ordinary photograph taken on a panchromatic plate recorded the drawing of a hand. Inspection by filtered ultra-violet radiations indicated that several drawings were still extant on the same page and it was possible, by photographing the page under ultra-violet illumination, to obtain a complete record of the drawings.
Charred documents. When a document has been carbonized by fire, it may still be possible to photograph the writing by daylight using a high-contrast blue-sensitive plate¹ or by ultra-violet or infra-red illumination.

In some cases the writing may be rendered sufficiently distinct to be read by immersing the paper in a 5 per cent. solution of silver nitrate for about three hours, when the writing will appear black on a grey ground.² The method which seems to have met with most success, however, is that of Taylor and Walls³ in which the document is given several applications of a chloral-hydrate solution (25 per cent. in alcohol), and dried at 60° C. between each application. It is then treated with glycerine (10 per cent.), dried as before, and photographed, using a contrasty non-colour sensitive plate. It is claimed that this method, even in the case of typescript or printing, has never failed to give a readable result. A comprehensive summary of the methods available has been published by the Istituto di Patologia del Libro, Rome.⁴

CHAPTER III

PRINTS, DRAWINGS, AND MANUSCRIPTS

AN OUTLINE OF METHODS OF CLEANING, REPAIR, AND MOUNTING

In the conservation of prints, drawings, and manuscripts the practical problems of cleaning, repair, and mounting are numerous and varied. While the more complex operations and delicate manipulative processes require the specialist, a number of simple methods remain that can be applied in an improvised laboratory by the amateur, and it is the purpose of this chapter to present a series of practical instructions for the collector who may be interested to carry out some of his own repairs.

Successful restoration depends on practice and experience. Old and worthless prints and drawings should be collected for experimental purposes, and repeated trials made in order to acquire dexterity in handling delicate material, and to discover the merits and limitations of the various processes described. Reagents should be tested on inconspicuous parts of the work before applying them generally, the milder processes being chosen initially and the more drastic ones only when necessary, and after carefully considering the risks involved. Thus weak solutions are tried before strong, cold solutions before hot, and mild reagents before those known to be more powerful in their action. Although technical details are given, the various processes may require modification according to the type of material undergoing treatment, and it is to be understood that the methods only apply, unless otherwise stated, to the more stable forms of art—prints, engravings, etchings, and drawings in carbon ink or in pencil when the design is not affected by immersion.

A large shallow sink provided with hot and cold running water is
an essential for print cleaning but, beyond this, no elaborate apparatus
is required. The sink should have a draining rack on one side, and be
flanked by a fair-sized table on the other, the table having half its
surface covered with ¼-inch plate glass held permanently in position
by a wooden moulding of the same thickness. It is convenient to
have heating facilities for paste-making in the same room. The room
should be well lit with both natural and artificial lighting. Beyond
storage accommodation for papers and chemicals the following com-
pletes the list of requirements—

Double saucepan and jug.
Measuring glasses.
Sponge and glass cloths.
Large photographic trays (porcelain and enamel).
A quantity of good-quality white blotting-paper.
Drawing-instruments, rules, squares, &c.
Drawing-boards and drawing inks.
Architect’s soft erasers.
Flexible paper-knives.
Print-trimming knife and oilstone.
Paired sheets of plate glass of varying weight and size.
A few heavy weights.
Brushes for dusting, retouching, and applying paste.
Electric iron.
Adjustable reading-lamp.
Copying-press and a capacious fume cupboard—optional.

EXAMINATION BEFORE TREATMENT

The paper. Examine by transmitted and reflected light, using a lens
when necessary. Test the crackle on shaking.

(a) Is the paper very porous, soft, or spongy? It may be so soft that
it would be unwise to use immersion methods of cleaning (see
Japanese Prints, p. 82). When paper is moistened it expands, the
effect of any size or binding material is weakened, and, if carelessly
manipulated in this condition, it may easily be torn.

(b) Is the paper hard or brittle, or the surface pitted or rotten? The
hardness due to size must not be mistaken for strength—test in a
corner with water. Papers which have been in the tropics are frequently so brittle that they will not stand manipulation without fracture. When the surface is pitted, ink may be inclined to flake, and immersion methods should be avoided lest the ink or paint should float away. Take note of weakened areas, creases, tears, wormholes, &c.

The drawing or impression. To what category does the print, drawing, or manuscript belong? What is the nature of the ink (pp. 63 seq.)? Is the technique simple or compound? Is any water-colour present? Does the picture bear traces of having been retouched or treated previously?

The object to be treated must be carefully examined in every detail before a course of treatment can be decided upon.

REMOVAL OF MOUNTS AND VARNISH

1. Removing cardboard backing

The card is always removed from the print: never the print from the card.

Cardboard is composed of laminated sheets, and the first operation is to insert a knife at a corner to determine the number of layers. A long flat paper-knife may then be pressed in and the laminations successively removed from the back, leaving about two of the constituent sheets attached to the print. In this condition the print is held with the back in the steam of a boiling kettle until the card becomes quite soft. It is then laid face down on dry clean blotting-paper and the card drawn slowly away by dragging gently on a corner across the part as yet undetached. In this way, if the card has been sufficiently steamed, it may be removed without straining the print. The print is now laid face down on fresh blotting-paper, and any residual adhesive carefully removed with a clean sponge. If this operation is omitted, the paper will eventually cockle. It is then allowed to dry between fresh sheets of blotting-paper under a weight such as a sheet of plate glass.

When, as sometimes happens, the backing is an inferior brown
strawboard, nothing but prolonged washing will soften it. This type of board is not laminated and can only be removed by rubbing it gradually away with the fingers while wet.

2. Removal of paper backing

When completely pasted down on paper, a print cannot generally be detached by steaming. Lay it face down on glass, sponge the back of the mount with warm water. Now float the picture face upwards on lukewarm water and allow plenty of time for the adhesive to soften before attempting to detach the mount. When starch paste is hard and slow to respond to such treatment, softening may be facilitated by floating the drawing face upwards on a warm aqueous solution of an enzyme (Appendix I. 3). Remove residual adhesive and complete operations as in 1.

3. Removal of canvas backing

When a print has been pasted down on canvas fixed on a stretcher, a sharp knife is inserted at one corner and drawn along the edges to remove the canvas and print from the wooden frame.

The back of the canvas is then sponged with lukewarm water, and laid against a wet sheet of glass, the fabric being made as wet as possible and the print kept as dry as possible. After some time when the adhesive is sufficiently soft, clean blotting-paper is laid on the print, and the blotting-paper, print, and canvas inverted and placed for support on another sheet of glass. The canvas is now worked back from a corner by gently pulling it across the part as yet undetached. If the paper starts to split, and shows signs of partly coming away on the canvas, stop immediately, apply very hot water to the canvas, and wait for some time before proceeding.

Complete the operation as in 1 above.

4. Removal of varnish

A print varnished with an oil varnish may be irretrievable as this type of varnish becomes insoluble with age. The removal of spirit varnish is possible, but is rather a specialized operation requiring
considerable care. The treatment given below may have to be modified to suit certain types of print.

Rub a tuft of cotton-wool in the palm of the hand (to give it a smooth surface), damp it, and lightly rub over the varnished surface. After drying, repeat the operation using turpentine instead of water. This clears the picture, and a record should be made of any colours before proceeding further.

In order to determine the most suitable solvent for removing the varnish, test a corner of the print with some methylated spirit. If this is not effective, try liquid ammonia (0.88) diluted with water (1:50). Support the print on a glass plate face up, and flood it with the chosen solvent, the action of which may be assisted by a flat camel-hair brush. Fresh solvent should be added repeatedly until it no longer becomes stained with the varnish. The print is now free from varnish although still more or less tinged faint brownish-yellow because the cellulose fibres have been stained with the darkened varnish. The print must be rinsed with water, then bleached (vide infra).

Plate 11 illustrates how, in the case of a printed book, it was possible to remove intensive varnish staining by the solvent action of alcohol followed by bleaching.

Old spirit varnish may be very brittle and may resist the above-mentioned solvents. However, hot water will sometimes soften it so that it flakes away. Pour boiling water into a large enamel photographic dish and immerse the print. As this method cannot easily be controlled where colours are concerned, it is risky with anything other than an engraving.

CLEANING: DRY AND WET METHODS

1. Dry methods

Although dry cleaning is not always necessary, pencil marks, &c., may become fixed in the paper if this is omitted before wet treatment is carried out.

If mildew is present, pick off the fluffy surface growths with a soft
camel-hair brush, care being taken that the spores are not scattered about in the process. Many are unavoidably rubbed into the tissue of the paper.

Use an architect's soft eraser for dry cleaning, or, if the dirt is too deeply ingrained, gently rub the surface of the print with a piece of stale (day-old) bread, using a light circular motion and taking the greatest care not to raise the surface of the paper. Change the bread for a fresh piece when it shows signs of dirt. The back of the print should be cleaned as well as the front. The print may be sterilized afterwards with thymol (p. 56).

Dry cleaning may be all that is required. The use of organic solvents (petrol, &c.) is not recommended except for removing specific stains. If dry cleaning in itself is not sufficient, two courses are open—either to clean the print as a whole by immersion in water, or to apply local treatment to remove specific stains.

2. Immersion methods

The print requires support when immersion methods are employed. Some workers use a sheet of plate glass, others a flexible support of stiff paper or Polythene. The print is never lifted from the water by its corners or handled while wet. It is the support that is handled, and when the support is slowly raised out of the water the print rests upon it and adheres so that it may be safely moved and if necessary turned over on to another support without being strained in any way.

A good soaking in cold water will always freshen up a print. After an hour it may be placed in a bath of hot water. Most fly-marks and mildew stains will respond to such treatment alone.

3. Cleaning with soap

In cases where further general cleaning is necessary, soap may prove to be a satisfactory cleaning agent. Carefully test the effect in a corner first. Lay the print face downwards on plate glass, damp it by contact with wet blotting-paper, then apply a little good-quality soap foam to the back with a large camel-hair mop. In larger prints
a very soft badger shaving-brush is a convenient tool, and if the tips of the hairs only are used without pressure, there should be no strain on the surface of the damp paper. If the results are satisfactory on the back, the face of the print may be treated similarly. All trace of soap must be removed afterwards by thorough washing; if this is omitted, the paper may go yellow.

Alternatively the print may be soaked in a dilute solution of a non-ionic wetting-agent (e.g. Lissapol\(^1\) or Nonex\(^2\)) for half an hour and then washed.

Washing is so important, especially when soap and chemicals have been used, that it must be carried out with something of the ritual of the photographic studio.

Lead a slow current of water by a rubber tube to the bottom of the dish containing the print, and allow the water to emerge below the support. The duration of washing will depend on the nature of the foreign matter to be removed and on the type of print under treatment, but at least an hour should be allowed where bleaching solutions have been employed.

4. **Drying and removal of creases**

In order to dry the print, it is laid face downward on a polished glass plate, and pressed on the back with a pad of blotting-paper to remove excess moisture. It is then set up to dry slowly in a gentle draught of air. Contraction of the paper on drying against the glass will remove most creases and marks of folds (see also p. 85).

If the print has lost its mechanical strength, it would be resized at this stage (p. 87).

The processes that have been described so far will freshen up a print and in many cases this will be all that is required. It is only when staining is persistent that it is necessary to have resort to chemical methods of cleaning—bleaching, and the action of specific solvents.

---

\(^1\) I.C.I. Ltd., supplied by Hopkins and Williams, Freshwater Rd., Chadwell Heath, Essex.

BLEACHING PROCESSES

Stains that have survived ordinary washing may often be removed by the process known as bleaching. This involves either treating the stain with a substance that will break down the colouring matter by oxidation into simpler colourless compounds that may be washed away, or treating the stain with a substance that will reduce it to a colourless compound that remains in situ. The most effective and permanent bleaching agents are in the first category (oxidizing bleaches): chlorine dioxide, hypochlorites, sodium perborate, hydrogen peroxide, and potassium permanganate. Examples in the second category (reducing bleaches) are sodium hydrosulphite and sodium formaldehyde sulfoxylate. While reducing bleaches are often effective in treating dye-stuff stains that withstand the action of oxidizing bleaches, they sometimes merely change the colour of the dye, but even if they go a stage farther and decolorize the stain completely, there is always the possibility that colour will eventually reappear owing to the subsequent oxidizing action of the atmosphere.

The simplest form of bleaching by oxidation is exposure to air and sunlight (or ultra-violet light) and this has given remarkably successful results in the case of Egyptian papyri where the writing material is a stable carbon ink.

The danger in using bleaching processes of any kind lies in the possibility of a loss of brilliancy in the inks and pigments and, if bleaching is overdone, the fabric of the paper may itself be attacked and weakened, especially when hypochlorites are used. For this reason bleaching agents must be applied under strict control and for the minimum time necessary to achieve the required results, and any excess of reagent is then decomposed with chemicals or removed immediately afterwards by thorough washing with water.

1. Use of hypochlorites

The traditional method of bleaching paper depends on the action of chlorine generated by calcium or sodium hypochlorite. The calcium compound known as bleaching powder is largely employed for bleaching the raw materials from which paper is made, and while
it may also be employed for bleaching prints, sodium hypochlorite is generally preferred as it is easier to prepare. Sodium hypochlorite is known, variously, as Eau de Javelle and 'chlorinated soda'. The stock supply of sodium hypochlorite should be the commercial preparation marked '10% w/v available chlorine' and it should be stored in a coloured bottle in a cool dark cupboard, otherwise it soon loses its strength. The solution must be diluted with water before use, generally one volume being added to twenty volumes of water; in no circumstances should the strength of the bleaching bath be greater than 6:20. The chlorine generated by this liquid has a powerful bleaching action, discharging the colour from dirt, mildew stains, fly-marks, and remains of varnish. If the action is prolonged unnecessarily, the paper itself will become a staring white and there is evidence that the cellulose may be degenerated and permanently weakened. The solution, which has an alkaline reaction, tends to soften the paper, but softening may be mitigated to some extent by having at hand a bath acidified with hydrochloric acid into which the print is transferred for a few moments now and then during the bleaching. Such a bath may be prepared by adding about a teaspoonful of concentrated hydrochloric acid to a quart of water. As in all processes involving the manipulation of wet paper, the use of a back-sheet or support is essential.

During the bleaching process, any writing in iron-gall inks will disappear unless protected beforehand. This is done while the paper is still dry, by a local application of a solution of nitrocellulose (5 per cent.) in a mixed solvent consisting of equal volumes of acetone and amyl acetate. The nitrocellulose can be removed by a wash of acetone at the conclusion of operations when the print is dry.

Bleaching is only allowed to proceed until stains become faint. The print is then passed through a bath of sodium thiosulphate (photographic hypo) of 2 per cent. strength to remove the residual chlorine, and washed thoroughly, when it will be found that the remaining marks gradually disappear. The hypo functions as an 'antichlor' and its employment is always desirable when hypochlorites have been used for bleaching.
2. Use of chloramine-\textit{T}

Chloramine-\textit{T} is a much milder form of bleaching agent than the hypochlorites mentioned above and its use is recommended next to that of sodium chlorite for routine bleaching. Chloramine-\textit{T} possesses the unique advantage that, when applied to a print, its bleaching properties are soon lost and nothing of a corrosive nature remains on the paper. Washing may thus be reduced to a minimum or entirely dispensed with. The process is particularly suitable for water-colour drawings, coloured subjects generally, and drawings in bistre and sepia, as the reagent may be applied locally to those parts of the work that are stained without exposing the whole work to the action of the bleaching reagent.

Chloramine-\textit{T} is obtainable commercially in the form of a fine white moderately soluble powder and must be kept in a well-stoppered bottle on account of its instability. Dissolve, only immediately before use, 2 gm. in every 100 ml. of water. Apply the solution to the stain with a soft camel-hair brush, cover with a pad of blotting-paper, and place under a sheet of glass. After an hour the print should be examined. Further applications may be necessary since the reagent is very mild in its action.

3. Use of sodium chlorite (chlorine dioxide)

This is the safest bleaching agent to use as there is no chance of its chlorinating the cellulose fibres of the paper; it is effective and easy to use but requires specialized equipment. The simplest method, which may be applied where there are fume cupboard facilities, has been described on p. 60.

4. Bleaching coloured and delicate prints

Many prints and drawings cannot be immersed in hypochlorite solution without damage, notably when colours are present. In the case of a stained coloured print, the best that can be done is to lay the print face down on glass; place a pad of wet blotting-paper on the back of the stain for a few minutes and then follow this with the
minimum of bleaching solution which is applied to the moistened back of the print with a camel-hair brush. It permeates the paper, having a mild bleaching action on the front. The stain should be kept under observation through the glass. When the result is satisfactory, flood with 2 per cent. hypo solution, then wash the print in situ, holding it against the glass all the time and at an angle to the running water. Now reverse the print, using a flexible support, and lay the support against the glass with the print face upwards, allowing a gentle stream of water to run over the picture for a short time. The print is then allowed to dry in this position.

An alternative method of application has been suggested for cleaning India-proofs and very thin papers. A sheet of blotting-paper is impregnated with a rather stronger solution of chlorinated soda, say 4:10, and allowed to become almost dry before being folded around the print and then sandwiched between two sheets of plate glass. The greatest care must be taken when removing the blotting-paper, and if there is any tendency for it to stick, it must be damped, otherwise it may bring away some of the ink and damage the print. When prints are of such a nature that they cannot be washed after this treatment, they should be exposed freely to the air for at least 2 days before being returned to the collection. It should be noted that where ‘antichlor’ cannot be employed, the method is not without danger to the paper, even though, judging by appearance, the result of operations seems to be entirely satisfactory.

5. Use of specific bleaching-agents and solvents

(a) Oil, fat, and tar stains: Pyridine. Pyridine, only the purest form of which should be employed, is an invaluable solvent for old partially oxidized oil, and for asphaltic stains, being decidedly more effective than benzene.

(b) Wax and candle-grease stains: Petrol. Some of the grease can generally be removed with a paper-knife. The whole print is then immersed in a bath of petrol. After soaking a few minutes the stain is rubbed gently with a camel-hair brush and soon disappears.

(c) Fly stains: Hydrogen peroxide, &c. Stippling the spots with
hydrogen peroxide in an equal volume of alcohol is often effective. If this fails, try stippling with 2 per cent. aqueous chloramine-T.

(d) Tea and coffee stains: Potassium perborate. Damp the area. Stipple a 2 per cent. aqueous solution of potassium perborate on the stain and expose to sunlight for an hour or so. The bleaching action is slow and as the reagent is alkaline it is not without danger to the paper. If the paper seems to be softened unduly, the action should be stopped at once by flooding the affected part with water. The final bleaching may be done by using ethereal hydrogen peroxide after the paper has been allowed to dry (see (f) below).

(e) Ink stains. Owing to the great differences in iron-gall inks, and even in modern inks of the blue-black type, no single process can be advocated as certain of success. A number of methods are available. Some of these may be found to bleach most of the stain, leaving a yellow tinge on the paper which, in turn, may be discharged by an entirely different process and reagent. Some of the possibilities are detailed below:

Brush with freshly prepared 2 per cent. aqueous chloramine-T. If not completely effective after two or three applications, and if the subject allows, try 5 per cent. oxalic acid or 10 per cent. citric acid, and then wash thoroughly. No tannate of iron can survive any one of these methods without being bleached, and if coloured matter remains, it is likely to be carbon or the residue of some dye-stuff. Always stop the action short of complete bleaching, and take no risks as regards the possibility of rubbing up the surface of the paper.

Should a coloured stain still persist when dry, a stronger bleaching agent will be required. Damp the stain and cover with powdered sodium formaldehyde sulphoxylate. This bleaches most ink stains, iron stains, and many dye-stuffs used in coloured inks, but thorough washing is essential after such treatment.

One further method may be mentioned, strictly as a reserve process to be used in case of emergency. It must be regarded as a last resort, as it is definitely deleterious to the paper; at the same time it seldom fails. An aqueous solution (0.5 per cent.) of potassium permanganate is painted over the stain where it forms a brownish-red
blotch of manganese dioxide. After about five minutes cover this
with a 2 per cent. aqueous solution of oxalic acid. The brownish-red
colour soon disappears, and the paper will usually be found to bear
no trace of the original stain. Thorough washing after treatment is
essential.

(f) **Blackened white lead and red lead: Hydrogen peroxide.** White lead
(basic lead carbonate) is readily converted to the black sulphide by
the action of sulphuretted hydrogen gas present in industrial atmos-
pheres. Red lead, an oxide, is similarly affected and in each case
stains are caused which are very disfiguring. The black sulphide may
be readily oxidized by hydrogen peroxide to lead sulphate which is
white and thus it is possible by simple treatment to remove the
staining and restore the brilliancy of the white pigment. In the case
of red lead that is superficially blackened, treatment by hydrogen
peroxide results in the formation of a thin white veil of lead sulphate
covering, but not concealing, the red pigment, and as it is unusual for
red lead to be converted more than superficially to sulphide, the
hydrogen peroxide treatment is generally all that is required to effect
restoration.

The direct application of a commercial solution of hydrogen
peroxide is sometimes recommended, but this is undesirable as it is
liable to contain corrosive impurities. The following methods are
designed in order to prevent contact of the corrosive impurities with
the paper. The hydrogen peroxide solution should be poured on a
stucco plate or porous tile, which is then fixed over the print about
\( \frac{3}{4} \) inch above it and left for a few hours. The vapour will clear the
blackened pigments. Similar results are obtained by using a shallow
bleaching-box with a false bottom of stucco that can be moistened
with hydrogen peroxide. The print is placed in the box, face upwards,
and left there until the stain has disappeared.

A method of more general application is to employ an ethereal
solution of hydrogen peroxide prepared as follows. Equal parts of
hydrogen peroxide (20 vols.) and ether\(^1\) are shaken together in a

---

\(^1\) Ether must be used only in a well-ventilated room free from naked lights as it is
highly inflammable.
(1) Text obliterated by a darkened resin varnish
(2) Extraction with alcohol removes resin leaving yellow stain
(3) Bleaching with sodium chlorite and formalin restores legibility

(4) Dark residue of resin obtained by evaporating the alcoholic extract
(5) The book, rebound

II. PRINTED BOOK (ITALIAN, 16TH CENT.)
A. Before treatment, showing stains due to the action of sulphur compounds on white lead.

B. After treatment with hydrogen peroxide

12. WATER-COLOUR DRAWING BY GUARDI
glass-stoppered bottle. The liquids are immiscible—the aqueous layer containing any impurities remains at the bottom, the ether layer rises to the top and this contains sufficient hydrogen peroxide for bleaching purposes. A glass tube (diameter ca. ½ inch) plugged with cotton-wool may be used for dabbing on the reagent. The tube is held in the hand, the cotton-wool dipped into the top layer only, and the ethereal solution is then applied to the stained areas by dabbing. Lead sulphide stains are bleached very readily by this method (Pl. 12 A and B). The action of the vapour may be prolonged by having a piece of blotting-paper at hand to slip over the print immediately after applying the reagent. The bleaching action may be intensified, if necessary, by adding a drop of perhydrol to the ether.

It is not unusual to find painted miniatures on paper or parchment heavily stained in places where white or red lead has been used, the stains being black or deep brown and having sometimes a silvery sheen. These can be freshened up very easily with hydrogen peroxide, and for this type of work it is best to employ the ethereal solution which may be applied locally with a small brush. If there is no immediate response to the action of hydrogen peroxide, it may be that the stains have arisen from the action of hydrogen sulphide on metallic silver. Silver was often used in Persian miniature landscapes to represent water, and when once blackened by sulphide, no satisfactory method can be recommended for its restoration.

Occasionally, a black smudge of lead sulphide will be found unexpectedly on a drawing, and closer inspection will reveal it to be a residue from a previous restoration that has been carried out either in white lead or in coloured pigments that have been mixed with white lead. In such cases the hydrogen peroxide treatment works satisfactorily as, even in the case of a pigment mixture, the original hue is restored by the conversion of lead sulphide to the white sulphate.

After hydrogen peroxide treatment there is much less chance of the white lead being blackened in future, because the sulphate is less susceptible to change than the basic carbonate.
SPECIAL CASES OF CLEANING AND RESTORATION

1. **Japanese prints**
   Few of the ordinary methods of treatment apply in the case of Japanese prints on account of the soft texture and quality of the paper. It is fortunate that this paper does not appear to be greatly attacked by foxing or mildew and the dirt is generally superficial. To clean a Japanese print lay it face down on glass and cover it with a rather larger sheet of tissue-paper (or preferably Japanese tissue which is thinner), in such a manner that one end of the tissue projects and can be held against the glass. Apply plain water all over the tissue in a series of light parallel strokes using a tuft of cotton-wool; sufficient will soak through the tissue to make the dust on the back of the print adhere and on carefully folding back the tissue this dirt will be removed. If not entirely successful, a very dilute gelatine solution (half strength, see page 87) may be substituted for the water, a fresh operation being conducted on the same lines. After cleaning the back treat the front similarly. As mauve and heliotrope pigments on Japanese prints are generally of vegetable origin and easily damaged, they should **never** be damped.

2. **Japanese vellum mounts**
   The so-called Japanese vellum forms an ideal mount for Japanese prints. Although it is a hard paper, the surface will not bear cleaning by the ordinary dry methods as it is very easily rubbed up. A clean soft sponge or, preferably, a tuft of cotton-wool is dipped in dilute starch solution, squeezed to remove excess liquid, and passed lightly and rapidly over the surface. By this means the dirt is picked up and the mount greatly improved in appearance.

3. **India proofs**
   The cleaning of an India proof is an extremely delicate matter on account of the ease with which the India paper becomes detached from its mount. When once detached the difference in expansion and contraction between the thin paper and its thicker mount renders it almost impossible to regain the original tension between the two
when they are remounted and dried. It should be noted also that the sharpness of detail in an India proof is bound to be lost to some extent if it is necessary to wet the paper for cleaning. Stains should therefore be removed, for preference, by dry cleaning or by local treatment if possible.

Reference has already been made (p. 78) to the method of cleaning using blotting-paper impregnated with chlorinated soda. This method, as has been suggested, is open to criticism, but alternative methods milder in action would necessitate longer contact with the damp bleaching-paper and result almost inevitably in the separation of the print from its mount. If the two sheets are only partially separated, they had best be completely detached from each other by soaking and floating them apart, and then chlorine dioxide (p. 60) or a chloramine-T bath may be used to clean both the print and the mount.

In order to remount the dry print it is laid face down on plate glass; the back is pasted all over, taking care to keep the paste off the glass, and the glass is then turned over and placed against the mounting paper which has been previously damped so that the India proof adheres in its correct position. The mounted India proof is dried superficially with blotting-paper, and after about ten minutes removed from the glass. It is then allowed to dry slowly under a weight protected by sheets of blotting-paper on each side. Should the blotting-paper adhere, it may easily be removed by damping with water.

For the preparation and use of flour paste see Appendix III.

4. Pastel and chalk drawings

Pastel drawings, especially those on vellum, should be frequently inspected as they are liable to become infected with moulds. Moulds may attack certain pigments (Indian yellow; bile yellows) or they may grow on the binding medium of the powder colour—generally gum tragacanth—or on the sizing of the paper.

When mildew is found growing on a pastel it must be picked off with a fine camel-hair brush slightly moistened with pure alcohol.
The picture is then sterilized with thymol. If the mould growth has caused permanent staining it may be possible to hide this by spreading the pastel pigment with a stump. Pastels should always be kept in a dry place and they should be framed in contact with a sheet of paper impregnated with thymol or Santobrite (see p. 57).

The treatment of pastels by amateurs should be restricted to the above.

When certain types of chalk drawings (outline drawings on inferior brown paper) are mildewed they may become stained in a manner which is very disfiguring owing to the sparsity of pigment. The stains cannot be bleached as the paper would be bleached also. There is a tendency in old pastel and chalk drawings for the colour to be fixed in the paper as a result of the prolonged effect of traces of moisture upon the gum of the pastel and size of the paper; these adhesives bind the particles of pigment that are in direct contact with the cellulose tissue. Water may actually be used for cleaning drawings of this nature, as follows: float the drawing on the surface of cold water face upwards (with a paper support immersed below), and rub the spots gently after some minutes with a fine brush. If this is only partially effective after ten minutes, remove the drawing, using the support, and float it on to very hot water in the same way. This treatment is only possible with old drawings when the pastel pigment has become fixed in the tissue of the paper. The mould growths are mainly on the surface of the paper and are removable after softening on water. By this treatment the worst of the staining is dispersed.

5. Bound papers

It is sometimes necessary to remove a page from a book for treatment by one or other of the methods described above. This is done as follows—insert a wet string against the page near its hinge and close the book. After a minute the page may be torn out. When the page is ready to be returned, the torn edge is pasted and the page aligned with the fore-edge and head end of adjacent pages; this will bring the pasted edge automatically into register with the hinge from which it was torn and it is caused to adhere by placing the book in the press.
REPAIR AND RESTORATION

1. Removing creases from paper. When not very bad the creases should be damped and a warm iron should then suffice to remove them.

Reference has been made above to the removal of creases by drying the print against glass. In the case of very bad creases the print is damped and laid face downwards on glass. Strips of soft white paper are then pasted all round ¾ inch on the glass and ¼ inch on the print, so that the tension may be greater as the print shrinks on drying. When the print is dry and flat, a little paste should be rubbed along the line of the crease and, if necessary, a strip of similar or thinner paper applied as a patch and allowed to dry before releasing the print from the glass. In carrying out such operations the guard papers must not be heavier or stronger than the paper to be flattened, otherwise tearing may occur.

2. Removing creases from parchment or vellum. With parchment or vellum the problem of creasing and warping is much more complicated and even prolonged treatment in a press is seldom effective.

A practical method of flattening parchment is first to relax it by sandwiching the membrane between two sheets of damp blotting paper and pressing lightly for a time. The blotting-paper should be uniformly damp but not too wet. The time required for this operation will be determined by the nature and thickness of the parchment. When the membrane is limp, it is placed on a glass that has been previously polished with French chalk to remove grease, covered with a sheet of dry blotting-paper of the same size, and held against the glass by weights placed around the edges. Strips of ¼-inch lead are ideal for the purpose and they may be as broad as a ruler. Weights should not be so heavy as to immobilize the parchment, the process requiring that, as the parchment dries and contracts, it will slide over the glass and take the weights with it, the centripetal pull being rather greater than the restraining action of the weights. By this means, even long-standing creases may be made to disappear. The most difficult types of membrane are those cut from near the edge of the skin, where thickness and grain are inclined to be variable, and
these may need localized treatment in some form of stretching frame in which the forces can be adjusted to the requirements of the membrane. Such an apparatus has been described by Cockerell¹ and there are cases in which the frame is an invaluable adjunct in removing creases from parchment.

If such operations are practised on worthless documents it will be noticed that care is necessary to prevent ink spreading and it is a wise precaution to fix ink beforehand with nitrocellulose as mentioned on p. 65. When colour is present, and especially gold lettering, the membrane must on no account be relaxed. Restorations of this nature can only be conducted satisfactorily by an experienced bookbinder.

3. Tears in paper. If the quality of the work allows, immerse the paper in water face downwards on a glass support, and when the torn pieces have been carefully floated into their correct positions slowly raise the support and paper out of the water. When half dry, tap along the joints with the back of a spoon in order to weld the fractured surfaces together. Paste may be added as a reinforcement and also a patch if necessary.

When machine-made paper is being repaired, the grain of the patch must be aligned with that of the paper. The direction of the grain of a machine-made paper may be discovered by moistening two edges at right angles; the paper swells in the direction across the grain, and acquires a crinkled appearance along the edge.

Hand-made paper should be patched with another piece of thinner hand-made paper using rather dry paste. Press very lightly thereafter.

When an insertion is required, it should be of the same age, thickness, and surface appearance as the paper being repaired. The restorer must have at hand a reserve collection of sample papers, engravings, &c., from which to select repair material when necessary. Even when the greatest care is taken in matching the papers, some allowance may have to be made for variations of sizing, and differences of shrinkage on drying. The tendency is to make the patch too damp with paste. The edges of the insertion and the paper are chamfered as described

¹ Cockerell, D., Bookbinding and the Care of Books, pp. 329 seq. 1901.
below in the repair of parchment, and after the edges have been pasted and the insertion stuck in position the whole is flattened in the press.

When a Japanese print is broken or much torn, it should be completely pasted down on another Japanese paper of similar quality and strongly pressed.

4. *Tears in parchment and vellum.* When creases are absent, there is no need to soak parchment or vellum. A simple chamfered tear may be repaired by sticking the edges together and pressing under a weight till dry. Paste, however, is not a very good adhesive for parchment. The replacement of torn corners requires a much more powerful adhesive, and stronger joints are obtained when the edges are painted with the minimum quantity of dilute acetic acid (say 10 per cent.). This gelatinizes the fibres so that the torn edges can be pressed intimately together.

Where there are no natural chamfers, as in a plain cut, the edges must be artificially chamfered by rubbing with an abrasive stick of chisel shape. This enlarges the cut, making it a V-shaped slit, and a patch of similar appearance but rather greater in size and thickness is cut to shape and then chamfered down to fit. It is inserted after applying adhesive to the edges, dried in the press, and then brought to the exact thickness by rubbing with a block of fine pumice. When such work is carried out carefully and the sheet rubbed over with a little pumice flour at the end of operations, the repair may be so perfect as to be almost beyond detection.

5. *Sizing and retouching.* The paper will require to be sized afresh if any retouching has to be done. Parchment size is sometimes recommended for this purpose but it has no advantages over gelatine. A good gelatine size is made by dissolving one sheet of clear gelatine in a quart of water (not more than 1·5 grams per litre). Size should always be freshly made immediately before use, and brushed thinly and lightly over the paper. Thick papers may be sized by immersion and hung on a line to dry away from radiators and draughts.

If any retouching is necessary, this should not exceed the absolute minimum required to conceal disfigurements due to tears and
abrasions of the paper. The method should generally be by stippling save where broken lines have to be joined up.

6. Toning. When one page of a book has been bleached or when one of a uniform series of engravings has been similarly treated, it may appear blanched and uninteresting beside its fellows. This may be remedied in a rule-of-thumb fashion by staining the paper with stout or a decoction of tea or coffee. Preliminary trials are made on similar paper in order to decide which stain gives the best results and to determine a suitable concentration.

MOUNTING

1. Parchment and vellum

Great care is required in mounting parchment and vellum as these are generally mounted in the damp relaxed condition and contraction on drying is considerable. If the paste is not uniformly applied around the edges, distortion will result. Assuming that there are no illuminations present, the parchment is relaxed in water and placed in sheets of dry blotting-paper between glass plates until it appears almost dry, while still retaining its suppleness. A thick mounting board is selected and four guide-marks made to show where the print is to be permanently fixed. The membrane is laid face down on glass and pasted half an inch all round the edge two or three times. It is then turned over and placed in its permanent position on the mount, and the edges pressed down very thoroughly with clean blotting-paper. It must be remembered that the edges will have to withstand a fair strain as the parchment shrinks on drying. Should the parchment have been too wet when mounted, the edges will pull away, dragging off part of the mount with them. Before drying, remove any marks with a damp sponge. After covering with two or three sheets of blotting-paper, weights are applied over the four corners and the membrane is left to dry, when it will become as tight and smooth as a drum.

Vellum is treated in similar fashion. The thinnest and finest specimens do not require preliminary soaking. It is sufficient to leave them
for a short time in damp blotting-paper pressed between glass plates before mounting.

When there is colour or gilding present, the membrane must on no account be relaxed, and manipulative processes are best left for the expert.

2. Papers

In the absence of special laminating equipment (p. 63), manuscript material is commonly mounted in guard-books (Pl. 8).

Prints and drawings are never ‘laid down’ (pasted down all over) on card, but attached to the card by hinges of thin paper folded back on themselves and known as guards. When only one such hinge is used the drawing can be turned over to inspect the reverse and watermarks can be studied by transmitted light. More usually four guard papers are employed and there is then less chance of accident by careless handling.

Only the best white or ivory cards should be used for mounts. These are cut to size and increased in thickness as required by pasting cards together. The mounts should be strongly pressed after pasting.

All mounts should be finished with rounded corners to avoid the chance of damage should they be accidentally dragged over the surface of adjacent prints. This can be effected by smoothing the edges and corners with glass-paper.

Three types of mount will be described—the solid, the overthrow, and the window mount. These are illustrated in Fig. 4 (p. 90).

(a) The solid mount. This type of mount is the one most commonly used, when there is nothing of interest on the back of the print. The print or drawing is attached by pasted paper hinges in a ‘box’ prepared by pasting a cardboard (3 sheet) frame to a backboard (4 sheet) of the same size as the frame. In the solid type of mount the cardboard frame does not cover any of the print. The backboard must always be thicker than the frame, the relative thickness depending on the size of the print. A comparatively large print would require a strong backboard for support. Deep ‘boxes’ must always be employed in mounting pastels.
(b) The overthrow mount. This type of mount is in general use where prints are hinged by one guard paper only so that the back can be inspected. It is also used when a print has a torn edge. Such a print is mounted on a backboard by pasted guards of paper and a frame is
prepared of such a size as to conceal the damage. This frame is hinged to the backboard by a strip of linen so that it may be opened to the left to expose the whole of the print for examination. A hinged arrangement of this kind is known as an overthrow mount.

(c) *The window mount.* When drawings occur on both sides of the sheet of paper the paper may sometimes be split so that the drawings are released and they may then be mounted separately. As the operation of splitting paper is always attended by risk, it is generally preferable to adopt instead the window type of mount so that both pictures remain visible. In this case the paper is attached only by its edges between superimposed rectangular openings of appropriate size in two cards. As the drawing might easily be damaged if left unsupported it is usual to mount a sheet of cellulose acetate\(^1\) at one side so that it forms a transparent support. If the paper is very frail or where much ink is present that may have rotted the tissue, it may be necessary to give it additional support by adding a similar sheet at the other side.

It is a general principle in mounting a collection to endeavour to make mounts in a limited number of sizes as this is convenient both for storage and for exhibition purposes. The card surround should be of such a thickness as to protect the picture from contact with the glass when framed. If the picture is in contact with the glass, condensation effects may encourage the growth of mildew.

3. *Documents with Seals*

When documents bearing appended seals are to be framed, they should first be inlaid in a solid card mount of such a size and thickness that the seals are backed by, and nested in, the card, so that their weight is supported. This protects them from the possibility of accident should the fabric by which the seals are attached become tendered and no longer able to sustain their weight.

Wax seals are easily broken and, if not repaired at the time, fragments tend to get lost and reconstructions become more complicated. A mixture of equal parts of rosin and beeswax is usually recommended

\(^{1}\) e.g. Celastoid. British Celanese, Ltd., Hanover Square, London, W.I.
as an adhesive for wax and, if it is a case of dealing with a simple fracture, this adhesive may be spread in the form of a hot liquid on the joints, pressing them together immediately till the parts are held securely. With heavy seals it may be necessary to insert one or two dowels before making the joint. The holes for these are made by probing with a hot needle, and the actual dowels can be made from fragments of broken needles of appropriate length and thickness. It is sometimes possible to reinforce a heavy seal by attaching a thin adhesive tape around the circumference in such a manner as to be unobtrusive, and it may be painted thereafter to match the colour of the wax.

When a seal is splintered and parts are missing, the main fragments should be brought together as above and any lacunae made up with plaster of Paris containing a little ochre and black as a rough match for the natural wax colour. Green and red colours are sometimes required. The tinted plaster should extend slightly above surface level, and after it is quite dry it is carved to match the design. It is then impregnated with thin celluloid lacquer to stop the suction and painted to match the original. While a repair executed in this manner should not be obtrusive or out of colour harmony, there should be no effort to conceal the fact that the seal has been restored.
CHAPTER IV

TEXTILES

The spinning of threads and the weaving of textiles come very early in the history of craftsmanship. Linen in southern Europe can be dated back as far as the Stone Age; in northern Europe wool was in use in the Bronze Age, and silk, which originated in China, can be traced back for more than 5,000 years. Embroidery, i.e. the decoration of the finished textiles, did not appear till much later.

Natural fibres may be of animal origin (wool, silk), or of vegetable origin (cotton, linen, hemp). Animal fibres contain keratin, a nitrogenous compound, and, when burnt, they contract and smell of burnt feathers due to the decomposition of the keratin. Vegetable fibres are composed of cellulose; they burn easily and give off a characteristic odour of burnt paper. Thus, it is possible by a burning test to make a broad distinction between animal and vegetable fibres, but, for specific identification, individual fibres must be examined under a microscope at about 100/150 diameters magnification when, with a little practice, the main types can readily be distinguished.¹

As textiles are of an organic nature and subject to attack by moulds and bacteria, the commonest factors that promote decay are those that favour the growth of these organisms, viz. a damp heat, lack of ventilation, and contact with decaying animal and vegetable matter. However, even under such unfavourable conditions ancient textiles may not be entirely destroyed. It is common, for example, to find fragments of textile surviving when they have been in contact with corroding copper, the corrosion products having acted as sterilizing agents. Textile remains are often found attached to the patina of

Chinese bronzes, in such a condition that the fibres can be identified and the weave recorded. Damp causes vegetable fibres to swell and soften, but the fact that animal fibres are not necessarily destroyed by immersion in water is proved by the survival of woollen material in peat bogs and lake dwellings. Excessive heat causes desiccation and embrittlement, and exposure to bright light and noxious gases cause the type of deterioration known as tendering. The astonishingly fresh appearance of some Egyptian mummy wrappings may be due as much to the presence of salt as to the absence of light and dryness of the surroundings.

TEXTILE ANALYSIS

Before beginning restoration work on textiles the first step is to carry out an examination with a pocket lens or binocular microscope, making a record of the following particulars.

The nature of the fibres of both warp and weft threads.
In spun threads, the direction of twist—S (') or Z (').
The type of weaving—tabby, twill, figured.
The count per centimetre of both warp and weft threads.
The presence of selvages.
Dye-stuffs or applied decoration—paint, gold leaf, metal threads, or embroidery.
The presence of sewing or stitch holes.

(If the material is very dirty, making analysis difficult, gentle blasts of air from a blow-ball or bellows may be used to remove loose dust, or in some cases a soft brush may be used.) It may be impossible to record all particulars until the textiles have been washed, but it is important to get as much information as possible in the first place in case any evidence should be lost during treatment. The preliminary examination will decide what are the safest methods to adopt for cleaning and restoration.

1 Henshall, A. S., Miss, and Maxwell, S., 'Clothing and other Articles from a late 17th-Cent. Grave at Gunnister, Shetland'. R. Soc. Proc. (Scot.) 1931–2, 86, pp. 30–42. This grave was found in peat, and when excavated, scarcely anything of the body was left, but all the woollen clothing in which it had been buried was well preserved. This clothing is now exhibited in the National Museum of Antiquities of Scotland, Edinburgh.
CLEANING OF MUSEUM TEXTILES

Textiles are easily contaminated with soot and dust, and being absorbent they are readily stained by contact with substances in solution, or decaying organic matter, or by the coloured substances produced by moulds in the course of their growth. Material that is dirty or stained, but which is sufficiently strong to be manipulated, may be washed or dry-cleaned, and specific stains can be dealt with locally. It must be understood, however, that the processes recommended may require modification depending on the condition and type of textile that has to be treated, whether carpets, tapestry, lace, &c. Complications may also arise from the presence of a dye, or of gold thread, or because of the vagaries of the weave or make-up of the fabric.

1. Washing

The first requirement for washing textiles is a supply of soft water. Rain water or distilled water may be used, or water that has been softened (demineralized) by natural zeolites or by special synthetic ion-exchange resins. Old textiles are washed preferably in flat vessels. Small samples may be conveniently dealt with in photographic developing dishes, but for washing large textiles the most practical arrangement is to use a wooden tray lined with Polythene. Such a tray is constructed with sides about 4 inches high, in one of which a V-cut is made for emptying; a sheet of Polythene is spread in the tray and made to take the shape of the vessel, and is then held in position around the edges by spring clothes-peg. By this means a waterproof tank of any size can be readily improvised. A very large tank may be emptied by siphon, or preferably hinged to the table along one of its longer edges, so that the water can be run out into an adjacent sink by tipping the vessel, the action being controlled by a long screw such as that commonly used for opening and closing hinged window-lights. In emptying the tank, the Polythene is pressed into the V-cut so that it forms a gutter between the table and sink.

A loose sheet of Polythene should be placed in the bottom of the
tank to act as a support for the textiles when they are lifted out of the water.

In the case of textiles of open weave, drawn-thread work, or lace, these should be pinned down or tacked to a sheet of Polythene, for safety during washing.

If textiles are coloured, a spot test with water will have to be applied to each colour before washing to determine whether the colours are fast. Fugitive colours may be fixed either by treatment with a 5 per cent. solution of common salt or acetic acid of the same strength. A stronger solution of acetic acid up to about 20 per cent. may be used if necessary. A spot test should be made, and the spot absorbed in white blotting-paper to ascertain whether the salt or the acid fixes the colour. The whole textile should then be immersed in the appropriate fixing solution.

When everything is in order, the tank is filled with soft water to a depth of about 3 inches, and the textile lowered gently into it, keeping the material flat and well spread out; it is kept immersed for an hour, changing the water as required, perhaps every twenty minutes. If the mains' supply of water is soft, a gentle stream of running water may be used. This is conducted to the tank by a rubber or Polythene tube and allowed to emerge beneath the Polythene support. When the running-water method of washing is adopted (in the case of a tank larger than the sink), a siphon draining system will be required.

Some of the dirt will be soluble in water, and insoluble sandy matter may be released by gently tamping with the fingers after the textile has been soaking for some time. When the washing is finished, the water is run off, and the Polythene support, to which the textile adheres, is raised and maintained at an angle to allow the material to drain. While still on the support the textile is pressed gently with absorbent towelling and it is then transferred to a backing of warm absorbent material—towelling or flannel.

When half-dry, the textile is turned over on to a Polythene sheet and straightened out, the weave being adjusted so that the warp and weft threads are at right angles to each other. Fine brass pins are inserted vertically at intervals through the textile into the Polythene
A. Photograph on Panchromatic Plate

B. Photograph on Infra-Red Plate

13. Fragment of Chasuble (13th Cent.) from Lincoln Cathedral. Silk with Gilt Parchment Appliqué from which all Gold has Disappeared
A. As seen from front

B. As seen from back

14. MANiple OF St. Cuthbert FROM durham cathedral (10th CENT.)

After cleaning and mounting prior to framing
sheet. The pins may require adjusting from time to time as the textile dries. Drying should be carried out in a warm well-ventilated room, and it may be facilitated by using a hair-dryer.

Use of detergents. While washing in plain water can bring about a remarkable improvement in the appearance of certain textiles, the use of a detergent is necessary if textiles are contaminated with grease. Soaps are not used as they tend to form an insoluble scum difficult to remove, but certain synthetic surface-active agents or wetting-agents may be employed. These are of two kinds, ionic and non-ionic, the latter type being the safer for textiles. Examples are Lissapol-N¹ and the preparation Igepal CA Extra.² These substances can be used with confidence, but a warning is necessary against using any of the commercial cleansing powders and liquids that are based on them, because patent cleaners may contain in addition soap powder, soda, or other materials that might be deleterious to fine textiles. There is the added complication that the formulation of any commercial product may be altered without changing the name and without any warning being given by the manufacturers.

It cannot, of course, be assumed that wetting-agents are necessarily safe to use in the presence of dye-stuffs, and preliminary tests must always be made by applying a spot of the concentrated detergent to all the dyes on the material. If any of the dyes 'bleed' under this test, they must be fixed as described above with salt or dilute acetic acid.

It is characteristic of wetting-agents that they are used in extremely dilute solution (as specified by the manufacturers), quantities of the order of 1 fluid ounce to 5 or even 10 pints of water, i.e. 1 per cent. to 0.5 per cent. strength, being adequate for most purposes. In using detergents, they should be added in the requisite amount to the water and thoroughly stirred so as to form a solution of uniform strength before the textile is placed in contact with the liquid. During washing, the wash-waters are changed at least three times, and a final bath of fresh soft water completes the operation. In cases where lukewarm water may be used, the cleaning will be even more effective,

¹ Imperial Chemical Industries, Ltd., loc. cit.
² Marketed by General Dyestuff Corporation, 435 Hudson Street, New York 14, U.S.A.
but the water should never be used hot as this has a softening effect on some fibres.

*Use of saponins.* Saponins are widely distributed in nature among the higher plants, and their virtues as substances having mild detergent properties have long been known. They are neutral in action, froth readily with water, and their cleansing properties are due to the ease with which they form emulsions with substances of a resinous or oily nature. Plant extracts containing saponins are largely employed in the East for washing clothes. They do not seem to have any deleterious effect on delicate fabrics, and are safer to use with coloured textiles than soap. The precaution should always be taken, however, to test the colours first.

A cleansing fluid may be made by infusing Saponaria (Soapwort) with water; or saponin can be obtained commercially as a white powder. To use the powder a little is switched to a froth with water and this is applied with a soft brush of the shaving-brush type. To clean upholstery the furniture is supported upside down or at an angle so that when the froth has done its work and is soiled it will fall away from the textile rather than sink in. The froth is lightly worked over the surface of the textile using only the tips of the bristles of the brush, excess froth being removed with a soft rag, and the textile dried thereafter with absorbent towelling. By this method excellent results are obtainable without rubbing and with the minimum of exposure to moisture. For cleaning damask that is frail with age, the process may be used with appropriate variations. It will usually be found that in cases where it is safe to use water, dirt can be removed more readily with the aid of saponin, and that colours are improved in appearance and enlivened by the treatment.

2. *Dry-cleaning*

Dry-cleaning may be carried out by mild suction, in conjunction with brushing, by using organic grease-solvents, or by dry steam, the nature and size of the textile being the determining factors in deciding which procedure is the most suitable.

*Vacuum-treatment and brushing.* Old carpets and curtains that are
very dusty may be improved in appearance by brushing with a soft brush (in the direction of the pile) towards a vacuum cleaner, or, alternatively, a suction nozzle may be used if the fabric is strong enough to stand it. It may be added that it is sometimes possible to improve the appearance of a rug by gently rubbing fuller’s earth or even baking soda into the pile, leaving it there overnight and then extracting it next day with a suction nozzle. In dealing with old rugs care must be taken to preserve the direction of the pile; if this is upset by violent local cleaning, the distorted fibres will scatter reflected light and the rug will have the appearance of being stained.

Brushes should be selected with care to suit the job in hand. The bristles should be white for preference and generally long, and the degree of softness is also important. Stiff brushes do have their place among the cleaning equipment but should be used with reserve.

Organic solvents. The term ‘dry-cleaning’ is usually employed to denote cleaning by anhydrous solvents. These are less likely to affect any batter in the cloth than water, but as they may well have a greater action upon dye-stuffs it is essential that all colours on a textile be tested individually by spotting with solvent before immersing the textile in the cleansing solution. Dry-cleaning is only given preference in cases where the presence of water is undesirable. The solvent most commonly employed is trichlorethylene (Westrosol), a non-inflammable liquid of high volatility (B.P. 88° C.). The Ciba Experimental Dye-house Laboratories, Basle, recommend that it should be used pure and cold, and that textiles should be immersed for from ten to thirty minutes at the outside. It is further recommended that where there is a danger of dyes running, dichlorethylene (B.P. 55° C.) should be selected as the cleaning agent in preference to trichlorethylene.

It is well established that when garments are soiled with grease and perspiration they are more prone to be attacked by moth than when freshly laundered, and cleaning with solvent has the same effect as laundering in discouraging attack. It goes without saying that garments should always be cleaned before they are incorporated in the collection.

Steam cleaning. For certain kinds of textiles, cleaning by steam (if properly controlled) may be much less drastic than either soaking with
water or cleaning with organic solvents. Steam may sometimes be used for softening and removing stains even in the presence of fugitive colouring matter that would spread if wetted. A steam-gun apparatus that will provide wet or dry steam at option is now regarded as an essential in modern laundry practice, and it could be applied with advantage to cleaning certain kinds of museum textiles—costumes, for example, or ethnographical specimens—as it makes possible a variety of cleaning operations that would not be warranted by any other process.

3. Removal of stains

It is not always advisable to attempt the removal of stains from old textiles. Stains of long standing may have undergone a chemical change with the formation of insoluble matter that can only be discharged by bleaching, and this process would be likely to weaken still further an old textile that had already become tendered. In certain cases, however, the removal of stains is to be recommended. In the unfortunate event of a textile being stained by accident, the staining should be dealt with quickly before it has had time to become fixed in the fibres. Iron stains commonly rot vegetable fibres, and these should be removed if the fabric is strong enough to withstand treatment. With frail old textiles, removal of stains may be a very delicate operation requiring experience and some knowledge of chemistry, but with modern fabrics, including many textiles in ethnographical collections, there is less risk of damage and, even for the amateur, a reasonable prospect of success.

In the removal of stains the first step is to ascertain the nature of the textile (e.g. wool, silk, or linen), and it is also a help to know the nature of the substance responsible for causing the stain; this may be obvious, but sometimes it will be necessary to make tests before the appropriate reagent can be selected. Even so, all sorts of difficulties are liable to be encountered. The textile itself may be decorated with paint susceptible to solvents; the colours of a dyed fabric may bleed in contact with liquids; or a particular lining may cause trouble by staining the material to which it is attached. In all cases the selected

1 Moss, A. J. E., Stain Removal, 1950 (Iliffe & Sons, Ltd.).
reagent must be tested on an inconspicuous part of the textile to see whether it is safe to use before applying it to the stain.

The success of the operation often depends on how the reagent is applied. The general tendency is to use too much solvent and so risk spreading the stain. The smallest quantity of the reagent should be applied on a small screwed-up tuft of cotton wool, and with a dextrous touch the stain should be lifted off rather than rubbed in. Several applications may be necessary, but if a stain is very stubborn, it is better to leave it rather than risk damaging the fabric.

In certain cases a different technique is required, known as ‘ringing the stain’. It is obvious that the application of any organic solvent to, for example, a grease spot, would simply result in the grease being spread over a larger area of the textile. The procedure adopted here is to stretch the textile, stained side downwards, over a glass plate covered with blotting-paper and apply the reagent to the back in drops from a small pipette so that it forms a ring around the stain. In this way the solvent bears in upon the stain from all sides at once, the grease is dissolved without spreading, and the resultant solution is absorbed in the blotting-paper.

An alternative method for removing grease or wax stains is to cover both sides of the stain with clean blotting-paper and apply a warm iron. This is the traditional method for removing ‘candle grease’; the grease melts and most of it runs into the warm paper. Any residue is then eliminated with benzene, turpentine, or trichloroethylene applied by ringing as described above. A blob of hardened paint or a blob of grease or mud should be carefully removed as far as possible with a scalpel or razor blade before the solvent is applied.

In using solvents for the removal of stains it must be borne in mind that animal fibres are very sensitive to hot water, which causes them to shrink and lose their lustre, to excessive rubbing, and to the use of those bleaching agents which depend for their effect on the evolution of chlorine. Vegetable fibres are generally more robust, but after bleaching or using acidic reagents it is essential to wash the treated part of the textile until it is neutral. These observations are summarized below in Table I.
### Table I

<table>
<thead>
<tr>
<th>Reagents</th>
<th>Animal fibres (silk and wool)</th>
<th>Vegetable fibres (cotton and linen)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Strong acids or alkalis</td>
<td>Never used</td>
<td>Never used</td>
</tr>
<tr>
<td>Dilute acids</td>
<td>Never nitric acid</td>
<td>Harmful</td>
</tr>
<tr>
<td>Dilute alkalis</td>
<td>Permitted</td>
<td>Permitted</td>
</tr>
<tr>
<td>Bleach containing chlorine</td>
<td>Never used</td>
<td>Use weak only</td>
</tr>
<tr>
<td>Hydrogen peroxide</td>
<td>Permitted</td>
<td>Use weak only</td>
</tr>
<tr>
<td>Very hot water</td>
<td>Never used</td>
<td>Permitted</td>
</tr>
</tbody>
</table>

A list of specific reagents that are commonly employed for removing stains is given in Table II which has been compiled with the object of systematizing procedure. The nature of the textile and of the stain being known, reagents are applied to the stain one after the other in the order given: thus, for example, the symbol GLA for the removal of mud stains on silk is to be interpreted as follows—hydrogen peroxide (G) is applied to the mud and, if no appreciable change is seen to take place in the course of a minute or so, very dilute ammonia (L) is applied. The stain should lessen in intensity and the mud be released so that it can be sponged off with hot water (A), taking care that it is not hot enough to destroy the lustre of the material.¹

### Table II

**Procedure for the Removal of Stains**

**List of Reagents**

- **A** Soft water (Hot)
- **B** Methylated spirits
- **C** Spirit soap (B.P.C.)
- **D** Trichlorethylene
- **E** White spirit
- **F** Hydrochloric acid (2 per cent.)
- **G** Hydrogen peroxide (10 vols.) made alkaline with ammonia
- **H** Potassium permanganate (1 per cent.)
- **I** Sodium hydrosulphite (5 per cent.) made alkaline with ammonia
- **J** Oxalic acid (1 per cent.)
- **K** Acetic acid (0.5 per cent.)
- **L** Ammonia (0.1 per cent.)
- **M** Sulphurous acid
- **N** Pyridine
- **O** Morpholine
- **P** Hydrofluoric acid* (1 vol. + 3 vols. water)
- **Q** Chloramine-T (fresh, 2 per cent.)
- **R** Tartaric acid (fresh, 5 per cent.)

* Rubber gloves should be used.

¹ For more extensive treatment see Holden, J. T., and Fowler, J. N., *The Technology of Washing* (British Launderers’ Research Assoc.) and Moss, A. J. E. *op. cit.*
<table>
<thead>
<tr>
<th>Nature of stain</th>
<th>Wool</th>
<th>Silk</th>
<th>Cotton or linen</th>
</tr>
</thead>
<tbody>
<tr>
<td>1. Mud</td>
<td>GLFA</td>
<td>GLA</td>
<td>GLA</td>
</tr>
<tr>
<td>2. Grease</td>
<td>D or E</td>
<td>D or E</td>
<td>D or E</td>
</tr>
<tr>
<td>3. Iron rust</td>
<td>PA or JKA</td>
<td>JKA</td>
<td>PA or JKA</td>
</tr>
<tr>
<td>4. Red ink</td>
<td>ABL</td>
<td>GJF</td>
<td>Q or LB</td>
</tr>
<tr>
<td>5. Blue-black ink</td>
<td>GFKA</td>
<td>GKFA</td>
<td>Q or HMA</td>
</tr>
<tr>
<td>6. Copying ink</td>
<td>BA</td>
<td>BA</td>
<td>BAI</td>
</tr>
<tr>
<td>7. Marking ink</td>
<td>CD</td>
<td>LG</td>
<td>AI</td>
</tr>
<tr>
<td>8. Fresh oil-paint</td>
<td>BCE or JA</td>
<td>BC</td>
<td>JA</td>
</tr>
<tr>
<td>10. Lipstick</td>
<td>GLKA or RA</td>
<td>GLKA or KA</td>
<td>KA</td>
</tr>
</tbody>
</table>

**THE TREATMENT OF FRAIL TEXTILES**

Textiles are liable to be presented for conservation in all stages of deterioration, but perhaps the most difficult problem arises in the case of material that is freshly excavated, possibly from a tomb. The textiles may be hard and brittle or so decayed as almost to resemble a spider’s web, or they may be damp and the pattern and colours concealed beneath a layer of dust and insect remains. But no matter what condition they are in, provided they have not completely disintegrated, they will always repay careful examination, and it is surprising what may be done in the way of preservation.

The first stage of treatment is always superficial cleaning, so as to enable the textile to be examined. Loose dust may be removed by bellows, but on no account must the material be brushed. Insect remains often adhere firmly and these have to be removed with scalpel or forceps working under a binocular microscope. If a textile has been damp, it may gradually develop a white bloom as it dries owing to the crystallization of salts, and this should be held in check by spraying occasionally with a fine mist of distilled water—not enough to soak the fabric and make it limp, but merely to prevent

---

1 When fatty acids are present the textile may be difficult to wet. If there is any difficulty in wetting the textile with the water spray, a little wetting-agent (Lissapol) should be added.
the surface from becoming white and so obscuring the structure. When extraneous matter has been removed as far as possible, reverse the textile and deal with the back in the same manner.

It is nearly always possible to wash frail textiles provided they are manipulated on a support and handled as little as possible while wet. As dirt and dust have a deleterious effect, washing is always desirable, but where there are decorative attachments, e.g. metal threads or leaf, leather, &c., that might be damaged by water, washing must be avoided. In such cases the textile should be photographed after preliminary cleaning, and by using an infra-red plate it may be possible to record details of ornament that are either concealed or that are difficult to interpret by ordinary photography (Pl. 13 A and B).

Sometimes textiles are rigid and crumpled or folded. When in this condition they should be laid on a glass plate or Polythene sheet and the wrinkles and folds sprayed so that they gradually become relaxed. If the material can be washed, no attempt should be made at this stage to stretch the fabric out. The material is floated into shape after it has been immersed in water.

Details of the washing process have already been described. A modification in the case of frail textiles is to use a porous support instead of the Polythene sheet, so that water with dirt and sandy material can drain through. Alternatively, the material may be sandwiched between two sheets of Terylene net fixed in a frame so that handling is reduced to a minimum. After washing, the Terylene is pressed between warm dry towelling to remove excess moisture, and when partly dry the frail material is transferred to a sheet of Polythene and pinned out as described on p. 96.

The greatest advances in the technique of handling ancient textiles have been made in the Textile Museum, Washington, D.C., and are fully described and illustrated in Workshop Notes published half-yearly by this institution. In order that the routine of cleaning may be visualized, a short summary of the operations that apply in the

1 Mrs. F. S. Greene (vide infra) recommends Lumite, a stiff plastic material having an open cloth weave. An English equivalent is Lumarith manufactured by British Celanese Ltd., Hanover Square, London, W.1.
case of woollen textiles is appended. For full details the reader is referred to *Workshop Notes No. 1.*

1. Textile analysis and scrutiny.
2. Test the chosen detergent on colours and fix them if necessary.
3. Support the textile between open-work screens.
4. Degrease in Stoddard Solvent (a mineral spirit) and allow to dry overnight.
5. Digest in a warm concentrated solution of Parazyme (a starch and protein digestive enzyme mixture), to render stains soluble, and flush with water at the same temperature.
6. Bathe in water fortified with detergent. (*Igepal-CA Extra is recommended, one part in two hundred parts of distilled water.*) Tamp, if necessary, through the open-work screen to remove residual earthy matter, and finally rinse—the last rinse in 10 per cent. glycerine.
7. Dry at first through the screen, latterly, if practicable, with a paper handkerchief placed next the textile.
8. Transfer to a plastic backing that will hold small blocking pins, and straighten the warps and wefts.
9. Mount the dry cleaned textile on a thin cellulose acetate sheet by occasional stitches of thread of the colour of the warps, inserted from the back between two warps, and passing over only one warp so that the stitches are eventually concealed by the wefts.
10. Frame the mounted textile passe-partout between two sheets of \( \frac{1}{16} \) in. Plexiglass (Perspex).

* Note: Woollens shrink and harden if exposed to changes of temperature while wet.

In comparative experiments conducted at the British Museum Laboratory with Miss Louisa Bellinger, Curator-Analyst of the Textile Museum, Washington, D.C., it was decided that Lissapol-N (I.C.I.) gave results comparable with those obtained in using the American preparation Igepal-CA Extra. Solvent Naphtha or white spirit may be used in place of Stoddard Solvent, and Diastase instead of Parazyme.

---

2 Cutilin Ltd., 17 Victoria Street, London, S.W.1. The manufacturers recommend
REPAIRING AND MOUNTING

Even when in a fragmentary condition, textiles must never be stuck down with adhesives. These harden and damage fabrics, they are also nutrients for moulds, and when a material has once been pasted down the reverse side is not available for study. Fragments may be preserved in plastic envelopes, but in order to prevent damage by handling they should be tacked to a sheet of cellulose acetate before being placed in the envelope. If textiles are too frail or brittle to be sewn, they can be exhibited by placing them on a sloping surface in a dustproof exhibition case; the surface should be faced with rough silk or velvet with the pile facing upwards so that the textile fragments adhere in position.

It is often desirable to drape tissues for exhibition, and when these are frail or tattered they must be given some form of support so that they can bear their own weight. It was formerly the practice to use a silk net for this purpose, but silk loses its strength in time, and it is subject to attack by insects and micro-organisms, whereas materials are available today that are not only stronger than silk but likely to be more permanent. A notable example is Terylene which is unshrinkable and rotproof, and so strong that a very fine quality can be used in repair work. Unfortunately there is some difficulty at the present time in applying dye-stuffs to the finished Terylene (though it may be dyed during manufacture) and this limits its use for the repair of coloured textiles, such as flags and banners. The traditional method of mounting old flags is to sandwich them between two pieces of silk net of wide mesh, and when the sewing is finished the threads of the net are tinted with water-colour to match the colours of the flag. The flag is then displayed with the pole in a horizontal position so that the strain on the textiles will be, as far as possible, equally distributed. Terylene would be a more suitable material for that Diastase should be used in 0.2 per cent. solution in water containing 0.3 per cent. of common salt. Stained textiles are soaked for about an hour in lukewarm water (without wetting-agent) and then placed in the Diastase-salt solution at pH 6.8–7.1 (see Appendix V) until the starch or protein is rendered soluble, when the treatment may be completed by rinsing and drying or washing with a detergent as in 6 (above). See also Appendix I.3.
this purpose than silk if the difficulty of tinting the threads could be overcome. But this is not the final solution of the problem, and the method is likely to be superseded. It would seem to be unwise to expose a valuable flag where it may be attacked by moth and harbour dirt; the most satisfactory solution seems to be to have a copy of the original made and hung, and to exhibit the original under glass nearby, as has been done so successfully with the relics of the Black Prince in Canterbury Cathedral.1

As a general principle, modern synthetic fibres should be used for the repair of old textiles whenever possible, as this makes it easy at any time to distinguish between what is old and what is new.

Terylene net was used in repairing and strengthening the fifteenth-century Pyx Cloth (Pl. 15), a frail textile of drawn-thread work which was required not only to support its own weight, but that of four wooden balls, one at each corner of the square. The balls were tasselled and covered with gesso and gold leaf. A Terylene net reinforcement was sewn on the reverse side around the central hole where the strain was greatest, and some reinforcement was also done at the corners; since both the textile and the Terylene were of a pale ivory colour the repair work was invisible at a distance of a few feet. All the sewing repair work was done with Terylene thread.

GOLD EMBROIDERY

Gold threads have been found to vary considerably in structure and quality. Sometimes they were made by winding a high-carat gold around silk thread (as in the case of the Durham vestments) and it was the custom to flatten such threads by hammering in order to enrich the appearance and convey the impression of a continuous sheet of burnished metal. At other times the gold alloy was much debased and became so tarnished that it will not respond to cleaning. Gold was sometimes wound on hair or on parchment, and even gilt paper has been used as a winding over a thread core. Gold threads

1 Restored by The Tower of London Armouries with the co-operation of The Royal School of Needlework. The original jupon has been strengthened with Terylene net, dyed to match the old material by The Fibres Development Dept. of I.C.I. Ltd.
should always be examined closely before any attempt is made to clean them as the poorer qualities would not withstand mechanical strain of any kind, and would be ruined by contact with water.

An unusual form of cleaning and mounting was carried out in the case of the gold-embroidered stole, maniple, and girdle, dating from the tenth century, that form part of the St. Cuthbert relics at Durham Cathedral.\(^1\) The work of cleaning involved removal of a hard buckram backing to which the frail old embroideries had been pasted at some time in an effort to preserve them. The buckram had shrunk, and caused the embroideries to cockle, and they were in a very brittle condition possibly due to excess of adhesive or glue from the buckram. The first step was to reinforce the embroidery by facing it with Japanese mulberry tissue. For this purpose it was necessary to choose an adhesive that would not be softened by water and that could easily be removed at the conclusion of operations without affecting the dye-stuffs. Polyvinyl acetate (dissolved in a mixed solvent consisting of toluene and acetone 95:5) satisfied these requirements, and by using this adhesive the thin tissue paper was attached as intimately as possible and the surface fixed securely. After the adhesive had dried, the embroideries were laid face down on glass, and relaxed under sheets of wet blotting-paper until the paste had softened to such a degree that the buckram could be detached. Residual lumps of softened paste that were adhering among the embroidery threads had to be picked out with forceps, and the back of the embroidery was then cleaned with saponin froth. What remained of the original lining was left in situ. The textiles were dried between sheets of warm blotting-paper and flattened under glass plates.

The next stage was to release the mulberry paper by the application of the mixed solvent, and it was found on removal to bear an imprint of the gold embroidery in dust.\(^2\) To clean the gold still further, fresh mulberry paper was applied with polyvinyl acetate, and after one or two applications the richness of the gold threads was

---

2. The dust contained silver sulphide. This arose from tarnishing of the silver constituent of the gold alloy that had been used in making the thread.
restored to a remarkable degree. The silk threads, however, were still very dirty. These were cleaned by several applications of saponin froth, and by this means it was possible to recover the natural sheen of the silk and to reveal colours that had been concealed by the grime of ages. The embroideries were then sterilized in the thymol chamber preparatory to mounting.

The Durham embroideries are composite in structure; in the main length of the stole and maniple there is embroidery on the face only, but in each case there are terminal panels with embroidery back and front and all the sections are edged with gold braid. For the purpose of mounting, the gold braids were removed and the back and front of the terminal panels separated. As the original foundation material of the embroidery had perished it had to be strengthened with a silk gauze backing. A fine silk gauze was stretched in an embroidery frame; the main section of the maniple was laid on this face upwards, and brown silk retaining threads were passed across at intervals. The four sections of the terminal panels were sewn down in the same way, separately, and the silk gauze reinforcement was cut all round each section, close to the embroidery. The mounting of the maniple was carried out as follows: a stronger and more open silk gauze was stretched in the embroidery frame and the main section of the maniple and the two front terminal panels were sewn on this in their correct relative positions. The frame was inverted, and a maroon silk lining similar to the original was sewn on the gauze along the back of the maniple, and the back sections of the terminal panels added. After this the edging braids were replaced (see Pl. 14 A and B).

The stronger gauze with the maniple in position was then stretched tightly in a thin slip frame and this was inserted, for exhibition purposes, in the channelling of a heavy wooden frame glazed on both sides.

This manner of mounting was devised by the late Mrs. Guy Antrobus. The reinforcement of silk gauze gives adequate support to the frail needlework, and the textiles are sealed from the air and out of contact with the glass. The Durham embroideries are the oldest and finest extant and it would appear that everything possible
has been done to ensure their preservation. Had the work been done today the only improvement that might have been introduced would have been to use Terylene gauze and thread in place of the silk.

ACTION OF LIGHT ON TEXTILES

No chapter on the conservation of textiles would be complete without reference to the damage that is done by exposure to light in causing tissues to become weakened and dye-stuffs to fade.

All natural fibres, animal and vegetable alike, gradually lose their strength on exposure to sunlight. This is illustrated in the following table published by Miss Elizabeth Stromberg,¹ which, even in the absence of experimental details, gives some idea of the rate of decay and the relative resistance of the various types of fibre:

<table>
<thead>
<tr>
<th>Sunlight Time to reduce Tearing Strength 50 per cent.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Silk, cultivated</td>
</tr>
<tr>
<td>Jute</td>
</tr>
<tr>
<td>Cotton</td>
</tr>
<tr>
<td>Flax</td>
</tr>
<tr>
<td>Wool</td>
</tr>
<tr>
<td>Wool, chrome-dyed</td>
</tr>
</tbody>
</table>

Actually, the rate of deterioration varies with the intensity of the light and the temperature and humidity of the atmosphere, but it is, above all, the invisible ultra-violet constituent of sunlight that is the prime cause of the weakening of textile fibres. In the case of dyed textiles, it is well known that certain dye-stuffs have a protective action in slowing down tendering while others accelerate this weakening of the tissue on exposure to light.

In the fading of dyed textiles the factors mentioned above are all of importance but the nature of the complex formed between the dye-stuff and the substrate must also be taken into account. When exposed to light, fading will occur if this complex is capable of

absorbing radiant energy which brings about a molecular change. It should be noted that the spectral absorption of the complex in the dyed textile need not necessarily be the same as it is for the dye-stuff in isolation. These considerations apply with equal force to any form of lighting, natural or artificial, but owing to the relatively low intensity of artificial light sources, the rate of change of colour may be so reduced that it can only be assessed after a prolonged period of time. Miss Stromberg has drawn attention to the cumulative effect upon museum objects of exposure to weak daylight and to artificial lighting, pointing out that, in certain cases, the result of long-term exposure may be just as serious in causing fading as exposure to sunlight for a correspondingly shorter time. However, in making comparisons of fading, it must be remembered that the relative humidity has been shown to have an important bearing on the fastness of dye-stuffs to light.

A detailed examination of the tendering and fading of textiles when exposed to narrow isolated bands of radiation in the extreme visible and near ultra-violet is being carried out at the present time under the auspices of the International Institute for the Conservation of Museum Objects. In a preliminary examination of tungsten and fluorescent sources, filtered and unfiltered, comparative tests have given an assurance\(^1\) that dye-stuffs suffer no abnormal fading when exposed to fluorescent lighting. For the internal lighting of cases, however, fluorescent tubes are to be preferred to tungsten lighting as they are cooler in operation, but it is essential that auxiliary equipment (chokes, &c.) should be mounted outside the case where, incidentally, it will be readily accessible for servicing.\(^2\)

While there is little to choose between tungsten and fluorescent lighting from the point of view of tendering and fading, the intensity of the light source is an important factor. Some suggestions are now discussed to indicate how textile collections may be protected from the deteriorating action of light, both natural and artificial.


\(^2\) For practical advice on museum lighting see *Use of Fluorescent Lighting in Museums* (1953), I.C.O.M., UNESCO House, 19, Avenue Kléber, Paris (16e).
When the museum or gallery is closed, collections should be kept in darkness unless all materials in storage. Permanent exhibition of valuable textiles is inadvisable unless special precautions are taken to reduce the period of exposure to light as much as possible; temporary exhibitions are to be preferred. Sunlight should be prevented from falling directly upon valuable textiles by the use of opaque blinds or curtains. The intensity of lighting, natural or artificial, should not be excessive. Roof lighting should be controlled by louvres or adjustable blinds, or it may be desirable in some cases to cut down the intensity of the lighting by colour-washing the glass in summer. Where there are vertical windows the lighting may be diffused by introducing blinds of the Venetian type, having a system of translucent opal plastic slats in place of the traditional wooden members; these are light in appearance and easy to adjust and clean. In certain cases protection may be afforded by fitting windows with special glass\(^1\) that has the property of excluding the rays that are most harmful. Such glasses cut out a portion of the spectrum and so must restrict the overall lighting to some extent, and they are necessarily slightly tinted, but they are of value for roof lighting and for lights in churches where, for example, it is desirable to prevent strong light from falling upon tapestries or altar cloths. Special glasses materially reduce the tendency to fading, but it should be added that no glass is available that will altogether prevent the fading of dyestuffs.

**ACTION OF SULPHUR DIOXIDE**

Sulphur dioxide is an impurity in the atmosphere resulting from the combustion of fuel, and it readily forms sulphurous acid which in turn is oxidized to sulphuric acid. Mention has already been made of its action in causing the deterioration of organic materials such as leather and paper, and it is the most potent cause of the tendering of

\(^1\) Pilkington Bros. Ltd., St. Helens, Lancs., make several such glasses. The most appropriate for window glazing is called Antisun, which at a thickness of 3/16 inch has a light transmission of 75–80 per cent., greatest in the middle of the spectrum where vision is best. This glass absorbs both heat and ultra-violet rays.
15. PYX CLOTH FROM HESSETT CHURCH, SUFFOLK. (LATE 15TH OR EARLY 16TH CENT.)

Repaired and strengthened with Terylene net
textiles. Cotton seems to be particularly prone to decay from the action of sulphur dioxide. Textile collections in cities or near industrial undertakings are likely to be best conserved when protected under glass. When the small concentration of sulphur dioxide within a case has been absorbed, no further contamination will take place until the case is opened and a fresh supply of polluted air is admitted.

Iron catalyses the change of sulphur dioxide to sulphuric acid, and it is a common observation that where iron nails or tacks have been used to fix textiles, the material has rotted more intensively where it is next to the iron, as, for example, around the tacks used to attach a canvas painting to its stretcher. For this purpose copper tacks should be used. But ordinary dust usually contains an appreciable quantity of iron in one form or another, and this is an additional reason for adopting some form of protection for materials that are exhibited in galleries in which the air is not fully conditioned.

FUNGOID AND INSECT ATTACK

In common with other organic materials, textiles are susceptible to attacks by moulds, but if the textiles are clean and conditions are reasonably dry, there is not likely to be any trouble in this connexion. In the event of mould growth being discovered, it can usually be arrested by airing the textiles, and any surface growths can be removed with a soft brush. If the outbreak is extensive, the textiles should be sterilized with thymol vapour (p. 56) but sterilization may be omitted if the textiles can be washed by an immersion process. If textiles are damaged by water, as might arise from a burst water-pipe or leaking roof, fungoid attack is likely to ensue, and the most effective first-aid measure is to expose them in a current of warm dry air. For emergencies of this kind the most suitable equipment is the electric hair-drier. The use of a heater without ventilation must be avoided as it would be likely to encourage and intensify mould growths.

Insect pests are a more frequent cause of damage to textiles than fungoid attack. Animal fibres, wool and silk, provide foodstuffs for
several varieties of insects, and dry conditions are no serious deterrent to their activities. Linens and cottons are normally immune from attack, and they should be segregated from the wools and silks to facilitate inspection of the more vulnerable materials. But textiles of all kinds are liable to be damaged if they should happen to be contaminated with substances that are attractive to the pests, and it is for this reason, as much as for any other, that garments are washed or cleaned before incorporation in the collection. Textiles should be inspected at regular intervals according to a routine, involving unfolding, dusting, airing, and sunning, and specimens should be scrutinized to discover any traces of insect attack—holes in the material, presence of grubs, cocoons, or loose silken threads.

There are three general ways of protecting textiles from insects. The first is to isolate the textile—to parcel it up so that insects cannot get at it. The second is to use insecticides, and the third is to use an inhibitor that is either so distasteful to the insect or so toxic that protection is assured. This last method involves the use of a proofing reagent that can be permanently fixed on the textile fibre. This necessitates the use of special apparatus and is a technique that can only be applied to new material in the course of manufacture.

Isolation of the textile. This is only a practicable proposition in the case of individual specimens, and, even then, there is always the possibility of infection being introduced in the parcelling and of activity proceeding unrecognized. It is advisable to include a quantity of volatile insecticide in the parcel. For wrapping textiles, paper is as effective as transparent plastic, but the latter is stronger, and has the added advantage that the contents are more readily inspected.

When Polythene bags are used, it is most important that the textiles are quite dry, as moisture cannot escape through Polythene, and if the textiles are damp when packed, they are almost certain to become mildewed. The proofed bags sold for the storage of furs are also satisfactory for textiles, but it is as well to put about a cupful of dichlorobenzene crystals in the bottom of the bag to ensure against the risk of moth eggs being present. Protection by sandwiching material passe-partout between glasses provides an alternative method
of isolation, and insecticides are normally not required here as the textile is exposed continuously to view.

Use of insecticides. Apart from vacuum fumigation that can be applied on a large scale to protect costumes or ethnographical material in bulk (see Wood, p. 123), the most effective protection is given by the presence of a volatile insecticide such as dichlorobenzene, which, though perhaps a trifle too volatile, is otherwise satisfactory and efficient. It can be filled into cotton bags and hung in wardrobes, or it may be scattered in drawers and chests between layers of paper. This substance is very effective against the chief insect pest, the clothes moth, and is also effective against dermestid beetle. It has, however, a rather overpowering smell which causes headaches, and as it is a liver poison it is important to avoid working for long over an open case in which this chemical is used.

For materials that cannot be protected in this way the alternative is routine spraying using, preferably, a power spray with a fish-tail nozzle, and working at a pressure to give fine atomization. The gun is held at an angle of about 45° to the textile, so that a cloud of insecticide floats on to it, rather than a direct blast which might be damaging. Many different insecticides are available for this purpose, the most satisfactory being those made up in a colourless and odourless petroleum distillate. Pyrethrum extracts have an immediate action upon moths, DDT is slower to take effect, and insecticides of a mixed type containing both ingredients are probably best (see p. 29). Very little of the spray material condenses on the textile, and there is no evidence that the regular use of the spray results in any staining or change in colour. Insecticidal dusts are liable to contaminate the material unnecessarily and are not recommended.

1 In Moth-proofing from the Entomological Point of View, Vezelinstituut T.N.O., Delft, 1951, Dr. A. D. J. Meeuse has shown that even fumigation with hydrogen cyanide is no guarantee against survival of moth eggs, and that camphor, naphthalene, and dichlorobenzene act on moths as respiratory poisons when used in high concentration, but have no repellent action in low concentration, though there are indications that dichlorobenzene is more effective than the other two. In 'Insect Pests of Books and Paper', Archives, 1952, 7, p. 19, A. W. McKenny Hughes recommends using dichlorobenzene crystals in an air-tight cupboard at a concentration of 1 lb. to 10 cub. ft. of cupboard space.
CHAPTER V

WOOD

Wood has been exploited by man since palaeolithic times. Man has done almost everything with wood—made his home of it, made fires with it to cook his meals, used it for utensils, weapons, and implements, built ships and bridges of it, used it for making vehicles, furniture, art objects, and musical instruments, and, in modern times, transformed it into paper and even clothing. It is not surprising, therefore, that wood forms a large part of the collections in museums—particularly in ethnographical and folk museums—and its conservation is a matter of considerable importance.

Being of organic origin, wood normally decays under combined biological and chemical attack when buried in the ground, but in exceptional circumstances it has been found to survive prolonged exposure to extremes of dryness or wetness. Worked timbers exist today in the dry tombs of Egypt that date back to the early dynasties, and it is somewhat of a surprise to find that these are still sound and often so fresh in appearance that they might be taken for timbers of modern origin. Timber that has been buried in wet peat bogs for a long period of time may retain its general shape and size. The absence of air inhibits fungoid attack, but profound changes may have taken place in chemical composition and microstructure, resulting in a loss of strength. Such timber can be dried, however, and its original appearance restored by laboratory treatment. In sharp contrast, wood completely decays when buried in damp sand that is well aerated. In this case decay is due in the main to fungoid attack. The excavation of the seventh-century ship burial at Sutton Hoo yielded many notable treasures (now in the British Museum) but the ship itself had succumbed, leaving only rusty iron nails and a stained impression in the sand to mark where the wooden planks had rested.
WOOD

No such extreme changes are to be expected in wood kept indoors. Under such circumstances the commonest form of deterioration, in temperate climates at all events, is due to the dimensional changes accompanying variations in the relative humidity of the atmosphere. These result in warping and splitting, features that are related to the hygroscopic nature of the material, and to the fact that it has a grained structure.

DIRECTIONAL PROPERTIES OF WOOD

In order to study the conservation of wooden objects it is necessary to be familiar with certain characteristics of the raw material. Wood is an anisotropic substance, exhibiting different degrees of hardness, toughness, &c., in different directions. It has an organized cellular structure, and as the fibres are for the most part orientated in the same direction, grain is a distinctive feature. This varies in appearance according to the species of tree and the cut of the timber.\(^1\) In a longitudinal median plank, two main zones are to be distinguished, namely heart-wood and sap-wood, the latter being situated towards the edges adjacent to the bark. In practice the direction of the cut is not necessarily chosen as being the most economical nor as giving the strongest timber, though both these considerations may apply; it is often selected in order to give some effect of texture or grain for ornamental purposes.

SEASONING OF WOOD

The sap-wood of a tree has a higher moisture content than the heart-wood, and when the timber is sawn into planks, the distribution of moisture in the planks varies accordingly. If left to dry out naturally, the moist parts will shrink more than the drier parts and the planks will warp. It is for this reason that they have to be submitted to a process of controlled drying known as seasoning. This consists in allowing the timber to dry out slowly under conditions

\(^1\) In *Museum*, 1955, 8, 3, p. 139, the relationship between cut and grain is illustrated in an admirable series of drawings.
in which distortion is prevented. But wood never loses all its moisture; the cell moisture\(^1\) remains, and, when seasoning has been completed, the moisture content of timber is found to vary with the relative humidity of the atmosphere surrounding it. The equilibrium state of air-dry timber that has been seasoned under the most favourable conditions varies in different regions. Desch\(^2\) quotes a moisture content of 12–15 per cent. as typical of timber seasoned in the United Kingdom and most regions of the United States, whereas in the more humid tropics, e.g. Malaya, 14–18 per cent. is characteristic, and in hot arid regions the moisture content may be as low as 8–12 per cent.

In the early stages of seasoning sap-wood is particularly vulnerable to attack by fungi and insects, not only because of its higher moisture content, but because it contains a greater concentration of nutrients. Kiln-seasoning is general today instead of air-drying, and as it speeds up the process there is less chance of infection; it has the additional advantage that the residual moisture content of the wood can be adjusted to suit the particular purpose for which the wood is required and the climate in which it is to be used.

**WARPING OF WOOD**

The most important factor in preserving seasoned timber is to avoid exposing it to wide variations in atmospheric humidity. Under such conditions it alternately absorbs and gives up moisture. While this causes little dimensional change along the grain, it causes expansion and contraction across the grain with consequent warping. If one side of a plank is protected from the atmosphere, the other side will absorb or give up moisture more quickly than the protected side and warping will be intensified. Humid conditions will cause expansion across the grain on the exposed side, so that the protected side becomes concave. Conversely, where conditions are very dry the protected side will become convex. This explains the ‘cupping’

\(^1\) i.e. the moisture contained within the cells as distinct from the moisture held between the cells.

that takes place in the constituent planks of a wooden panel painted only on one side, when exposed in an atmosphere that is too dry. Varying relative humidity tends to cause back and forward movements which are a significant factor in the life of panel paintings (p. 4). If a panel is restrained in a frame by glue or nails, changing humidity may cause the wood to crack or become distorted.

The elimination of warping is a lengthy and uncertain process, although at times it has to be attempted in the interests of conservation. The operation may take several weeks or even months. The treatment consists in moistening the concave side so that water is slowly absorbed; this swells the tissue and encourages the wood to return to the flat condition, and it is then kept flat under weights until dry. But even if this operation appears at first to be successful the tendency will be for the wood to return gradually to the warped condition unless some form of cradling can be used to confine the stresses to one plane—see pp. 161–2.

A simple form of metal reinforcement that is often quite effective and may be applied where the wood is of reasonable thickness is to screw a series of angle irons to the back of the panel at right angles to the grain by countersunk screws passing through slots (not holes) in the metal. The edges of the slots should be slightly chamfered. This arrangement admits of lateral movements of the wood and prevents the panel from cupping. The metal and screws must be of such a nature as not to loose freedom of movement by corrosion or otherwise.

When, as in certain types of furniture, the wood is painted all over or French-polished, all the surfaces are equally protected and movement of the constituent timbers is reduced to a minimum. In the case of synthetic grainless boards that have no directional properties, or plywood having an adequate number of layers, a balance of forces is achieved structurally, and warping is prevented. Large thin boards may buckle under their own weight, but this is purely a mechanical matter which can be dealt with by arranging for suitable supports.

1 When wood is old and has been in a warped condition for a long time, a modification of cell structure may have occurred which precludes the possibility of a successful restoration. Wood in this condition is said to have acquired a 'permanent set'.
ATTACK BY FUNGI

A certain degree of dampness is required before fungi can grow on materials supplying the necessary nutrients. This has already been referred to in the case of paper and leather, and it is equally true of wood. Structural timber is most prone to attack because it may become wet owing to the inadequate provision of damp-proof courses, and large wooden objects, such as canoes, totem poles, and carvings exposed out of doors, are much more likely to be attacked by microfungi than objects kept indoors. The fungi attacking such timbers may be of the virulent 'dry rot' group that is so difficult to eradicate without wholesale sacrifice of the infected material. Preventive measures take the form of isolating the wood, if possible, by the introduction of a damp-proof course of slate, lead, &c., and by protecting any end-wood by impregnating it with a waterproofing agent such as linseed oil, wax, or a water-repellent silicone varnish. When timbers are exposed in the open, advantage should be taken of any possibility of deflecting water by dripcourses or courses. When wood is in contact with the ground, it should be impregnated with a fungicide, and any part under ground level encased in waterproof cement.

The following fungicides are colourless and odourless and will not stain wood, brickwork, or plaster.

(A) 3–6 oz. of commercial sodium fluoride dissolved in one gallon of water and applied cold by brush (two coats), or

(B) Half a pound of commercial acid magnesium fluoride dissolved in one gallon of water.

For details of these processes and of other methods of dealing with an outbreak of dry rot, the reader is advised to consult the publications of the Forest Products Research Laboratory. No satisfactory account of methods of conserving timber would be possible without frequent reference to the valuable publications of this laboratory,

1 Midland Silicones, Ltd., 19 Upper Brook Street, London, W.1.
2 A wooden tub must be used in making this solution as it attacks metal.
which are based upon wide experience and research, and cover all aspects of the subject.

When moulds are growing on art objects in the museum, ethnographical specimens, &c., it will usually be found that they are feeding upon oil, grease, or the binding medium of gesso, rather than on the wood itself, and growth may be arrested by cleaning and applying fungicide. But the treatment should not stop here. It is equally important to take measures to prevent a recurrence of the outbreak by improving ventilation.

**ATTACK BY INSECTS**

Insect pests of timber are a greater menace to museum objects than moulds, because the possibility of infection is ever present. The nature of the damage will depend on the type of infestation, which varies in different parts of the world. The commonest pests in Great Britain are various types of wood-boring beetle, and the damage is done by their grubs which tunnel into the wood, converting sound tissue into powder. The grubs are popularly known as ‘wood-worm’.

If the attack is not checked at an early stage, the wood may fall to pieces and may be found to be riddled with insect channels filled with powder and frass resembling fine sawdust. It is usually the presence of powder of a light colour that provides the first indication of attack; if structural timber is inaccessible, the first indication may be that a piece of timber gives way and falls to the ground, as happened in the case of a wooden rib from the roof of the North Transept of York Minster in 1934. When this occurs the attack will have already proceeded to great lengths, and it can only be dealt with by an expensive operation involving wholesale impregnation of the old sound wood with insecticides, and by making extensive replacements of the infected wood.

Furniture should be inspected regularly to ensure that it is free from ‘worm’, paying special attention to plywood back-boards and also to any sap-wood that may be present. It is not uncommon to find the first evidence of a wood-beetle infection in the ply-backing
of a framed picture, or in the plywood that has been used to repair the back of a chest or a bureau. When discovered, infection should be dealt with at once, either by burning the plywood or by treating it with a reliable insecticide (see p. 125).

The discovery of worm holes in wood is not necessarily an indication that insects are still active. Sometimes they die off before all the wood has been destroyed, but it is never an easy matter to be certain that activity has ceased. Clean wood dust falling from an old hole may suggest activity which, in fact, does not exist; but if an object that has been lying undisturbed with accumulations of dirty wood powder is seen to have some clean wood dust upon it, this will be a clear indication that insects are still alive in the wood. In all cases where there is any doubt, it is safer to assume the worst and the object should be lifted carefully into the open air, brushed clean, and treated with an insecticide.

Before it can be taken for granted that sterilization has been completely effective and that no viable eggs remain, the wood must be kept under regular observation during at least one complete life-cycle of the insect, but as insects develop at different rates this involves identification. For example, the common powder-post beetle (Lyctus) breeds through from egg to adult in three months to a year, whereas in the case of the death-watch beetle (Xestobium)—the major pest of inaccessible structural timbers—development is much slower, taking possibly two years or more to complete. The life cycle of the common furniture beetle (Anobium) may also run to two years or more. Wood-boring insects may be identified by the size of the holes they leave on emerging to lay eggs, or by the materials attacked, or by the nature and extent of the outbreak, age of the timber, &c., but the most certain method is a microscopic study of the grub or full-grown insect and comparison with photomicrographs and drawings in the standard works of reference.¹

As an aid to long-term inspection and sterilization all surface

¹ Fisher, R. C., ‘Household Timber Insects, Recognition, Prevention, and Control of Infestation’, The Sanitarian, November 1949; see also the leaflets of the Forest Products Research Laboratory devoted to specific pests.
powder and the frass should be removed and the old holes filled with soft wax (see p. 125) so that any fresh ones will be at once apparent. In adding to a museum collection, the greatest care must be taken to exclude material that might introduce infection until such time as it has been sterilized. When wood is found to be attacked, treatment should be carried out as soon as possible; the longer the delay, the more difficult it will be to bring the infestation under control.

METHODS OF STERILIZATION

Insects may be killed by rise of temperature, by reduction of pressure (vacuum treatment), and by poisoning with gas (fumigation), or liquids (spraying; impregnation). It may be stated in general that fumigation methods are immediately effective but confer no lasting protection, whereas impregnation methods may be slower to take effect, but the poison remains and confers protection over a period.

Sterilization by fumigation. Sterilization may be carried out in a sealed gas chamber, and with suitable equipment it is possible to reduce the pressure to a partial vacuum before admitting the gas, which ensures that the fumigant will penetrate in a reasonable time into the objects under treatment.

Hydrogen cyanide gas is used in such a chamber at the British Museum (Pl. 16) for dealing with ethnographical material in bulk, mainly textiles, skins, feathers, and small wooden objects, for which a twenty-four hours' exposure has been found adequate. With denser materials such as heavy timber, methyl bromide would be likely to give better penetration, but methyl bromide has the disadvantage that, in dealing with ethnographical specimens which may contain feathers, leather, &c., or with upholstery containing horse hair, it tends to form compounds having a very unpleasant odour. This rules it out for museum work where objects in storage have to be frequently examined. When large objects have to be sterilized by fumigation and no special plant exists for the purpose, it may be possible to arrange for this to be carried out professionally by a firm specializing in such work.¹

¹ London Fumigation Co., Marlowe House , Lloyd's Avenue, E.C. 3.
Carbon disulphide is a reliable insecticide that can be used for the fumigation of valuable objects of all kinds as it does not harm the most delicate material. It is a clear liquid at normal temperatures, having a low vapour pressure (B.P. 46° C.) and an unpleasant odour. As it evaporates readily to form explosive mixtures with air, rigid precautions must be taken to exclude naked lights and prevent smoking in any room in which this fumigant is used. Some museums are equipped with special vacuum plant for fumigating with carbon disulphide vapour. The special plant is enclosed and therefore safe and assures good penetration. It is possible to fumigate with this insecticide at normal temperature and pressure with improvised equipment. This type of fumigation is conducted as follows. Dishes of the liquid are exposed above worm-infested wood in an air-tight cupboard or box which is then sealed up, the doors being pasted round with adhesive paper or Polythene tape. A Polythene bag may be used instead of a cupboard provided it can be adequately sealed with tape. Liquid carbon disulphide should be used in the proportion of 1 ounce to about 8 cub. ft. of air space in the chamber. The liquid volatilizes in the course of a few hours, and the heavy vapour pours down over the wood and tends to accumulate near the base of the cupboard. The time of exposure required for fumigation is from 14 days to three weeks, and a fresh amount of liquid carbon disulphide should be added at the end of the first week to take the place of that which has evaporated. After fumigation the unpleasant smell disappears on exposing the wood to air for a short time.

It will be obvious that fumigation by carbon disulphide is particularly effective for sterilizing boards that can be treated in a horizontal position in a shallow box, and as this fumigant is without action on paint or varnish the method is very useful for the treatment of worm-eaten panel paintings.

Liquid carbon disulphide should be stored in a cool place and protected from daylight as this causes the liquid to develop acidity.

For the sterilization of wooden objects that are not decorated with

1 Carbon disulphide is a dangerous chemical and fumigations involving its use are best left in the hands of trained personnel.
fugitive colours, a non-inflammable solution of one part of carbon disulphide in four parts of carbon tetrachloride may be recommended. While this is not quite so toxic to insects as pure carbon disulphide, equally satisfactory results are obtainable either by using a greater concentration or extending the period of exposure.

Sterilization by impregnation. Impregnation with liquid insecticides may be carried out by using a pipette or a syringe to inject the liquid into the holes. Alternatively, the insecticide may be brushed in. Large undecorated timbers may require to be drilled in inconspicuous parts to enable the insecticide to penetrate.

Suitable liquid insecticides are those containing DDT, BHC (gammexane), pentachlorophenol and its derivatives, chloronaphthalenes, or the metallic naphthenates. Many reliable mixtures are available commercially. If applied thoroughly to all insect holes, the impregnation method may be as effective as fumigation, but before using an insecticide it should be tested to make sure that it is non-staining and that it does not soften paint or damage any decorations on the wood with which it might come in contact.

For filling the holes after treatment, the best material is usually a soft wax to which an insecticide has been added. A suitable preparation for filling the holes can be made by stirring DDT powder into molten beeswax, bleached or appropriately coloured to match the timber. When solidified, the wax paste may easily be spread into the worm holes with a blunt knife. If the wood requires strengthening (see below) this ought to be attended to prior to waxing.

Sterilization by Spraying. While spraying is not very effective for killing pests that have already bored their way into wood, there is evidence that spraying with a 2 per cent. aqueous emulsion of DDT can prevent the spread of an outbreak of Lyctus.²

Proofing of wood against insect pests. Proofing is essential in the case of timber that may be exposed to attack by termites (‘white ants’) — the most injurious wood-feeding insects known. Creosote conform-

---

² Forest Products Research Laboratory, Leaflet No. 43, 1948.
ing to a standard specification\(^1\) is used for this purpose but to be entirely effective the wood should be impregnated under pressure, and the creosote retained by the timber in certain minimum amounts. Metallic naphthenates are also used for pressure impregnation. For the protection of museum objects of wood such as furniture that cannot be impregnated under pressure, the only alternative is to saturate the wood as completely as possible by dipping or brushing, using a non-staining insecticide that will penetrate deeply into the timber. Protection is limited in proportion to the depth of penetration of the preservative. Such an insecticidal mixture may be prepared by dissolving pentachlorophenol in a petroleum distillate. This leaves the wood clean after treatment, but it is emphasized (F.P.R. Leaflet 38) that a certain minimum period of immersion is necessary if this treatment is to be entirely effective. For thin wood where glue is absent a hot 4 per cent. aqueous solution of zinc chloride or sodium fluoride may be used, the object being dipped into the solution; but with aqueous solutions penetration is less effective and the degree of protection usually less satisfactory than that obtainable by the use of petroleum solutions or chloronaphthenates.

**STRENGTHENING TIMBER**

Wood that has been weakened by the attack of fungi or insect pests may be strengthened by impregnation with a consolidating agent or it may be reinforced mechanically. The nature of the specimen and the condition of the timber will determine which of the two methods is likely to be the more satisfactory, but in many cases it will be found that both mechanical strengthening and consolidation are required in order to restore solidity. When this is the case impregnation may sometimes precede reinforcement, but sometimes it will be more appropriately applied after the wood has been mechanically strengthened.

\(^1\) British Standard Specification Number 144. See also Forest Products Research Laboratory Leaflet No. 38 which gives an account of the habits and habitat of termites. References are given in this leaflet to the problem as it presents itself in Australia, India, Malaya, West Indies, and U.S.A., as well as in Great Britain.
WOOD

Consolidation by mechanical reinforcement

Of the mechanical methods for strengthening timber, the follow-
ing are the most important.\footnote{This matter, in so far as it concerns painted panels, is given detailed attention in \textit{Museum}, 1955, 8, 3, pp. 139 seq.}

1. Dowelling either with metal or wooden pegs, and refacing if necessary with wood.

2. Inlaying across cracks with solid \(X\)-shaped wedges to prevent the cracks from opening; or covering the joints by ‘buttons’ of wood, i.e. small palettes \textit{ca.} 3 in. by 2 in. glued across the cracks.

3. Reinforcing with wooden splints, glued and/or screwed to the old wood. Special bracket irons, angle irons, \&c., may be useful in repairing furniture.

4. Stopping irregular cavities with a gap-filling cement (see repairs, p. 132).

Consolidation by impregnation

Methods of consolidation by impregnation may be applied to all porous materials and are particularly suitable for dealing with intricate shapes, carvings that have suffered erosion by exposure, or wood that is riddled with worm holes. Impregnation is carried out by immersion, injection, or the solutions may be applied with a brush.

Where the surface is covered with loose \textit{gesso} and gold leaf it can only be satisfactorily consolidated by impregnation with a substance that penetrates easily, and synthetic consolidating agents dissolved in organic solvents are recommended in this case. Painted \textit{gesso} is in the same category, but the paint must be tested to ensure that it will not be softened or dissolved by the solvent. If there is any sign of its being attacked it may be possible to change the solvent for one less active, or waxing may be the preferable technique.

\textit{Impregnation by wax}. When mechanical repairs are necessary they are always carried out before the wood is waxed. For many purposes the wax-bath is a most effective form of treatment, but for large objects special equipment is required. The bath may be a capacious,
rectangular iron tank, heated electrically and thermostatically controlled so that the temperature of the wax is not allowed to rise above predetermined limits. There should also be an independent safety device to switch off the electric supply in case of failure on the part of the thermostat. Block and tackle will be required for raising and lowering heavy objects, and with this equipment it is possible to deal conveniently with life-sized wooden sculptures.

The composition of the wax mixture may be varied but the main ingredient is usually unbleached beeswax to which has been added a resin in quantities not exceeding 50 per cent.

The object to be impregnated, which must of course be quite dry, is lowered into the molten wax and kept beneath the surface by weights or otherwise. As the temperature gradually rises the air is expelled in a stream of bubbles and wax enters the pores of the wood. Any traces of moisture are driven off by maintaining the temperature at about 105° C. till bubbling ceases, and it may then be allowed to rise to about 120° C. The time of immersion depends on the porosity and bulk of the timber. When the impregnation is considered to be complete the heating is shut off, the object raised from the tank, and the hot wax allowed to drain away. Finally, surplus wax is removed from the surface with turpentine.

Waxing by immersion has been developed on a large scale in Baltimore by Rosen. The process is also applied with much success in the Central Laboratory of the Belgian Museums, Brussels, as a routine method of treating church furniture, architectural material, wooden carvings, and the like (Pl. 17).

For the impregnation of small objects the wax immersion process can be carried out using improvised equipment, but a note of warning is necessary about a potential source of danger when the tank is heated from below. If moisture has been introduced it will accumulate


2 Private communication from Dr. Paul Coremans, Director of the Central Laboratory of the Belgian Museums, Brussels.
17. VAT FOR IMPREGNATING LARGE WOODEN FIGURES WITH MOLTEN WAX (LABORATOIRE CENTRAL DES MUSÉES DE BELGIQUE)

A polychrome oak figure being lowered into a vat of hot wax
18. SPECIMENS OF WATERLOGGED WOOD AFTER DRYING

Adjacent sections of ash originally of the same diameter. *Left*: Untreated, showing considerable shrinkage. *Right*: Pre-treated to prevent shrinkage on drying.
at the bottom of the tank as the wax solidifies; on reheating, steam pressure may be generated beneath the semi-solid wax and may cause an explosion. This danger can be mitigated by placing a metal rod upright in the wax as it cools; the rod is withdrawn and most of the water decanted away before the wax is reheated. It must also be remembered that the molten wax mixture is inflammable, and that serious burns can be caused by splashing. When wax is being melted over an open flame, an asbestos board should be at hand to place over the vessel in the event of the wax catching fire.

Wax-resin mixtures are inert and very stable, and being waterproof they afford protection against damp. They are thus of value in preventing movements in wood due to changes in the relative humidity of the atmosphere. Waxing has certain disadvantages, however, unless the objects are kept in cool surroundings. Even if the surface of a waxed object has been cleaned after immersion in the bath so that only the thinnest film of wax remains, there is a tendency for wax to creep out gradually from the inside, and in a matter of time the surface will become sticky and tend to collect dirt. In addition, if a waxed object is exposed to the sun or exhibited in a glass case that is lit internally, the specimen may develop an objectionable gloss with rise of temperature. A further disadvantage of wax mixtures is the fact that they have a high refractive index and lower the tone of colours. The optical qualities of light colours are best conserved by using materials having a lower refractive index than wax, and in certain cases colourless synthetic resins may be preferred.

Impregnation by synthetic materials. Thin varnishes are used as impregnating agents to consolidate wood either before or after mechanical repairs have been carried out. By making a test with coloured ink it can be shown that wood absorbs liquids much more quickly in the direction along the grain, and this fact is borne in mind when impregnation is carried out by brush or by injection. The thin liquid is fed into the end-wood so that it permeates the timber as far as possible, and the work is completed by coating the surfaces; on volatilization of the solvent the resin remains to consolidate the tissue. The following colourless
synthetic resin varnishes are suitable consolidants for wood—polyvinyl acetate dissolved in a mixed solvent of 9 vols. of toluene and 1 of acetone, or Bedacryl 122 X\(^1\) diluted with toluene to a suitable consistency. Other materials that have become available recently for the impregnation of dry wood are the polyester resins such as Marco S.B. 26C\(^2\) or Bakelite 17449.\(^3\) These are supplied as mobile liquids which set in the pores of the wood to hard insoluble solids, as a result of a catalysed chemical reaction. One of these resins was used with success by Dr. G. A. Horridge for the consolidation of the delignified wood of Cody's Tree at the Royal Aircraft Establishment, Farnborough.\(^4\) An object that has been tunneled by wood-worm may be strengthened by running the solution into the worm holes with a fountain-pen filler or drawn-out glass tube as is done in impregnating with insecticides.

In order to strengthen dense timber the impregnation should be conducted in an apparatus from which the air has been withdrawn. For this purpose a metal vacuum tank is used. If the tank is much larger that the object, a vessel of smaller dimensions is used as the container in order to economize in the use of solvent, and this vessel is placed inside the larger tank which is then evacuated. When the pressure has been reduced to the required degree, as recorded by the pressure gauge, the impregnating solution is admitted gradually through a tube to the smaller vessel until the object is completely covered with the solution; it is allowed to remain there for about an hour, so that the solution is absorbed as far as possible by the wood. Air is then admitted and this drives the consolidant farther into the wood. When the tank reaches atmospheric pressure again, the object is lifted out of the container and allowed to drain; it can be dried without acquiring a gloss by leaving it in an atmosphere of the solvent. This process, which is decidedly the most effective method

---

\(^1\) Imperial Chemical Industries Ltd. Supplied as a 40 per cent. solution in xylene which is too viscous for most purposes and requires dilution.


\(^3\) Messrs. Bakelite Co. Ltd., 12 Grosvenor Gardens, S.W. 1.

\(^4\) Royal Aircraft Establishment, Farnborough. Technical Note 1236, 1954.
of strengthening wood, is of general application to all kinds of porous material whether of organic or inorganic origin.

Apparatus for vacuum impregnation can be set up on a small scale using ordinary laboratory equipment. All that is required is a vacuum desiccator provided with a two-way tap, one outlet of which is sealed until such time as it is used to admit the solution, and the other is connected via a trap bottle to a suction pump operated from the main water-supply. Alternative arrangements are shown in Fig. 5, p. 152.

REPAIRS AND MAINTENANCE

The first requirement for wood repairs is a good adhesive that is easy to apply and which will make a strong joint. Calcium caseinate is a powerful adhesive for wood; it was used in the Middle Ages for furniture-making and for joining planks together to make into broad panels as supports for painting. A good-quality hide glue is more convenient in practice, and can yield joints that are strong enough for most purposes. The tendency nowadays, however, is to choose one of the modern synthetic resin adhesives. These set by chemical action when mixed with a hardener to give very strong joints. Four main types\(^1\) are recognized, namely:

- Urea Formaldehyde (U.F.) adhesives.
- Phenol Formaldehyde (P.F.) adhesives.
- Resorcinol Formaldehyde (R.F.) adhesives.
- Epoxy resin adhesives.

The first three tend to shrink on setting but the epoxy resins are characterized by the fact that they set without undergoing any appreciable contraction in volume.

The application varies according to the manner in which the hardener is mixed with the resin before use.

\(^1\) These are marketed under a variety of trade names by different firms. Details may be obtained from the following: British Industrial Plastics Ltd., 1 Argyll St., London, W. 1; Aero Research Ltd., Hinxton, Duxford, Cambridge; Leicester Lovell & Co. Ltd., St. Christopher’s Works, North Badgesley, Southampton.
(1) The hardener is mixed with the resin in the stated proportions before the adhesive is applied to the joint.

(2) The hardener is applied to one side of the joint, the resin applied to the other side, and the two held together till the adhesive sets, or

(3) the adhesive is supplied as a dry powder consisting of resin plus hardener to which it is only necessary to add water before use.

Detailed instructions are supplied by the manufacturers in each case. In making a choice of adhesive it may be of significance that the R.F. hardener is neutral, whereas the others are acidic or alkaline.

If, in addition to repairing a joint with an adhesive, it is necessary to fill irregular cavities in such a manner that the filler will itself confer strength, a modern gap-filling adhesive must be used, one of the most satisfactory being Aerolite 300, a U.F. resin. If it is merely a question of filling up a hole, choice may be made from a large selection of materials. The following are available—ordinary putty (whiting and linseed oil); Alabastine (a fine hard-setting plaster of Paris); and a viscous celluloid syrup in which has been incorporated a porous filler such as sawdust or pumice powder; if a black filler is not out of place, Chatterton’s Compound may be used, applied with a hot knife. Glue and sawdust may sometimes be all that is required, and awkward holes may be filled in with shaped balsa wood inserted with an adhesive. In pegged repairs of this nature, extra wood must always be allowed and the excess cut or filed away after the adhesive has finally set.

Veneers and inlays tend to suffer from desiccation of the glue which causes them to lift; in this condition they are easily damaged with a duster and they should be temporarily secured with adhesive tape. This is a case where early attention is amply repaid. The process of laying back damaged veneer is a matter for the specialist—amateur repairs are seldom very permanent or satisfactory—and for the job to

---


² Celluloid fillers must be applied in thin layers allowing time for the material to set between each application, otherwise shrinkage may be considerable.
be done properly, it requires the use of a flexible glue\(^1\) and the appropriate irons, the veneer being flattened and maintained under pressure until the adhesive has hardened.

Marquetry can be kept from drying up by rubbing occasionally with a little non-drying oil. Olive oil answers well for the purpose. Almond oil is used for cleaning the wooden parts of violins and other musical instruments that are decorated with inlays. Wooden furniture having a good surface, whether French-polished or wax-polished, should be kept in condition by rubbing at intervals with a soft cloth. The original polish is always worth preserving whether applied to veneer or solid timber and it improves with time. Furniture polish coats the surface with a thin film of wax. This is a good thing, but wax should not be allowed to accumulate in cracks or crevasses as it will detract from the appearance and collect dirt. For cleaning polished furniture, of whatever kind, a suitable emulsion can be made by shaking together, vigorously, half a pint each of linseed oil, turpentine, and vinegar, to which is added a small teaspoonful of methylated spirit. This mixture is comparatively inexpensive; it removes dirt and polishes in the same operation, and is harmless if applied in moderation.

**BASKETRY**

Problems in preserving basketry are essentially those caused by mould growths and insect attack, and treatment is conducted on the same lines as in the case of wood similarly affected. Basketry tends to become brittle, however, and very dusty. It may be cleaned by Lissapol froth, dried, and then impregnated with bleached beeswax dissolved in benzene. Celluloid in acetone forms a satisfactory adhesive for repair work.

Bark-cloth garments disintegrate owing to the rotting of the stitching and they are then unable to sustain their own weight. Restoration is carried out by couching the bark to a Terylene lining, and an improvement in appearance results from spraying with a

\(^1\) Animal glue can be rendered flexible by the addition of starch derivatives, e.g. sorbitol.
dilute methacrylate solution or with polyvinyl acetate spray solution (p. 178).

WATERLOGGED WOOD

When wood is exposed for long periods in water or in wet clay or peat, it undergoes a form of decomposition arising from degradation of the lignocelluloses of which it is composed; much of the finer cellulose tissue disappears, but the thicker lignin structures remain and these preserve the shape of the wood and often the general surface appearance. It is not uncommon to come across wooden objects from lake dwellings or from wells that appear to have survived for generations with little disfigurement, dark in colour but preserving their shape and ornament in minute detail. The loss of the finer cellulose tissue does not cause much alteration in the gross volume of the wood, but the porosity is much increased and the wood is able to absorb water like a sponge. When filled with water it is very heavy, and in this waterlogged condition it may be unable to bear its own weight so that if carelessly handled it may fracture across the grain. The preservation of waterlogged wood is one of the major problems for the archaeologist. Objects in this condition must be lifted on a rigid support or splint and packed at once in wet moss, cotton-wool, newspapers, textiles—anything that will keep them thoroughly wet—so that they can reach the laboratory in the condition in which they were found. They must then be transferred immediately to a bath of water, when they can be cleaned and examined at leisure, measured, and photographed. So long as the objects are kept wet they will keep their shape, but if allowed to dry without being specially treated, the weakened cellular tissue that is held apart only by water will collapse and the specimens will then be damaged beyond recovery.

It goes without saying that waterlogged wood can be preserved in water (containing preferably a disinfectant such as 2 per cent. w/v of carbolic acid), but it is much more interesting if it can be dried without becoming deformed in the process. Two general methods are available for accomplishing this. The first is to replace the water
in the pores of the wood with a liquid that will solidify there and make contraction impossible as the wood dries, and the second method is to fix the tissue so rigidly that it will not collapse after the water is removed. The alum process is an example of the first method, and the alcohol-ether-resin process an example of the second.

1. The alum process

The alum process depends on the fact that potash alum is a crystalline substance freely soluble in hot water, but only soluble to about 10 per cent. in cold. If a hot concentrated solution can be made to permeate the tissue by soaking the wood in the liquid for sufficient time, the mass will solidify as soon as the specimen is removed from the hot solution and no shrinkage will be possible. In practice, glycerine is added to the alum solution to facilitate impregnation and it may help also to preserve the colour of the wood.

The composition of the alum bath is important. Alum is dissolved in hot water in such proportions as to give the strongest concentration consistent with reasonable fluidity at a temperature near the boiling-point of water. The solution is prepared in a copper or an iron vessel—not galvanized iron as the alum would dissolve the zinc. From two to three parts by weight of potash alum are dissolved in one part by weight of boiling water to which has been added one part by weight of glycerine. The wood is submerged in this solution, using lead or iron weights if necessary to keep it below the surface, and the bath is maintained at 92–96° C. for at least ten hours, a longer time being allowed when the timber is bulky or has a close grain. Rosenberg\(^1\) recommends up to thirty hours—from which it will be seen that the treatment of heavy timber may take a long time. The level of the liquid should be maintained during heating operations by adding hot water as required.

When it is considered that impregnation is complete, the wood should be lifted out, washed quickly with warm water to remove superfluous alum from the surface, and set aside to cool. The residual white deposit which invariably remains is brushed off when dry, or removed

with a cloth wrung out in hot water or preferably in hot glue size, which, with the alum, will form an insoluble skin. After a few days, when quite dry, the surface is sealed as described on p. 143.

There is evidence that the best results are obtainable with hardwoods rather than softwoods and that success depends on the use of a sufficiently concentrated hot solution of alum, and on securing adequate penetration. An example of the satisfactory application of the process to thin slats of yew is described in detail by Moss¹ in the preservation of an Anglo-Saxon wooden bucket.

2. The alcohol-ether-resin process

The process of drying tissues in successive baths of ethyl alcohol to remove water and following this by immersion in successive baths of ether to remove alcohol is a well-known biological technique. If the final ether bath contains a resinous material in solution, the resin will remain in the wood on the evaporation of the ether, and the wood structure may thus be fixed rigidly.

This has been found to be a very successful method of dealing with small objects. Werner Kramer has obtained excellent results by the alcohol-ether process, impregnating the wood finally under vacuum with a concentrated solution of dammar resin in ether (Pl. 18). His success appears the more striking in that he bleaches the wood as a preliminary by immersing it in a 5 per cent. aqueous solution of hydrogen peroxide for a week before commencing the alcohol treatment, and by this means the black slimy surface characteristic of waterlogged timber is converted to a pale slate or biscuit colour. The natural appearance of the wood may thus be restored and the grain revealed.² Sometimes, as in the case of birch poles, it is even possible to recover traces of the silvery bark, the presence of which was unsuspected prior to treatment. Stone axe handles from a prehistoric site in Switzerland responded most successfully to the method and emerged as if they had been newly fashioned.

² The method was developed at the laboratory of the Swiss National Museum, Zurich, under the guidance of Professor Emil Vogt.
In recent work at Copenhagen, Christensen\(^1\) has found that it is quite possible to dry waterlogged wood if the water is first replaced by alcohol, and this, in turn, by a liquid of low capillarity such as ether, which is then dried out very quickly by sudden exposure in a vacuum. This is done by having the object in a closed vessel connected by a stop-cock to a larger vessel that has been previously evacuated: the opening of the stop-cock reduces the pressure in the smaller vessel very quickly, and causes rapid volatilization of the ether.

Such vacuum-drying prevents shrinkage, but in thin woods may cause some warping. If a little shrinkage is permissible, the ether may be allowed to evaporate in the ordinary way without employing a vacuum technique and there is then no warping.

The scheme advanced by Christensen is briefly as follows:

1st Bath. Alcohol—in volume equal to five times the bulk of the wood.
2nd Bath. Repetition of No. 1.
3rd Bath. Fresh dry ether—Duration 2 days.
4th Bath. Repetition of No. 3.

Alcohol of at least 95 per cent. strength is used and great care is taken to ensure that the time of soaking will be adequate to remove all water. For timbers such as oak and ash which are close-grained the time may be calculated at one day for every centimetre length of wood, but it is emphasized that the soaking in alcohol cannot be overdone.

In normal cases the ether may be evaporated very quickly as described above, but when the object is thin and easily warped, the safer procedure is to allow the ether to evaporate naturally. Objects are then strengthened by impregnation in a 3 per cent. solution of polyvinyl acetate in pure benzene. The object is dipped in this solution for a moment only, and when dry, it is coated with a 10 per cent. solution of dammar resin in petroleum ether (B.P. 80–100° C.).

\(^1\) Christensen, B. B., *Om Konservering af Mosefundne Trægenstande*, 1952 (National Museum, Copenhagen).
This must be applied sparingly. It is unwise to attempt to cut short the process by going directly from the alcohol bath to impregnation with polyvinyl acetate, so dispensing with the ether, as this leads to unsatisfactory results.

Both the Kramer and Christensen processes have been tried in the British Museum Laboratory and found satisfactory. The hydrogen peroxide method of pre-bleaching had been used for a number of years to reveal wood grain but always in a restrained fashion. More recently, following a demonstration at Zurich, the bleaching was carried out in the laboratory to the limit and the specimen, a Roman scoop of hard wood, was restored to what must have been its original appearance. In this case the final bath was a strong solution of dammar resin in ether, applied under vacuum. It was found that a piece of softwood treated in the same fashion could not be bleached beyond a pale slate colour, and it seemed to become more porous than the hardwood, but even in the softer wood the fineness of detail was well preserved: chisel marks survived and actually a series of fine saw-marks, no more than 1 millimetre apart. This bears striking testimony to the delicacy of the method.

While alcohol and ether are expensive and involve a serious fire risk, they are certainly to be preferred for the treatment of small objects, and there seems no doubt that this treatment coupled with hydrogen peroxide bleaching would be justified in all cases where it is essential to preserve fine detail of workmanship.

RESTORATION OF THE STANWICK SCABBARD

An unusual type of restoration was successfully accomplished in the case of the iron sword discovered in its wooden scabbard by Sir Mortimer Wheeler in the excavations at Stanwick, Yorkshire, in 1952. This was a unique object dating to the first century A.D. (Pl. 19). As it was waterlogged, it was dispatched by the first train to London packed in a mass of wet newspapers, and it was this prompt action that made its preservation possible. Reference is made later (p. 282) to the corrosion of the iron sword by sulphate-reducing
bacteria and to the preservation of the weapon. The following
details relate to the preservation of the wooden scabbard.

The scabbard was made from two slats of ash of a maximum thick-
ness of \( \frac{3}{8} \) inch and tapering towards the edges. These were held to-
ether by a series of bronze rings of symmetrical eye-shape set at
intervals, and decreasing in size from the mouth-piece to the chape.
As the sword could not be withdrawn from the scabbard, in which
it was firmly held by its iron sulphide corrosion products, the alum
method could not be used for preserving the wood. This acidic sub-
stance in reacting with the sulphide might have caused disintegration
beyond control. It was decided, therefore, to remove the scabbard
from the sword by dissection under water. The varying width of the
bronze fittings afforded the possibility of their removal from the
chape end; this proved to be a delicate but not a difficult task, and
the bronze mouth-piece was finally released from its own end.

After the removal of the bronze fittings the wood was detached
with the aid of scalpels from one side of the sword-blade and floated
on to a glass plate cut to the length of the scabbard. The blade was
then reversed and the remaining wood removed and floated on to a
second glass plate.

The specimens were then treated by immersion in a series of
alcohols of increasing strength in order to eliminate the water. The
treatment was carried out in boxes made specially from tin-plate.
The alcohol in the wood was eventually replaced by amyl acetate,
the final immersion being carried out in a solution of celluloid dis-
solved in equal volumes of amyl acetate and acetone to a strength of
about 5 per cent. Throughout these operations, the wood remained
on the glass-plate supports and was not handled until it had been
strengthened. This work took about fourteen days, and on the
evaporation of the solvent, the thin porous wood was found to be
toughened quite satisfactorily.

For the reassembly of the scabbard it was necessary to prepare
a core of such dimensions that, with the old wood in position, the
bronze rings would slip on from the pointed end, and the recon-
structed scabbard would conform to original dimensions. But here a
difficulty arose due to minor irregularities in the shape of the metal fittings. This necessitated the use of a resilient core which was made as follows: a piece of wood was cut slightly thinner and narrower than the sword, and to this was fixed along its length a piece of thin steel cut from a flexible steel rule, the attachment being made with a bandage soaked in shellac. This core was then impregnated with a solution of nitrocellulose and the old wood of the scabbard was attached to it on both sides with Durofix, the contact being made as intimately as possible. The rings could now be slipped on from the pointed end, there being some slight elasticity in the core in consequence of the action of the spring in its interior. When the adhesive hardened the core became rigid and some minor repairs with a mixture of celluloid and sawdust completed the restoration.

It was, of course, never intended to return the sword to its scabbard; the two were subsequently mounted side by side on a wooden baseboard to form an exhibit, interesting in itself and convenient for purposes of study.

HEAVY TIMBERS. DUG-OUT CANOES, ETC.

The preservation of heavy waterlogged timbers, such as dug-out canoes, presents a special problem because of the difficulty of transport and the expense of the large quantity of materials required for treatment.

The wood is cleared of mud and drawn out of the water or moist ground on an improvised stretcher or sledge preferably of sheet metal, and it is set in such a position that it can be washed down, measured, and photographed. The usual procedure is then to cover it under a good heap of earth and to leave it for at least a year but preferably longer so that the water will drain gradually from the timber. When this takes place slowly the heaviest oaken timbers do not warp, although the surface may, to some extent, tend to open up. In the case of a dug-out canoe, the sides, which are much thinner than the base, will tend to warp inwards and these must be supported before covering with earth by inserting wooden cross pieces padded with sacking, and lashing them in position. These should be readjusted
when the earth is finally removed and before the vessel starts to undergo acclimatization to atmospheric conditions. The acclimatization process may take an additional year or longer. During this time, the wood must be protected from the weather, particularly from frost and direct sunshine. Towards the end of the acclimatization period, the surface may be sealed by painting it with a solution of sodium silicate of high silica content diluted with distilled water, e.g. P 84 brand of sodium silicate\(^1\) diluted with six times its volume of distilled water. Such treatment has proved to be satisfactory in several cases. Sometimes where the wood has appeared to dry well impregnation is unnecessary, but with large thin timbers the element of chance is considerable and much depends on the grain of the wood and its freedom from knots. All would agree, however, that the slower the process of drying the better the chance of achieving a successful restoration. It is, of course, fatal to bring such an object indoors, at least until it is in equilibrium with conditions out of doors. The proximity of radiators, for example, can undo the work of years expended in caring for the object in the early stages.

The remains of two Iron Age boats were recovered from the River Humber at North Ferriby in 1947. These consisted of keel planks and major portions of the lower timbers of the vessels sewn together with thin yew branches and caulked with moss. The boats were salvaged by lashing them to metal skids at low tide and drawing them up to the river bank out of the mud. As these early relics were of unique interest, they were taken at once to the National Maritime Museum, Greenwich, where they underwent a course of treatment to dry the wood with the minimum of distortion.

Previous investigations by the finders\(^2\) had yielded several old spars, some of which had been placed in glycerine and some allowed to dry naturally, thus providing useful material for comparative study. While the air-dried wood showed little deformation, there was a definite tendency for surface layers to develop a *craquelure*?

---

1 Imperial Chemical Industries Ltd.
which on the thinner portions formed a system of raised flakes. A long-term experiment with a waterlogged spar showed that glycerine could be introduced to replace the water in sufficient amount to slow down the rate of drying, leaving a surface that could eventually be sealed with shellac; by this means the bulk of the water was eliminated without the formation of craquelure.

Galvanized iron tanks of ample size to accommodate the timbers were constructed by bolting smaller units together and these were provided with drain holes and plugs so that the final de-mudding could be carried out in the tanks. The wood was arranged in a single layer supported above the bottom on bricks. After washing and draining thoroughly, the plugs were inserted and the wood was covered with a dilute glycerine solution which was replaced three or four times over a period of two years, the progress of operations being followed by stirring up the solution at intervals and testing the density of the liquid with a hydrometer. When the glycerine liquors corresponded in strength to those from which the test spar had been taken, they were run off and the timber was allowed to dry in air for more than a year. The pieces were then removed and treated individually with a solution of shellac in alcohol. With heavy timbers there must always remain an element of doubt as to when they are dry enough to be considered stable, and while in the present instance there appear to have been no complications that were not immediately overcome, it is recognized that it may take years for equilibrium to be established and it will be necessary to keep the work under observation and to apply more shellac if the wood requires it. In the present instance even the caulkling and yew-stitching survived treatment and all the timbers were decidedly improved in appearance.

The largest water-logged vessels that have been preserved are undoubtedly the Viking ships at Oslo.¹ There are two superb examples,

the Oseberg ship and the Gogstad ship. The third, the Tune ship, which is in a fragmentary condition is still of much interest in showing clearly how such vessels were constructed. These ships, and many of the carved wooden objects that they contained, were preserved mainly by the alum method. Before beginning treatment exact sizes and shapes were recorded by taking plaster impressions while the objects were still wet. The alum-treated wood was finally coated with a mixture consisting of equal volumes of turpentine and linseed oil with the twofold object of improving the appearance and water-proofing the surface.
CHAPTER VI

BONE AND IVORY

From earliest times bone and ivory have exerted a peculiar fascination for the craftsman. Bone was the more easily come by and, as it was available in a variety of shapes and degrees of hardness, selected pieces could be fashioned into all sorts of useful objects—fish-hooks, arrow-heads, tools, and implements. Ivory, though of standard shape, was the more suitable material for fine work and lent itself to decorative treatment. It could be carved, etched, stained, painted, gilded, inlaid with metals and with precious and semi-precious stones; it could be used, moreover, for inlaying in wood, and for veneering. When a material was required for the finest carving, the choice fell on ivory rather than on bone. Thus, ivory was in demand for the plaques and reliefs that were used for diptychs and bookbindings. It was also ideal for carving figurines and statuettes, and specimens are extant that date back to the earliest dynasties of Egypt. A great wealth of ivory-carving has survived from early Christian times in the West, and many fine works in ivory have also come down to us from remote civilisations in China and the Near East. Although bone was more generally used for utilitarian purposes, it was often brought into service for ornamental purposes as well, owing to the relative scarcity of ivory.

When worked and decorated, the two materials may be so much alike that they cannot be distinguished by mere inspection, and identification is then a task for the specialist. It is not possible by chemical means alone to distinguish bone from ivory. In both materials the main inorganic constituents are the same, namely calcium phosphate associated with carbonate and fluoride, and the organic tissue of both is ossein; while the latter varies in quantity, it is usually present to at least some 30 per cent. of the total weight.
A. Sword and waterlogged scabbard cemented together by corrosion products

B. After separation and restoration

19. IRON AGE SWORD AND WOODEN SCABBARD FROM STANWICK
20. DECORATED IVORY PLAQUE FROM NIMRUD. (9TH CENT. B.C.)

Before treatment

See frontispiece for conditions after treatment
Microscopic examination will, however, permit of a differentiation, since bone and ivory both have a cellular structure, and the microstructure of each is definitive. In cross-section, bone shows a rather coarse grain with characteristic lacunae, whereas ivory, being composed of the hard dense tissue known as dentine, is more compact and is characterized by the presence of a network composed of tiny lenticular areas resulting from the intersection of systems of striations that may be seen radiating from the centre of the tusk. In favourable circumstances this can be observed without sectioning by studying the surface with a pocket lens.

The main factors bearing on the conservation of bone and ivory may be deduced from a consideration of their physical structure and chemical composition. Bone and ivory are anisotropic having directional properties and for this reason they are easily warped upon exposure to heat and damp; furthermore, they are decomposed by the prolonged action of water due to hydrolysis of the ossein, and the inorganic framework is readily disintegrated by acids. Being porous and of light colour, bone and ivory are easily stained. They tend to become brittle with age and they lose their natural colouring when exposed to sunlight. When burnt, they become grey or blue-black, and when buried in the ground for prolonged periods of time they are greatly weakened and may be disintegrated either by salt incrustations or by water; in the latter case they are converted ultimately to a sponge-like waterlogged material similar to waterlogged wood. Under other circumstances they may become fossilized. With the onset of fossilization the organic content gradually disappears and the remaining calcareous matter becomes associated with silica in the form of quartz and with mineral salts derived from the ground. When bone and ivory are in good condition it is possible to preserve them by taking very simple precautions, but when they are in a fragmentary state, waterlogged, or fossilized they can only be cleaned, strengthened, and stabilized; a satisfactory restoration may be impossible.

Two general features should be considered before discussing

1 Bone-black and ivory-black are two well-known pigments.
detailed methods of preservation and restoration. These relate to size and colour. Bone and ivory are usually only available in pieces of limited size, so that large objects made from these materials will of necessity be composite in structure. In the case of ivory, for example, when a large object does not conform to the shape of the tusk, the artist may add portions by dowelling or jointing (tongue and groove, half-lap, &c.) and, when any warping of the component pieces occurs, the greatest strain will tend to be imposed on the joints. This is particularly noticeable in the case of books that have covers made of ivory; a clasp or even a pair of clasps may have been fixed to keep the covers shut and the pages flat, but should swelling of the pages occur (and this is by no means unusual if they are of parchment, see p. 47), the strain on the ivory covers may be considerable and the bindings may open along the joints. This type of accident is caused by the cumulative effect of damp which causes the pages to swell, but similar accidents may be the direct result of exposing the ivory to sunlight or to heat. Joints may open, veneers may warp, and if warping is serious, it may be impossible to restore the original shape.

Old ivory often has a yellow colour and this is accepted as a form of natural patination which may help to enhance the appearance. On the other hand, coloration may have been artificially applied, as is often done by the Japanese carver of netsuke, who displays the greatest originality in arriving at a surface effect that will show his handiwork to the best advantage. The greyish colour of burnt ivory has already been mentioned, and in some of the Nimrud ivories there seems reason to suppose that the burning was intentional and was carried out with the object of achieving a particular colour effect. Staining may be due to a residue of some former embellishment executed in a fugitive dye-stuff that has decomposed; in this case the residual colour may show some signs of regularity or of pattern work which may be of interest and value to the specialist as evidence of a decoration that once existed but no longer survives. There also arises the problem of adventitious staining which may be due to a variety of causes. When there is no sign of regularity in the staining its origin may be obscure; it may possibly be due to some photochemical
effect or to contact of the bone or ivory with another material. Rust stains and copper stains can be helpful in the reconstruction of a broken object by showing which pieces originally fitted together, or in calling attention to the previous existence of hinges or metal attachments. In the case of the Franks whalebone casket in the British Museum, staining of this kind was particularly valuable to those engaged in its reconstruction. Finally, there may be disfiguring stains arising through use or accident. Stains may be removed by appropriate chemical treatment, but since there are so many possible causes of coloration in bone and ivory objects, the greatest caution is necessary before undertaking any treatment that might result in the loss of important characteristics; only in exceptional cases can the removal of stains be justified, because although chemical treatment may discharge the stain, it may result in damage to the texture and patina of the ivory, and thus magnify the disfigurement.

METHODS OF CONSERVATION

Removal of surface dirt

Having referred to the damage that can be caused to bone and ivory by exposure to water, it may seem illogical to assert that washing with soap and water can be carried out without danger. This can, however, be done if the objects are in good condition, and if they are dried with soft towelling immediately after treatment, so that the wetting is confined to the surface. The use of water containing a detergent (soap, or one of the new surface-active agents) is indeed essential for the removal of soot and grease, but the object must not be allowed to remain in contact with the aqueous solution longer than is absolutely necessary, and for this reason the washing is carried out with a brush rather than with cotton-wool or a sponge. The hardness of the brush and the length of the bristles is also important; too soft a brush would only prolong the time necessary for cleaning. The brush must therefore be selected with some care to suit the job in hand, taking into account the strength of the object and the amount of grime to be removed.
A more complicated problem arises in the case of partially decayed ivory or bone, especially when the polished surface skin has deteriorated and there is evidence of cracking. In this case the ivory is washed as before, but instead of being dried with towelling it must be passed immediately through two or three changes of 95 per cent. alcohol before being dried by contact with absorbent material such as a paper handkerchief or blotting-paper.

Removal of salts

One of the most difficult problems has been that of cleaning salt-incrusted and semi-fossilized ivories from Egypt. These may be very absorbent and, if placed in water even for a few minutes, they tend to split and to warp, providing an almost impossible task for the repairer. However, the cleaning of such ivories ceased to present any real difficulty when it was discovered that it was not necessary to remove all the salt but only the surface deposits, and that success depended on the speed of operations. In a typical instance a cracked ivory, disfigured by a salty incrustation, was placed in a shallow earthenware dish, and covered with distilled water. This water was discarded and replaced with fresh distilled water at intervals of five seconds, four or five washings being considered to be sufficient. The ivory was then immediately washed in 80 per cent. alcohol for thirty seconds, followed by two washings in 95 per cent. alcohol or industrial methylated spirit for a similar period. Finally, the object was immersed in ether for one minute, and dried in air. By this procedure the entire cleaning was completed within three minutes, the ivory having been exposed to water for a much shorter time, and it survived without any apparent change of shape or extension in the cracking. Such treatment may seem to be too quick and easy to have any permanent effect, but, in fact, it has been applied on many occasions in dealing with salty ivories from Egypt and Palestine. The results have been entirely successful, and the times have been cut down well below those stated where specimens were particularly frail or fragmentary.

Incrustations of carbonate provide yet another complication. In
this case, the use of acid is essential to decompose the carbonate, but
during decomposition the effervescence due to the evolution of
carbon dioxide gas would inevitably result in disintegration of the
ivory. This difficulty is overcome by applying the acid locally only,
_i.e._ no more than a square centimetre is dealt with at a time. Ivories
incrusted with carbonate are cleaned under a binocular microscope
at about 10 diameters magnification, applying a 1 per cent. solution
of hydrochloric acid with a small water-colour brush repeatedly. As
effervescence ceases, the excess of liquid is mopped up either with a
second brush or with shreds of blotting-paper applied with forceps.
The acid must be used in quantities just sufficient to soften the incrus-
tation, which is then removed with a needle. Finally, in order to
eliminate the last traces of acid, the ivory is washed in several changes
of distilled water for a few seconds at a time, and then dried by
the alcohol and ether method already recommended for delicate
material.

Occasionally one comes across an ivory that is incrusted with
calcium sulphate (selenite). This incrustation cannot be removed by
chemical means owing to its comparative insolvability; any attempt
to do so would involve a long soaking process which would cause
the ivory to disintegrate. The only procedure that can be recom-
mended for removing calcium sulphate from ivory is to reduce the
incrustation mechanically by the use of dental equipment—inverted
cones and burrs. Needless to say, such treatment can only be under-
taken if the ivory object is sufficiently robust to withstand the
mechanical stresses involved.

In describing the different methods of cleaning bone and ivory,
emphasis has been laid on the importance of cutting down the time
during which they are exposed to the action of chemicals and even
of water, but in an exceptional case it may be necessary to break
these rules in order to save a fine specimen. Such action is only justi-
fiable when the risks to be taken are fully understood, so that treat-
ment can be suitably modified to reduce these to a minimum, and
when there is a reasonable hope of success. An interesting example is
provided by the restoration of a well-known Egyptian ivory figurine
in the British Museum, once thought to represent Menes of the First Dynasty.\textsuperscript{1} This ivory had been broken in antiquity; the parts were incrusted and held together with calcium carbonate which, in crystallizing, had opened up the structure to form a series of strata interspersed with carbonate, so that the object was both swollen and deformed. In order to restore the object to its original shape it was necessary to clear away all the carbonate from the cracks; therefore the cleaning could not be confined to the surface as is normally the case. Actual soaking in dilute hydrochloric acid followed by washing was necessary. It was foreseen that effervescence during the acid treatment would probably cause the specimen to disintegrate, and this eventuality was prevented by previously impregnating the ivory under vacuum with a solution of celluloid (2 per cent.) and allowing it to dry. The celluloid solution had the effect of slowing down the penetration of the acid, thus holding the effervescence in check. When the calcium carbonate had dissolved, the constituent pieces fell apart. These were washed in distilled water, dried through alcohols to benzene, and then impregnated with a solution of dammar resin dissolved in benzene, using the vacuum technique already described. Reassembly of the fragments, using a viscous solution of dammar resin in benzene as the adhesive, presented no problem and the original shape was recovered. After restoration the figurine was found to be more complete in regard to its ornamentation than had been previously suspected. Only one major gap remained and this was filled in with a mixture of beeswax and carnauba wax and tinted with thin oil-paint to match the biscuit colour of the ivory. It should be noted that when cracks in ivory arise from warping it is not desirable, in general, to fill them in, because under different environmental conditions they may tend to close up, and if this natural tendency for the cracks to close is prevented by the presence of a filler, it may cause the ivory to rupture. However, in the present instance a small portion of ivory was missing, leaving a fissure that caused a structural weakness. This weakness was overcome by the filling which, at the same time, had an aesthetic value in minimizing the disfigurement.

\textsuperscript{1} Figure illustrated in Plenderleith, H. J., \textit{The Preservation of Antiquities}, 1934.
Consolidation

When it is merely a question of consolidating a dry powdery ivory, several suitable transparent media are available, notably polyvinyl acetate and the polymethacrylates. Polyvinyl acetate lacquer was recommended by the present author as long ago as 1934 as a medium for impregnating and strengthening bone and ivory and also in stronger solution as a cement.\(^1\) Todd\(^2\) has described how he consolidated carved ivory spatulas of the Shang dynasty with Gelva (a polyvinyl acetate) by impregnation under vacuum. Similar materials have been used for strengthening bone.\(^3\) The use of an aqueous emulsion of polyvinyl acetate for hardening bone that has been excavated from bogs has been described by Purvis and Martin.\(^4\) Waterlogged bone provides a much easier problem than waterlogged wood, as it can be strengthened directly by vacuum impregnation with dilute polyvinyl acetate emulsion in an apparatus such as that illustrated in Fig. 5. Purvis and Martin state that the individual globules of such an emulsion have a diameter of about 0.01 μ. When the pressure is released, the smallest interstices of the bone are filled with the emulsion and it only remains to drain the bones on a wire frame. The treated material is described as being robust and as having recovered its original colour variations and minute surface architecture, and as being capable of satisfying the requirements of the most exacting osteologist. McBurney\(^5\) speaks very favourably of the excellent qualities of polyvinyl acetate (presumably the emulsion) for hardening bones during excavation both before removal from the ground and immediately after, especially when the bones are very wet and soft. Bedacryl L\(^6\) (a polymethacrylate emulsion) also works well, and this may be readily transported as a concentrate for field work and diluted with water to the required consistency before

---

6 Imperial Chemical Industries Ltd.
use. These emulsions are stable so long as they are protected from frost.

Although nitrocellulose lacquers and adhesives suffer from certain minor defects, these are outweighed by the advantages that they offer for strengthening and repairing bone and ivory—ease of application, reliable adhesive properties, and ease of removal. The latter characteristic is important as it allows adjustments to be made in the course of repair work, and makes it possible at any time in the future to remove the adhesive, should this be desired. By using Durofix, a complicated repair like that of the St. Cuthbert comb, which had split into countless fragments, could be tackled with confidence and with the knowledge that under museum conditions the restoration would be, to all intents and purposes, permanent.

MOULDING OF IVORIES

Particular care must be taken in handling ivories when there are features of interest that must be preserved; for instance the carved ivory plaques of the Christian era that have prayers written on the
back in ink, and others that are coated with wax as a ground for writing with a stylus. There are cases where damage has been caused in the moulder’s shop when making plaster replicas of ivories. Moulding is not necessarily dangerous provided the moulding materials can be used cold. The temptation to use a warm jelly mould or warm rubber latex, which is so convenient for dealing with undercuts, must be resisted, as such treatment may cause cracking or staining. A safer procedure is to piece-mould the ivory or, preferably, to employ a cold jelly mould prepared from alginate; but all materials that require to be used wet necessitate pre-treatment of the ivory to protect it from possible damage by water. This is generally done by giving the object several thin coats of nitrocellulose lacquer diluted with acetone. Such a thin coating need not affect the sharpness of the replicas, and may easily be removed afterwards by swabbing with solvent. Perfect replicas may be obtained by the use of alginate and, when expertly handled, this material imposes no strain on the ivory.

RESTORATION OF AN INLAID PHOENICIAN IVORY

During an excavation at Nimrud in 1952, Professor Mallowan recovered a number of magnificent ivories from the bottom of a well, dating from the period of Assur-naṣir-pal II (883–859 B.C.) and, by drying them slowly and uniformly, he was able to preserve their shape and prevent serious cracking. It may be of interest to describe the subsequent treatment that was applied to one of them—a masterpiece of exquisite carving, embellished with gold and incrusted with lapis lazuli and carnelian (Frontispiece).

When this reached the British Museum Laboratory it was covered with fine clay (Pl. 20), and an X-ray examination showed the presence of deep cracks which were widest at the back, or external side, of the tusk, but, fortunately, scarcely apparent on the decorated side. The clay was carefully scraped from the back and, as this revealed a surface without decoration, it was rigidly secured (as a first-aid precaution to prevent any further opening of the cracks) by backing

1 Alginate Industries Ltd., Dipple by Girvan, Ayrshire.
it with layers of broad adhesive tape, the equivalent of surgical strapping. The front was then dealt with, working under a binocular at a magnification of 10 diameters. First the upper layers of clay were removed with needles to expose the sculpture. Many fragments of gold leaf were recovered from the clay; these were washed with 1 per cent. nitric acid and then with water, and set aside for subsequent replacement. The ivory was further cleaned with pellets of blotting-paper held between pointed forceps and moistened with detergents in order to peptize and release the clay with the minimum of strain, and later the surface was washed and polished by the same technique, doing small areas at a time. The final stage in the restoration was the replacement of the loose fragments of gold leaf in their correct positions on the cleaned ivory, using Durofix as the adhesive.

The background of this superb object—thought to be part of a throne—is decorated with an all-over floral pattern consisting of alternating flowers and seed capsules, the stems and sepals being covered with fine gold. The forms are deeply carved and the thin walls that outline the flowers are also gilt so that they appear like metal cloisons framing the lapis lazuli inlays of the petals. The seeds (or buds?) are represented by polished carnelians of dome shape, serrated at the base to engage with the gilt ivory calyces. This rich background forms a canopy for the main carving below, a scene in high relief depicting a lioness in the act of killing a Nubian. The ivory body of the animal is unadorned, and powerfully modelled. It stands out in sharp contrast to the relaxed human victim with his gleaming golden loin-cloth, and spikelets of crisp curly hair, an effect obtained by fixing gilt-topped pegs into the head which was stained black beforehand. In spite of the loss of much of the gold overlay and the blue and red incrustation, this ivory carving still gives the effect of a faceted polychrome jewel.

The following technical points relating to the construction of the ivory were noted during the cleaning and are of special interest. The lapis lazuli inlays are of varying thickness, but the recesses in

1 Lassoband, Smith and Nephew Ltd. (Technical Tapes Division), Welwyn Garden City, Herts.
the ivory are of uniform depth. Consequently, it was found necessary to bed the inlays in a cement so that their polished surfaces would be level with the cloisons. This is the only case known to the author where a blue cement has been used for inlaying lapis lazuli. It is composed of lime putty coloured to match the lapis lazuli by the addition of copper frit and where inlays are missing the ivory is often stained by a residue of this frit cement. Another interesting observation was made on examining the back of fragments of gold leaf from the ivory. Traces of a brownish film were clearly visible on the gold under the microscope. This material which was of organic origin and swelled in water, becoming very sticky, appeared to be the original adhesive employed in laying the gold leaf. It is difficult to believe that a reversible colloid could have survived in such circumstances but apparently this was the case, and it was due no doubt to the protection from air and moisture afforded by the coherent film of gold.

HORN, ANTLER, ETC.

Many of the methods that have been described for the treatment of bone and ivory apply with slight modification to kindred organic materials that are frail or porous, such as tortoise-shell, horn, and antler. These may be washed in the same way as ivory.

In the case of horn, which is usually thin and hollow, the surface may be strengthened and prevented from flaking by impregnation with polyvinyl acetate and it may be possible to confer added strength by attaching a plastic tape or bandage to the inside surface of the horn. Only part of the interior can be covered, if a horn is intact, but this will be the thinnest part and the part, therefore, that has the greatest need of reinforcement.

Antler usually has a spongy central tissue covered with a continuous outer rind. Under conditions of prolonged burial the spongy tissue generally decays and only the outer rind survives. Thus the antlers of the Sika deer that once crowned a Chinese wooden deity were found to have lost all their spongy tissue; they were hollow and partially collapsed. The surface skin had survived, however, and it
was possible to support this with a wire core bound with cotton-wool and reinforced where necessary with a filling of nitrocellulose and cork dust. The antler picks or levers found in flint workings in the chalk are usually dry, and in this case may be treated by pouring dilute polyvinyl acetate lacquer through the porous tissue. It should be noted that this may leave an unpleasant shine on the outside of the antler unless the surface is swabbed with toluene and the antler allowed to dry in toluene vapour. This can easily be done by suspending it in a tall vessel containing a couple of ounces of the liquid solvent.

Although jet and amber are not chemically related to bone and ivory, their treatment may be referred to here as they are washed according to the methods outlined above for ivory. While fossil amber is insoluble in organic solvents it is not easily distinguished in appearance from other resins that are soluble and for this reason it is advisable to avoid the use of alcohol and ether altogether in cleaning objects of resinous origin.

Finally, a special warning is necessary in dealing with jewellery that may be inlaid with materials such as shell, mother of pearl, and coral (which consist essentially of chalk). These substances are decomposed by acid; where there is any doubt about identification, cleaning-agents of an acidic nature must at all costs be avoided.
CHAPTER VII

EASEL PAINTINGS

Easel paintings are defined as paintings on panel or canvas in tempera or oils, and they present unique problems in conservation because of their variety and complexity of structure. The pictorial image is, however, the important thing—colour and optical quality have to be studied in relation to conservation—and it is for this reason that the care of valuable paintings is regarded as being the most exacting and responsible of all museum curatorial activities. Restoration is a matter for the expert as it involves an appreciation of aesthetic quality as well as a specialized training lying outside the field of ordinary museum work; but the curator must himself be familiar with all aspects of his subject, and the present chapter is designed to provide such technical information as may help him to appreciate the fundamental factors involved in the care of paintings whether in the gallery or in the restoration studio. It deals with the structure of paintings, with the ills to which they are subject, and with methods available for their conservation.

STRUCTURE OF PAINTINGS

An easel painting cannot be regarded merely as a coloured surface. Although painting techniques may have changed through the centuries, the basic fact remains that, from the physical point of view, a painting is a stratified structure built up in a series of layers each of which may exhibit complexities of its own as shown diagrammatically in Pl. 21. There is first of all the support, a prepared panel or canvas; on this is spread the ground, which consists essentially of a white inert substance in a glue medium. On Italian panels the inert substance was usually calcined gypsum (Ital. gesso), but Northern
painters favoured the use of chalk. For the application of gold leaf, a special ground was used in which the inert constituent was a species of clay known as bole. The paint layer overlies the ground and it consists of an aggregate of pigment particles in a binding medium. In tempera painting the medium was egg yolk or the whole egg, but later, drying oils were more commonly used. Finally, there is usually a layer of varnish and this surface coating serves a twofold purpose: it protects the paint layer against atmospheric contamination and also confers upon it an enhanced brilliancy. When a picture is in a sound condition, its various layers are fused and interlocked, the closest study revealing no sign of cleavage.

The character of the painting is mainly influenced by the nature of the binding medium and, as the medium carries the whites and blacks and other pigments, its stability is crucial to the life of the picture. By the test of time, egg-tempera and oils have proved themselves satisfactory media for easel paintings, but they suffer in varying degrees from a tendency to shrink and to become brittle with age. When a picture reaches a certain age, its surface usually becomes patterned with a series of micro-cracks which have picked up foreign matter and reveal themselves as dark hair-like lines most readily discernible in the lighter passages; these lines intersect and form a complicated network known as craquelure. Many factors contribute to the general appearance of craquelure, but in the main it is due to the natural shrinkage of the medium on ageing and the inability of the brittle aged film to withstand the slight rhythmical movements of the support. The forces thus imposed on the paint layer may cause a craquelure which extends through both the painted layer and the ground; different types may be recognized as characteristic of painting on wood panel or on canvas.

While superficially the craquelure on a painting may often appear

1 Cennino Cennini has described the whole process of tempera painting in his famous Treatise published in 1437 (date of earliest surviving text). Cennino D'Andrea Cennini: Il Libro dell' Arte (The Craftsman's Handbook), translated by D. V. Thompson, Jr., 1933.

2 Movements in response to changes of the relative humidity of the atmosphere (see later).
to resemble the intentional crackle of a glaze on oriental pottery or porcelain, it differs in this respect that it is an acquired characteristic and develops slowly; it is not regarded as a disfigurement, and rarely is it of such a nature as to introduce instability.

Two subsidiary types of *craquelure* may be mentioned. One is due to faulty technique, such as the application of a layer of quick-drying paint over a slow-drying paint; this type of so-called ‘youth’ *craquelure* is usually restricted to certain passages of the painting. The other condition is due to a faulty choice of material and is known as ‘alligating’, a term used to describe the system of wide depressions in the paint layer resulting from the use of bitumen or asphaltum, a material that is soluble to some extent in the binding medium and has notoriously bad drying properties.

In contrast with *craquelure* which is a more or less normal condition in old pictures, cracking and cleavage are serious forms of deterioration which affect canvases as well as panel paintings. A crack has the appearance of a crevasse in the paint layer, running inwards at right angles to the surface, and it usually penetrates through all the layers of the picture. It is caused by flaws in the ground or support. Cleavage, the more insidious form of deterioration, is due to loss of adhesion between the superimposed strata of the picture; it manifests itself in the formation of blisters which, if they be allowed to fracture, will lead eventually to the loss of paint.

While cracking and cleavage may be found in pictures of any age, they are to be seen most frequently in old pictures where the binding media have lost much of their original elasticity as a result of chemical changes, and are no longer able to adapt themselves to movements of the support occasioned by exposure to environmental changes. Both these conditions are revealed particularly when there are sudden extreme changes in the relative humidity of the atmosphere. The conservation of the paint layer depends in the first instance on the condition of the supports, whether panel or canvas, and their reactions to the environment.
PAINTINGS ON PANEL

Church furniture of painted wood was in common use in Italy from very early times, and an obvious development was the use of flat painted panels for wall decoration and for altar pieces. When large panels were required they had to be made by joining planks together, and for this purpose a cement was used made from lime and cheese which is, essentially, calcium caseinate. If the planks were thick enough, they were pegged as well with iron or wooden dowels. Very heavy panels were often additionally reinforced with cross-bars, either inserted wedge-fashion into a converging channel cut across the grain, or fixed in some other way to the back. Such reinforcement was not possible in the case of thin panels especially when, as in the side leaves of a triptych, they were painted on both sides. Panels completely protected on both sides by paintings have proved to be very durable, whereas thin panels painted only on one side have been less able to withstand the influences that cause warping.

The mechanical behaviour of painted panels is closely related to the environment to which they are exposed:

1. When conditions of storage or exhibition are kept constant as regards temperature and relative humidity, a panel tends to attain a state of equilibrium and no warping, cracking, or movement of any kind need be anticipated.

2. When a picture is transported to a locality where the environmental conditions are different from those to which it has been accustomed, a certain amount of movement will take place. In a composite panel that is unprotected at the back, each constituent plank will tend to cup, and at this stage cracks may make their appearance on the painted surface opposite the joints.

3. In a picture gallery where the air is not conditioned, day-to-day variations in temperature and relative humidity are to be expected, but normally these do not impose any great strain on the support. However, in addition, there are the long-term variations of a seasonal nature to be considered. When artificial heating is brought into action in the autumn and discontinued in the spring,
Schematic representation of the stratified structure of a panel painting (vertical section)
A. Encrusted with a corroding copper alloy which has afforded it cathodic protection

B. After treatment

22. SILVER LAMP FROM UR OF THE CHALDEES (c. 3000 B.C.)
the environmental changes may be considerable. These changes impose
strain on the support and for this reason care must be taken especially in
the spring and autumn to look for any signs of cracking and blistering
or weaknesses in the paint overlying the joints in composite panels.

Temperature and humidity are the most important factors in the
life of a panel painting. Exposure to extremes of temperature will
cause desiccation and shorten the life of the picture, but extremes of
humidity are the more potent and insidious cause of trouble, wood
being a bad conductor of heat but responding quickly to damp and
dryness. In the conservation of panel paintings, however, it is, above
all, variation of relative humidity that has to be guarded against as
this causes intermittent movements of the support that give rise to
cracking and cleavage of the ground and paint layers.

Easel paintings have been carried out on several other types of
support—metals, slate, marble, &c. Of these the metals are the most
important despite the fact that ground and paint layers do not adhere
well to the non-porous surface. Metals are, moreover, good con-
ductors of heat, and for this reason paintings on metal are more prone
to damage by variations of temperature than by variations of the
relative humidity of the environment.

Panel paintings: mechanical damage and treatment

1. Cracking. The most sensitive kind of easel painting is un-
doubtedly the large wooden panel, especially when it is made up
from thin planks and painted only on one side. Such a panel is prone
to mechanical damage. It may be fractured even by lifting or moving
it carelessly. A less obvious cause of trouble may be the position of
the picture on the wall, where it is perhaps exposed to a mild draught
of air from a door or window, and the changing humidity will
ultimately cause it to crack. This tendency to cracking explains why
so many panel paintings have been treated in the past with a cradling
or parquetry reinforcement at the back. The cradling has been applied
with the twofold object of giving mechanical strength to the panel
and of confining its movements to one plane so as to prevent twisting
or warping. Cradling consists of a series of wooden slats ('stretchers')
fixed rigidly at intervals in the same direction as the constituent planks, i.e. along the grain of the panel; these stretchers have slots to accommodate another series of wooden slats (‘runners’) arranged at intervals at right angles to the grain and in contact with the panel. While the stretchers are fixed, the runners should have freedom of movement to allow for slight expansion or contraction of the panel, but so often the runners themselves become warped and immobilized, with the result that the cradling becomes the prime mover, and the panel has to adjust itself as best it may. For this reason, if a cradling has to be applied to strengthen a picture, it should be kept as simple as possible.

Other ways of reinforcing a panel are by inserting X-shaped wedges or inlays across the joints of planks in order to prevent their opening, or to trust to a series of ‘buttons’ glued across the joint to hold the crack. Buttons are palettes of wood about 4 in. by 2 in. by \( \frac{1}{2} \) in., the size varying with the conditions. A common type of reinforcement is that achieved by gluing textile to the back of the panel, and this is often found where a panel has been enlarged by the artist in the course of painting the picture. Rubens often varied the proportions of his smaller panels, and reinforced the backs in this way. When a thin panel has been enlarged it is seldom that the different members are in harmony, and warping often occurs. Some of these enlarged panels are so warped that it is difficult to contain them in their frames. Not infrequently textiles were fixed on to the face of panels before applying the painting ground, and this is a sound technique because it not only strengthens the panel, but at the same time provides a good key for the painting ground.

When a panel painting is observed to develop a convexity on the painted side, or a series of convexities or undulations corresponding to its constituent members, this should be regarded as an emergency requiring immediate attention. The trouble may arise from one of two causes:

Firstly, it may be due to the fact that the picture has recently been hung in a dry room or exposed to heat, e.g. by sunning or by being near a calorifier.
Secondly, if a picture has attained a state of equilibrium under conditions that are rather damp, say in a large private house, and has been lent to an exhibition, its exposure in the drier conditions of the gallery may cause dehydration at the back, where unprotected by paint, with consequent cupping of the panel (see Wood, p. 118).

In both cases the picture should be taken at once to a cool basement room where conditions are known to be damper and laid flat, face downwards. The picture may usually be left in its frame, but any nails or metal cleats around the edges that might restrain the natural tendency of the panel to revert to its original shape should be removed. A restorer with experience of panel work should then be consulted. He may consider that it will be sufficient to leave the panel to recover over a period of time, or that some special form of treatment is required. Recovery, which in the natural course of events may take many months, can be assisted by the judicious application of pads of damp blotting-paper to the unpainted side.

After the panel has recovered its shape and been given any necessary reinforcement, cracks may remain on the painted side, and these will require to be filled with stopping. Various materials may be used for stopping, one of the commonest being a putty made from plaster of Paris and glue to which has been added a little stand oil. After the crack in the painting ground has been filled in, the surface of the stopping is painted in the colour and texture of the original, taking care that the new paint is level with the adjacent surface (see Inpainting, p. 175).

When a panel painting is being sent from its normal environment to an exhibition, it is a wise precaution to protect it on the back with a material that is impervious to moisture, e.g. Polythene sheet, fixed with Polythene tape. Protection of the back in this way tends to counteract the effects of changes in humidity and warping is prevented. This precaution is only necessary when the panel is removed from the environment to which it has become accustomed. On its return home the sheet should be detached as there is no advantage in covering the back when the panel is known to be stable in its home surroundings.
2. Cleavage and flaking paint. Cleavage may take place between any of the adjacent strata of the painting. When it occurs in isolated areas it will commonly result either in the appearance of a blister or in paint becoming loose on the surface.

When cleavage is widespread, the condition is more serious. Extensive cleavage is an indication that the panel is no longer able to fulfil its function as a support, and consideration has then to be given to the possibility of its replacement, either by a fresh panel, or by some structural material that may serve the purpose more suitably. This type of operation, known as 'transfer', is one of the greatest delicacy, and is only undertaken as an extreme measure by an expert.

When loose paint is found on the surface of a picture, action must be taken at once to prevent it falling off. If movement of the picture is likely to result in loss of paint, emergency measures should be taken while the picture is still on the wall by sticking mulberry tissue over the damaged areas using as the adhesive dilute gelatine. If the glass cannot be detached from the front, the picture will have to be taken down carefully and removed from its frame. It is then laid face upwards in preparation for treatment. The cause of flaking may be due to the breakdown of an old repaint or to tension in the panel resulting from an advancing crack in the wood, but there is no doubt that, fundamentally, such instability is caused by exposure to uncontrolled humidity changes (see p. 4).

The flaking paint is reattached to the picture by injecting an adhesive under the raised paint—usually either a dilute solution of gelatine or a mixture composed of beeswax and resin—and applying gentle pressure and heat so as to flatten the paint. This work must be carried out with dexterity as the glue so quickly becomes tacky and the wax-resin adhesive shows a tendency to set prematurely. Operations can be greatly facilitated by using the appropriate tools. Fine brushes and spatulas are required, and an electrically heated steel spatula having a gently curved blade is an invaluable instrument in preventing the adhesive from setting too soon. The degree of heat should be controlled within narrow limits so that the instrument can
be employed to soften and flatten the paint without changing the
shape of the fractured edges.

It is important to draw attention to the danger of fingering a
blistcr before treatment, as the paint is brittle and under compression,
and it may easily fly to pieces at a touch.

PAINTINGS ON CANVAS

The advantages of a textile support for easel painting came to be
recognized in the fifteenth century with the discovery of the possi-
bilities of oil painting, and from this time onwards canvas was
destined to become the principal form of painting support. It had
the merit of lightness and flexibility. A painting on canvas could be
carried as a banner and, within limits, it could be rolled, so that the
large easel painting was no longer static in the sense of being an
elaborate piece of furniture that must remain where it had been
erected.

Canvas is the fabric in general use today for easel painting. It is
woven from spun threads of cotton, linen, or hemp, in tabby or twill
weaves, and the fibres are protected from direct contact with the oil
by a sizing of glue. The canvas is then primed in such a manner as to
preserve the texture by covering it with a thin ground of linseed oil
and white lead. This painting ground provides a uniform surface to
work upon and one that will reflect light, so that the artist can exploit
to the full the translucent quality of his paints. The canvas is stretched
in a wooden frame provided with wedges in the corners so that it can
be tightened or slackened at will.

A stretched raw cloth canvas tightens when damp and expands
when dry, but the effect of glue sizing and of priming and painting
reduces such movements to a minimum, and may even cause them
to operate in the reverse direction, some expansion taking place in
damp conditions and some tightening in dry conditions. Very large
thin canvases are those most liable to sag in their stretchers and, if
allowed to do so, they may easily take on a permanent warp, espe-
cially noticeable towards the corners, where undulations appear that
soon collect dust. It is not a good thing, however, to tamper much
with the stretcher keys after they have once been adjusted to a reasonable mean. It may be that the site occupied by the picture is exposing it to too high a temperature or relative humidity. The picture should be rehung elsewhere for a year and kept under observation. Should there be no improvement it may be necessary to have the canvas removed from its stretcher and replaced in contact with a stretched raw cloth canvas. The additional canvas at the back will give support to the picture and slow down its response to atmospheric changes in humidity.

It is clear that in the case of canvas, the environment plays a part in conservation second only to its influence on panel. While the oil ground and the superimposed paint layers on a canvas are more adaptable to movements of the support than the gesso ground and paint layers on wood, the elasticity of the oil is gradually diminished with the passage of time; the paint layer becomes brittle and is easily cracked by movements of the support, due to humidity changes.

An accidental knock at the back of an old canvas may cause local stretching, and result in the formation of a series of concentric cracks in the ground and paint layer, a disfigurement not necessarily very serious, unless accompanied by cleavage and blistering of the paint.

It is characteristic of canvas that it tenders with age, and gradually becomes weakened so that it is easily punctured or torn. Tendering may be due to a variety of causes but a major factor is certainly the cumulative action of sulphur dioxide absorbed from the atmosphere and its conversion to sulphuric acid by the catalytic action of iron. A canvas may become so rotten with age as no longer to be able to bear the strain where it is held to the stretcher by tacks. In this case it may be sufficient if the edges only are strengthened, and this is done by a process known as strip-lining. Eventually there will come a time when the whole fabric gets so frail that it can no longer be regarded as a satisfactory support, and then consideration has to be given to the possibility of backing it with fresh canvas, a process known as lining or, more generally, relining.

1 In tacking a canvas to a stretcher copper tacks should always be used.
Canvas paintings: mechanical damage and treatment

1. Patching holes. Holes in canvas are repaired by backing the canvas with a patch, and filling either with stopping or with an insertion depending on the size of the hole. If the canvas is puckered it will have to be relaxed with moisture and flattened, and it may be necessary to remove hard paint from loose threads with a paint remover in preparation for the repair. In the case of a small hole the picture is laid face down on glass with the damaged area in contact with an oiled paper. A patch is prepared from a piece of raw cloth canvas larger than the hole, and it should be 'chamfered' by pulling several threads from the weave all round, thus thinning the edges so that the shape of the patch will not appear on the front of the painting. A suitable adhesive is a mixture of warm beeswax and resin, the patch being fixed with a warm iron; or a commercial rubber-paste cement¹ may be used or a glue-paste mixture. The picture is then turned over, the oiled paper removed, and the hole stopped with putty made from whiting and linseed oil in preparation for inpainting.

If the hole is large, it is filled in by inserting a piece of primed canvas of the same weight and weave as the original canvas. The weave of the inserted piece is aligned with the threads of the canvas, the contour traced, and the piece cut to the exact size and shape of the hole; it is then fitted in position, and secured temporarily on the front by Sellotape. A patch is applied to the back as already described, and stopping is used on the front if required before inpainting.

2. Mending tears. When tears in the canvas have to be mended, the torn edges are first brought together working from the back so that the weave can be distinctly seen, the broken threads being lined up, and the tear secured with Sellotape. The picture is turned over and the tear secured on the front. Then working from the back again, the Sellotape is removed and a patch applied as described above. In applying a long and awkward patch with a glue-paste adhesive, care must be taken not to stretch the material too tightly as on drying it may shrink and pucker the canvas. In using aqueous adhesives puckering of the joint due to differential shrinkage is a frequent

¹ e.g. Grade 753: National Adhesives Ltd., Slough, Bucks.
cause of embarrassment. Polyvinyl acetate emulsion\(^1\) or epoxy resins\(^2\) are less liable to cause trouble in this respect; they have the merit that they are used cold, and no great pressure is necessary in order to make good the joint. Wax-resin adhesives may also be used, and in this case the patch is applied, as mentioned above, with a warm iron.

3. \textit{Strip-lining}. When the canvas support is in good condition generally, and only the edges are weakened where they have been attached to the stretcher, these can be strengthened by applying strips of canvas to the back of the painting. Raw cloth canvas is used of the same weight and texture as the picture. The strips are cut to fit, allowing slightly more material at the outside edges for fitting over the stretcher; the inside edges must not extend beyond the width of the frame, and these edges are ‘chamfered’ as in preparing a patch. A strong adhesive is necessary for strip-lining, and the glue-paste mixture may be used, but the best cement for this purpose is probably Araldite 101 as this does not shrink on setting, and there should be no tendency for the canvas to cockle. Should its removal ever be necessary, however, this could not be accomplished by heat or solvent action, but only by rubbing down from the back with glass-paper. Whatever adhesive is used, the repaired canvas must be put in a press or kept under weights until the joints are secure.

4. \textit{Relining}. Relining is for canvas paintings what transfer is for panels: it is a method that can be used for saving a picture that is threatened by the failure of its support. Most old paintings of any importance have already been relined.

In relining, the painted surface is first reinforced with tissue paper applied with an aqueous adhesive; then, any previous relining canvas is removed. If a painting is being relined for the first time, the old canvas is smoothed down before a new canvas is applied. Relining is a major studio operation, but it is only by virtue of the fact that it can be expertly accomplished that so many Old Master paintings survive in such excellent condition today. This work is never attempted unless it is deemed to be essential for the survival of the picture.

\(^{1}\) e.g. Texibond \(\text{VaN}\): Messrs. Scott Bader & Co. Ltd., 109 Kingsway, W.C. 2.
\(^{2}\) e.g. Araldite 101: Aero Research Ltd., Hinxton, Duxford, Cambridge.
and then it is only undertaken by a qualified expert. Apart from the general risks involved in relining, the chief danger is that characteristics of the artist’s technique may be impaired; for example the *impasto* may be flattened and a tonality introduced that is foreign to the work, but in recent years the technique of relining has been improved and brought more under control. When heat is required, it is applied either by electric irons thermostatically regulated or by the use of a hot table,¹ and much ingenuity has been devoted to evolving methods of protecting the *impasto* during relining.

The glue-paste adhesives that were originally in universal use have largely been replaced by various wax-resin mixtures containing inert materials such as China clay, powdered chalk, &c. Wax relining has several advantages: the wax consolidates the paint layer by impregnation through the cracks from back to front, it avoids the complications that may result from using aqueous cements, and it can be easily removed at any time without strain when further relining is required. It is possible to carry out relining without heating by using a cold-setting emulsion adhesive such as Texibond V₄N and this has the advantage over glue that it remains permanently flexible. It is only applicable, however, in cases where the paint layer is in such good condition that it does not require to be consolidated by impregnation with wax.

In the preceding paragraphs treatment has been prescribed for dealing with certain types of deterioration arising primarily from failure of the panel or canvas support. It falls now to consider in how far the paint and varnish layers may suffer damage in themselves, or from what may be regarded as surface agencies.

**THE PAINT LAYER**

The paint layer is composed of insoluble particles of pigments mixed together in the greatest complexity, each particle being surrounded by an impervious film of medium. In egg-tempera and oils

¹ Ruhemann, H., ‘The Impregnation and Lining of Paintings on a Hot Table’, *Studies in Conservation*, 1953, 1, p. 73.
the medium isolates the pigments from moisture and noxious gases. This is not so in the case of water-colours or gouache, in both of which gum Arabic is the essential binding medium. White lead, which in water-colour is so readily blackened by the action of sulphuretted hydrogen in the atmosphere (p. 80), is immune when ground in oil or egg-tempera.

Pigments that are chemically incompatible in water-colour may often be mixed in oils with much less chance of their interacting with each other, and when resin is added to the oil it is even possible to use colours made from mixtures of pigments containing sulphur and copper without fear of subsequent chemical interaction which would lead to loss of brilliance. This has always been well known, and El Greco, for example, mixed such potentially incompatible pigments freely by employing isolating resin varnishes. It would seem therefore that in easel paintings there is no problem as regards the preservation of the pigment constituent of the paint layer. Whenever chemical changes have occurred, they are usually irreversible. Two of the commonest examples to be found on Old Master paintings are associated with ultramarine and copper resinate green.

Ultramarine is occasionally found to have lost its brilliance and to have become mottled white in appearance. This is due to the action of acid, probably arising as a legacy from some ill-advised process of cleaning in the past. In water-colours this so-called ‘ultramarine sickness’ has even been traced to acidity in the paper support.

In many early paintings it has been observed that areas that were originally a brilliant green have now assumed a dirty brown colour. This is due to a chemical change which copper resinate greens undergo, particularly in alkaline conditions. Since alkaline reagents (e.g. soap, wood, ash, ammonia) were used in the early days for cleaning paintings, these may be responsible for the browning that is so often observed. Alkalis can also destroy the colour of Prussian blue, a pigment widely used in the palette with yellows in composing mixed greens, and in this case also there is a tendency for the greens to go brown and there is no possibility of restoring the original hue.

It is questionable whether the amount of sulphurous gases present
even in industrial atmospheres would be enough to affect pigments bound in oil or tempera, but it is a different matter when soot, which is greasy and contaminated with sulphuric acid, is allowed to accumulate on a painting: it becomes ingrained in the *impasto* and *craquelure* and the weave of the canvas, and causes deterioration of all the materials with which it is in contact. Dust in any form is harmful and for this reason paintings that are of any value should never be hung against the chimney-breast of an open fire, or above a radiator; apart from the undesirable temperature and humidity conditions in these situations, the dirt problem is intensified by convection currents. The main reason for keeping pictures under glass is to protect the paint layer from noxious gases and from dust more effectively than can be done by the varnish itself. When a picture gallery is air-conditioned glazing is unnecessary.

As an oil painting matures there is a tendency for it to go down in tone; the medium becomes more translucent and darker in colour. The increased translucency is due to an increase in the refractive index of the oil on ageing, and this cannot be prevented. The main form of deterioration in the paint layer, however, is that caused by the darkening of the oil medium. Linseed oil is a complex mixture of fatty acid glycerides, certain of which (e.g. linolenic) tend to become dark in colour on ageing. When an oil painting is kept in a dull light the rate of darkening is intensified, and although the main cause of such darkening is usually due to the deterioration of the varnish (see below) the change in the medium itself is a factor of importance. The pernicious practice of rubbing over the surface of a painting indiscriminately with linseed oil under the mistaken idea that this will help to preserve it merely aggravates the tendency to darkening and cannot be too strongly condemned. Darkened oil paint may be restored in large measure by exposing the picture to sunlight which bleaches the discoloured ingredients of the oil. The medium of tempera painting is in a different category. It has no tendency to darken or lose its optical qualities, and as it has a comparatively low refractive index, delicate nuances of colour can be appreciated in this medium to a degree impossible in oil painting.
THE VARNISH LAYER

The final coat of varnish is applied to a picture to enhance its appearance, to give depth and luminosity to the colour, and unity to the composition. At the same time it gives the picture some mechanical protection and shields it from direct contact with the atmosphere. But varnish is impermanent, and as it decays and is removed and replaced many times during the life of an Old Master painting, it is necessary that it should be of such a nature that it can be easily removed, even in the decayed condition, without exposing the paint layer to the action of solvents that might soften it, and without involving any mechanical strain that might weaken it. It is hardly necessary to say that a varnish must also be transparent and remain so, and it must be tough; and to be of maximum optical value it should have a high refractive index approximating to that of the dried oil films (ca. 1.53). Further qualities are required, but the few enumerated above are sufficient to restrict the field to what are called the spirit varnishes, i.e. those made from resins that are directly soluble in volatile solvents such as turpentine or white spirit, and that on evaporation of the solvent leave behind a film of resin having the desired qualities. Of the natural resins, mastic and dammar come nearest the mark, and in spite of certain deficiencies are in most general use today. Certain synthetic varnishes have been recently introduced and these are discussed in detail later (p. 178).

Varnish deteriorates by blooming\(^1\) and by becoming brittle and opaque.

(a) Bloom. This word has been used to describe the dull bluish-white appearance that develops in a transparent film and leads to a cloudiness. The precise cause of blooming is not known, and while many factors may be involved, damp is certainly a major factor in the blooming of varnish. Dammar and mastic are sensitive to moisture which affects the varnish film after it is dry, causing it to become clouded with a bluish-white mist. The blooming of varnish may also be caused by the use of inferior materials or incorrect solvents in its formulation.

If dealt with in the early stages, bloom can be removed from the surface of the picture by rubbing with a soft silk pad, either dry or charged with a little wax polish, but if left unattended, the bloom tends to work its way into the film and it can then no longer be removed by surface treatment.

In the case of pictures that are prone to bloom, a palliative may be found in the use of a microcrystalline wax polish (see Appendix XII). For the precautions necessary to prevent bloom forming during revarnishing, see pp. 176–8.

(b) Embrittlement and loss of transparency. So far as our present knowledge goes, all spirit varnishes based on natural resins become brittle, disintegrate with age, and develop a yellow colour. The yellowing is most pronounced in varnished pictures that are exposed to a bright light. While daylight prevents oil paint from yellowing it has the opposite effect on dammar and mastic varnishes. Daylight is essential for the life of a painting in the oil medium, but exposure for long periods to direct sunlight is not recommended as this causes premature ageing of the varnish and hastens the day when its replacement will be necessary.

In old pictures that may still retain a coating of oil varnish, e.g. an oil-run copal, the decay of the varnish is accompanied by crazing which may communicate itself to the underlying paint and cause serious damage to the picture.

Whatever the nature of the varnish the breakdown of the film exposes a much larger surface area for the condensation of moisture and accumulation of dirt, and when the varnish has become badly discoloured and porous and is no longer protective its removal is imperative.

STANDARD STUDIO TECHNIQUES

1. Removal of varnish

Films of soft resin varnishes (mastic and dammar) are removed from a painting by applying solvent on swabs of cotton-wool to small areas at a time, the dissolved varnish being absorbed by the wool and the swabs changed frequently during the course of the
work. The choice of solvent is crucial as it must dissolve the varnish without affecting the underlying paint. Many solvents for resin are available—alcohol and acetone are the two most generally used—but they must be diluted with a restrainer (turpentine or rectified petroleum) in such proportions as to ensure that the liquid has no action on the paint layer, but dissolves the varnish without undue friction. In order to adjust the proportions of solvent and restrainer, cleaning tests are first made on the edge of the picture. It may be found that a mixture of alcohol and turpentine in the proportion of 4 vols. to 20 vols. removes the varnish without affecting the paint; a 2/20 mixture may be equally satisfactory, whereas a 1/20 mixture may be found to be effective only on continued rubbing and this is undesirable as it would be likely to damage impasto. In such a case the 2/20 mixture might seem to be the best to employ. This would then be tested on varnish overlying small areas of bright colour, e.g. vermilion (which is usually most sensitive), using for the purpose of the test a wisp of cotton-wool twisted round a match-stick. The swab is closely examined in order to discover whether, as a result of cleaning, any pink colour has been communicated to the cotton-wool. If so, the cleaning solution is too strong and it must be further diluted with turpentine. When the mixture has passed these tests satisfactorily, parts of the painting are selected and rectangular areas cleaned of varnish in order to form an estimate of the condition of the underlying paint and the change of tonality that may be anticipated as a result of cleaning. This preliminary routine is a necessary prelude to the cleaning of any important picture.

In stripping a picture the restrainer must always be readily accessible. In practice, the operator can hold the cleaning swab in one hand and the restraining swab in the other, so that should there be any indication of the softening of the paint, as by the appearance of a trace of colour other than brown on the cleaning swab, he can flood the area with restrainer and arrest the action. The areas are finally wiped over with the restrainer to remove the last of the solvent and any softened residues of varnish that may remain on the picture. In the case of high impasto or resolute brush-work, such as is found in
paintings by Rembrandt, hollows may be filled with dark varnish while the raised portions are light and possibly abraded. The cleaning of such paintings requires time and patience as the work may have to be carried out very largely by using small wisps of cotton-wool on match-sticks as described above, rather than larger swabs held in the fingers.

The experienced restorer will be able to estimate beforehand what the paint layer can stand in the way of cleaning. He will recognize the existence of any soft resins used by the artist in the course of painting the picture, and ensure that such a painting is not damaged by his cleaning solutions. He will consult with the curator regarding the removal of any repainting that dates from a former restoration. In cleaning a picture it is often found that previous repairs have been concealed by unnecessarily large areas of repainting, and the removal of such restorations, followed by inpainting of the stopping only, will usually be desirable.

2. Inpainting

The term ‘inpainting’ is used in picture restoration to emphasize the fact that the original paint is sacrosanct and must not be covered with modern additions of colour. The only exception is where the picture is abraded and part of the design rubbed away; here it may be necessary in the interests of aesthetics to retouch the paint layer with colour—a delicate operation for which paints are employed that can readily be removed if necessary. Where inpainting is required, either in oil or tempera paintings, the medium that is chosen is egg-tempera. This is applied irrespective of the original technique because it is not subject either to darkening or to increase in transparency; the oil medium, as has been mentioned, is subject to both. Tempera has the one disadvantage that it becomes insoluble, but this is no serious inconvenience when it is applied to areas of stopping only. In case it should ever be necessary to remove the modern paint, it may be applied over an isolating varnish of dammar in turpentine and this is the technique generally adopted, each coating of tempera being given a thin coating of dammar varnish before the next is
applied. As the dammar can always be softened by the use of resin solvents, this makes it possible to remove the retouchings at any time.

In order that the quality of the inpainting should be in harmony with the rest of the picture, it is necessary that the transition from stopping to finished painting should follow the same course as that adopted by the artist. Thus, the stopping should be coloured to match the ground of the painting; if drawing or under-modelling is present, this should be reproduced on the stopping, and the underpainting and glazing should be built up as nearly as possible in the manner adopted by the artist himself. The inpainting should be carried out using stable powder pigments ground in the medium and it is important that the finished restoration should be of slightly lighter hue than the adjacent passages of paint, a dark retouching being much more obtrusive than a light one. The aim should be not to deceive; it should be to conceal forms of damage that would distract the attention and by their presence make it impossible to appreciate the picture as the artist intended. A trained eye can usually detect restorations but, should the matter be in doubt, suspicions can often be confirmed by inspection of the painting in a darkened room under ultra-violet radiation, or by infra-red photography, or X-radiography.

After the painting is dry the picture is ready for revarnishing.

3. **Revarnishing**

As already mentioned, mastic and dammar varnish are the picture varnishes in most common use. It is difficult to decide which is the better as the resins themselves are complex mixtures\(^1\) and subject to much variation in quality. Mastic seems to give the more brilliant finish, but it requires a turpentine solvent which may be one reason why it becomes so yellow on ageing. Its use was abandoned by the National Gallery in 1949 in favour of dammar varnish,\(^2\) which, in

---


The tail of the dragon is an ancient repair in gesso covered with gold leaf.
24. Gold head of a bull from Ur of the Chaldees (c. 3000 B.C.)
contrast to mastic, may be dissolved in white spirit and is believed to be less prone to yellowing.

Dammar picture varnish is made from selected dammar resin by dissolving it in white spirit without the aid of heat. The resin is suspended in a gauze bag in the solvent contained in a covered jar, and the liquid shaken or stirred at intervals over a period of some weeks. A suitable concentration would be about 4 oz. of dammar in 1 pint of white spirit. A little stand oil is added, finally, to act as a plasticizer to toughen the film and to help brushing-out qualities, but never in excess of 5 per cent., i.e. 1 fluid ounce to 1 pint of varnish.

A picture should never be varnished until the paint is quite dry. The surface must be free from dirt, grease, and wax, and to ensure this it is rubbed over lightly with white spirit and allowed to dry before beginning operations. The varnish may then be applied by brush or spray. The work should be carried out in a warm dry room free from dust and draughts; care should be taken that the varnish is not chilled but is several degrees above the temperature of the room.

If a brush is used, it should be a special varnish brush made with bristles of the finest quality and kept exclusively for the purpose. The varnish is applied to the picture thinly in a series of overlapping squares, say 8 in. by 8 in. (depending on the size of the picture), using first horizontal strokes and then immediately afterwards vertical strokes. In this way any excess of varnish is removed and a film is left of uniform thickness free from brush marks. A second coat may be applied if considered necessary after the first has dried, but the aim should be to apply the thinnest varnish film consistent with realizing the desired optical effect.

When the spray is used—and spraying has much to commend it especially when there is high impasto—the fan-shaped type of nozzle is best. The pressure and the distance of the jet from the picture have a bearing on success and will be determined to a large extent by the viscosity of the varnish. In using the spray the jet must not be too near the painting, because it will then deliver varnish that is too liquid to control; on the other hand, if it is too far away, it will tend
to give a matt effect by covering the surface with minute spots or scales of resin that are too dry to coalesce, thus affording no protection.

In theory, the thinnest uniform coating should be obtainable by spraying, but in practice a uniform coating is difficult to achieve without building up a film that is at least comparable in thickness with that obtained by brushing.

When a picture has been varnished it is allowed to remain on the easel till the varnish is dry; it should then be kept under glass for a few weeks until the varnish becomes really hard, when there will be less tendency for the surface to bloom.

**Synthetic resin varnishes.** Various types of synthetic resin are now available and research is constantly proceeding with the aim of producing a synthetic varnish suitable for paintings that will have all of the required characteristics (p. 172) and none of the demerits of natural resins.

One of the first synthetic resins to be tested was polyvinyl acetate which has been used for many years on quite an extensive scale in the U.S.A. The choice of solvent is important as there are many solvents for this substance that are too active for application to oil paintings. Alcohol only should be used for preparing a varnish for brushing purposes and the standard solution is made by dissolving 20 grammes of polyvinyl acetate in 100 ml. of ethyl alcohol\(^1\) without the application of heat. This may take a long time but solution is facilitated by stirring and shaking. The standard solution may be diluted, if necessary, to half or quarter strength, by adding ethyl alcohol. This formula is given by Bradley\(^2\) together with a reliable spray formula for a polyvinyl acetate picture varnish composed as follows:

<table>
<thead>
<tr>
<th>Standard solution of polyvinyl acetate in ethyl alcohol</th>
</tr>
</thead>
<tbody>
<tr>
<td>(20 per cent. w/v)</td>
</tr>
<tr>
<td>Ethyl alcohol</td>
</tr>
<tr>
<td>Cellosolve acetate (or cellosolve)</td>
</tr>
<tr>
<td>Diacetone alcohol</td>
</tr>
</tbody>
</table>

\(^1\) Of strength 96 per cent. It should be noted that polyvinyl acetate does not dissolve in absolute alcohol.

Attention has also been devoted to the esters of polymethacrylic acid, which have the advantage that they are soluble in hydrocarbon solvents.

These two types of resins are linear polymers of relatively high molecular weight. Hence, for a given solids content, the viscosity of their solutions is high and this gives rise to practical difficulties in their application. These can be overcome by using a varnish based on polycyclohexanone\(^1\) which is a polymer of low molecular weight similar in physical properties to the natural resins. This is made up in white spirit and yields a relatively non-yellowing film which, un-plasticized, is rather glossy and rather brittle. The glossy nature of the film is no disadvantage as the optical quality can easily be adjusted after the varnish has dried, by the application of wax polish (see below). The brittleness, however, is a problem that is still engaging the attention of manufacturers, and several formulations are already on the market. It seems certain that with the discovery of a suitable plasticizer, this varnish will become a serious competitor to varnishes of the conventional type.

4. Wax varnishing

Wax varnishes are merely polishes, because even if applied thinly to the painting by brush, the wax residue that remains on the picture after the solvent has evaporated is polished by using a soft long-bristled brush. The resulting gloss may be enhanced, if desired, by further polishing with a soft cloth. The surface of a wax varnish even when polished is duller than that given by resin or synthetic varnishes, and wax may be applied over these to reduce their high gloss. Resin and synthetic varnishes are never applied over wax.

A common formula for wax varnish contains 4 oz. of white beeswax dissolved in one pint of distilled turpentine with the aid of a water-bath maintained at about 60° C. An improved form due to A. E. A. Werner is the wax salve made from microcrystalline paraffin wax and Polythene wax (see Appendix XII). This leaves

\(^1\) This material is available commercially as Resin AW\(_2\), manufactured by Badische Anilin u. Soda Fabrik, Ludwigshaven; an English equivalent, Resin MS\(_2\), is manufactured by Howards of Ilford Ltd., Ilford, Essex.
a surface that is relatively non-sticky and the salve may be used either as a polish or, in the case of smooth painting, for eliminating ingrained dirt.

In order to wax-polish a picture, the surface is first rubbed very gently with a piece of soft silk to remove any dust. Some of the salve is then placed in the palm of the hand, where it quickly softens. A piece of cotton-wool wrapped in thin silk is charged with the softened wax, and used to polish the picture with the minimum of pressure. In the case of a picture ingrained with dirt, the polish is applied direct, using the cotton-wool without the covering rag. The waxed cotton-wool collects the dirt, and finally the surface may be polished (using a new covered pad) by the method described initially. Large canvases may require some support from the back (by a pad of papers or the like) during cleaning and polishing.

EXCEPTIONAL FORMS OF DETERIORATION

1. Insect attack on panels. Painted panels are sometimes found to be attacked by wood beetle (Anobium), the larvae of which tunnel into the wood to emerge eventually from the surface as fully developed insects. Activity may be recognized by the appearance of holes or frass and, if action is not taken in time, the support may become so weakened that cleavage of the paint results.

When insect pests are found to be active, the painting should be at once isolated to prevent adjacent pictures becoming infected. It must then be sterilized, using an insecticide that will not damage either paint or varnish, the most satisfactory form of treatment being fumigation with carbon disulphide (see p. 124). To confirm that sterilization has been successful, the panel should be inspected at intervals of six months after fumigation. If the old insect-holes are filled with wax at the time of fumigation, the appearance of any new ones will be readily detected.

2. Fungoid attack on canvas. A canvas that has been relined with a glue adhesive is prone, under conditions of damp heat, to be attacked by moulds. The appearance of fluffy white growths, visible especi-
ally on dark passages of paint, is an indication of fungoid activity, and if observed through a reading-glass will likely be found growing on the glue where it is exposed in the craquelure. To arrest and prevent further growth, the surface of the painting should be rubbed over lightly with soft pads of cotton-wool and the picture exposed to sun and air. This may be all that is required. The glass, however, should be sterilized, before it is replaced, by cleaning with cotton-wool that has been moistened with formalin. In very bad cases it may be necessary to use formalin on the actual painting, but this should be left to the discretion of the restorer who will know from his experience or can discover by testing whether the picture is in a condition to be treated with aqueous solutions. The back of the canvas must also be considered and may be given permanent protection against mould growth by the use of a non-volatile fungicide such as Santobrite (p. 28).

For Further Reading

PART II

METALS
PART II

MATERIALS
CHAPTER VIII

GENERAL INTRODUCTION

The conservation of museum objects of metal is a study in itself. Metals form a heterogeneous though well-defined group of materials, almost all subject to corrosion—that is, to the loss of metallic properties with the formation of mineral incrustations. This is due to a series of chemical or electro-chemical reactions, and disintegration may be slow or accelerated depending on the nature of the metal and the conditions to which it is exposed. That metallic corrosion is accompanied by a change in appearance is fortunate, as this calls attention to the fact that chemical change is taking place, and the sooner the object is treated the better its chance of survival without loss of character.

In order to arrest corrosion, it is necessary to discover and remove the cause. This may be, and often is, the presence of some active element such as chlorine that has entered into chemical combination with the metal, in which case it can only be removed by a process of counter-attack involving other chemical changes that are applied in the laboratory under scientific control and that can be relied upon to achieve this end. The effect of such treatment may be striking, as when an object concealed under a gross incrustation of minerals is so dealt with that its original metallic quality is restored; but such an extreme change in appearance is not the inevitable result of chemical treatment. It is often possible by the exercise of ingenuity to arrest corrosion without sacrificing the mineralized or patinated surface, and there are many cases where this is desirable. The nature and condition of the specimen will determine what freedom of action exists and to what degree stability can be regained with the minimum of sacrifice.
In dealing with corroded metals there are, in fact, three possible methods of treatment: the use of solvents, the use of chemical and electro-chemical reduction, and mechanical methods; and while it will be convenient to study these methods individually, it should be understood that they are not mutually exclusive, but are applied as occasion demands in combination. Thus, the use of solvents and reduction methods are each dependent for ultimate success upon the application of certain forms of mechanical treatment (brushing, washing, &c.) and there are many cases where the best results can be obtained only by a combination of all three processes. Metals have individual characteristics; they are susceptible to different chemical reagents, they corrode in different ways, and in doing so they form different kinds of incrustations that must be recognized and studied before it is possible to decide upon the best form of treatment. Chemical methods involving the use of specific solvents can only be described in dealing with the metals themselves. By contrast, the methods of reduction and the mechanical methods are of general application to all corroding metal, and a description of these processes is introduced, therefore, as a preamble to the study of the individual metals of antiquity.

In considering what treatment to apply, a physical examination is of the greatest importance. This is carried out with the aid of a lens and, if necessary, by exploring with a needle or in the case of ferrous metal by using a magnet. It is necessary to form some estimate of the thickness and regularity of the encrusted layer, strength of the residual metal, presence of ornament, taking special note of fine detail and of any cracks. To reveal the presence of hidden ornament in cases where the metal is heavily corroded, it may be necessary to use X-rays. Methods of treatment are based upon this preliminary assessment, and tables are given at the end of each chapter (save in the case of gold), to assist in deciding what may be the best procedure. For the application of the methods recommended reference must be made to the text, but whether the suggested methods are, in fact, the best to apply in the circumstances can only be decided by the knowledge, experience, and resource of the operator.
If metals are considered as raw materials, it is noteworthy that few are found in nature in the free or uncombined state. They occur in combination with non-metallic elements in the form of minerals, and these provide the ores from which the metals are won by smelting. Minerals, rather than metals, are thus the stable forms under natural conditions, and it is to these minerals that metals tend to revert, especially when they are buried in the ground for prolonged periods of time. It has been observed that, other things being equal, the more easily a metal is won from its ores, the greater is its stability. Thus, metallic tin which is obtained with the minimum expenditure of energy from the oxide (cassiterite) shows less tendency than copper to become reoxidized. Because of its relative stability, tin has been used as a coating for protecting copper vessels from at least the beginning of the Christian era, and even in this attenuated form it is often found to have survived in excellent condition.

When metals are buried in the ground, the rate of corrosion is intensified according to the degree of acidity of the soil, its porosity, and the presence of naturally occurring soluble salts. These substances in the presence of moisture conduct electricity and are known as electrolytes. Metallic corrosion is an electro-chemical phenomenon. The absolute strength of electric currents flowing over corroding metal can be measured, and by comparing such measurements with the rate of corrosion, change of weight, &c., it has been shown that the whole complicated cycle depends on electro-chemical principles.

Metals that have been buried and have already suffered some degree of corrosion have a comparatively porous surface and are thus liable to retain traces of salts. Often these salts are sealed up in what appears to be a stable incrustation or patina, but should they be exposed to moisture and oxygen they are likely to give rise to fresh activity, with consequent pitting of the surface and possibly serious disfigurement. Metals behave differently, however, in their reactions to non-metals. Sometimes there is a progressive growth of the surface minerals which increase at the expense of the metallic core; in others, the mineralization is compact and stable and, after preliminary development, it tends to inhibit further change. Where development
has been slow and uniform the intricacies of shape and ornament are preserved and the appearance of the object may even be enhanced by the colour and texture of the patina. A patina that has developed slowly may sometimes be evidence of antiquity and a fine patina will determine the market value of an object. It is necessary, therefore, in considering the preservation of metal objects to consider also the preservation of their patinas when these may be taken as evidence of age or are in themselves of aesthetic value. The patinas of Chinese bronzes, for example, which have matured through the years in ground substantially free from soluble salts, have an irresistible appeal for the collector; but the danger is that mineral patinas which are apparently quite stable may become unstable if exposed to damp, and when corrosion becomes active it is sometimes a most difficult task to arrest it without serious loss of surface appearance.

Burial in the earth, however, is by no means essential for the corrosion of metals. Even when objects are kept indoors, moisture and oxygen are enough in themselves to cause dulling of the surface due to the formation of a layer of metallic oxide. The results of attack by sulphurous gases that are present in industrial atmospheres are even more readily apparent, yielding the darker films of metallic sulphide known as tarnish; but disfigurements of oxide and sulphide are purely superficial and readily amenable to treatment.

The individual metals of antiquity are few in number—gold, silver, copper, lead, tin, and iron, but they have been mixed together as alloys, intentionally or otherwise, from remote periods. Thus, electrum (an alloy of gold and silver) is much more common than pure gold, and ancient silver almost invariably contains copper. Copper and tin were alloyed intentionally to make the more useful metal called bronze, and lead was often added to the mixed metal. But there is evidence that tin was not always distinguished from lead by the early metallurgists; analyses sometimes reveal the presence of both or even a preponderance of lead when this could only yield an alloy that was less serviceable for the purpose intended. Some of the earliest examples of bronze from Ur of the Chaldees contain such a small
quantity of tin that it is doubtful whether the composition was intentional and it may therefore be misleading to use the term ‘bronze’ in this connexion. Lead and tin have themselves been alloyed together from remote periods. Copper and zinc were alloyed intentionally in Roman times for coinage, but here again, there are isolated examples of brass occurring in earlier civilizations (China), and to what extent this was intentional can only be determined by further study. The manufacture of steel is a comparatively modern process, but the carburization of iron was discovered many years previously at the armourers’ forge, where iron and slag were hammered in the presence of carbon and wrought, possibly by chance, into steel, an alloy superior to all others for the manufacture of sword-blades. Such a chance occurrence might possibly provide a basis for the celebrated sword-blades of fable and legend!

Metallic corrosion has been described as an electro-chemical phenomenon. This action can be demonstrated where two dissimilar metals are in contact in the presence of a conducting solution of a salt—the essentials of a simple electric cell; under these conditions the baser metal is corroded preferentially as long as the two are in electrical contact, the more noble metal surviving by what is called ‘cathodic protection’. By conducting such experiments with different pairs of metals it is found that the metals can be arranged in order of their nobleness from gold downwards, through silver, copper, lead, tin, and iron, the baser metal always being sacrificed in the presence of an electrolyte where it is in contact with another metal higher up in the series. An interesting example of survival by cathodic protection is that of the frail but perfect silver lamp discovered at Ur of the Chaldees among a mass of copper lamps (Pls. 22 A and B). The fact that only the silver lamp survived was significant, and the condition of the copper lamps was such as to confirm that they had been preferentially corroded in accordance with this theory. For the same reason, bright inlays of copper or its alloys are sometimes found

---

1 It should be noted that when metals are alloyed, as in stainless steel, the position of the alloy in the electro-chemical series cannot be determined even approximately.
embedded in heavily rusted iron swords and knives, iron in relation to copper being the baser metal.

It is not only the conjunction of two different metals that promotes electro-chemical decay. Where two or more metals are alloyed together as tin and copper are in bronze, the susceptibility to corrosion is greater than in the case of an unalloyed metal and, when once it has set in, the decay of bronze proceeds more quickly than is the case with copper alone. In the same way base silver is corroded more intensely than pure silver. Indeed, it is no uncommon occurrence to find on treating an object that has all the appearance of being a corroded bronze, that it is in fact made of base silver, the baser constituent (the copper) having corroded preferentially and covered the white metal with a heavy green deposit, thus concealing the true nature of the alloy. In such cases treatment can be applied to remove the green incrustation, revealing the underlying silver enriched in appearance in proportion to the quantity of copper that has been leached from the alloy in the process of corrosion.

From these considerations it follows that different metals are at different electric potentials relative to a surrounding electrolyte, and it is then but a step to the idea that if different potentials could be established on one and the same metal (e.g. by local inclusions in the surface, varying porosity and concentration of the electrolyte, &c.), an electric current would flow leading to an electrolytic form of decomposition. This has proved to be the case, and in some measure may serve to explain the more catastrophic forms of corrosion such as the rusting of iron and also the behaviour of metals that have been subjected to the periodic action of saline water.

There is, however, a brighter side to the picture, and it may be introduced by reference to a minor catastrophe which took place, not in the earth, but in a museum. A painted zinc label had been left on a corroded silver dish in a cupboard. One must assume that the cupboard was damp and that the corrosion layer on the silver was able to act as an electrolyte. After the lapse of some years it was discovered that the label had gradually subsided into the dish and the remains stood in a clear liquid to which part of the label had apparently de-
composed. The liquid proved to be zinc chloride and on clearing it away it was found that the silver at the point of contact had been reduced from dull horny silver chloride to bright metal. Now, metallic zinc is a baser metal than any of the others to be preserved in museum work, and this simple reaction may actually be put into practice in modified form in the laboratory, for the treatment of metal objects that have been disfigured by corrosion. The method is known as electro-chemical reduction.

**REDUCTION METHODS**

1. **Electro-chemical reduction**

   The standard method involves the use of zinc and caustic soda, and the reaction is carried out in an iron or enamelled container with the aid of heat. Heat promotes chemical action, and thus enables results to be obtained reasonably quickly.

   **Materials.** The zinc is used in granulated form or in the form of a coarse powder, so as to present a large surface of contact with the object under treatment. It may also be used in the form of zinc wool and wrapped around the object. Zinc dust is not as a rule so effective, as it tends to clog together.

   Granulated zinc may be purchased as such, or zinc may easily be prepared in this form by heating the metal in an iron ladle (it melts at 419° C.), and pouring the molten metal from a height into a bucket of cold water. If required as powder, the granulated zinc is pulverized in an iron mortar to the requisite grain size, and it may then be passed through a sieve of about 30 mesh to the inch, which yields a granular powder sufficiently fine for the most delicate operations. The advantage of using zinc wool is that wrapping ensures that electrical contact will be satisfactory; on the other hand, the object is concealed during treatment by the wool, and the progress of the reaction more difficult to follow.

   Commercial flake caustic soda is dissolved in water and employed as the electrolyte. The strength of the solution should be not less than 10 per cent. (2 oz. to 1 pint of water). When in contact with excess of
zinc this solution will be 'spent' after half-an-hour's heating, i.e. evolution of hydrogen which is the potent factor in effecting reduction will then have ceased. It is usual, therefore, to employ stronger solutions, say up to 20 per cent., for objects that are heavily corroded. It is better to change such solutions occasionally rather than to work with solutions in excess of 20 per cent. concentration.

Method of procedure. Heavy objects such as celts, spear-heads, and the like are buried under a heap of granulated zinc in an iron basin, covered with the caustic soda solution, and the liquid boiled for an hour or so over a gas-ring or Bunsen burner, preferably in a fume cupboard as the vapours evolved are irritating. During this time, distilled water is added, as required, to maintain the volume of the solution at the original level. Delicate objects such as fibulae are buried in zinc powder, covered with a 10 per cent. solution of caustic soda, and heated over a steam bath for an hour, taking the same precautions to maintain the volume of the solution.

The result of such treatment is to soften the incrustation. In the case of bronzes, the green colour will disappear, and it will be found possible by brushing under a stream of water to remove much of the brown, muddy deposit, revealing a metallic surface upon which are retained any details or ornament. If the metallic surface is not entirely clean, it will be necessary to repeat the reduction with fresh zinc and caustic soda. It is not permissible to leave vestiges of oxide unreduced as they seal in traces of chloride, and the only satisfactory method of eradication is by further reduction followed by brushing and washing. If traces of chloride are allowed to remain, a further outbreak of corrosion is inevitable, sooner or later. In some cases the vestiges of oxide may be found to be covered by a film of metallic copper which will make them more difficult to remove. It is most important, however, that they be removed. This copper plating is not so liable to occur when the reduction is carried out in boiling solutions, as the plating is discouraged by the mechanical agitation.

In cases where residual incrustations cannot be removed even with a steel or glass brush they will have to be picked or scraped off. This often requires the exercise of much patience, using a magnifier and
25. CROSS-SECTION OF A CORRODED BEAD OF BASE SILVER FROM UR OF THE CHALDEES, SHOWING STRATIFICATION

26. CROSS-SECTION OF PYRAMIDAL OUT-GROWTH KNOWN AS ‘WARTY PATINA’ FOUND ON A CORRODING CHINESE BRONZE

The light areas are tin oxide and the dark areas mainly cuprite. The multi-layered structure indicates that the incrustation is of great age.
A. As excavated—the silver, inlaid with gold and niello, completely covered with corrosion products

B. After treatment and restoration

27. THE BULL CUP FROM ENKOMI (c. 1400 B.C.)

28. FRANKISH BUCKLE (7TH CENTURY)
Left: Before cleaning. Right: After cleaning by the thiourea method
mounted needle, and going over the whole surface a small area at a
time, picking off residual lumps of incrustation to expose the pockets
of greenish-white pasty material (cuprous chloride) that underlie
them. A subsidiary reduction is then required to ensure that salts
exposed by this treatment are rendered soluble so that they can be
washed out.

Where equipment is available for electrolytic reduction it is con-
venient to give the finished object half an hour’s treatment in the
tank (Fig. 6, p. 194) as this facilitates the removal of the last traces
of salt, thus cutting down the time required for washing. This
operation must be conducted with thoroughness and is described in
detail on pp. 197–9.

In the course of reducing objects with zinc and caustic soda, the
soda is spent and must be discarded, but a quantity of zinc remains,
and, as it is rendered largely inactive due to the formation of a film
of oxychloride and carbonate, it tends to accumulate in the labora-
tory. It may be recovered for further use by washing it with water
acidified with hydrochloric acid, then with distilled water, drying,
and, if necessary, melting in the iron ladle, and regranulating.

Variations in procedure. The electro-chemical reduction process as
outlined above admits of several useful variations:—

(i) If the object is encrusted with lime or chalk in addition to
metallic oxides, it may be that the reaction will proceed more regu-
larly if the reduction is carried out using a 10 per cent. solution of
sulphuric acid instead of caustic soda.

(ii) Where local treatment only is required, as in exposing an
inscription on the base of an Egyptian bronze statuette, reduction
may be done by repeatedly applying a paste of zinc powder and
sulphuric acid of 90 per cent. strength. Alternatively the paste may
be made in situ on the surface of the object by applying a few drops
of the acid and mixing in a small quantity of the zinc powder with
a glass rod or a tuft of glass bristles held in a piece of glass tubing.

(iii) Silver objects which are not heavily coated with horn silver
respond well to reduction with zinc and hot formic acid, and even
more readily when aluminium granules are used instead of zinc.
The main advantages of electro-chemical reduction are that it requires no apparatus beyond what is readily available or can be easily improvised, and with the modifications described above it provides the means of reducing metal objects locally when desired. It is, however, subject to the limitations that apply generally to all methods of reduction. See p. 217.

2. Electrolytic reduction

As an alternative to the method already described, it is possible to effect reduction by using an electric current, see Fig. 6. The corroded metal object is made the negative electrode (cathode) in a suitable electrolyte such as caustic soda solution, the positive electrode (anode) being of sheet iron; under these circumstances the reducing action is then dependent on the application of an external e.m.f. as, for example, by the use of an electric battery. This method is sometimes referred to as electrolysis.

When current passes, hydrogen is evolved at the cathode with the result that the incrustation is gradually reduced and saline matter decomposed. As the reduction progresses, chlorides are transferred from the cathode to the iron anode. This method can be very effective in cleaning badly encrusted metals of all kinds, but unless conditions are made to conform to accepted standards (as described below), complications in the form of secondary reactions may vitiate results.
Equipment. The electric supply must be direct current, either from a secondary cell or accumulator, or it may be A.C. stepped down from the mains to about 12 volts and rectified (see Diagram, Appendix XIII). It is the amperage in the circuit that is the most important factor, however; this depends upon the resistance of the electrolyte and the size of the electrodes, but will vary during the process as the resistance of the incrustation on the object alters under treatment.

In practice, caustic soda of about 5 per cent. strength is chosen as the electrolyte, and two iron anodes are hung equidistantly, one on each side of the cathode, and only a few inches from it. Under such standardized conditions the rate of reduction will then depend mainly on the current density which can be regulated by a variable resistance to approximately 10 amps per square decimetre of cathode area, so as to give a steady vigorous evolution of gas. This figure is not critical for iron or steel. For copper and silver, however, the current density should not be allowed to fall below 2 amps per square decimetre, otherwise there is a tendency for a film of metal, usually salmon-pink copper, to be deposited on the surface of the object in the region of the chloride-containing incrustation, and this is difficult to remove. At first the resistance of the incrustation may be considerable, but in the course of treatment, after perhaps an hour or more, a rise in amperage will indicate that the resistance of the incrustation has fallen, and this should be compensated by increasing the external resistance.

Method of procedure. A convenient arrangement is as follows:— A glass tank is filled to a suitable depth with 5 per cent. caustic soda and brass bars are laid across the top from which the electrodes are suspended by copper wires, thus allowing of easy adjustment. The object is attached to the negative pole of the electric supply, whilst the iron anodes are wired together and joined to the positive pole through an ammeter and adjustable resistance.

The iron anodes are heavily attacked during electrolysis. They tend in time to deposit flakes of metallic oxide, &c., in the solution, a state of affairs that may be prevented by tying them in nylon bags. But, even so, it is necessary to remove the anodes occasionally and
scrub off the products of chloride attack. Graphite and carbon electrodes have also been used, but these break down in alkaline solutions and deposit a black film on the object. Attempts to brush this off only burnish it into the surface. An advantage of the iron anode is that when iron is plated on to the object, as often happens when prolonged electrolysis is necessary, it can be easily removed in the subsequent washing process. The chance of complications taking place is reduced by using anodes of stainless steel. Nylon bags may be tied round the objects being treated if additional support should be required, or if there is any chance of metal inlays, &c., falling off into the solution.

Another refinement is to cover the surface of the electrolyte with floating glass capsules or short lengths of Polythene tube, called croffles, in order to minimize the escape of caustic spray which, though not excessive, tends to irritate the throat. The glass croffles are preferable to those of Polythene, and are used in the proportion of about 1 lb. of croffles to 90 square inches of surface.

Instead of using iron or steel anodes, the electrolytic tank itself may be of iron instead of glass, and made to serve as anode, but if originally galvanized it will be necessary to remove all traces of the galvanizing metal which otherwise might plate on to the objects. This may be removed by running the tank as anode for a considerable period with a temporary iron (tin-plate) cathode, the caustic soda solution being maintained at the highest possible level in the tank until the last of the zinc has been stripped from the metal and the tank surface is clean. After washing and making up with fresh electrolyte, the tank is ready for use.

In reducing a corroded object the duration of treatment will depend upon the nature of the incrustation, but its removal can be facilitated by taking out the specimen under treatment occasionally and brushing and scraping the surface. The object should be removed while the current is still running. In no circumstances should it be left in the tank after the current is switched off, otherwise it may become plated with any metallic impurity present in the electrolyte.

Caley\(^1\) has referred to a complication that is liable to interfere with the cleaning of bronzes that contain much lead. The lead is distributed in the alloy in the form of globules, and as these are gradually dissolved by caustic soda, the bronzes become pitted. Also the electrolyte tends to build up a concentration of lead plumbite. Lead may then be replated on the objects as a grey coating. Usually this is easily brushed off, but an electrolyte contaminated with lead might well plate this metal on to other specimens treated subsequently, with undesirable results. He recommends that, when necessary, the lead be removed from the electrolyte by plating it on to a large sheet of copper attached as a temporary cathode to the cathode bar which normally carries the objects under treatment. The lead coating is easily stripped from the copper sheet later, by passing it through a bath of 10 per cent. nitric acid, and after washing in water it is ready to be used again should occasion require.

This serves to draw attention to the importance of keeping the electrolyte clean. Lead can easily be detected in the electrolyte by withdrawing a small sample in a test-tube and adding dilute sulphuric acid, until it is slightly acid. If a white precipitate of lead sulphate is formed, this is an indication that the electrolyte is contaminated with lead, in which case the lead may be removed by the auxiliary electrode method described above; alternatively a fresh lot of electrolyte may be made up to replace that which is contaminated.

**INTENSIVE WASHING TECHNIQUE**

When an incrustation has been broken down by reduction there will remain on the surface of the metal a sludge of insoluble oxides and metallic powder; this will also contain chlorides, rendered soluble by the treatment, as well as a residue of the electrolyte employed in the process. The sludge is removed by brushing the object under running water, and while this will remove most of the superficial soluble impurities at the same time, it is not by any means sufficient

to deal with all. It will be readily understood that when an incrustation has grown upon metal, the metal underlying the incrustation will be microporous, and when the incrustation is removed, the surface layers of the metal will behave as if they were a mass of capillary tubes, and these will retain a residue of chlorides. This corrosive matter is not easy to remove. It can be eliminated by prolonged soaking in changes of distilled water, or its removal may be speeded up by using hot water, but unless the capillaries can be flushed with the distilled water, there will always be the chance of chloride being retained in cracks or pits sufficient in concentration to cause a fresh outbreak of corrosion at a later date.

Such considerations have been examined theoretically by R. M. Organ,¹ and as a result of practical tests carried out under controlled conditions, using a conductivity meter, he has been able to recommend a form of intensive washing and also a particularly sensitive method of carrying out the silver nitrate test, i.e. the test that is used in practice to determine when objects are free from chloride. In this method it is recommended that the washing should be carried out with pure (distilled) water, by alternate heating and cooling, at least in the early stages, to ensure permeation. The heating causes any air or liquid in the capillaries to expand and partly leave the metal, and on cooling fresh water is sucked in. In dealing with a batch of bronzes, it is convenient to use a steam oven, in which are placed Pyrex beakers of distilled water containing the objects. The oven is kept at about 98° C. during the working day, the temperature falling overnight. This alternate heating and cooling ensures that the capillaries in the metal are flushed with water, and the progress of the washing is followed by testing the conductivity until it reaches a minimum.

The procedure described above is of wide application to metals and many siliceous materials, notably to baked cuneiform tablets, but this method is not suitable where leaden objects are concerned. Hot distilled water reacts in a few hours with lead to form lead hydroxide which would cover the metal with a milky white film. A special procedure is required for washing leaden objects (see p. 260).

¹ Organ, R. M., Museums Journal, 1955, 55, p. 112.
The standard test for chloride is carried out by adding silver nitrate solution in presence of nitric acid, when the formation of a white flocculent precipitate (of silver chloride) indicates that chloride is present in the solution that is being tested.

In the final stages of washing, when testing is of the greatest significance, it is essential to be able to detect the presence of the merest traces of chloride remaining in the water. Organ has emphasized the importance of applying the test under optimum conditions. The volume of soak-water should not be much in excess of that required to cover the object and the water should be in contact with the object for some time (until the maximum electrical conductivity of the water is attained). Then the test is carried out under the following standard conditions which give the most sensitive results. About 10 ml. of the water to be tested are poured into a tall narrow cylinder standing on a black surface. Several drops of dilute nitric acid are added (exact concentration immaterial), a stopper is applied, and the vessel inverted several times in order to mix the solutions. At this stage the liquid should be clear when examined in a good side lighting. Five drops of dilute silver nitrate (17 grammes dissolved in a litre of distilled water) are then added, mixed as before, and time allowed for any opalescence to appear. Under these conditions one part of chloride per million of solution may easily be detected in a column 10 cm. high.

Drying metals

Two methods are commonly employed for hastening the drying of metals—heating and desiccation. Heating in an oven at 105° C. is the simpler process, and is satisfactory for iron objects of all kinds. It is satisfactory also in the case of silver and copper alloys that are smooth and non-porous. Porous metal, however, takes longer to dry, and base silver and copper alloys that are at all porous may acquire a disfiguring film of oxide in the oven. Although oxide films may be removed by glass-brushing, this introduces another operation, and

1 The use of a glass or rubber stopper is preferable to using the finger, as chlorides from the skin would be liable to upset the results.
one that can be avoided if drying is done in a desiccator. Drying can be accelerated by passing an object through a bath of acetone and then placing it in a desiccator from which the air can be evacuated. A capacious vacuum desiccator is a very useful piece of apparatus to have in a conservation laboratory. It is charged with silica gel (see Appendix XI), and evacuated as required by means of a vacuum pump attached to the main water supply (see Fig. 5, p. 152). The vacuum desiccator is used not only to dry objects, but also to store them until such time as they can be lacquered or protected with wax.

**MECHANICAL METHODS**

When a metal object is showing signs of active corrosion, there can be no question that the only way of checking the activity and effecting a permanent cure is to employ chemicals, but this involves using certain ‘mechanical’ operations as well, in order to facilitate the action of the chemicals and their removal at the conclusion of the treatment. Mechanical methods may be employed directly for cleaning oxidized or tarnished metals, where it is merely a question of removing surface staining; they may also be employed in dealing with rust fragments that are completely oxidized, and no longer subject to chemical change.

Even with the most primitive workshop facilities it is possible to make a simple kit of tools that will be adequate for most purposes.

*Making needles and chisels*

The commonest tools are needles and small chisels, and the most satisfactory are those that one makes oneself. Needles may be mounted in wooden or metal handles, or in a small chuck or mandrel. They should be short and stiff and of varying thicknesses. An etcher’s needle, as sold for dry-point work, is a useful instrument for general purposes.

A series of chisels and scrapers can be made from old files that have been annealed in the fire, or from pieces of silver-steel rod of round, square, or rectangular section. The cutting edges are made first of all
by grinding, and then the tools are given heat-treatment to adjust the hardness prior to sharpening. The heat-treatment is conducted in two stages as follows:

**Hardening.** Heat about 1 inch length of the chisel at the sharp end to bright redness, avoiding over-heating, then immediately quench the metal by dipping the point in water and moving it about rapidly. This will make it dead-hard and brittle, and it will be dark in colour. Polish the metal on a fine emery paper until it is silvery white in order that the temper colours may be observed in the subsequent operation.

**Tempering.** To toughen the hardened metal, heat the polished chisel \(\frac{1}{2}\) inch or even farther from the cutting-edge—very gently and under strict control—and watch the temper colours moving towards the cutting-edge in the sequence—yellow, straw, purple. Aim to arrest operations by throwing the tool into cold water when a middle-straw colour extends from the cutting-edge some distance up the chisel. If the colour of the cutting-edge has gone to the purple stage, the chisel has been overheated, and the hardening and tempering operations must be done again.\(^1\)

Finally, clean and sharpen on a carborundum slip or oilstone.

Mechanical methods may be defined as picking, chipping and scraping, grinding, cutting, brushing, shot-blasting, polishing, and burnishing.\(^2\) These will now be considered *seriatim*:

**Picking**

Picking is a technique apparently so commonplace as to seem unworthy of further description, but there is much more to it than would at first appear. It is not often realized that a force of \(1\) lb. exerted by means of the point of a needle \(1/5000\) inch in diameter is equivalent to a pressure of several tons to the square inch.

---

\(^1\) A pale yellow colour represents razor hardness and the metal is brittle; a blue or purple colour implies flexibility as required for springs and hack-saws. Intermediate straw or brown colours are an indication that the steel will be tough enough for use as a chisel and will keep its cutting-edge.

A needle is thus a very potent tool and the correct method of handling it is shown in the diagram (Fig. 7). If pressed near the edge of a brittle surface, the edge will chip off, and the nearer to the edge it is pressed, the less the pressure required to remove tiny flakes from the rust escarpment. Considerable masses of rust can be eliminated, grain by grain, by using a needle in this manner; this is a labour of patience and the operator must be content to advance very slowly, using the minimum of pressure. Any attempt to hasten proceedings by pressing the needle too far from the edge, places unnecessary strain on the object and may end in disaster. The needle may be used on occasion as a tiny lever or even as a scraper.

**Chipping and scraping**

When metallic excrescences have to be removed this can be done by using a small chisel and jeweller’s hammer, or a chisel mounted for use as an engraving tool. Mechanically operated needles and micro-chisels are available, but, owing to their vibration, these are only useful for heavier work.

When the object is very thin and brittle, chisels are not employed as they are liable to fracture the specimen: it is safer in such a case to use the needle or fine engraving tool, but dental burrs may at times be useful to help to reduce thick lumps of oxide.

When incrustations have been softened by reduction, it is useful to be able to remove them in the wet condition, and in cases where it would be unsafe to use metal tools, lest they should cut into the
surface, a sharpened stick may be employed, cut to a chisel edge. Cane tools are also useful as scrapers, and these may be used to rub the surface of an object undergoing treatment with zinc and caustic soda, in order to observe the progress of the reduction.

**Grinding**

When excrescences are exceptionally hard, it may be preferable to grind them away. For this purpose a small wheel or inverted cone may be used on a dental drilling-machine; alternatively, the grinding may be done with slips of carborundum which can be obtained in various shapes and degrees of roughness. Grinding, however, must not be lightly undertaken, as it can destroy markings in the rust—textile marks, wood grain marks, &c., which may be of significance. On the rusted blade of a sword, for example, they may indicate that a scabbard had been present and decayed *in situ*; or markings on the tang of the weapon may be all that remains to show how the hilt was constructed. It is also necessary to add a warning that grinding may give iron oxide a dull metallic polish so like that of metallic iron itself that it becomes difficult to distinguish between the two.

**Cutting**

Cutting by using a hack-saw is only employed as a means of separating two objects that are corroded together into a solid mass, and which could not otherwise be separated. Such a mass may be cut as a preliminary to chipping and picking. Slit-saws are sometimes useful, or fret-saws, or even a copper wire hardened by stretching, fixed in a frame and fed with carborundum powder in oil. The quillon of a sword that had slipped along the tang in antiquity and was rusted firmly into position in the wrong place was released by working a small hole through the rust between quillon and tang, inserting the copper wire and 'sawing' a slit through the rust along the tang. The wire was found to be much more effective for the purpose than the finest fret-saw since it was more flexible.

**Brushing**

Brushing is a frequent accompaniment of all other mechanical
processes, its object being to eliminate foreign matter and reveal the progress of the work. Where a good solid iron core remains and objects can be dealt with chemically, occasional brushing in a stream of water is essential to eliminate the by-products of the reaction, which tend to adhere to the specimen in the form of a sludge. Different kinds of brushes can be selected to suit the type of work in hand. Hard nylon tooth-brushes are invaluable for use on bronzes after reduction. If a steel brush is used, it must be dried thereafter or it will rust. A motor-driven brush can lead to a great saving of time when objects are strong enough to stand this treatment. When wire brushes of appropriate coarseness are used, either hand or motor-driven, they will help to scrape away loose rust, but care must be taken that the bristles do not penetrate too deeply into softer parts of the incrustation. Rotating wire brushes, if soft enough,\(^1\) smear the products of reduction over the metal and make the surface appearance more uniform. In some cases it is safer to use graded steel wool rather than brushes. The finest grades are very gentle in action, especially when used with a little oil. Glass brushes are used for fine work such as jewellery or for polishing silver inlays in iron—they are indispensable in the laboratory. For removing hard rust from soft metal inlays, there is no more effective tool than a carborundum pencil, or a slip of this abrasive that has been rubbed down to a point. It enables the rust to be pushed off without rubbing the surface of the inlay.

**Shot-blasting**

One of the most effective methods of dry-cleaning is to use a jet of particles of fine grit blown from a modified form of spray gun, as in the shot-blast cabinet. Several types of cabinet are available commercially. Fine work may be cleaned in the dental cabinet, using a spray of bauxite grit, and at the other extreme, heavy metal castings are dealt with by using various grades of powdered cast iron, but bauxite is suitable for most museum purposes. It is possible by varying the grade of bauxite, the pressure, the distance of the nozzle from

---

\(^1\) A warning is necessary against the indiscriminate use of brass brushes as they tend to impart a yellow brass colour to other metals.
the object, the angle of attack, &c., to carry out a variety of cleaning operations quickly and efficiently. This type of cleaning is never used where soft metals are present as it would give them a matt surface. It is useless, therefore, for gold, silver, and lead alloys, but it can be very useful for the removal of mud and soft corrosion products from bronze and iron. Indeed it can safely be employed for the removal of siliceous deposits and dirt from certain kinds of smooth bronze patina without harming the surface.

Shot-blasting has also proved to be the best means of removing surface incrustations of various kinds from baked clay tablets (p. 321).

Polishing and burnishing

Polishing is done with the finer grades of abrasives such as emery-flour, rouge, diamantine. These are used either as powders or made up with tallow in paste form. Special cloths impregnated with such materials can also be obtained. Abrasives are available in emulsified form as metal polishes, or compounded with paraffin and cotton-wool as polishing wools. The degree of abrasion depends not only on the size of the particles, but on the nature of the abrasive—carborundum powder, for example, being harder and having a greater cutting action than pumice powder of the same degree of fineness. Levigated alumina is a finishing abrasive by means of which a high polish can be obtained on metal that has been previously rubbed up with a series of coarser grits. The final effect of brilliance can be obtained on steel only by burnishing to eliminate all porosity from the surface. This is done either by using a cloth wheel on a buffing-machine or, in the case of fine work, a hard polishing-stone, e.g. haematite, the so-called bloodstone. A chain burnisher is very seldom used, and then only for coarse work in exceptional cases. Burnishing and polishing by abrasives, although of great importance in cleaning jewellery, or in removing rust spots from otherwise bright armour and accoutrements, are, however, techniques of limited application. In cases where the rust is not of a spotty nature, but present as a continuous layer of fine dust on a coat of mail, for example, it should be removed as far as possible with a stiff brush, if the metal is strong
enough to stand this. No attempt should be made to attain a bright finish by burnishing with an abrasive, as the links will almost certainly have been weakened by rusting. In such cases sufficient protection is given, after cleaning, by the application of a thin film of wax, which, when brushed, leaves a dull shine of satisfactory appearance that acts as a foil for any embellishments. This type of treatment and finish, by brushing, waxing, and polishing, has been applied with considerable effect in cleaning sword guards and t'suba where the steel is covered with fine ornament and often inlaid with gold and silver.

The most perfect examples of the armourers’ craft in existence are, without question, Japanese swords, and if these become spotted with rust, the use of tools or of etching or abrasive materials in an effort to remove the rust does more harm than good. All that can be done is to try to minimize the disfigurement caused by the dark spots of rust on the mirror-like surface of the steel by alternate applications of kerosene and lanolin, and by rubbing with silk or a Selvyt cloth. The surface may be protected thereafter with a thin film of a neutral microcrystalline wax (see Appendix XII). Care is necessary to avoid possible accident, as the swords are often exceedingly sharp.

Summary

The general methods of treatment that are applied in dealing with corroded objects have been described, namely, reduction methods (electro-chemical and electrolytic), mechanical methods, washing, testing for chloride, and drying. The action of chemical solvents in arresting corrosion and cleaning and restoring metal objects is of equal importance as a general method of treatment, but, as solvents are selective in action, they are dealt with in the specialized chapters that follow.
CHAPTER IX

GOLD AND ELECTRUM

Gold is found in nature in the metallic condition as a rich yellow, soft metal, commonly associated with quartz and certain sands. It does not corrode, it is not dissolved under natural conditions, but it is often found alloyed with the baser metals silver and copper. The presence of silver in gold gives pale-coloured alloys in which the degree of paleness is in proportion to the amount of silver present, and such alloys have been known since the time of Pliny as 'electrum'. Both gold and electrum have been used for coinage and jewellery from the earliest times.

When gold is found alloyed with copper, there is usually some silver present as well, and although such alloys may be very pale yellow or even greenish yellow when freshly cast or cleaned, they lose their pallor after years of burial in the ground and acquire a warm yellow hue. This is known as surface enrichment and is due to the action of salts in removing the baser metals from the surface of the alloy, leaving a film of pure gold.

It is usually desirable to preserve the rich colour, even though it may only be a surface enrichment, and in cleaning, care should be taken not to rub the surface because its quality is easily lost. Incrustations of lime can be removed from the gold by applying dilute nitric acid (1 per cent.) locally with a match-stick, a small brush, or a capillary glass tube. Siliceous or muddy incrustations on gold may be removed by soaking in a 2 per cent. solution of a surface-active agent such as Lissapol. Organic remains are softened by immersion for a few minutes in caustic soda (2 per cent.) and removed with a match-stick. If incrustations resist these attentions, they may sometimes be disintegrated by using a fine jet of steam.
Gold is sometimes found having a clear purplish red or rose-pink colour. In the eighteenth to twentieth dynasties of Egypt, a striking rose-pink colour was imparted to gold used for personal ornaments, horse-trappings, and sequins. On opening a box of royal robes in Tut-ankhamûn’s tomb, for example, it was observed that the garments had been decorated with sequins in a variety of shapes and, although the textiles had perished, it was clear that in one instance at least a garment had borne a stitched panel of massed sequins regularly positioned, rose-pink alternating with yellow gold, to give a diaper effect, thus proving that the pink colour was not an effect of age. A pink sequin from the Tut-ankhamûn robe was analysed and found to contain in addition to silver and some copper, 0.85 per cent. of iron which was the colouring constituent. The pink colour was confined to the surface of the sequin back and front.

Attempts to reproduce this coloured alloy were successfully carried out by R. W. Wood,¹ who fused gold with iron and obtained an alloy having the same colour and micro-structure as the sequins. If this were the technique applied in the early dynasties of Egypt, it would be necessary to suppose that the iron was introduced in the form of a mineral from which it could be released when in the crucible, as the melting-point of iron is 1,535° C. An independent examination of the sequin problem made at the British Museum Laboratory has shown that when gold containing silver and copper is fused with iron pyrites and soda, some of the silver and copper combines with the sulphur in the pyrites and rises to the surface of the molten gold as a dross, leaving the gold alloyed with iron. When this gold is hammered into sheets and given heat treatment, it develops a superficial colour characteristic of the Egyptian sequins. Iron pyrites (M.P. 1171° C.) has about the same melting-point as pure gold (M.P. 1063° C.) and often has the yellow metallic appearance of gold, for which it might easily be mistaken; indeed it has been called ‘fool’s gold’.

If by chance some of this material were inadvertently mixed with gold scrap for the crucible, it would be possible for the rose-pink colour to be developed accidentally. If such were the case, it

29. Roman beaker of dark blue glass blown into a perforated silver framework (1st cent. A.D.)

A. Before treatment—corroded coins fused together as the result of a fire

B. After separation in citric acid and cleaning by reduction

30. HOARD OF ENGLISH SILVER PENNIES (EDWARD I AND EDWARD II)
would explain why this particular technique appeared sporadically in these dynasties and was later forgotten.

Gold coins sometimes have what appears to be a variety of this same rich red patina, but here it is an acquired characteristic, the colour having developed by contamination and possibly heating during the life of the coins. At its best this adventitious red can be very handsome, and is regarded by collectors of coins as a valuable patina worth preserving, but it is easily rubbed off and when once lost it is virtually impossible to restore the rich colour by artificial means.

Although gold is so durable it is a very soft metal and can be beaten out to form the thin leaves used for manuscript illumination and for applying to gesso. It can, moreover, be dissolved in mercury fairly easily to give gold amalgam. This substance has the consistency of butter and when rubbed over clean silver or copper, it adheres. The mercury may be readily volatilized by heating and the silver or copper thus becomes covered with a continuous layer of gold. The process is known as ‘mercury gilding’ or ‘fire gilding’.

It was interesting to find both techniques used together on the dragon heads that ornamented the periphery of the shield from Sutton Hoo (A.D. 650) now in the British Museum. The original heads were executed in bronze that had been gilt by the mercury gilding process, but certain repairs had been made in antiquity and some of the heads had been replaced by models in gesso covered with gold leaf. The same type of repair was also evident on the large dragon shown in Pl. 23.

Gold is likely to be found as decoration on armour and accoutrements, on buckles, on bronze statuettes, especially in the eyes of the figures, and on silver dishes, in which case it may be limited to the embellishment of reserves of ornament. Silver so decorated is said to be ‘parcel gilt’. This type of decoration was exploited in Persia during the Seljuk period for the embellishment of horse-trappings, rose-water sprinklers, incense-burners, vases, dishes, and caskets. When gold is present in reserve ornament on silver, special care is required in cleaning because the gold is thin and the surface soft,
and the gold reserves could be easily damaged by the cleaning materials used to remove tarnish from the surrounding silver.

CLEANING GILT METAL

Metals that are gilt should not be reduced because of the danger of losing the gilding, but if the gold overlies solid copper or bronze, it is sometimes safe to apply the Rochelle salt process (see p. 239) for cleaning the base metal. If the base metal is encrusted with corrosion products, however, and it is known that a film of gold exists beneath or among them, the only possible line to take is that of mechanical cleaning, using needles under a binocular microscope at a magnification of some ten diameters. Very dilute nitric acid (1 per cent.) can be used to clean the surface of the gold when it is exposed to view, but acid should not be used to soften the base incrustation, otherwise the ornament may become detached. To reveal the gold can be a long and arduous task, but the results are rewarding if the shape and decorative appearance can be recovered.

When gilding is present on bronze statuary that is exposed to soot deposits and accumulations of dust, it requires occasional cleaning, and as dirt of this nature is usually acidic, it responds readily to washing with dilute ammonia. The statuary should be well washed with water thereafter. This simple type of cleaning was applied to the royal effigies in Westminster Abbey, and the results were very satisfactory.

CRUSHED GOLD OBJECTS

Objects of gold are often found in a distorted condition when they are dug from the ground, and there may be a strong temptation to try to fold the metal back into shape. This must be resisted as it is not the easy task that it appears to be. Pure gold is readily malleable, but when it is alloyed, as it usually is to some extent, the metal becomes brittle on ageing, and mechanical manipulation should only be undertaken after scientific examination has shown that the metal is of such a composition that reshaping can be considered a reasonably safe procedure in the hands of an expert.
GOLD AND ELECTRUM

Many examples of the restoration of crushed golden objects, by careful annealing and tooling back to shape, are provided by the material excavated at Ur of the Chaldees by Sir Leonard Woolley. One of the finest specimens is the wig-helmet of Mes-Kalam-Shar, made from a single plate of 15 carat gold, and weighing 2 lb. 4½ oz. A more intricate reconstruction was that of the gold bull’s head forming the frontal ornament of a lyre (Pls. 24 A and B). This was brought to the Laboratory as a complex, held together by wax and bandages, so that the relative position of the various pieces could be studied at leisure. The gold mask, ears, and horns were first recovered, and as the gold, though badly crushed, was much thinner than that of the helmet, it was correspondingly easier to manipulate. The original shape of the animal’s head was re-established, and having done this it was possible to reset the eyes and reassemble the various pieces of carved lapis lazuli forming the beard.

The special tools used for such work are metal and wooden stakes, levers of various kinds, mallets of box-wood, hide and horn, and shaped sand-bags of leather.¹

¹ Maryon, H., op. cit.
CHAPTER X

SILVER

Silver is a soft white lustrous metal which fuses at about 960° C. It is found locally in the metallic condition (as native silver), but is more generally distributed in mineral form, two of the commonest minerals being argentite (silver sulphide), and cerargyrite (silver chloride), commonly known as 'horn silver'. The sulphide is black in colour, and the chloride a dirty white or slate grey. The pure metal is malleable and ductile, and is capable of taking a high polish, and for this reason has always attracted the craftsman. It has enjoyed a deserved popularity in most countries of the world for objects of personal adornment and decorative metal-work. Gold and silver have been used for coinage throughout the ages, and for this reason the ratio of values has always been a significant figure, the relationship fluctuating with the scarcity of one or other metal. Partington\(^1\) quotes a number of examples illustrating how the relative values changed at different periods and in different localities, observing that in the earliest periods of civilization gold was the commoner metal, because it could be obtained from rivers, whereas silver had to be obtained from the hills where it occurred, not as a surface deposit, but buried, often deeply, in the ground.

Both gold and silver were in general use at Ur of the Chaldees in the Early Sumerian Period, at a time when silver was a rarity in Egypt. Indeed, silver was never available there in any quantity until the Graeco-Roman period, and it is not surprising, therefore, that the robbers who entered Tut-ankhamûn's tomb were found to have concentrated first upon the silver.

SILVER

TARNISH

On exposure to city atmospheres, silver is readily tarnished by the formation of a thin surface film of silver sulphide, and for this reason polished silver requires regular cleaning. The sulphur which tarnishes the silver may be carried in the air in the form of sulphuretted hydrogen from flue gases either industrial or domestic, its action being particularly noticeable in foggy weather. There are other, perhaps not so obvious, sources of sulphur contamination. It may be impossible to keep silver bright in places where there are vulcanized rubber floor-coverings or where a case has a rubber seal of the draught-excluder type. Cheap paints may contain volatile sulphur impurities that will tarnish silver, and even certain types of dry paint film may be potential sources of trouble. Casein is a common constituent of water paints and this is subject to bacterial action that releases sulphur compounds that will tarnish silver. Emulsion paints should not be used to decorate cases that are designed for the exhibition of polished silver unless they can be guaranteed as free from casein. Paints based on lithopone are also suspect. The safest paints to employ are either cellulose paints or good-quality oil-paints based on titanium white. These should contain enough lead-driers to act as a sulphur barrier and prevent any sulphur compounds that may be present in the oil film contaminating the air in the case. A more insidious source of trouble is contact of the silver with textiles that have been ‘finished’ by treatment with chemicals containing sulphur. Such textiles may contain as much as 200 parts per million of reducible sulphur, whereas all that is required to cause visible discoloration is 2 parts per million.

Tarnish is easily removed with plate powder or rouge cloths, but for antique silver which requires regular cleaning such abrasives are not recommended as they would tend in time to cause damage by wearing down stamp marks and fine ornament. The same applies to Sheffield plate or silver-plated ware, where drastic cleaning or the use of chemicals might result in exposing the underlying base metal. In these cases it is safer to confine routine cleaning to rubbing with a soft cloth and a little French chalk, moistened to a paste, if desired, with methylated spirit containing a few drops of ammonia.
For silver that has been neglected and is heavily tarnished, the following simple electro-chemical method of treatment may be used. Place the silver in contact with aluminium, cover it with a dilute solution of caustic soda or sodium carbonate (washing soda)—a solution of 5 per cent. strength is ample—and allow it to remain until the stain has disappeared. It can then be washed in running water and polished with a soft cloth or cotton-wool. Alternatively a chemical solvent for tarnish is available called Silver Dip,¹ and this gives excellent results provided the silver is not left in the solution longer than necessary and is washed thoroughly afterwards.

Tarnishing may be prevented by the exclusion of sulphur in any form. This is easier to achieve with silver in storage than when silver is on exhibition, as protection may be afforded by wrapping in several layers of soft tissue paper, and if the silver has been polished and washed beforehand it will be likely to remain bright for a long time. Alternatively, the silver may be kept in Polythene bags having an air-tight seal. A recent advance has been the introduction of an anti-tarnish paper—a soft velvety paper that has been impregnated with chemical substances (copper compounds, chlorophyll, &c.) so that it will absorb hydrogen sulphide from the air, thus leaving correspondingly less sulphur to attack the silver. Anti-tarnish papers should not be in actual contact with the silver or there may be staining by transfer of the sulphur, but as external wrapping papers they cannot be other than advantageous.

Silver should be exhibited in a dust-free case, provided with a breathing channel to allow for the expansion and contraction of the air that occurs with change of temperature. This channel which is usually filled with cotton-wool should contain in addition a loose plug of anti-tarnish paper. The efficiency of such papers when placed actually inside a vitrine has been found to vary, however, because they tend to become easily desiccated. Actually they are more effective when freely exposed to the varying temperature and humidity of the atmosphere; nevertheless, when a bulk supply of anti-tarnish paper

¹ Design and Research Centre for the Gold, Silver and Jewellery Industries, Bull. 15, 1953.
can be included in a closed case, it reduces the rate of tarnishing even under dry conditions. This has been established in a series of tests in which the protected silver was found to remain bright for up to a period of several months after unprotected silver had visibly blackened.

Where silver is displayed in a closed glass case on shelving draped with Shantung silk that has been passed through a 10 per cent. solution of lead acetate, dried and ironed, it may remain bright for many years, as demonstrated by Bengt Bengtson in the Nordic Museum, Stockholm.

With certain designs of vitrine it is possible to arrange a tubular filter-vent, through which sulphur-free air can be introduced by blowing from outside to replace the contaminated air in the case. The joints of the vitrine are then sealed with Polythene tape. A suitable blower can be improvised from a vacuum cleaner, the bag being replaced by a filter tube charged with material that will absorb sulphuretted hydrogen, e.g. pumice granules that have been previously treated with copper or lead acetate and glycerol.

It will often be more convenient to protect silver objects by a thin coating of impervious lacquer and if this is applied with care it is practically invisible (p. 229).

PATINA AND CORROSION

In silver objects there is generally a clear-cut distinction between surface deposits that are disfiguring and those that are thin and stable and which improve the appearance. Tarnish is a disfigurement, but one occasionally comes across specimens of ancient silver having a thin, uniform, shiny black patina of silver sulphide, and this is worth retaining for its decorative quality. It has not been possible by laboratory methods to reproduce such a black surface, which is probably to some extent an effect of ageing.

A thin film of silver chloride can itself form the basis of an agree-

1 This method was first suggested by Professor J. Tandberg; the lead acetate solution should contain 2 per cent. of glycerol and about 1 per cent. of ammonia.
able patina, as in the case of silver objects excavated from soils containing little chloride. T'ang silver from the loess has sometimes a beautiful patina of a chalky texture and pale lilac colour, stained pink and green with cuprous oxide and malachite respectively. As this patina is stable and enhances the appearance of the object, it is desirable to preserve it. If the patina is concealing fine ornament, a compromise is necessary, however, and it may then be permissible to expose the ornament by rubbing with a mild abrasive. The impregnated cloths that are sold for polishing metals are suitable for this purpose (Dura-glitz, &c.) provided the appropriate grade of cloth is used, i.e. one for cleaning silver: those sold for cleaning copper and brass are too drastic in action and would abrade the metal unnecessarily.

When chloride is present but sulphide preponderates, the silver is dark in colour and has usually a stained and corroded appearance. It may sometimes be improved by applying selective solvents (p. 219) to remove the staining, and by picking or brushing as required to release any incrustation. Such treatment will restore the natural whiteness of the metal, but care must be taken that ornamental details are not sacrificed in the removal of the incrustation. It is better to take no action than to risk damaging the object; specimens in this condition, although unsightly, are generally stable and treatment is only warranted if it improves the general appearance and reveals ornament more distinctly. Corrosion by sulphide formation may sometimes be found to have proceeded to the limit, as in the case of a Persian silver anklet examined by the author. In such cases the metallic quality is entirely lost, the material is black and brittle and has the appearance of old leather. In this condition the silver is beyond restoration.

When silver has been buried for long periods in salty soil, as in the case of the silver vessels from the Ziggurat at Ur, it is largely transformed on the exposed surfaces into stable silver chloride (horn silver), which has a grey earthy appearance of no aesthetic value. Deformation may be considerable, owing to the great expansion accompanying the change from metal to mineral, but as it is stable, treatment is only required to improve the appearance of the specimen, to expose details of ornament, or to recover an elegant shape. Horn
silver is tough, and the object is robust so long as the incrustation of chloride is in position, but if removed or reduced, it should be borne in mind that the vessel may be much weakened. Reduction tends to give a discontinuous surface film, lacking in coherence and rigidity. Another factor of importance is that stratification is a feature of the incrustation resulting from long-continued corrosion, and it is likely that some of the segregated constituents are in the form of discontinuous strata. This is clearly seen in the sectioned silver bead from Ur (Pl. 25), where, in addition, the residual uncorroded metal in the heart of the object is of varying thickness and porosity. Accordingly, when an object of high silver content is so heavily corroded that little silver remains (as, for example, in the case of a thin metal jug or bowl), treatment, if applied at all, should be carried out with a view to exposing surface features and removing irregularities. Electro-chemical reduction would be likely to be too drastic in its effects. On the other hand, there may be every justification for reducing a massive silver object, where it is clear that a solid core of silver remains to ensure the mechanical strength of the specimen after treatment. The strength of such a silver core is, however, not always easily assessed—old silver is invariably much more brittle than silver that has been recently cast. The brittleness is caused by a change in the microcrystalline structure with the passage of time, a change that takes place irrespective of whether the metal is superficially corroded or has remained bright and free from chemical attack.

TOUGHENING FRAIL SILVER

Brittle silver may be toughened either by heating to a dull red heat, or by maintaining it at a somewhat lower temperature for some time. The operation requires dexterity and experience, because silver melts at about 960° C. and there may be impurities present that lower the melting-point of the metal considerably. The metal begins to glow at about 600° C. If there are any mineral compounds of silver and copper present, such a temperature cannot be attained without fusion. Silver chloride, for example, melts at 455° C. and
cuprous chloride (nantokite) at 422° C. Where doubt exists, or where the silver is very thin, the minimum of heat should be used. Toughening actually begins to take place at about 250° C., and if the silver is placed in an oven, the temperature of which rises in about two hours to 400° C., this should suffice for most purposes.

In the case of sound metal, free from salts, the silver may be heated in an electric furnace, the temperature of which is adjusted to 600–650° C. The best results are obtained in the electric furnace: if a gas furnace is used, it is difficult to prevent the silver from blackening unless all fumes can be excluded during heating. When silver is debased with copper, heating will cause the formation of a black surface film of copper oxide. Staining due to oxidation is, however, easily removed in a bath of sulphuric acid (5 per cent.).

**CRUSHED SILVER OBJECTS**

In the case of silver that has been crushed out of shape, heat treatment is always applied before any attempt is made to bend the metal. When once toughened the metal may be worked back gradually to shape with the fingers, or with wooden tools covered with chamois, a process requiring much patience because the softened metal hardens again on cold working, and repeated heat treatment may be required as the work proceeds. When dents have to be removed from modern silver, there is more latitude, and it is usually possible to rub out small deformities, without preheating, by the use of shaped tools manufactured for the purpose. The most useful are bloodstone burnishers mounted in brass sockets on stout wooden handles. There is no question of using these on old silver, however, until it has been previously toughened.

The largest of the silver salvers from the Sutton Hoo Ship Burial was excavated in a crushed and very brittle condition. It was toughened by heating it repeatedly with a large blowpipe flame and restored to shape with the fingers, and later with the help of a hide mallet. When a free flame is used in this way heating should be carried out in a darkened room, as it is impossible otherwise to detect
when the metal begins to glow, and part of the object might easily be inadvertently over-heated.

**Copper Incrustations on Silver**

Reference has already been made (p. 189) to the cathodic protection of silver in the presence of corroding copper. A common result of such action is the formation of a heavy deposit of copper corrosion products on the silver that is itself uncorroded, and this may be difficult to remove. The usual procedure is to get rid of as much as possible of the incrustation by mechanical methods (picking, grinding, &c.) first of all, and then to endeavour to eliminate the rest by the action of selective solvents. Some examples will now be given to illustrate the treatment of silver objects by the application of solvents, selected because of the specific properties they possess.

(i) *Ammonia*. In the presence of air ammonia dissolves copper and it may be claimed that no reagent is more effective for cleaning silver that has become encrusted with a massive deposit of copper by long contact with the baser metal. Ammonia, however, must be used with restraint where silver chloride is present, because it dissolves this substance and, if applied to corroded silver, would leave the metal in a much weakened condition. It softens organic remains such as residues of wood, skin, textiles, &c., and is thus a general cleaning-agent where such types of incrustation are concerned. It is used in concentrated form (density 0.88), preferably in a fume cupboard.

(ii) *Potassium cyanide*. This is highly poisonous and seldom required, but it is a very potent stain remover. It is not applied in the form of a solution and it should on no account be used where acid is present or be allowed to come in contact with the skin. For removing stains from silver, a small pellet of cyanide is held with forceps and rubbed on the wet silver, and in dissolving a little of the silver it will at the same time remove most stains.

(iii) *Formic acid*. This has the particular merit that it may be used to dissolve copper compounds without affecting silver or silver chloride.

(iv) *Silver nitrate*. For removing a thin film of metallic copper
from silver, the use of a solution of silver nitrate is preferable (see p. 224).

In certain circumstances it may be necessary to use more than one solvent. Thus, for example, in the case of the silver lamp (Pls. 22A and B) the bulk of the copper incrustation was cut away but a resis-
tant layer of metallic copper remained firmly adhering to part of the frail silver. The removal of this necessitated prolonged treatment in ammonia followed by local treatment with potassium cyanide. After all traces of the incrustation had been removed the lamp was dark in colour; it was given a bath in warm formic acid (about 30 per cent. strength) for a short time (twenty minutes) to brighten the metal and, after washing and drying, the silver was toughened in the furnace as described above.

A striking illustration of the use of formic acid was in the treat-
ment given to a silver cup from Enkomi, Cyprus (fifteenth cen-
tury B.C.), which arrived at the Laboratory covered with a green incrustation. This incrustation had originated partly from the copper constituent of the silver alloy and partly no doubt from contact with base metal in the ground, and it gave the impression that the cup was made of bronze (Pl. 27A), concealing the fact that it was actually of silver elaborately decorated with niello and gold. The true nature of the object having been discovered by X-ray examination, formic acid was selected as the cleaning agent, and after a successful experiment on one of the fragments, the cup was immersed for twenty minutes in a boiling solution composed of one volume of commercial formic acid and two volumes of water. By such treatment the green incrustation was removed, leaving patches of cuprite and re-
duced copper which yielded eventually to local applications of ammonia (0.88). The ornament around the cup was revealed in surprising freshness, the niello for the most part being in excellent condition and the gold inlays rich in colour without showing any evidence of blanching (Pl. 27B). The silver proved on analysis to be alloyed with 3.4 per cent. of gold and 9 per cent. of copper.

Niello, a black decoration, has been used for ornamenting silver

from very early times. Pliny\textsuperscript{1} refers to the technique having been employed in Egypt, though few examples have survived. There are, however, black decorations still extant on the dagger-blades from Mycenae dating to 1500 B.C. and the niello technique is well established in Roman, Byzantine, and Anglo-Saxon art. Examples of black inlays from many sources have been subjected to laboratory examination by Moss\textsuperscript{2} who has developed micro-tests for their identification and shown that prior to the eleventh century niello corresponded with the mineral acanthite (silver sulphide), whereas subsequently a fusible mixture of silver and copper sulphides was used that corresponded to the mineral stromeyerite. Lead sulphide (galena) is sometimes present in addition.

Niello ornament might be difficult to discern on heavily tarnished silver, or might even be eclipsed by the spread of incrustation; it is necessary, therefore, to add a warning that the application of any electro-chemical process would reduce the black surface of the niello to metal, and thus destroy the decoration. Should doubt exist, electro-chemical methods are best avoided; but if the reduction has been brief, the black colour of the niello can be exposed again by the judicious use of metal polish, the reduction effect being purely superficial owing to the high electrical resistance of the niello relative to that of the silver.

(v) Ammonium thiosulphate and thiourea. A useful process for the removal of thick horn silver (silver chloride) is to glass-brush\textsuperscript{3} the surface with a 15 per cent. aqueous solution of ammonium thiosulphate containing 1 per cent. of Lissapol. The thiosulphate may be replaced by 5 per cent. thiourea although this requires to be rubbed more vigorously and is best reserved for cast silver such as buckles and ornaments that are rigid enough to withstand the mechanical stresses involved. The thiourea method is, however, very mild in its chemical action and can be perfectly controlled. It can be used where

\textsuperscript{1} Natural History, Book xxxiii. 131.


\textsuperscript{3} If the brush is provided with a metal holder, it should be washed frequently otherwise the holder will soon become corroded.
niello decoration is retained in horn silver, and for this reason alone it forms a very useful addition to the methods available for conservation. The silver buckle (Pl. 28) responded well to treatment by the thiourea process.

(vi) Citric acid. This is a convenient cleaning-agent for corroded copper, and hence for base silver covered with green corrosion compounds. It has the merits of being non-poisonous and odourless, but it must be used with a certain reserve as it is a non-selective stripping agent, and even in 5 per cent. aqueous solution may tend to over-clean. It is used as a pickle for separating objects such as base silver or copper coins that have become corroded together (see below).

ELECTRO-CHEMICAL REDUCTION OF SILVER OBJECTS

Electro-chemical methods may be applied to remove corrosion from silver within the limits already described in this chapter (p. 217). Besides the aluminium and soda method referred to for removing tarnish (p. 214), a number of other methods are available that give satisfactory results in dealing with corroded silver. For example, it may be that in using 30 per cent. formic acid as a solvent the cleaning has not been entirely effective; in this event, zinc or aluminium powder may be added and the mixture heated for a time. An alternative to the formic acid and zinc is to use zinc and caustic soda, which has a more pronounced softening effect on the corrosion. The horn silver becomes slimy after a time, and the object considerably weakened; brushing during reduction must be gentle and frequent, especially towards the latter stages. The reduced silver chloride incrustation takes the form of a white spongy deposit which is easily removed, and the residue can be consolidated by burnishing with a glass rod to restore the metal-like quality. After electro-chemical reduction the specimen may be hung in an electrolytic tank for ten minutes or so and boiled afterwards in several changes of distilled water. The alternative is to allow it to remain under running hot water for several hours, but in any event the washing is completed in several changes of distilled water until all chlorides are removed. The intensive method of washing (p. 197) can also be applied to silver.
After treatment, the silver is dried in an oven at 105° C. and it should be kept there for some hours, as the surface of reduced silver tends to be porous. It may sometimes be preferable to dry in a vacuum desiccator (see p. 200), as oven-drying tends to produce a film of tarnish. Where corrosion has been heavy and deep-seated it may be unwise to attempt to toughen the silver as the necessary high temperatures would tend to cause distortion of the metal.

**ELECTROLYTIC REDUCTION OF SILVER OBJECTS**

Silver may be reduced in the electrolytic tank in the usual manner using caustic soda as the electrolyte and iron anodes.

A variation, particularly to be recommended in the case of this metal, is to substitute formic acid of 15 per cent. strength for the soda electrolyte, this being the concentration giving highest conductivity. This method was very useful in dealing with the following rather unusual specimen. The silver casing that was attached to a Greek vase of blue glass was badly corroded, and as the vase was of such a shape that the silver could not be removed for treatment it was necessary to select a method that would clean the silver without harming the glass. Choice fell upon electrolytic reduction in 15 per cent. formic acid, using anodes of stainless steel, and the result was considered to be entirely satisfactory (Pl. 29). A caustic soda electrolyte could not have been used here as it would have been likely to attack the glass.

Stainless steel anodes are not an essential in using formic acid—indeed prolonged electrolysis in formic acid leads to the steel being attacked and there may then be a deposition of iron on the silver. With iron anodes in formic acid this deposition would take place even more quickly. There are no such complications when using anodes of graphite or carbon with formic acid. Good results are obtained, for example, using carbon anodes with an e.m.f. of 12 volts and a current density of 1 amp. per sq. dm.

During electrolysis of a base silver object the copper minerals in the incrustation are reduced to metallic copper, and when using the formic acid electrolyte there is a tendency for a film of copper to
be left on the object. This film is removed (after heating in an oven to 110° C. to eliminate formic acid) by immersing the specimen in 20 per cent. silver nitrate made up in distilled water. By this means the film of copper is replaced by silver powder which is easily brushed off.

CLEANING SILVER COINS

Old silver coins that have been found in the ground provide examples of decay which will vary according to the quality of the alloy and the nature of the soil. Identification of the coins is a primary requirement, and treatment is carried out with a view to making the ornament distinct and the inscription, if any, readable. It is sometimes the case, however, that an inscription is more legible in the corroded state, and for this reason it should be studied before treatment. After such preliminary study, a hoard of coins is assorted into groups according to the different forms of treatment required. In some cases corrosion may have proceeded in such a manner that the surface of the coin is black and wax-like, and very little metal remains. This is characteristic of a sulphide incrustation, and such coins had best be treated only by washing and drying. When the surface is cracked or the coin is very porous it may be reinforced by impregnation with a nitrocellulose solution.

More usually, a preponderance of metal remains, and the incrustation is due to silver chloride; but silver coins are generally debased and the incrustation will be found to be mixed with the products of the corrosion of the copper present in the silver alloy. The coins may then be adhering firmly together in masses. A conglomerate of this nature may be broken down by formic acid treatment or by soaking in 5 per cent. citric acid, although sometimes the individual coins are more easily released by reducing the mass with zinc and caustic soda. The alkaline Rochelle salt process (p. 239) has advantages where the silver is greatly debased with copper, but in this case the coins must be carefully brushed from time to time since the cuprite which cements them together is dissolved by the Rochelle salt exceedingly slowly and then only in the presence of air.
31. THE EMESA HELMET (1ST CENT. A.D.)

Left: As received in the Laboratory. Right: After treatment and reconstruction
32. CHINESE MIRROR OF HAN DYNASTY

The hair-like cracks result from stresses in the brittle speculum metal set up by oxidation.

33. CHINESE MIRROR OF HAN DYNASTY FROM THE HUAI VALLEY

The oyster-shell pitting is characteristic of advanced spotty corrosion on a brittle speculum metal.
In the British Museum Research Laboratory, where coins frequently arrive for treatment in hoards, they are dealt with in a machine constructed on the principle of the ball mill. This apparatus takes the form of a deep rectangular tank made of polyvinyl chloride mounted at an angle of about 45° from the vertical ("tumbler tank") and arranged to revolve slowly about its longer axis. It is driven by an electric motor. For cleaning base silver coins the tank is filled with an appropriate reagent such as alkaline Rochelle salt, and a few loose coins are added to act as hammers when the tank revolves and so help to separate other coins from the conglomerate mass. As the tank revolves the contents fall four times from the flat walls during every revolution and these movements assist the chemical action so that the coins are soon separated. Individual coins are washed with water thereafter and dried in hot sawdust in the tumbler tank; they emerge clean and with a slight polish which seems to heighten the contrast of light and shade and thus enable lettering and ornament to be more easily studied.

Before the installation of the tumbler tank, electrolytic reduction was employed, and while this procedure was found to give equally satisfactory results, it involved more labour. It was applied, for example, in cleaning a hoard of 15,000 silver coins from Dorchester. During electrolysis the coins required continuous attention and had to be frequently removed and brushed in order to prevent the reduced incrustations from being deposited as a resistant metallic film on the coins. The small mass of corroded silver coins shown in Pls. 30 A and B had become agglomerated as a result of fire. The mass was loosened to a considerable extent by soaking for some hours in a bath of citric acid (5 per cent.), final cleaning being done by placing the coins in stirrups of copper wire suspended from the cathode bar of the electrolytic tank.

COMPOSITE OBJECTS

Silver has been used so widely in decoration that all sorts of problems are presented in the conservation of objects of a composite nature containing the metal in conjunction with other materials.
In dealing with silver inlays it is safest to adopt the mechanical methods which have already been described (p. 200). If chemical treatment is essential, it may be necessary to isolate the inlay with a wall of Plasticine or clay, but there is always the danger of chemicals seeping beneath the inlay and presenting a problem in washing. When washing after treatment is impossible, as with a fretted silver ornament set in lacquer, cleaning can only be satisfactorily accomplished by the use of abrasive cloths. The problem of revealing silver inlays in rusted iron generally resolves itself into adopting methods for removing the rust which tends to overrun and obscure the silver.

RESTORATION OF THE EMESA HELMET

An unusual and very interesting problem involving the conservation of silver in juxtaposition to iron was provided by the Emesa helmet, a treasure from Syria (Pl. 31), and as it presented certain novel features its restoration will now be described in detail. When received, this fine helmet, with its silver visor in the form of a human face, was at first taken to be a ceremonial piece, but, as pointed out by Seyrig, the protection afforded by the iron skull-cap was continued by a lining of iron of substantial thickness 1–6 mm. underlying the visor and conforming exactly to its shape. The probability is therefore that the helmet was intended as head armour to protect the wearer under war conditions; the iron cranium was dented as if by a heavy blow. The silver was found on examination to be very brittle and cracked in many places; the major cracks had been filled with a dark stopping material in an effort to reinforce the silver as shown in the photograph. The iron backing of the visor had rusted and the swollen metal pressing behind the brittle silver had been responsible for distorting the features and forcing the cracks open, in some cases to as much as 4 mm. A state of strain had set in which, if not released, could only lead to further deterioration as the silver itself was only 0.5 mm. in thickness. A necessary preliminary to treatment was the removal of the visor from the helmet; this proved to be easy because

---

1 Seyrig, H., 'Le Casque d'Émèse', Annales, Arch. de Syrie, 1952, 2, Nos. 1 and 2.
the hinge, although no longer movable, was already detached from
the head-piece and the two parts were merely held together by wire.
After temporary consolidation by applying transparent Sellotape
across the cracks in the mask, the visor was bedded in a thick pad of
cotton-wool, and the layer of rusty iron was cut away from the back
in the region of the mouth and lower jaw. It was in this region that
the silver was weakest and most distorted through contact with the
corrod ing iron. It was clear that no satisfactory restoration would be
possible unless the silver could be toughened. A pilot experiment
was carried out on a tiny detached fragment and as this gave re-
assuring results the visor was heated in the electric furnace for three
hours, the temperature rising slowly during this time to a maximum
of 310° C. Heating was arrested at this point. The silver was now
darkened, partly by the burning of the stopping of rust and wax that
had been used in a previous restoration to fill the cracks and partly by
the blackening of residual rust that remained attached to it. The rust
was removed by brushing with 9 per cent. oxalic acid (see p. 278)
before the final heat-treatment as additional heating would have made
the iron compounds too insoluble to remove later. Further heating for
eighteen hours at 600° C. and thirteen at 650° C. was necessary before
the silver was considered to be sufficiently toughened to stand up to
the manipulation that would be necessary in closing the cracks. In
order to strengthen the visor, where iron had been cut away, the
underside of the silver was cleaned and given temporary reinforce-
ment by applying patches of silver gauze across the cracks with soft
solder. The gauze took the shape of the silver very readily and could
be fixed without involving any pressure. The front of the visor was
then carefully cleaned with Dura-glit cloths to remove all of the
dark oxidation products and it was then in a condition for the restora-
tion work to begin.

The first operation was to restore the shape of the lower jaw by
dealing with the cracks, one series at a time. The temporary patches
of silver gauze were removed, and the silver was manipulated to get
the edges of the cracks to meet exactly, then fresh patches of gauze
were applied, and fixed permanently in position across the cracks on
the underside, using soft solder with a non-corrosive flux. This was a long and exacting process but well worth all the care expended upon it because, as the work advanced and the cracks disappeared one by one, the visor became noticeably stronger and more rigid, and the likeness nearer to that shown in the earliest photographs.

It should be emphasized that it was the toughening of the silver that made this type of restoration possible; if for any reason it had not been possible to toughen the silver by heat treatment, it could not have been manipulated. In its brittle condition the only possible action would have been to try to reinforce the inside, and fill the cracks from the front. It is fortunate that the more permanent type of restoration could be carried out on the Emesa visor, and that it was not necessary to use filling material, as this tends in time to become a conspicuous disfigurement. Care was taken to leave no solder on the front of the mask; it appeared inevitably as thin lines in the repaired cracks, and these were concealed by the local application of a surface coating of silver, applied by ‘ragging’ with a pointed silver anode tipped with cotton-wool moistened with potassium cyanide (p. 289).

It now only remained to replace the rusty iron that had been removed from the back of the visor. This material, after cleaning, was still in the form of sizeable fragments that could be returned to position in contact with the silver, where they were consolidated with Durofix.

The major fractures that remain in the silver after restoration no longer weaken the structure, nor do they cause serious disfigurement. It would have been possible to deal with all the cracks, but this would have meant removing practically all of the iron from the back of the visor, and, as much of the iron was in sound condition, this large operation was not considered justified. The interest and value of the object depend as much on the variety and perfection of the craftsmanship displayed in its construction as on its actual appearance. With the removal of the loose and swollen rust, where the strain was greatest, and the consolidation of the iron and silver in the lower part of the face, the helmet is now considered to be in a stable condition.

The small holes beneath the eyes in the visor had been designed so that the wearer could look down to the ground in front of him, but
evidently they were not enough, and it is amusing to find that they had been enlarged, in each case, by a narrow notch cut in the central lobe, no doubt in an emergency as this adjustment is not the work of a craftsman.

**LACQUERING SILVER**

Lacquers that are of the right quality and consistency are almost invisible on silver if they are skillfully applied. Many types are available, and some of the best are those based either on polyvinyl acetate or the polymethacrylates. A lacquer that has been found in practice to have excellent protective properties, and that is almost invisible when sprayed, or carefully brushed on silver, is the commercial product known as Ercalene. Frigilene is also very good.¹

Before lacquering, the silver must be cleaned and degreased, and while cleaning can be done by any of the methods recommended, a word of warning is necessary as regards the washing process to remove grease. Many proprietary detergent preparations are available today that are excellent for household cleaning, but some are entirely unsuitable for silver, as they contain phosphates which react with the silver to form a resistant brown stain. It is better to degrease the polished silver by washing it in hot soapy water, or the pure detergent, Lissapol, may be used; the silver is then rinsed in running water and dried. After degreasing, care should be taken not to spoil the surface by touching it with the fingers. When quite dry the silver is ready to be lacquered.

Lacquer may be applied by spraying, dipping, or brushing. The brushing technique is probably the simplest. A soft clean brush is used and the lacquer should be sufficiently thin to spread without leaving brush marks. It must not be allowed to collect as a thick film in any hollows where it would catch the light and reveal its presence by a shine foreign to that of the silver. It is useful to have a solvent at hand which can be used, if need be, to remove any surplus. Lacquer can be prevented from collecting in hollows by stippling with a brush or by using a gentle puff of air from a blow-ball.

¹ Both preparations are marketed by Messrs. W. Canning & Co. Ltd., London.
TABLE III. SILVER ALLOYS

Silver alloys

Uncorroded

- Solid, Substantial
  - Bright
    - 1; 2, 8a, 7, 11
  - Oxidized, tarnished
    - 3a, b, e, f, 4g, 8a, 7, 11

Part of object corroded, or having attachments of corroded base metal

- Macrocristalline, brittle, or only seriously deformed
  - Bright
    - 6, 7, (9), 11
  - Oxidized, tarnished
    - 4g, 8a, 6, 7, (9), 11

Incrusted with green copper compounds

- 1; 4d or e, 8a or b

Thinly covered with silver chloride

- 1; 4h or i, 8a; 5c or e, 8a

Corroded

- Heavily incrusted with horn silver, &c.
- Mineralization complete
  - 1; 2, 9a, b or e

Corroded material to be removed (extraneous)
- Either 10 or 4 or 5 followed by 8a, 7, 11

Corroded material to be retained (structural)
- Either 1 or 2, 3, or 4d, 8a, 7, 11

For method of use see p. 186.
1. No treatment.

2. (a) Warm soapy water; (b) Lissapol froth.

3. Mild abrasives: (a) French chalk; (b) Rouge; (c) 'Plate powder'; (d) Pumice or emery powder; (e) 'Metal polish'; (f) Dura-glit; (g) Special cloths.

4. Solvents:
   (a) Hot concentrated aqueous ammonia.
   (b) Potassium cyanide.
   (c) Nitric acid (5 per cent.).
   (d) Hot formic acid (50 per cent.).
   (e) Alkaline Rochelle salt.
   (f) Sulphuric acid (5 per cent.).
   (g) Silver Dip.*
   (h) Ammonium thiosulphate plus Lissapol.
   (i) Thiourea plus Lissapol.
   (j) Citric acid (5 per cent.).
   (k) Silver nitrate (20 per cent.) for removing a copper coating from silver.
   (l) Concentrated hydrochloric acid plus ferric chloride (10:1) for soft solder on silver.
   (m) Glacial acetic acid plus '20 vol.' hydrogen peroxide (3:1) for ditto.
   (n) Oxidizing Rochelle salt. (See p. 241.)

5. Reduction:
   (a) Aluminium and caustic soda.
   (b) Zinc and caustic soda.
   (c) Zinc and formic acid.
   (d) Zinc and dilute sulphuric acid.

6. Annealing and reshaping.

7. Protective finishes: (a) Lacquer: Ercalene, Frigilene, polyvinyl acetate, or a polymethacrylate, e.g. Bedacryl 122X.
   (b) Paraffin wax dip.
   (c) Microcrystalline wax polish.

   (b) Sodium sesquicarbonate (5 per cent.).

9. Consolidants: (a) Nitrocellulose (Durofix).
   (b) Bedacryl 122X.
   (c) Polyvinyl acetate.

10. Mechanical: (a) picking, (b) chipping, (c) grinding, (d) brushing, (e) grit spraying, (f) polishing, (g) burnishing.

11. Storage and exhibition. Dust-free case with sulphur-free atmosphere.

* Silver Dip that has been used will contain silver in solution and will deposit this upon copper. For this reason it is safer not to use Silver Dip for cleaning silver inlays in bronze. For such a purpose mild abrasives are probably most effective.
CHAPTER XI

COPPER AND ITS ALLOYS

Copper derives its name from Cyprus which was the Roman source of the metal, *aes cyprium* being the 'bronze from Cyprus'. It is, however, an element, and it occurs in nature in the metallic condition as well as in the form of many minerals, chief of which are cuprite (cuprous oxide), chalcocite (cuprous sulphide), chalcopyrite (copper pyrites), and the basic carbonates, malachite and azurite or chrysocolla.

Metallic copper resembles silver in being sensitive to sulphur which causes it to become covered with a film of tarnish consisting of copper sulphide. It differs from silver, however, in being sensitive also to oxygen; pure copper oxidizes very readily when exposed to moist air, whereas, under similar conditions in air that is free from sulphur compounds, silver remains unstained. Thus it happens that while silver objects in the museum tend to tarnish, copper objects are more usually covered with a thin film of oxide which confers on them a dull appearance. The oxide film does not increase appreciably in thickness with time and for this reason it may be regarded as protecting the underlying metal; oxidation may be so slight that the metallic appearance of the metal is maintained and there is no disfigurement; but if the metal is alloyed with tin, lead, or zinc and the constituents imperfectly mixed, the oxidized surface may be patchy and give rise to a rather unpleasant appearance. The restoration of the metallic condition is not, however, a major problem as the thinly oxidized metal may be cleaned with metal polish or by immersing the object for some hours in dilute sulphuric acid (5–10 per cent.), washing afterwards, in each case, drying, and rubbing up with a soft cloth. But the brilliancy is soon lost and to preserve a burnished surface, even under museum conditions, it is usually necessary to cover it with an impervious film of lacquer.
When buried in damp soil, copper soon loses its metallic appearance. The oxide layer increases in thickness, and cuprous oxide becomes compacted into the purplish-red mineral known as cuprite; this, in turn, may become encrusted with basic carbonates that are green or sometimes blue in colour and that correspond to the minerals malachite and azurite. Such incrustations are stable when free from chloride and they protect the underlying metal from further corrosion.

It does not follow, however, that all green incrustations are stable. Appearance has little to do with stability, even though it be true that a thin coherent deposit is more likely to be stable than a heavy porous incrustation that would tend to behave like a sponge and absorb moisture and soluble salts from the soil. When more than one metal is present, as in the tin bronzes,¹ it is only to be expected that incrustations will be more complicated both in composition and structure and therefore more liable to contain and retain salts. Contamination is the rule rather than the exception because soluble salts are widely distributed in nature, in the soil as well as in the sea, and for this reason they are usually present in excavated material.

It was noted in the case of silver (p. 216) that when this is buried in salty ground it tends to become covered with an insoluble shell of stable silver chloride. In the case of copper and its alloys, however, the presence of chlorides in the incrustation presents an acute problem from the point of view of conservation because an unstable cuprous chloride is formed; this cuprous chloride continues to react and there is progressive corrosion, even under museum conditions, with the result that the surface becomes powdery and spotty. So common are these features in corroding bronzes that the appearance of characteristic pale green powdery spots is referred to as 'bronze disease' whether occurring upon copper or on any of its alloys.

The spots grow as a result of the nantokite (cuprous chloride) being transformed into atacamite (basic cupric chloride) by the action

¹ Gettens, R. J., *J. Chem. Education, 1951, 28*, pp. 67–71. In this paper the reactions that take place during corrosion of a tin bronze are described and the morphology of the incrustation illustrated in a series of photomicrographs.
of oxygen. This change is facilitated by the presence of moisture and it is not surprising that ‘damp’ has been mistakenly credited with being the cause of corrosion since serious outbreaks of active corrosion have been observed to take place in damp weather. But as the features of bronze disease cannot be reproduced even under damp conditions when chlorides are absent, it is clear that moisture is only a subsidiary factor in the corrosion of copper and its alloys. The main objective, therefore, in conservation is to get rid of chlorides. Here we are faced with two difficulties. Cuprous chloride cannot be removed by mere washing with water. Not only is it insoluble but it is inaccessible as it occurs in greatest concentration, not in the surface spots, but in the deeper layers of incrustation. In actual practice, chemical methods can be applied to convert insoluble chloride complexes into soluble salts that can be removed by washing, but it is the inaccessibility of the nantokite that introduces the serious problem. A corroded bronze has a banded structure (Pl. 26), the metal core being surrounded by layers of massive cuprous oxide interspersed with powdery tin oxide, and these layers underlie an external shell of basic carbonates. Chlorides may be present in the green surface shell, but in a heavily corroded bronze concentrations of cuprous chloride are found in the absorbent layers of tin oxide, and even in the micro-cracks and fissures that can be shown by metallographic technique to exist in the interface between metal and cuprous oxide.

In view of these facts it would seem to be outside the bounds of possibility to remove all chlorides from a mineralized bronze without at the same time decomposing the incrustation, and, indeed, this is the only certain way to stabilize a bronze that is heavily corroded. There are many laboratory methods by which this may be accomplished, and they are generally referred to as stripping methods. But there are times when it would be quite unthinkable to sacrifice the patina of a bronze and when, if corrosion has set in, ways and means must be found to deal with the attack while preserving, as far as possible, the appearance of the specimen.

In the study of ancient metals it will be noted that bronzes have survived for thousands of years even though their porous incrusta-
tions have been impregnated with salts. The explanation must be that an equilibrium had been established between the corroded bronze and its environment in the ground, and it is to this that we must attribute survival. Excavation destroys the equilibrium, however, and exposure to new and very different environmental conditions provokes changes that are often profound. It may sometimes happen that a bronze can undergo a second acclimatization to the museum environment, without visual change, but it is more usual for further corrosion to take place in the process, and if neglected, equilibrium will in many cases only be re-established at the expense of complete disintegration. All metal objects should, therefore, be sent for laboratory treatment as soon as possible after excavation.

When electro-chemical methods can be adopted, the objects will be restored to a stable condition and to something resembling their original uncorroded appearance. But it will not be possible to apply electro-chemical methods in every case. Reduction methods are only applied when a substantial metallic core remains and when its mechanical strength is beyond doubt. In the absence of a metallic core the aim will be to arrest decay by the use of selective solvents. When electro-chemical or electrolytic methods are out of the question, as in the case of a finely patinated bronze, the prospect of arresting decay without destroying the patina will depend upon the nature and distribution of the corrosion, and also upon what success can be achieved in localizing the action of the chemical reagents during treatment. When conditions are dry, chloride activity will be at a minimum and if bronze disease should break out and can be dealt with in its early stages, there will be the less chance of its getting out of hand.

In this type of problem the following points are important—(1) to treat the spots in the early stages, (2) to dry the bronze well thereafter, and keep it dry, and (3) to be alive to the necessity of giving further treatment in good time should further spots make their appearance. Methods of treatment for bronzes where patina has to be preserved are given on p. 242, but it will be realized that these methods do not eliminate all chlorides from within and beneath
the incrustation. Such treatment must therefore be empirical to a
certain degree, and there can be no absolute assurance that preserva-
tion will be permanent, although many cases are on record where the
treatment has proved to be successful. By these processes, chlorides
are removed from the surface where the shell of incrustation is most
porous and weakest, leaving the more solid crust of corrosion to seal
up the remainder. It is a common error to believe that a short cut is
possible—that wax or other film-forming substance may be used to
seal in the decay and so act as a protective layer. Experiments are
described below which show this to be a fallacy. No impregnating
medium has been found to be of any permanent value in arresting
the progress of ‘bronze disease’, unless the spots are excavated in the
first place or treated by chemical means to enable the chlorides to
be extracted.

Before proceeding to describe and recommend processes of treat-
ment, it may be useful to refer to some observations made during the
examination of a mineralized ‘bronze’ spear-head of rectangular
cross-section from Ur of the Chaldees that could be sacrificed for
experimental purposes. Although covered with a heavy green in-
crustation, the object was apparently stable under museum condi-
tions and there was no evidence that corrosion had taken place in
recent times. Chemical analysis showed the metal core to consist of
97.2 per cent. copper with less than 1 per cent. tin.

The spear-head was sawn into a number of fragments roughly
cubical in shape. The cross-section showed the structure to consist of
a metallic core enveloped in massive purple-red cuprite, on the sur-
face of which had grown a green patina or incrustation of about
3 mm. in thickness, mainly malachite (Fig. 8). The test pieces were
dealt with variously as follows:

1. Controls. Some cubes were kept as controls, and these broke into
active corrosion within forty-eight hours of being cut. On examining
the section, pale green powdery material was observed growing
thickly along the inner margin of the outer green patina. This was a
zone of nantokite (cuprous chloride) which had been sealed within
the incrustation, but on exposure to air it was converted to atacamite
(basic cupric chloride) and exhibited the typical appearance of bronze disease within this very short period of time.

2. Washing. Other pieces were washed in distilled water with the greatest care until apparently free from chloride and dried by passing through alcohol and ether, and maintaining at 105°C in an oven for several days. When these were exposed in a humid atmosphere bright green striae formed on the exposed edges of the outer green layer following certain lines of the strata, and these could be shown by micro-testing with silver nitrate to contain chloride. There was no indication of any chloride in the central metal core or in the cuprite immediately overlying it, nor could chloride be detected on a longitudinal external face of the specimen. The chloride was, therefore, located within the green rind of incrustation but not on the outside surface, and it could not be removed, even where exposed on the cross-section, by the most careful washing alone.

3. Reduction. A third series of test pieces was subjected to electrochemical reduction with zinc and caustic soda or to electrolytic reduction, washed free from chloride, and dried. Corrosion could no longer be induced to take place even when the samples were kept in a damp atmosphere. Reduction is therefore a cure for corrosion when applied for long enough in conjunction with washing to

---

1 Although there was no evidence of it here, it is quite common to find the attacking layer of cuprous chloride between the cuprite and the metal core.
eliminate all chlorides. But this process involves the loss of the green patina.

4. Acid treatment. Specimens were treated by soaking them in citric acid (5 per cent.) and sulphuric acid (15–20 per cent.). The latter was more effective. It seemed to go a long way towards removing the chlorides in the course of a few days. There are practical difficulties in using it, however (see p. 241), and it is not recommended as a standard process. These acids act as stripping reagents and the green surface is lost as in the case of No. 3 above.

5. Alkali bath. A test was made by immersing some pieces in sodium sesquicarbonate solution (5 per cent.). This did not remove the green incrustation or patina, but gradually extracted the chloride in the form of salts that could be dispersed and eventually eliminated by frequently renewing the solution. This operation, including washing with distilled water, took several weeks to reach the stage where chlorides could no longer be detected and the dried bronze was stable when tested under moist conditions. There can be no doubt that, after such treatment, much chloride must remain inaccessible beneath the green surface, but it seems that the sodium sesquicarbonate gives rise to basic copper carbonate which fills the pores of the incrustation, sealing off the residual chlorides from atmospheric moisture.

Assuming that the test-pieces are representative of the average conditions that obtain when heavily corroded bronzes are contaminated with chloride, the experiments indicate in which part of the incrustation chlorides are to be found in the greatest concentration, and how very sensitive they are to moisture. They prove the inadequacy of washing alone, the effectiveness of stripping methods, and the possibility of protection by removing exposed chloride, and sealing the remainder behind a moisture-barrier.

The choice of a method of conservation depends upon the physical and chemical condition of the specimen, the amount of chloride present, and the nature of the incrustation—its thickness, stratification, and porosity. If it is essential that the patina should be saved, it is clear that the operation may be a lengthy one, and the results inevitably fraught with some uncertainty.
METHODS OF TREATMENT

1. *When incrustation or patina can be sacrificed*

When the incrustation or patina can be sacrificed, corroded copper alloys may be dealt with by electro-chemical (p. 191) or electrolytic reduction (p. 194), or by employing alkaline Rochelle salt. The latter process has been found to be the one most generally useful. Preliminary treatment by Rochelle salt is recommended, therefore, for dealing with copper incrustations, even where apparatus is available for electrolytic reduction, though it will be found convenient to employ the electric current in the final stages, in order to facilitate the removal of salts and chemicals, and thus economize in the time required for the final washing in distilled water.

(i) *Use of alkaline Rochelle salt and dilute sulphuric acid.* Copper combines with oxygen in two different proportions by weight to form two definite oxides, cuprous oxide and cupric oxide, and these compounds are commonly present in incrustations, and give rise to two series of salts, the cuprous salts and the cupric salts respectively. These may be carbonates, chlorides, sulphates, &c., and the chemical nature and distribution of these compounds will depend on the conditions to which the metal has been exposed. Usually the cupric salts predominate in the visible incrustation, and they happen to be soluble in an alkaline solution of Rochelle salt. By dissolving them away the green encrusted surface of a copper alloy is broken down, and it then remains to find a method of eliminating the residual cuprous salts. This can be done with dilute sulphuric acid. There are two operations, therefore, and these are applied consecutively in the standard form of Rochelle salt treatment, the result of their application being to remove all the mineralization and to restore the metallic condition of the specimen.

The two solutions are prepared as follows:

*Solution 1. Alkaline Rochelle salt.* Dissolve 1 oz. of commercial flake caustic soda in a pint of cold water, and dissolve 3 oz. of Rochelle salt (sodium potassium tartrate) in this solution.

*Solution 2. Dilute sulphuric acid* (10 per cent.). Add 2 oz. of
concentrated sulphuric acid slowly to a pint of cold water in a porcelain dish, stirring continuously.¹

In order to carry out the treatment the encrusted metal is allowed to soak in solution 1, in a covered container of glass, porcelain, or stainless steel, either in the cold or, if it is desired to hasten operations, in a steam-bath until the incrustation is softened, when most of it can be removed by brushing. The colourless Rochelle salt solution becomes deep blue in reacting with the incrustation.

After the first part of the treatment, a brownish-red residue of cuprous oxide (cuprite) will be found firmly adhering to the metal, and it does not respond well to brushing. It is usually associated with pasty cuprous chloride. Sometimes there is also a layer of metallic copper which has been deposited within the incrustation during the process of corrosion. This copper (referred to as redeposited copper) tends to seal up the minerals beneath it, and protect them from attack by the reagents. If redeposited copper is observed, this complicates the treatment because it is insoluble in Rochelle salt and in dilute sulphuric acid, and it must be removed mechanically. This is not always an easy matter and it takes time, but unless all of the copper deposit is picked off at this stage, there can be no guarantee that the process will give permanent results.

The object is then placed in solution 2 in a container of glass, porcelain, or stainless steel.² It is set aside in a warm place with occasional brushing, and examination with a pocket lens, until as much as possible of the cuprite has been eliminated together with a sludge of copper powder.

A peculiar condition that sometimes arises and may be seen under the lens is one in which cuprite forms a crystalline mosaic with copper crystals on the apparently intact surface of the cleaned metal. As this film may have chlorides trapped beneath it, it must be broken

¹ The acid must always be added to the water, and not vice versa; water should never be added to strong sulphuric acid as this might cause serious accident due to spurt ing accompanying the sudden evolution of heat.

² If a stainless-steel vessel is used with acid, insulate the object in the vessel by placing it upon an old photographic plate or other resistant electrical insulator to avoid deposition of the copper on the steel.
34. EGYPTIAN BRONZE FIGURE OF ISIS AND HORUS (PTOLEMAIC PERIOD)

Left: As excavated (plinth is modern). Centre: After treatment with Rochelle salt and caustic soda. Right: After final treatment
35. BRONZE BUST OF AN ETHIOPIAN QUEEN (7TH-6TH CENT. B.C.)

Left: As excavated. Right: After restoration.
down in the final operation. It is in helping to break down this film that the method of electrolysis is so useful, and the electrolytic tank is thus a valuable adjunct to the Rochelle salt method of cleaning. Electrolysis is not essential, however, as brushing and pickling in dilute sulphuric acid solution can achieve the same result, although this usually takes much longer and is more laborious. The object is boiled finally in several changes of distilled water, cooling on each occasion in the water to flush the capillaries as described on p. 198. The steam-oven method of heating and cooling is more convenient when dealing with a batch of objects. Washing is continued until chlorides can no longer be detected in the water when the silver nitrate test is applied as described on p. 199.

(ii) Use of alkaline Rochelle salt with hydrogen peroxide. When bronzes having fine line ornament are heavily encrusted the sulphuric acid, in dissolving the cuprite, tends to leave any fine lines filled with copper powder, and in this event the copper is difficult to brush out. This complication can be avoided by using a single oxidizing bath made up by adding 100 ml. of hydrogen peroxide (20 vols.) to each litre of the standard Rochelle salt solution I. The object is then immersed and brushed from time to time and picked if necessary. The action of the hydrogen peroxide may be followed by observing the small bubbles that are liberated at the point of attack, namely, around the residues of cuprite; cuprous salts are oxidized and become soluble in the Rochelle salt solution. The hydrogen peroxide method is slower in action than when sulphuric acid is used, and the metal object may be slightly attacked in the final stages if the oxidizing solution is used for too long. But, under strict supervision, the oxidizing process is, on the whole, the more likely to yield a satisfactory restoration of fine line ornament, and it is considered therefore to be the preferable technique where fine decoration exists.

The beginner tends to seek for a simple course of treatment, and may be tempted to ask whether it is not possible to simplify operations by carrying out the whole of the cleaning with sulphuric acid alone. This is unlikely to be successful. Sulphuric acid does not remove redeposited copper, and in fact, owing to the length of time
required for pickling, using sulphuric acid alone, further copper is almost certain to be redeposited in a fine-grained adherent form in the neighbourhood of the chloride deposits. This would involve an inordinate amount of mechanical cleaning: the redeposited copper would have to be removed as otherwise salts might tend to be sealed in, with the risk of a further outbreak of corrosion.

Sulphuric acid is really a clearing reagent for cuprite. It is also useful for the preliminary treatment of heavy highly-corroded bronzes, such as a bunch of spears or tools that have become corroded together. In this case it is used at about 20 per cent. strength as a pickle to release the corrosion products so that the various objects can be separated and then given individual treatment.

Detailed instructions such as the foregoing tend to suggest that the methods are beset with complications, but in the application of stripping processes this is not really the case. Objects do tend to react differently under treatment, but this adds interest to the work, and at the same time provides a stimulus for ingenuity.

2. *When patina or incrustations should be preserved*

A most difficult problem is presented by the well-patinated bronze which is corroding in spots. Here the aim must be to arrest the corrosion while retaining the general character of the object. The method of treatment is determined in the first instance by the structure and appearance of the patina and by the distribution of the spots. The following are the three commonest conditions—isolated spots in thin soft patina; isolated spots on thick hard patina; and spots overlying a porous area. Examples will now be presented to illustrate the treatment applied to typical specimens in each category, but methods may have to be varied considerably to suit particular requirements.

An Islamic bowl of sound metal with chased ornament all over it had a thin green patina tinged with brown cuprite. The patina was spotty (indicating the presence of chlorides) and, moreover, it concealed important details of the ornament. It was desired to remove chlorides, get rid of the spots, and at the same time expose the
pattern-work where it was concealed, without giving an effect of
over-cleaning. The bronze was soaked in changes of sodium sesqui-
carbonate (5 per cent.) for some weeks, rubbing away the incrusta-
tion after it had softened in those places where it concealed ornament.
This could be done with the fingers or with a soft wooden scraper.
The bowl was washed until chloride was no longer detected in the
wash-water on testing with silver nitrate, and then boiled in water
for some hours so as to darken the metal where it had been exposed.
The bronze was dried and polished with a waxed brush and it still
retained the appearance of a patinated bronze of brownish colour
relieved by green carbonates in the deeper parts of the ornament.
This type of treatment is often applied and gives good results with
thin patinas if they are carefully handled while in the softened con-
dition but it must be noted that warming the sodium sesquicarbonate
solution would tend to dissolve the patina.

A fine Persian bronze vase having a comparatively thick hard
green patina was similarly infected with a few deep-seated pale green
spots. These not only detracted from the appearance of the vase but
were observed over some weeks to have increased in size and owing
to the thickness of the mineralized surface the condition was re-
garded as serious. In this case the spots were carefully cleaned of all
powdery matter, using a needle under a binocular magnifier; they
were then reduced locally, by applying to each individual spot zinc
powder and drops of 90 per cent. sulphuric acid from a fine glass
capillary tube. By this means the chlorides responsible for the cor-
rosion were rendered soluble and could be washed out in changes
of distilled water. Such treatment, which removes all the powdery
material and leaves the spots a dull brown colour, is most likely to be
effective when, as in the present instance, the general patina is hard
and shiny, but care must be taken in using acid to prevent it spreading
as it attacks malachite instantly. After treating a spot and allowing
effervescence to take place for perhaps half a minute or less, the
bronze is held in a stream of water, then mopped dry, and the area
inspected. It may need further excavation and further reduction—
experience will tell. In restoration work of this kind it is worth while
making experiments on worthless material before attempting to restore an object of value. It should be emphasized that strong sulphuric acid is a dangerous chemical (see Appendix II) and is only used in a room where a water-supply is available in case of emergency.

Incidentally, the method when applied over a larger area provides a means for the local treatment of ornament and has been found useful in rendering legible incised inscriptions that have become clogged with oxide, where it is desirable to avoid wholesale reduction. After such treatment, which, of course, removes the green incrustation, the metal should be washed locally by a water jet so arranged that the washings fall away without running over untreated parts of the object.

Egyptian bronzes are often heavily infected with salts and for this reason seldom possess good patinas. When a good patina does occur, therefore, every effort should be made to preserve it, but if the surface once becomes spotted, restoration may be a difficult matter. An Egyptian bronze cat about 10 inches high, having a patina that was exceptionally fine, persisted in corroding over a small porous area. As there was a natural reluctance to excavate the corroded area, it was treated locally with jets of distilled water, arranged to run continuously using a water circulator.1 This was continued for many weeks in the hope that prolonged washing might be effective. The bronze was then dried and the area painted over with lacquer. Since this treatment failed to arrest corrosion, the lacquer was removed and the washing repeated, but no matter how many times this was carried out, nor how many sealing agents were tried, it invariably happened that after some months of quiescence the trouble broke out anew. It was found necessary in the end to excavate the area where the salt was most concentrated and to make good with a little powder colour ground in Ercalene lacquer in order to preserve the appearance of the specimen; but such treatment has little chance of being effective unless the object is kept permanently in a dry case, or protected by some other means from damp.

The intimate connection between damp and corrosion is illustrated by the interesting case of a Greek bronze head of a Barbarian dating to 400 B.C., which had a very fine patina that had never given any trouble. Suddenly it showed signs of corrosion and the cause was traced to damp from a leaky roof that had entered the wall against which the bronze had been exhibited. The spots were excavated individually with dental tools and, when the bronze had been carefully washed and dried, the cavities were filled with lacquer. No further outbreak was detected until the following winter when a series of new spots had to be treated, but these were neither so intense nor so obtrusive as before. There has been no further trouble,
nor indeed is any expected, as the specimen has now been mounted in a case of the island type kept permanently dry by silica gel (see Appendix XI). Such an arrangement is shown diagrammatically in Fig. 9. In instances of this kind it is often better to accept the risk of further outbreak and be prepared to attend to the specimens again, rather than adopt drastic measures that would be certain to change the character of the piece.

It is obvious that the methods already given are capable of being varied in many ways to suit the problem in hand. Thus, the water-circulator might be used to circulate sodium sesquicarbonate or some other solvent, and the local reduction process might be carried out using caustic soda in place of sulphuric acid; or aluminium or magnesium powder might be substituted for zinc. Whatever treatment may be applied, the desiccator type of museum case, containing silica gel, is a useful stand-by in helping to arrest any residual activity.

The foregoing examples illustrate how the choice of method in dealing with corroded bronzes is limited by the presence of a good patina. It is sometimes limited by the presence of material of a non-metallic nature. Thus, the plaster and gold leaf that covers certain types of Egyptian bronze may provide a serious complication. Vestiges of the former ornament may remain, and the bronze cannot, therefore, be immersed in any liquid. In one such case, corrosion was arrested by daily applications of paper pulp (see p. 299) that had been soaked in a 5 per cent. solution of sodium sesquicarbonate. Many applications were made before the spotiness finally disappeared, and the metal surface, where exposed, was covered by a stable green film of basic copper carbonate.

Egyptian bronzes sometimes retain a siliceous core as a legacy from the casting technique, and when such bronzes are diseased it is usually a difficult matter to restore their stability. As a siliceous core is so much more porous than bronze, it contains proportionately more salt, and unless the core can be removed, any immersion process is likely to do more harm than good. Such bronzes can only be given local surface treatment.

The final example is of an Egyptian bronze having a core of a
different type. This finely chased specimen was a hollow casting of a falcon, and the bronze had a deep green patina which, however, was pitted and spotty. As various forms of surface treatment had been tried without success, it was eventually decided that the salt must be inside and that it would be necessary to remove the base-plate in order to gain access. When the base-plate was cut out the mummified remains of a bird were found in the interior, and after these had been extracted it was a simple matter to restore the bronze by the use of chemical solvents, immersing it first in citric acid and finally in sodium sesquicarbonate until stability had been attained.

3. Removal of calcareous deposits. Use of sodium hexametaphosphate (Calgon)

Bronzes are sometimes covered with calcareous deposits, and while these may be removed with dilute nitric acid this is undesirable where there is fine work, or where metal is very thin, as the acid treatment is rather drastic. In all cases it is preferable to remove the deposits by soaking the bronze in a 5 per cent. solution of sodium hexametaphosphate (Calgon),¹ which in time releases the deposits of calcium and magnesium salts by combining with them to form soluble salts. Stronger solutions up to 15 per cent. may be used. This hastens the reaction, and warming the solution hastens matters still further, but not without risk to the patina. On the other hand, where the patina is not of importance, Calgon solution can be used to remove the green incrustation completely, merely by continuing the treatment for long enough. Calgon has the merit that it may safely be employed for cleaning the surface of copper alloys that are in such an advanced state of decay that no metallic core remains. For this reason it is a useful material to have in the restoration laboratory.

CHINESE BRONZES

Chinese bronzes comprise alloys of the gun-metal and bell-metal class, containing tin in the proportion of 10–20 per cent. Such alloys were used for the massive funerary bronzes. The mirrors are of

¹ Farnsworth, Miss Marie, *Technical Studies in the Field of Fine Arts*, 1940, 9, p. 21.
speculum metal, a white bronze which may contain tin in quantities up to about 30 per cent. Lead is commonly present as a minor constituent in all Chinese bronzes; its presence seems to have little effect on stability, though when present in quantity it tends to lower the tone of the patina.

In alkaline soils which contain little chloride, the corrosion of copper and its alloys takes place more slowly and more uniformly than when much chloride is present, and the products of the reaction, as we have seen, are more likely to be stable. They may, in fact, constitute a beautiful patina well worth preserving, either for aesthetic reasons or as evidence of antiquity. Such are the patinas found on Chinese bronzes. They may be thin and enamel-like, moderately encrusted, or massive, and they may vary in colour from white or the palest turquoise to the positive greens and blues characteristic of malachite and azurite; a warmer tone is contributed by the presence of cuprite in the lower layers and when exposed at the surface may add a colour varying in hue from pale ochre to deep reddish-purple. In advanced stages of corrosion the remaining metal may be cracked and frail, but the progress of mineralization has not always been so destructive, and specimens have survived for 3,000 years that are beautifully patinated and can still be described as robust.

Chinese bronzes seldom require chemical treatment to ensure their preservation. They may sometimes be improved in appearance by mechanical treatment to remove mud or gross incrustations and reveal concealed features of ornament or inscription. Indeed, the bronzes that reach Western markets have sometimes a mineral surface suggesting that a form of mechanical treatment has already been applied to remove incrustations and expose to view the more subtle hues of the under layers and, in particular, the pale turquoise layer of tin oxide. This oxide, which, in the absence of chloride, is usually laid down as a thin coherent film, is normally white, but as it is very readily stained with copper compounds, it forms the basis of smooth shiny coloured patinas that are stable and much admired and sought after by collectors.

The speculum alloys are characterised by brittleness and this ac-
counts for some of the features of corrosion peculiar to the mirrors. When the brittle core is unable to adjust itself to the increasing strain resulting from expansion accompanying the change from the metallic to the mineral condition, cracks begin to make their appearance. Sometimes they are superficial and hair-like (Pl. 32), in which case they tend to become centres of corrosion, from which copper compounds spread out over the adjacent metal without necessarily attacking it. The rate of decay is slow, however, compared with that found in bronzes containing less tin. Sometimes corrosion takes place in isolated spots, and when this happens in the case of speculum the surface of the metal becomes cracked around the spots in a series of small concentric rings, so that each spot looks like a tiny oyster-shell, and further cracking may even lift the spots clear of the surface where they remain as hard excrescences (Pl. 33). Such pitting is characteristic of speculum metal that has at one time been highly polished; it may be taken as indicating considerable antiquity in the specimen, and there is no evidence of any change in the appearance of the spots under museum conditions.

DECORATION IN GOLD, SILVER, AND TIN

The conservation of copper alloys is often complicated by the presence of decorations in other metals, chief among which are gold, silver, and tin, and it may be as important to preserve as much as possible of these other metals as to preserve the copper alloy itself. Decorations may easily be lost in a heavy incrustation of corroding copper and it is very important, therefore, to be on the look-out for evidence of ornament, as this might be of great archaeological significance. The discovery of such decoration will obviously have an important bearing on the method of treatment that is employed in conservation.

Copper alloys may be gilded with gold leaf or gold amalgam. Whichever process has been adopted there is always a problem when the metal underlying the gold corrodes, because the gold becomes covered with copper compounds so that its presence may not at first
be recognized. Gold is insoluble in the chemical reagents normally
employed for cleaning bronzes, but if these reagents should attack
the corroding bronze lying underneath the gold, there is nothing
to prevent the gold being lost. If, from a preliminary examination,
the presence of gold is suspected, it is preferable to deal with the
gold first and the bronze, locally if necessary, afterwards. A useful
reagent for clearing the gold is weak nitric acid (1 per cent.), as this
softens carbonate incrustations and enables the extent of the gilding
to be assessed. Drops of the acid are applied with a match-stick to the
encrusted gold, preferably under a binocular magnifier, and absorbed
with the dissolved incrustation in filter-paper; to assist the action
of the acid, any insoluble minerals are cleared away, little by little,
with the help of a needle. Working in this manner it is possible, in
time, to expose all of the gilding. As a final operation, the gold may
be fixed with lacquer after washing and drying.

Examples are found among Chinese bronzes of the T'ang and
later periods where the film of gold is coherent and alloyed to solid
bronze by the mercury gilding process. In these cases, alkaline
Rochelle salt may be used as the cleaning-agent. Where there is gold
embellishment and the underlying base metal has expanded by oxi-
dation, however, the gold may be porous, and an object in such a
condition cannot safely be treated with Rochelle salt or, indeed, by
any other chemical process as any chemical action on the base metal
would be likely to loosen the gold which would float off. Mechanical
picking has to be resorted to and is carried out under the microscope,
particular attention being given to base metal deformations with a
view to recovering as far as possible the original shape and the
artistic appearance of the object.

Silver and tin decorations on bronze survive without corroding,
but in each case preservation may be difficult. Reference has already
been made (p. 189) to the cathodic protection afforded to silver by
contact with copper in presence of an electrolyte, and to the fact that
a base-silver alloy having a green incrustation may be mistaken for
bronze and be dealt with as such (p. 220). When white metal is
observed in treating an object having a green incrustation, the action
should be arrested by washing the object with water. The partially cleaned specimen should then be closely inspected with a magnifying glass to determine whether there is any pattern-work before deciding what modification of treatment, if any, is necessary in the circumstances. If the white metal proves to be silver (as can be determined by its being stained black by a drop of warm ammonium sulphide),¹ the probability is that warm formic acid, applied locally to the white metal, will be the most suitable treatment, but mechanical methods should be exploited as well. When a bronze is embellished with silver it is generally desirable to preserve the patina of the bronze and in such cases it may be better to switch over to sodium sesquicarbonate. If the bronze has been reduced, it should be boiled in distilled water for some hours at the conclusion of the treatment to darken it, if necessary, so that the dark metal will act as a foil for the silver.

Should the white metal prove to be tin it is likely to be in the form of a thin coating and in this case reduction methods would be inadvisable. Tin is found on bronze cooking-vessels, on ornaments such as fibulae and in association with red ‘sealing-wax’ enamels and inlays on the escutcheons of Saxon hanging bowls, while, in China, it was used for elaborate inlays on Han bronzes—chariot fittings, belt fasteners, and sword guards. On all of these objects a heavy incrustation of copper minerals is best removed from a film of metallic tin by the application of mechanical methods of treatment—by the use of needles working under a binocular microscope. On the other hand alkaline Rochelle salt may be made to give satisfactory results but there is then no guarantee that all of the tin will be preserved when it is attached to metal that has already corroded.

Extreme caution is necessary when beginning work on freshly excavated material. This is well illustrated by some of the objects from the ship burial at Sutton Hoo in East Anglia, a notable example being the large dragon from the shield (Pl. 23) which was

¹ A more precise test for silver is to place a drop of 15 per cent. sulphuric acid on the metal and a tiny crystal of chromic acid in the drop and wash off after one minute. A red stain (silver chromate) left on the metal indicates that silver is present. The stain may be easily removed.
concealed under a green muddy incrustation. The mud was removed with the aid of a small stencil brush moistened with a little detergent, and the object was found to be gilt-bronze having local areas washed with tin, and it was inlaid with garnets; in the course of operations it was revealed that the tail of the dragon was an ancient restoration composed of chalk and glue overlaid with gold leaf. Here any treatment involving immersion in an aqueous solution would have been disastrous as the chalk and glue ground would certainly have been softened or damaged.

When dealing with fine metal-work that is decorated with gold, silver, or tin, dry or semi-dry methods of cleaning should always be applied in the first instance. There remains the possibility, however, that wet methods may ultimately have to be used to ensure permanent conservation. Also, whatever standard form of treatment is applied, it may be necessary to introduce modifications at any stage of the work to suit the particular requirements of the specimen.

The restoration of an Egyptian bronze statuette of Isis and Horus (Pl. 34) may be cited as an example of how technique has sometimes to be modified to meet individual requirements. This bronze was heavily encrusted, the white patches showing in the first photograph being an indication of bronze disease. Preliminary examination revealed that the eyes were inlaid with thin sheet gold, but the gold was considered to be sufficiently firmly attached to remain unaffected by the reduction of the incrustation.

Treatment was begun by immersing the bronze in a solution of alkaline Rochelle salt. The solution was changed frequently with intermittent brushing, until the green coating had all disappeared and had given place to reddish-brown cuprite. This took three days and details of the features were now becoming apparent, as shown in the central illustration. Mechanical cleaning was then carried out very thoroughly prior to soaking in sulphuric acid (15 per cent.); the acid treatment went smoothly and seemed to remove all the cuprite. Thus, a remarkable improvement in appearance was achieved in a comparatively short time, but it was clear, having regard to the coarse incrustation that had been removed, that the metal must be in
a highly porous condition and the entire elimination of salt would be difficult.

The bronze was therefore placed in a nylon bag (as a precaution lest the gold should fall from the eyes) and reduced continuously for seventy-seven hours in the electrolytic tank using a solution of caustic soda as the electrolyte, with periodic brushing. This treatment disclosed the existence of an inscription around the base. The inscription was cleaned out as far as possible with a needle, and the bronze was then ready for washing. Washing was carried out by the method described on p. 197 and had to be continued for six months before the final traces of chloride were eliminated. The specimen was dried, and the gold on the eyes which had survived perfectly was burnished. Although the bronze was now stable and clean, the appearance was unsatisfactory, as the metal had a mottled pale straw colour, and this made it unattractive as a museum exhibit. This discoloration was dealt with by the device of applying warm Plasticine as a coating all over the bronze and leaving it there overnight. (Only some brands of Plasticine are effective, and these contain traces of sulphur compounds which impart just the right degree of tarnish.) The metal had now a rich brown-black colour. It was washed with methylated spirit to remove grease, and lacquered with Ercalene. Finally a little chalk was rubbed into the inscription so that the characters would be more apparent against the darkened surface of the metal. The restoration being completed, the object was remounted on its plinth.

An altogether simpler case was that of the heavily mineralized bronze head shown in Pl. 35. Although swollen and cracked, the metal was so substantial that it could be reduced by covering it with granulated zinc in an iron basin and boiling for two hours in strong caustic soda solution. The bronze was then removed and brushed with a steel-wire brush. This revealed ornament that was rendered more precise by further reductions in a fresh soda solution. Washing proved in this case to be a simple matter, and the specimen was eventually dried, and coated with a nitrocellulose lacquer. The discovery of cartouches is always interesting, but in this case the interest was intensified by the fact that the cartouches revealed by
the cleaning contained the names of a hitherto unrecorded Ethiopian king (Kheperkari Alentamani). The presence of the vulture head-dress identifies the head as being that of the queen.

CRUSHED COPPER OBJECTS

When thin copper alloys are buried in the ground, they are liable to be crushed by the weight of earth that covers them. Before any attempt is made to restore the shape, the metal must be softened by heating and plunging into cold water. Over-heating will cause extensive oxidation of the surface and it is better to soften little and often during the bending process rather than attempt to achieve too much in one single operation. Pls. 36 A and B show how a shield boss of interesting shape was restored and mounted. Where there are several dents, the order in which they are removed is sometimes of importance and the operator works, as a rule, from the main body of metal towards the extremities, endeavouring to recover the shape with the minimum of movements of the old metal. As for tools—pin vices, wooden cramps, clothes-peggs are all useful, but where the metal is thin enough, the most sensitive control is that given by the hands, the softened metal being pressed against shaped wooden stakes held firmly in a vice. Cracks may be reinforced at the back with soft solder (p. 286).

A novel type of problem is presented when only fragments of a metal object remain. If, as in the case of the Saxon hanging bowls (Pls. 37 and 38), the fragments are large enough to allow the original shape to be established with certainty, it may be worth while to construct a model from a transparent plastic substance following the contours exactly so that the fragments of bronze can be attached to it in their appropriate positions. The first requirement is to model the bowl in clay in the inverted position and upon this to make a strong plaster cast. This cast may then be used as a negative and coated internally with a thermoplastic material such as Perspex, a technique depending on the fact that sheet Perspex may be softened by heat and blown into the required shape by compressed air. Some
shapes can be made up more satisfactorily in the cold by turning a series of Perspex rings on the lathe and building them up from the base upwards, the parts being cemented together with Perspex cement or with glacial acetic acid. In working Perspex it is difficult to make long joints perfectly; when solvents are used, these evaporate and there is a tendency for air to be sucked in, thus making the joint conspicuous. The most perfect joints are made by firms specializing in heat-welding by supersonic diathermy. After the Perspex shape has been made, the old metal fragments are easily attached to it using either Durofix or chloroform.

When a large thin bronze vessel is patinated but disfigured by a hole, a very effective repair can be carried out by a method devised by Murray Pease of the Metropolitan Museum of Art, New York. According to this process, a piece of felt is blocked roughly to shape, soaked with polyvinyl acetate, and, when partially dry but still pliable, it is moulded to conform with the required contours. It is then fixed inside the vessel and after the adhesive has dried, the outside is made good to match adjacent metal in colour and texture.
A. As excavated

B. After restoration

36. EARLY IRON AGE BRONZE SHIELD-BOSS FROM ANGLESEY
37. BRONZE HANGING BOWL (SUTTON HOO)
Shape reproduced in Perspex by the blowing technique

38. BRONZE HANGING BOWL (SUTTON HOO)
Shape reproduced from a series of Perspex rings of appropriate radius
<table>
<thead>
<tr>
<th>No.</th>
<th>Treatment</th>
<th>Solvent(s)</th>
<th>Reducing Agent(s)</th>
<th>Protective Finish(s)</th>
<th>Method of Darkening</th>
<th>Consolidant(s)</th>
<th>Storage and Exhibition</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>No treatment</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>2</td>
<td>Warm soapy water</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>3</td>
<td>Mild abrasives; pumice or emery powder; metal polishes; Dura-glue; special inpregnated cloths</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>4</td>
<td>Sulphuric acid (50 per cent.)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>5</td>
<td>Nitric acid (10 per cent.)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>6</td>
<td>Alkaline Rochelle salt (5 per cent.)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>7</td>
<td>Sodium hexametaphosphate (5 per cent.)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>8</td>
<td>Alkaline Rochelle salt containing 10 per cent. of 20 per cent. hydrogen peroxide.</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>9</td>
<td>Zinc and sulphuric acid (60 per cent.)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>10</td>
<td>Electrolysis in caustic soda.</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>11</td>
<td>Zinc and ammonium sulphate.</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>12</td>
<td>Electrolysis in caustic soda.</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
CHAPTER XII

LEAD, TIN, AND PEWTER

Lead and tin are soft white metals, and though very similar in appearance, they are easily distinguished by their relative density—lead being almost twice as heavy as tin—and by the fact that clean lead makes a black mark when rubbed on white paper and tin does not. Lead is subject to corrosion in cases where tin is resistant and many organic acids which attack lead have no action on tin. Tin can be used with perfect safety for coating copper cooking-vessels and, in the form of foil, as a hygienic wrapping for foodstuffs.

Tin becomes oxidized, however, as a result of prolonged burial in the ground. In these circumstances the greyish film of stannous oxide which initially forms on the metal gradually gives place to white stannic oxide. The characteristic corrosion product of lead is the basic carbonate which is also white, but the two are readily distinguished by applying a drop of acid which causes the lead incrustation to effervesce, whereas the stannic oxide is unaffected.

Alloys of lead and tin are known as pewter, the hardest variety of which contains about 80 per cent. of tin and 20 per cent. of lead. Roman pewter contains about 80 per cent. of tin, but the proportions of tin and lead in pewter have been found to vary considerably throughout the centuries, owing to scrap metal having been so often melted down and refashioned. Modern pewters are in a different category. Lead is usually absent, its place being taken by antimony and possibly some copper, with the result that these alloys are much harder and less subject to oxidation than the traditional alloys.

LEAD

Museum objects of lead are normally covered with a thin film of oxide. When this has grown slowly in pure air, it has a dull grey
appearance, and acts as a protective patina: when the film of oxide has grown in impure air (air that is contaminated with traces of organic acids, paint fumes, &c.) it is discontinuous and non-protective; a certain milkiness may then make its appearance on the surface, and in time active corrosion breaks out with the formation of basic lead carbonate. The carbonate is loosely adherent and the corrosion is accompanied by a considerable increase in volume. It is for these reasons that leaden objects often suffer serious disfigurement unless the corrosion is checked in the early stages.

When leaden objects are excavated from the ground they are commonly found to have a white incrustation. This is composed of lead compounds formed by chemical action between the metal and saline matter, and also by the action of oxygen and carbon dioxide dissolved in the ground water. Even when such an incrustation appears to be stable, it is often so unsightly as to be unacceptable on specimens for study and exhibition.

The aim of laboratory treatment is therefore either to arrest corrosion, or to improve the appearance of the lead. Carbonates and other salts must be removed, and then the cleaned metal is preserved either by exposing it in an uncontaminated atmosphere so that it will build up its own protective film of oxide (see above), or, more usually, by sealing the surface with a film of wax. A serious complication arises from the fact that the expansion of the surface on corrosion causes micro-cracks to form in the underlying metal, and these have to be cleared of impurities if the results are to have any permanence.

**Electro-chemical and electrolytic reduction**

While the standard reduction methods apply in the case of lead, they are not the safest processes to adopt as lead is slightly soluble in caustic soda (unless it is cathodically protected) and prolonged immersion may result in the loss of fine detail. The same type of accident might happen during electrolysis if the electric current were switched off while the leaden object was immersed in alkali. Reduction is useful, nevertheless, for dealing with odd specimens where fine detail is absent, and especially with large objects that are heavily encrusted.
Special care is required to wash away the last trace of soda from the metal, as, if alkali is left in the porous surface or in cracks, the cleaned lead will eventually become white.

Washing lead free from caustic soda. Washing cannot be carried to completion by using cold water alone, nor is it possible to be certain that the lead is free from alkali unless indicators are used, the most useful in this case being thymolphthalein and phenolphthalein. The procedure is carried out in two stages as follows: First, place the object in changes of very hot tap water\(^1\) to remove the bulk of the alkali, and test the waters from time to time by adding a few drops of thymolphthalein.\(^2\) When the indicator no longer turns the water blue this is a sign that the first stage of washing is complete. The lead is then passed through a series of baths of hot distilled water (that has been freshly boiled to expel dissolved oxygen and carbon dioxide) and this is continued until the washings no longer turn pink when tested with phenolphthalein. The object is then removed from the hot water by tongs (it should not be handled until after waxing), mopped dry with a soft clean cloth, and passed through a bath of 95 per cent. alcohol (industrial methylated spirit). The lead is taken from the alcohol bath by tongs, shaken free from liquid, and placed in a bath of molten beeswax, or low-melting (49° C.) paraffin wax (which has the merit of giving a less brittle film at room temperature), and kept there for a few minutes at a temperature above 100° C. to remove residual traces of moisture. It is withdrawn from the wax, shaken, and placed on blotting-paper to drain. A domestic hair-dryer is very useful for keeping wax molten on the object so that any unnecessarily thick surface deposits may be wiped off.

\(^1\) Waters from peat contain humic acids, &c., and are unsuitable as they tend to dissolve lead. The water from a lead piping system may safely be used for washing.

\(^2\) Phenolphthalein is too sensitive for use with tap water, as some drinking-waters are faintly alkaline in their action, or develop alkalinity on heating, and in such cases the test would not be discriminatory. Thymolphthalein, on the other hand, is not so sensitive to faint alkalinity. The indicators remain colourless in acid and neutral solutions, but give the following colour reactions in the presence of alkali:

- **Thymolphthalein**: Blue (pH above 9·3).
- **Phenolphthalein**: Red (pH above 8·3).
thin film of wax that remains permeates the porous surface of the metal and acts as a protective coating.

*Treatment of leaden objects with acids*

As the white incrustation that develops when lead corrodes is essentially carbonate, it is readily dissolved by acids.

Dilute nitric acid was formerly recommended for removing incrustations of lead carbonate from antiquities, the cleaning effect being immediate. It seemed that excess acid could be neutralized with alkali, such as caustic soda, and the soluble residue removed with water, leaving a clean metallic surface which, on drying and sealing with wax, would remain permanent under museum conditions. This, however, has not proved to be the case. It appears that, after a lapse of several years, lead so treated acquires a milky appearance, and the method is no longer considered satisfactory.

Experiments using acetic acid have given even less permanent results judged on a time basis. This is not surprising as lead corrodes very readily in presence of acetic acid vapour and carbon dioxide. This, indeed, forms the basis of a commercial process for the manufacture of white lead.

Caley\(^1\) has reported favourably on a method of treating lead by hydrochloric acid and ammonium acetate, and this is of special interest as being in the nature of a long-term experiment. Fifty-six leaden objects from the Agora were dealt with in 1937, and seventeen years later these were found to be in perfect condition. For this method two solutions are required composed as follows:

*Dilute hydrochloric acid:*

100 ml. concentrated hydrochloric acid (Analytical Reagent, Sp. Gr. 1.19) in a litre of distilled water.

*Ammonium acetate solution:*

100 gm. ammonium acetate in a litre of distilled water.

Besides these it is necessary to have a quantity of distilled water that has been boiled vigorously a few minutes prior to use (to expel dissolved gases) and then protected from the air.

The procedure is as follows:

1. *Hydrochloric acid bath.* Soak object in some fifty times its volume of acid till effervescence ceases (time 1–2 hours, or overnight). Drain off acid and place object for a few minutes in some hundred times its volume of hot distilled water. Decant and repeat the washing twice.

2. *Ammonium acetate bath.* Soak washed object in some twenty-five times its volume of warm acetate solution till no more corrosion products visible on surface of lead. This should take about 1 hour and should not be prolonged beyond 2 hours at normal room temperature. Wash as follows: place object in some 100 times its volume of cold freshly boiled distilled water. Time ten minutes. Decant and repeat the washing three times.

3. *Drying.* Dry at room temperature without heat, or dry through alcohol in the usual way.

4. *Waxing.* Immerse the object for a few minutes in molten paraffin wax at 100° C.

The advantage of using ammonium acetate is claimed to be two-fold; it dissolves lead dioxide which is insoluble in hydrochloric acid, and it acts as a buffer to protect the lead from the action of any trace of hydrochloric acid that may remain from the first bath.

Note: This method is recommended for leaden objects only and not for tin, as tin is soluble in hydrochloric acid!

*Use of ion-exchange resins in the treatment of lead*

A notable contribution to the preservation of leaden objects was made in the British Museum Research Laboratory when it was discovered that lead carbonate and chloride could be removed from the metal by the use of ion-exchange resins.¹

The ion-exchange principle has long been known as a means of softening hard water, i.e. removing dissolved calcium salts which form insoluble compounds with soap. It was found that hard water is softened by passing it through a column containing certain naturally occurring complex sodium silicates (zeolites). More recently special synthetic resins were introduced which carried out the exchange

more efficiently. The function of the ion-exchanger is to abstract the calcium ions that are in the hard water and replace them with sodium ions. When the resin is saturated with calcium ions it can be regenerated by treatment with brine which, in turn, replaces the calcium ions with sodium ions so that the resin can be used again. It is claimed that the process of regeneration may be repeated indefinitely.

The above principles may be employed in cleaning leaden objects that are encrusted with carbonate. The process is very simple. All that is necessary is to place the corroded leaden objects in contact with granules of an appropriate ion-exchange resin,\(^1\) cover with distilled water and keep hot, changing the resin if necessary until the white incrustation disappears. Metallic lead is unaffected, but the incrustation of basic lead carbonate which gradually dissolves is removed from the system; the lead ions are taken up by the resin in exchange for hydrogen, and carbon dioxide is evolved from the hot liquid. The ion-exchange method is a great advance on previous methods that have been used for the treatment of lead, as no chemicals have been introduced into the system, and therefore subsequent washing of the lead is unnecessary. Although the initial outlay in resin is rather expensive, the method is economical because the resin may be regenerated after use by treating it with nitric acid to dissolve out the absorbed lead, and washing it with distilled water till the washings are neutral to litmus paper: the resin is then in a condition to be used again. The apparatus is shown diagrammatically in Fig. 10 and the column used in regenerating the resin may be seen (middle left) in the general view of the Laboratory, Pl. 39. The method has

\(^1\) ‘Amberlite IR 120’, British Drug Houses Ltd., Poole, Dorset.
the further advantage that, when specimens are badly swollen and the cracks filled with a resistant white deposit of basic carbonate, it is not essential to remove this, as in using ion-exchange resins the carbonate has not been contaminated with chemicals. Nevertheless it will usually be desirable, for the sake of appearance, to remove all of the carbonate if possible.

In applying the ion-exchange method the following points should be noted. Ion-exchange can only be conducted in the presence of water, and as distilled water has a certain solvent action on lead, the rate of cleaning should be speeded up whenever possible. To facilitate this, any hard incrustations of carbonate occurring in cracks should be opened up by picking with a needle. There is little likelihood of serious damage by water, however; at the worst there may be a tendency for ornament in relief to be rounded, but the effect is very slight and could only be of significance in cleaning leaden coins where the inscription is barely legible.

The use of ion-exchange resins is advocated for the treatment of small objects—coins, medals, weights, badges, sling stones, &c., but when the objects are heavily corroded, or when they have been treated with shellac or other varnish, it has been found advantageous to give them a preliminary treatment for a short time in the electrolytic tank.

Storage of leaden objects

The susceptibility of lead to attack by organic acids is one of its principal characteristics. The effect of carbonic acid in presence of acetic acid vapours and of humic acids has already been noted. Tannic acid is equally corrosive, and an exudation of tannic acid from oak cupboards and drawers is credited with causing the intensive corrosion to which leaden objects are subject when stored therein. Coins and medals, beggars' badges, leaden crosses, inscribed leaden scrolls, &c., have been found converted completely to amorphous masses of white powder as a result of their being kept during past generations in oak cupboards. Nor are other woods necessarily innocuous. To be harmless, the timber would require to be well seasoned, and if any
LEAD, TIN, AND PEWTER

doubt exists, it should be sealed with a resistant lacquer. This applies particularly to nests of shallow drawers where there is a large surface of wood and limited ventilation (Pl. 40). Well-seasoned mahogany is probably the safest wood to use for cupboards and cases designed for the storage of leaden objects.

TIN

Tin has been described as being very stable under normal atmospheric conditions, but when buried in the ground and exposed to moisture and oxygen over a long period it loses its lustre, becoming granular and grey in appearance. This change is due, in the main, to oxidation. The oxide layer may have a greenish hue when traces of copper are present. A more advanced stage in the corrosion is the conversion of the grey stannous oxide to a higher form, stannic oxide, which is nearly white in colour, but it is probable that this only occurs as a secondary reaction in the presence of soluble salts.

Tin is often found as a thin silvery layer on bronze, especially in jewellery and suchlike. This has been applied to the bronze in much the same way as a plumber applies soft solder. The bronze has been cleaned and fluxed to prevent it oxidizing, heated to a temperature slightly above the melting-point of tin (232° C.), and the molten white metal then wiped over it with a rag charged with tallow. Thin washes of tin are easily destroyed both by acids and alkalis, and when found on ancient metal objects these should be cleaned by mechanical methods if possible.

Solid objects of tin, e.g. plates, jugs, coins, and medals, are cleaned either electro-chemically or by electrolytic reduction in the usual caustic soda electrolyte. Corroded tin coins have been found to respond well to electro-chemical reduction in the cold, using either zinc, aluminium, or magnesium powder and caustic soda.

The problem of corroded tin is often confused with an allotropic change that takes place in the metal at low temperatures, known as ‘tin pest’. This is, however, a different phenomenon to that of metallic corrosion. It is speedier in action, and more catastrophic in its results,
for the metal changes its crystalline state and falls into a coarse grey powder. The destruction of everything of archaeological interest is thus complete, and restoration is impossible. It is fortunate that true ‘tin pest’ occurs very rarely, and only in exceptional circumstances has it been identified with any certainty as occurring in the case of museum objects. When an object of tin shows signs of decay these are almost invariably the result of metallic corrosion, and are therefore amenable to treatment.

Special care is required in dealing with heavily oxidized tin that bears incised ornament because in such cases the detail may be entirely in the oxide layer, which must therefore be preserved, and, of course, reduction would destroy this. A case in point is the Hartogs plate in the Rijksmuseum, Amsterdam. This famous plate was recovered after more than eighty years of exposure on the coast of Australia, and it had been supposed to be a victim of ‘tin pest’ because of its frail condition. It was shown, however, by micrographic examination that the weakness had resulted simply from prolonged metallic corrosion. Oxidation of the surface had been accompanied by a considerable increase in volume, and this had resulted in strains that tended to make the oxide layers split away from the core of metal, threatening the loss of the inscription. In such cases, reduction is out of the question, and the only possible treatment is to consolidate with a suitable adhesive such as Durofix. Fortunately, in the case of the Hartogs plate, there was no indication that corrosion was still taking place, and chemical treatment was therefore not required.

In the case of tin coins that have been struck and bear a raised inscription, however, conditions are different. During corrosion the inscription is preserved at the expense of the worked metal constituting the flat part of the coin. This is an interesting case of the debasement of a metal in those parts that have been subjected to the greatest mechanical strain. Thus in struck coins the inscription may be clarified by careful reduction as was done in the case of corroded tin coins from Malaya\(^1\) where good results were obtained by using magnesium powder and caustic soda.

---

Embedding lead and tin in plastic materials

When a lead or tin coin is of unique interest but is so weak or fragmentary that the remaining pattern is likely to succumb to any process of cleaning, the only way of saving it is to embed it in a transparent plastic. This can be done with Marco Resin\(^1\) S.B. 26 C. which sets at room temperature without the use of pressure. Purves and Martin\(^2\) have described the method of embedding biological material in this resin, and Miss Plesters\(^3\) has given full details of its use for embedding paint fragments that are to be prepared as cross-sections. When it is desired to preserve coin fragments, the most convenient shape of the polymerized resin block is one that may be kept in a coin cabinet, i.e. a disk, which should be highly polished to offer minimum interference with vision. As there is a certain shrinkage when the resin sets, it is very difficult to prepare such a polished disk by simply allowing the resin to set in a mould with polished plain faces. Hence it is usually necessary to work and polish at least one of the flat faces. This may be a lengthy operation. The Marco resin process is unsuitable where copper is present, as copper produces a green stain in contact with the resin.

If an electrically heated press is available, a thermoplastic resin may be used, and the embedding operation is greatly facilitated, since blocks may be prepared of uniform size that only require final polishing. A methacrylate resin called Transoptic\(^4\) has been found to give reproducible results under these conditions. The powdered plastic should first be spread on white paper and examined carefully, any dust or foreign matter being removed. The object is then washed in chloroform, air dried, and thoroughly dried over silica gel. (Avoid touching with the fingers after washing with chloroform.) A circular mould, \(1\frac{1}{4}\) inches in diameter, is then partly filled with the powder and the object placed gently on top. An equal amount of powder is added and the plunger of the mould inserted. The mould is then

---

placed in the press and heated gradually to 150°C. without applied pressure. Heating is discontinued, the pressure raised to 2,000 lb. per sq. inch, and the mould cooled to 40°C. while maintaining the pressure. Cooling may be hastened by a water-cooled jacket. The clear resin block containing the coin is then removed and may be finished off by polishing with I.C.I. Perspex Polish, grades 1, 2, and 3 being used successively on Selvyt cloths.

If at a later date it is desired to remove the embedded object, the block is soaked in chloroform. Transoptic swells and dissolves; Marco resin is crazed, and in each case the object may easily be removed without any risks of damage. The thinnest lead foil has been mounted in Transoptic and demounted without distortion.

PEWTER

Pewter vessels depend to a large extent upon their patina to give them an appearance of dignity and maturity; chemically cleaned pewter has the appearance of lead and so in the conservation of pewter objects it is essential to avoid overcleaning. Where the metal is merely stained or covered with a film of oxide and it is a question of improving the appearance, a rub with a mild abrasive such as rottenstone applied with an oily rag will be all that is required.

When pewter carries a fine design hidden by a thin deposit of lead carbonate, as in the case of the Bastille medal (Pls. 41 A and B), the best treatment to adopt is, without doubt, the ion-exchange process described (p. 263). This removes the corrosion product while preserving the finest details of the design.

Old pewter is occasionally found to have wart-like growths of incrustation on the surface, possibly arising as a result of localized contamination with salts. If these spots have a hard skin and are not showing signs of active corrosion, it may be safer to leave them alone than to attempt to get rid of them (e.g. by grinding) because the material underneath may be mainly tin oxide which is often very crumbly. If exposed to the air, moreover, the under-layers of such spots may begin to corrode actively. If corrosion is already active and
treatment essential, electrolytic reduction may be carried out. This will remove all corrosion products but the appearance of the object will suffer because the surface will be left full of small depressions or pock-marks. In such a case the only course that can be recommended is to fill the depressions and make good the surface with wax.

Reduction is the safest procedure, however, when searching for the original pewter marks under a layer of heavy incrustation. Such marks are easily damaged by scraping as the underlying metal may be softer than the incrustation. Tankards and flagons are often marked beneath the lip, and care should be taken with capacity marks on the side of vessels, and makers’ and owners’ marks under the base. When the site of the mark is known or suspected, local reduction with zinc and caustic soda may be all that is required (cf. p. 193).

An example of the unexpected discovery of a pewter mark occurred in the case of a plate brought up from the sunken galleon in Tobermory Bay. This had a thin greyish-white incrustation and was apparently devoid of documentary interest. A short reduction in the electrolytic tank, however, revealed a stamp on the under surface of the plate, and it was sufficiently legible to be recognized by the Duke of Argyll as indicating that the plate was of Portuguese origin.
TABLE V. LEAD AND TIN ALLOYS

Lead and tin alloys

Uncorroded

Stable

Appearance satisfactory

No metal remaining (worthless)

Corroded

Heavily corroded or laminated

Heavily corroded

Lightly corroded or spotted

Corrosion active

Metal remaining as a solid core

1. No treatment.
2. Washing: (a) Freshly boiled distilled water; (b) Freshly boiled potable water from a lead pipe system.
3. Ion-exchange method.
4. Solvents: Hydrochloric acid treatment followed by ammonium acetate.
5. Reduction: (a) Zinc and caustic soda; (b) Electrolysis in caustic soda. (Lead being soluble in caustic soda, these processes are applied for short duration only.)
6. Protective finishes: (a) Paraffin wax dip; (b) Impregnation with a plastic substance or embedding.
7. Mechanical: (a) picking, (b) chipping, (c) grinding, (d) scratch-brushing, (e) grit-spraying, (f) and burnishing.
8. Storage and exhibition. Avoid the use of oak furniture or unseasoned wood.
CHAPTER XIII

IRON AND STEEL

Iron occurs in the metallic condition in meteorites where it is associated with small quantities of nickel, cobalt, copper, &c., and the oldest iron objects that have been subjected to chemical analysis have been shown by the presence of these trace elements to be of meteoric origin. Terrestrial iron is of rare occurrence as the metal is so readily oxidized and converted to minerals, and these are abundantly distributed throughout the earth's crust.

Objects of iron and steel provide some of the most intractable problems for the conservator, because of the variety and complexity of their corrosion products. Iron corrodes easily, the corrosion products are unsightly, and the swelling and deformation of the objects may be severe.

Rusting. Iron is readily attacked by oxygen in presence of moisture to form rust—a name derived from the characteristic orange and red compounds that appear as the first products of corrosion. These consist at first of a mixture of ferrous and ferric hydroxides, but on further oxidation the rust becomes substantially a hydrated ferric oxide, in which some carbonate is usually present as well.

This might seem to be fairly straightforward, but when salts are present that can act as electrolytes, the chemical reactions that cause corrosion are reinforced by electro-chemical reactions, and mineralization is greatly accelerated. It was noted in introducing the subject of corrosion (p. 190) that when a metal was partially protected from aeration by patches of oxide, &c., certain areas became anodic and others cathodic so that, in fact, the corroding metal behaved in the presence of an electrolyte as if it were a number of tiny galvanic cells. This is a feature of iron corroding in the presence of sodium chloride, the anodic areas dissolving to form ferrous chloride whilst
the cathodic areas become alkaline due to the formation of sodium hydroxide. A film of hydrogen gas begins to accumulate on the cathodic areas, and as this has a comparatively high resistance to the passage of electric currents, its presence tends to slow down the reaction. In presence of oxygen, however, the hydrogen is continuously removed, the two gases reacting to form water or hydrogen peroxide. Thus the electrolytic action is enabled to proceed and it continues until the area is covered with a deposit of rust formed by the interaction of the ferrous chloride and sodium hydroxide. New areas then take on the roles of anode and cathode and since sodium chloride is regenerated at the same time as the rust is being deposited the cycle of reactions can continue. This is an important factor in the conservation of iron as it means that we are faced with the same conditions that confront us in the treatment of ‘bronze disease’, namely, that stability cannot be assured until all the chloride has been removed from the corroding object.

It has been mentioned that the accumulation of a film of cathodic hydrogen slows down the electro-chemical action, but that the action of oxygen in breaking down this film makes it possible for the corrosion to proceed. It does not follow, however, that, in the absence of oxygen, iron will necessarily remain uncorroded. The film may be broken down in another way, namely by bacterial action, and this introduces a novel factor that is important in studying the corrosion of ferrous metals. Iron and steel are often found to be very badly corroded when buried, under anaerobic conditions, in heavy clay containing sulphates. In this case corrosion has been traced to the presence of sulphate-reducing bacteria which act in a two-fold manner: firstly, converting the sulphates to sulphides which attack the iron, and secondly, breaking down the hydrogen film and thus making it possible for corrosion to continue. When corrosion takes place under such circumstances the surface of the iron is found to be covered with a black crust of iron sulphide, and the clay surrounding the object is also stained black. Should there be any doubt about the nature of the deposit, iron sulphide is easily detected by its reaction with acids to liberate sulphuretted hydrogen recognizable by its
40. PILGRIMS' BADGES, BULLAE, CLOTH MARKS, ETC., OF LEAD

These show various stages of disintegration characteristic of the change from metallic lead to a basic lead carbonate.
characteristic odour of rotten eggs. The first of the sulphate-reducing bacteria to be made the subject of laboratory experiment was *Vibrio desulphuricans*, but many new varieties have since been isolated and it is probable that there are several species. This type of corrosion is very prevalent and is a common cause of the destruction of iron pipes buried in clay.

Reference should also be made to the action of sheath-forming bacteria (e.g. *Gallionella ferriginea*) in promoting corrosion by differential aeration. These bacteria form a blistered structure of tubercles that is a common feature of heavily corroded iron, and sulphate-reducing bacteria may also be active in the anaerobic areas beneath this cellular structure.

PRELIMINARY EXAMINATION

When an iron object comes to the laboratory for treatment, a careful examination is necessary to determine the condition of the specimen before a course of action can be decided upon. If a massive incrustation of dry rust is free from chloride, it may be stable under museum conditions, or if rusting has proceeded to the limit and no metal core remains, even if the oxide is swollen and fissured, the specimen will also have reached a stage of stability and no treatment will be essential. On the other hand, if corrosion is still taking place, some process for the removal of chlorides will have to be employed in order to preserve the object. Therefore the first question to decide is whether there are any signs of chemical activity.

An examination is first made of the surface. Corroding iron does not display the range of colour that is apparent in bronzes that are ‘diseased’, but even so areas of active corrosion may often be detected by the fact that they have a slightly different colour and texture to those of the surrounding metal or incrustation. But changes of colour

---

and texture do not necessarily indicate in themselves that corrosion is active. Dampness on the surface of the object is, however, a sure sign, because when chloride is present in contact with iron it gives rise to corrosion products that are hygroscopic and beads of brown liquid may be found on the surface of otherwise dry rust; even a comparatively thin film of rust will show signs of dampness due to chemical activity if chloride is present.

For assessing the internal condition of an iron or steel object, the method of X-radiography is particularly useful. Iron oxides are relatively much more transparent to X-rays than the solid metal, and X-radiographs reveal the distribution and extent of oxidation more certainly and directly than any other method; they enable one to estimate the depth of pitting, and hence to decide on the best method of treatment.

But only in rare cases can radiographic methods of examination be considered an essential and their application in any case is limited by the thickness of the metal. Whether facilities are available for X-radiography or not, a careful examination is always carried out, first with a magnet in order to determine the extent of the metal core, and then by a needle or metal probe, used with the aid of a lens, in order to reveal the extent and condition of the rust layers and discover, if possible, the existence of any decoration hidden beneath the incrustation. Other tests are made instinctively, e.g. the balance of a rusty blade as an indication of the regularity or otherwise of the corrosion, and an estimate of weight for bulk (relative density) which, with the magnet test, should help in determining the extent of oxidation.

**Treatment of Iron Objects**

From the preliminary examination it will be decided in which category to place the specimens, and guidance as to subsequent procedure may then be obtained by consulting Tables VI and VII below (pp. 282 and 290).

Where a substantial core of metal remains in the heart of the rust, reduction methods can be employed without the cleaned object being
mechanically weakened, but if, as in the case of a thin and heavily rusted blade, the metallic core is discontinuous, reduction methods are best avoided.

When a small object no longer has a metallic core it is very brittle, and as any treatment would be likely to weaken the specimen still further, objects in this category are best left alone. Sometimes interesting ornament may be hidden in a mass of rust, and can never be revealed by cleaning because of the frailty of the specimen. In such cases photographic evidence may be made available by radiographic methods, using X-rays or radio-isotopes. At other times a little careful work with a needle under a binocular microscope may succeed in revealing something of special interest. When this can be done without undue risk it is worth while in specimens that are designed for exhibition in public galleries.

Laboratory attention may not be required in order to ensure the stability of a specimen, yet it may be desirable in order to improve the general appearance. Apart from the matter of ordinary repairs, treatment may be required for any of the following reasons—to reveal the original contours or shape; to expose ornament existing in the rust, or decorative inlays that are hidden by the incrustation; to study the method of construction, or, in general, to reveal evidence which, in the absence of such treatment, would be likely to remain concealed. Such evidence may sometimes be obtained by radiography, but more generally by the application of mechanical and chemical methods used either alone or together. Mechanical methods are of special significance in dealing with corroded iron and steel.

Reduction. The standard practice with iron objects is to apply reduction methods whenever possible, i.e. when a good continuous core of metal remains, and there are no complications in the way of inlays or non-metallic attachments. In cases where the surface is not much pitted, electrolytic reduction may be applied forthwith, but when pitting is at all extensive it is advantageous to start with electro-

---

chemical reduction, using zinc and caustic soda, as this treatment is more likely to get at the deep-seated corrosion in the pits and cracks; but a final electrolytic reduction is always worth while, as it has been found to remove most of the residual chloride, and thus cut short the time necessary for washing. This is clearly an advantage in dealing with metals that rust.

Difficulty is sometimes found in making a good electrical contact with corroded iron objects, and it may be necessary, by scraping away the rust, to expose some uncorroded metal for this purpose. Electrical contacts may have to be made in several places in preparing large objects for electrolysis.

As a matter of interest, electrically reduced iron often steams when first rinsed after removal from the tank. This is due to the rapid oxidation of pyrophoric iron, and a case is on record where a large piece of black reduced rust, when picked off, actually glowed and burned through the French polish on the bench! This, however, is very exceptional.

**Caustic soda treatment.** Where chemical corrosion is observed to be taking place on an object that is too frail for electrolytic reduction, it may be treated by boiling the object in several lots of dilute caustic soda solution followed by boiling in changes of distilled water. The results may be satisfactory when the treated object is kept in a dry place, but with this method there is a chance that chlorides may remain in the under layers, and the result of caustic soda treatment is therefore not necessarily permanent.

**Heat treatment.** The rough and ready method of separating a mass of corroded iron objects by the use of a blow-lamp is obviously attended by the danger that specimens may fly to pieces. This is particularly liable to happen in the case of an object having a bubbly surface of ferric oxide in which air and salts may be trapped. When iron is damp or badly rusted, heating is liable to cause a good deal of flaking from the surface, and although it loosens deposits of carbonate, it hardens any clay deposits and makes the rust impervious to the action of solvents. The method, therefore, cannot be recommended without serious reservations.
Use of rust softeners, solvents, and inhibitors. Thin rust spots on burnished steel may often be softened by keeping the steel under paraffin oil for a few hours, and then rubbing locally with worn emery paper of fine grade or a match-stick, but care must be taken to remove all the paraffin oil or fresh rust is likely to be formed. The paraffin oil is removed with a dry cloth, and replaced with a lubricating oil, and this is eventually removed by dry rubbing, any thin residual film of lubricating oil being harmless. The substance known as ‘Plus-Gas Fluid A’\(^1\) is even more effective in softening rust than oil. Unless rust spots are removed, they tend in time to bite deeply into the metal. At these points anodic attack is taking place, and this localized chemical action is a stimulus to greater activity. It is important, therefore, in maintaining collections of armour and the like, to be aware of the danger of rust-pitting, and to take action in good time to remove the rust. Where salts are absent, and atmospheric relative humidities are at 50 per cent. or under, rusting is not a serious problem, but moisture readily condenses on cold metal surfaces, and condensation is always a menace if there should be a sudden drop in temperature. Also in a city atmosphere, condensed moisture might well contain sulphur dioxide which would promote rusting. A smear of lanolin (wool fat)\(^2\) has proved to be a good protective agent for burnished steel, but as it leaves the surface of the metal sticky, it is only of practical value when the steel is protected by being enclosed in an exhibition case. A thin layer of non-sticky wax is to be preferred for museum objects (see Appendix XII).

Certain commercial rust removers come within the category of rust solvents,\(^3\) and these are of special value for derusting objects that are either too heavy or too large and unwieldy for treatment by electrolytic methods. Some have the added advantage of being rust inhibitors, i.e. they leave a protective film on the surface of the metal after treatment. Rust solvents may be applied by brushing, and when

---

\(^3\) e.g. ‘ACP Deoxidine No. 125’, Imperial Chemical Industries Ltd., Paint Division, Slough, Bucks., and ‘Jenolite’, Jenolite Ltd., 43 Piazza Chambers, Covent Garden, London, W.C. 2.
the solvent action has gone far enough, either the surface is wiped clean without washing, or in other cases any remaining chemicals are washed away. In using these commercial preparations the makers’ instructions must be followed implicitly.

In these rust inhibitors the active constituent is often a derivative of phosphoric acid which forms an inert film of ferric phosphate on the surface of the metal. In this connexion it is interesting to observe that objects of iron or steel are occasionally found in the ground in a substantially uncorroded state after years of burial. This may sometimes be explained when they are from a burial site, and have been exposed to the action of phosphates from adjacent bones. The protection given by tannates, which may have their origin in decomposing leather or oak bark, is even more marked.

It happens that caustic soda is also a rust inhibitor (p. 276), and this property gives it an added advantage over acid electrolytes in dealing with ferrous metals. The metal objects must still be washed after reduction to remove dirt and the last traces of chloride, but if the final wash water is rendered faintly alkaline with caustic soda (and the object dried thereafter), rusting will be inhibited.

Rust may be removed by the use of a solvent such as a 9 per cent. solution of oxalic acid, but it is often more convenient to use a complexing reagent, which has the property of combining with the iron dissolved from the rust; the iron is then said to be sequestrated. Some of the most effective complexing reagents are derivatives of ethylenediamine tetra-acetic acid (EDTA products), and examples are the Versenes,¹ a range of compounds that will complex iron under acid or under alkaline conditions. ‘Versene Fe-3 Specific’ is claimed to be the most powerful complexing agent for dealing with iron salts, but ‘Versene T’ is of greatest interest as it can be used in presence of caustic soda which slightly dissolves rust to form ferrites. This is extremely effective on freshly deposited rust, but less so on aged and especially on heated iron oxide. In these difficult cases it is a help to

add sodium hyposulphite. A solution which has been used with effect contained 30 ml. ‘Versene T’, 30 gm. caustic soda, and 6 gm. sodium hyposulphite per litre. The method of application is to keep the rusty object immersed in a hot solution until such time as the rust is dissolved to the required extent, the object being thoroughly washed thereafter and dried.

Protective finishes. Whatever chemical method is adopted in cleaning iron and steel, the subsequent washing is likely to be accompanied by a certain amount of rust discoloration, and when the metal is dry this is removed by brushing, but if washing has not been prolonged, the rust discoloration can be avoided by a final bath in acetone which removes the surface film of water. There then arises the question of what type of protective finish is likely to be most suitable. Choice may be made from a wide range of materials: there are sealing agents, such as oils and fats, waxes or lacquers: there are also chemical inhibitors, one of the most effective of which is sodium benzoate. This may be applied in either a water or glycerine solution of 1.5 per cent. strength for keeping steel bright in moist surroundings. See also Appendix XII.

Certain chemical substances have the property of preventing the corrosion of iron and steel in enclosed spaces without being in contact with the metal surface at all. These are known as Vapour Phase Inhibitors (V.P.I.), but it is not safe to use them indiscriminately, as, however effective they may be with ferrous metal, some have been found to attack copper alloys and others soft solder. If such deficiencies can be overcome, as seems not impossible, vapour phase inhibitors should have wide application in the museum world in helping to preserve objects of iron and steel, but meantime it is likely that the use of wax or lacquer will continue to be the most popular type of finish for archaeological specimens. In some comparative tests conducted at the Ancient Monuments Laboratory,  

1 Formerly known as hydrosulphite. This is not to be confused with photographic 'hypo', which is sodium thiosulphate.


dipping in molten bleached beeswax was shown to be the most effective method of preserving treated objects.

**PATTERNS IN SWORD BLADES, DAMASCENE WORK, ETC.**

A feature of the Japanese sword-blade is the pattern latent in the steel, arising from the method of forging by folding and twisting the red-hot billet of metal according to a pre-arranged formula. A series of intricate cloud, wave, and other patterns may thus be repeated along the length of the blade, and good examples are greatly esteemed by the connoisseur. Should the metal be allowed to become dull the pattern may become indistinct and finally disappear. Reference has been made (p. 206) to the conservation of such weapons. The following note refers to a method of restoring a lost pattern, using the reagent called ‘nital’ prepared by dissolving 1.5 ml. of concentrated nitric acid (Analytical Reagent, Sp. Gr. 1. 42) in 100 ml. of alcohol (industrial methylated spirit). First remove any grease with a soft rag moistened with benzene. Moisten a swab of surgical cotton-wool with freshly prepared nital solution and apply it uniformly to the blade, rubbing backwards and forwards from hilt to point. This is done beside a running tap under which the blade is plunged to arrest the etching of the acid. Sometimes this solution develops the pattern within a few seconds; at other times several short applications may be necessary before the desired result is obtained. The blade should be closely inspected after each washing; as an excessive use of the reagent might blur the pattern, it is very important to avoid carrying the action too far. The blade must finally be washed under the running tap for about half a minute and carefully dried with a soft cloth.

Japanese swords should be kept *slightly* waxed. If the waxing is overdone, there is a possibility that dust will accumulate, and as great care must be taken to ensure that the scabbards are kept free from any dust that might cause scratching, it is better not to wax at all than to apply wax too liberally. The formula using microcrystalline wax given in Appendix XII is suitable. The smallest quantity is brushed uniformly over the burnished metal with a moderately
stiff brush such as is used for cleaning silver. Then as much as possible of this is removed by brushing with a softer brush having longer bristles—the kind of brush that is used for silk hats.

The textured patterns in damascened blades often have the appearance of ‘watered silk’, an effect obtained by long heating and light forging, producing what is known as spheroidization of iron carbide in a pearlite structure. Damascened blades should be treated with the same care as is afforded to Japanese swords.

The coarser pattern effects sometimes found in swords of the eighteenth century, and typically in the Malayan kris, are obtained by variations in relief of the different elements of the pattern. This is attained by an etching process and in general such blades provide no complication as regards preservation.

PRESERVATION OF RUST

When iron is completely converted to massive oxide and no free metal remains, the specimen has been described as stable and no laboratory treatment is required for its preservation. When the rust is in a granular rather than a massive condition, however, there is always the possibility of disintegration due to the crystallizations of salts, and in this case the specimen will have to be washed free from salts before its stability can be assured; the surface may have to be consolidated with nitrocellulose solution before washing as is done in the case of brittle pottery and stone (see p. 299).

Rust should be preserved when its removal might be attended with the collapse of the specimen. It should also be retained if its removal would be likely to result in serious disfigurement. A common example is the spotty surface sometimes found on a reduced iron object; the spots are due to compacted masses of ferric oxide occupying pits in the metal, and if these were dug out, the smooth surface would be made unsightly, and they are, therefore, left in situ. The risk that corrosion might break out again is extremely slight under museum conditions, when salts have been removed as far as possible, and the object has been given a protective coating of wax or lacquer.
### TABLE VI

<table>
<thead>
<tr>
<th>Object</th>
<th>Features</th>
<th>Treatment</th>
<th>Result</th>
</tr>
</thead>
<tbody>
<tr>
<td>Coat of mail</td>
<td>Loose, adherent red rust.</td>
<td>Denaturing solutions considered dangerous and</td>
<td>The cleaned chain had a black appearance and dull shine.</td>
</tr>
<tr>
<td></td>
<td></td>
<td>lightly with wax salve.</td>
<td></td>
</tr>
<tr>
<td>2. Welwyn fire dogs</td>
<td>Large, heavy, and broken</td>
<td>Cleaning in part by electrolysis, in part by</td>
<td>The broken parts reassembled, and details of the structure revealed.</td>
</tr>
<tr>
<td></td>
<td></td>
<td>picking, and in part by pickling.</td>
<td></td>
</tr>
<tr>
<td>4. Stanwick sword</td>
<td>Corroded and encrusted with</td>
<td>Electrolysis.</td>
<td>Shape recovered; object fail but complete.</td>
</tr>
<tr>
<td></td>
<td>iron sulphide, due to action of</td>
<td>Mounted on a base-board having a recess</td>
<td></td>
</tr>
<tr>
<td></td>
<td>sulphate-reducing bacteria</td>
<td>Mounting the sword.</td>
<td></td>
</tr>
<tr>
<td></td>
<td>largely mineralized.</td>
<td>Mechanical treatment.</td>
<td></td>
</tr>
<tr>
<td></td>
<td>2. The nose of hilt largely</td>
<td>Reassembled in such a manner that each piece is</td>
<td></td>
</tr>
<tr>
<td></td>
<td>mineralized.</td>
<td>detachable.</td>
<td></td>
</tr>
<tr>
<td>6. Sutton Hoo shield</td>
<td>Encrusted with carbonate and rust concealing ornament in gold, bronze, tin, and garnet.</td>
<td>Dilute nitric acid (1 per cent.) locally, washing and drying through alcohol and other baths, and finally lacquering.</td>
<td></td>
</tr>
</tbody>
</table>

The original state recovered as far as possible, and much interesting ornament revealed. (Pls. 44 and b.)
7. Sutton Hoo helmet.* In hundreds of rust fragments with remains of ornament in tinned and gilt bronze and in silver. Detailed study and reassembly of ornament; recovery of crest and main shape of body of helmet. Discovery of neck piece and earlap appendages and their assembly on the body of the helmet. Fitting to model head made in plaster of Paris.

Reconstruction based as far as possible on an interpretation of the evidence. (Pl. 45.) Lacunae filled with coloured plaster.

8. South Shields sword. Fragmentary masses of rust, apparently insignificant. X-radiography to reveal inlay. Mechanical treatment to remove overlying rust.

Inlaid figures of Mars and of Roman standard revealed. The brass inlays exposed to view on either side of the blade. (Pl. 46 illustrates the progress of the work.)


Design and inlays revealed. (Pls. 47a and b.)


Silver pattern completely revealed on a ground of black iron. (Pl. 48.)


Contour established, and intricate ornamental inlays of noble metal recorded by radiography. (Pl. 49.)

12. Sutton Hoo axe. Fragmentary; massive rust; no metal remaining. Cross-sections of fragments indicated original shape of haft and head; also the existence of a ring at end of handle. Original shape recovered by picking and grinding.

Reconstruction based on evidence that was only discovered by patient microscopical studies.

---

5 While this material could not be sampled for analysis, a brass of similar appearance on a Roman scabbard in the British Museum, of the first century A.D. and bearing the image of Tiberius, was spectrographically examined and proved to be an alloy of copper and zinc containing a lesser quantity of tin; traces of silver, lead, and iron were present. It is possible that the two Roman alloys correspond with the metal known as orichalcum (*Plato, Critias*, 114 b) obtained from a naturally occurring mineral.
Iron rust may have an evidential value. It is never safe to proceed to reduce a rusty object without making a careful examination to ensure that there are no features peculiar to the rust that will be sacrificed in the process. Thus, a rusted shield grip may retain fragments of a leather belt, or the oxidized surface of a sword may provide the only evidence of the structure of the scabbard or the hilt. Textile imprints in rust may be important, and even the off-prints from reeds or grasses with which the corroding object was in contact. Compacted rust is to be regarded, therefore, as a possible source of evidence that might otherwise be unobtainable, and laboratory treatment, on occasion, may have to be directed to the preservation of this evidence, rather than to the restoration of the object as a museum specimen.

PRACTICAL EXAMPLES

In the absence of a personal examination, it is never possible to give more than an indication of what treatment would be likely to be the most desirable in any particular instance, but some further guidance may be obtained by studying case sheets. Brief practical notes are added relating to a dozen iron objects selected as being representative of different stages of mineralization, in the hope that they may illustrate the application of methods already described, and at the same time be useful for reference; see Table VI, p. 282.

RECONSTRUCTION WORK

Iron is unique among the metals of antiquity in that it may be decomposed to fragments of rust that are stable in themselves, and not subject to further corrosion. It is sometimes possible, therefore, by studying the fragments and fitting fractured surfaces together, to reconstruct an object, in part or whole, and by this means recover a specimen of archaeological importance from apparently worthless material.

Two of the tabulated examples, No. 7 (the helmet) and No. 12 (the axe), were received as a mass of fragments, some friable and en-
crusted with sand, others very hard, and partly transformed into limonite—a hydrated oxide of iron. In each case the laboratory task was to discover the shape of the object, to assemble the fragments and effect as far as possible a reconstruction. Such work is a variation of the familiar jigsaw puzzle, but it must be carried out in three dimensions, instead of two, and since many of the pieces are missing, every scrap of evidence has to be marshalled before any progress is possible. Thus no preliminary form of cleaning can be undertaken, as this would destroy the natural variation in the colour and texture of the rust that helps to indicate which pieces come together.

The first step is to set aside in groups all pieces bearing fragments of decoration, all pieces having distinctive grain markings or stains, and pieces that are of the same thickness and may belong together.

Maryon\(^1\) describes an early stage in the reconstruction of the Sutton Hoo helmet as follows:

When sorting the fragments, I found that to facilitate the handling of each piece, and to preserve any delicate edges, it was a good plan to provide each fragment with a piece of stiff card, upon which it could rest, and by which it could be lifted. On each card the outline of the fragment was drawn, and notes made of details of any ornament or other feature of interest which had been observed. Search through, and study of, the recovered fragments, continued for a long while, but gradually some more general observations became possible.

This method of dealing with the fragments on cards is invariably employed.

Attention is next devoted to fragments containing ornament, and when pieces are found that fit together they are stuck in position with Durofix. In such reconstruction work a box of sand is very useful for holding pieces at any required angle whilst the adhesive is hardening. In this manner and with the exercise of much patience forms begin to take shape.

There comes a time when no further pieces can be made to fit. At this stage in the reconstruction of the Sutton Hoo helmet, all

---

discarded rust, clay, &c., was re-examined, and it was found that certain shapes had been impressed on worthless material that had once been in contact with the helmet. Impressions of these shapes were taken in plaster of Paris, and the plaster models were found to be the equivalent of some of the pieces missing from the helmet. A fresh impetus was thus given to the work—and when, eventually, certain puzzling fragments were recognized as being the remains of cheek pieces and neck guard, it was clear that a satisfactory reconstruction was within sight. The original material when brought to the laboratory consisted of ‘a gilded bronze nose and mouth-piece, two gilded bronze dragon heads, parts of what once had been a silver crest, and three or four hundred fragments of sand-encrusted rusty iron’. The latter might long have remained of little significance had it not been for the development of the reconstruction technique that made possible not only the recovery of this helmet, but several other important objects from Sutton Hoo—notably the axe, the standard, the shield, and the drinking-horns.

REPAIRING METAL ANTIQUITIES WITH SOFT SOLDER

Metals that are uncorroded may be joined together by soldering, i.e. by alloying them with another metal of lower melting-point that can be made to flow through the joint and, on solidification, act as a bond of union between them. Full details of the process are given in textbooks on metal-working,¹ and it will suffice here to state some general principles that apply to soldering as a method of repairing antiquities.

There are two principal kinds of solder, hard solders which may contain many different metals and melt above 600° C., and soft solders that are composed of a mixture of lead and tin. It is only the soft solders that are of interest to us, and, in particular, the material known as Tinman’s solder. Tinman’s solder Grade K (B.S.S. 219) is of general utility and contains three parts of tin to two of lead. Its melting-point is so much lower than that of either tin or lead that it

¹ e.g. Maryon, H., op. cit.
can be used for repairing either of these metals as well as all other metals of antiquity.

The surfaces to be joined must be clean, i.e. free from oxidation and tarnish and any other impurity that will prevent the molten solder from coming into intimate contact with the metals with which it is to unite. But even the heat required in the process of melting soft solder would cause the metal to become oxidized unless it is protected; it is necessary therefore in the process of soldering to use an oxide-solvent, called a flux. Fluxes are of two kinds: the first contain chlorides such as zinc chloride and ammonium chloride, while the second include those based on resins and fats. The former are very effective but difficult to remove completely afterwards, and for this reason are likely to give rise to subsequent corrosion; the latter are less efficient but do not cause corrosion.

In repairing antiquities, the non-corrosive fluxes are to be preferred, but we are then faced with the difficulty that while the old-fashioned resin fluxes are adequate for work on new metal (as in joining electrical cables, where absence of corrosion is an important factor), they are intractable where there is intercrystalline oxidation as in the case of ancient metal. A compromise is thus necessary, and it is to be found in the proprietary materials sold as ‘non-corrosive’ or ‘activated resin’ fluxes. These still require the joints to be clean, but are easier to work than the simple resin fluxes.

COMMERCIAL FLUXES AND SOLDERING FLUIDS

As it is important to have a working knowledge of the properties of fluxes used in the repair of metals, the following notes relating to some of the commonest that are commercially available may be of interest. Comments are based upon workshop experience rather than laboratory testing.

Corrosive fluxes

1. Baker’s Soldering Fluid. This is recommended for general-purpose work, such as repairs to tin, brass, copper, or iron articles, and for joining and backing-up electrotypes. The joint requires thorough washing.
(2) Fryolux. This solder paste is very useful for tinning two surfaces that have to be closely fitted, such as a piece of brass tubing telescoped into another piece. Useful for 'sweating' screw heads into their sockets.

(3) Multicore, ARAX. This flux is very good for general-purpose work and can be used with stainless steel.

(4) Solderine. This does not have the same self-cleaning properties as 1, but is satisfactory on surfaces that can be very thoroughly cleaned before application. It is less corrosive than 1, 2, and 3.

(5) Coraline. This solder paste is similar in action to 4, but possibly rather less corrosive.

Non-corrosive fluxes

(6) Multicore Solder Paste, Cored Ersin Solder, and Liquid Ersin Flux. All three have been found satisfactory on cleaned surfaces, the Cored Ersin Solder being ideal for electrical contacts.

(7) Alcho-Re. This solder paste has been found to be satisfactory when applied to surfaces made spotlessly clean beforehand.

When ancient metal is found to be very difficult to solder, the best flux to use is probably a mixture composed of 70 parts of zinc chloride and 30 parts of ammonium chloride. This has the advantage of melting at 180° C., but it is very highly corrosive. When it is used in making a joint, the object should be washed immediately afterwards in water acidified with a few drops of hydrochloric acid in order to decompose any insoluble zinc oxychloride that may remain on the joint, and it should be thoroughly washed finally in water made faintly alkaline with a little washing soda.

Soldering is required most frequently in repairing brittle metals, e.g. speculum or ancient silver, or in mending fine work such as that on brooches, handles, &c., where leverage would tend to rupture joints made by adhesives. Soldering is often used as a temporary fixing in 'tacking' cracked metal together as is done in the reconstruction of crushed metal objects. When the original shape has been re-established, a permanent soldered joint can be made to take the place of the temporary fixings which are easily removed. (Cf. treatment of the Emesa silver helmet, p. 226.)
A. Before treatment

B. After treatment

41. PEWTER MEDAL OF THE CAPTURE OF THE BASTILLE
A. As excavated

B. After cleaning and restoration

42. ROMANO-BRITISH IRON STEELYARD WITH LEADEN WEIGHTS
(c. 3rd cent. A.D.)
(Avery Historical Museum)
In exceptional cases it may be necessary to apply a modern metal patch to an antiquity. This should bear an unobtrusive but distinctive mark to indicate that it is modern; a small depression, for example, such as can be made by using a spring-loaded centre punch.

Soft solder can be given a coating of copper by degreasing the solder with trichlorethylene and then rubbing it with a moistened crystal of copper sulphate. The copper surface thus obtained is less obtrusive than the white metal. Soft solder that has been applied to silver can be concealed by degreasing and then silver-plating. This is done by ‘ragging’, i.e. plating with a glass rod anode around which is wrapped silver wire in contact with a small piece of soft rag moistened with a cyanide electrolyte. The solder is rubbed with the rag whilst a small current is passing (at about 3 volts) and it gradually becomes covered with a thin coating of silver.

TABLE VII. IRON AND STEEL

Iron and steel

Uncorroded
1 or 5

Mildly corroded
gilt or inlaid
2, 5

Corroded

Stable

Appearance satisfactory
1

Appearance unsatisfactory

Heavily incrusted

Rust porous
3a and 2c

Rust compact
6 or 1

Lightly incrusted
3a or 3b; 6d and 5

Thin and frail
2; 3a, 6, 5

Unstable (corrosion active)

Corrosion localized

Stout and heavy
6, 4, 7

Rust spots on burnished steel
3, 2, 6f

Corrosion general

Several objects corroded together
6

No metal remaining
1 or 5d

Object in fragments
Reassemble

Metal remaining as a solid core
4, 7

For method of use
see p. 186.
1. No treatment.
2. Mild abrasives: (a) Fine emery powder. (b) Carbontanum powder. (c) Steel wool.
3. Rust removers: (a) Petroleum jelly. (b) Lubricating oil. (c) Clock oil.
4. Dorsamides: (a) Decotheine. (b) Decotheine. (c) Decotheine. (d) Electrolysis in caustic soda.
5. Protective finishes: (a) Bituminous paint. (b) Paraffin wax. (c) Micromeraldine wax polish.
6. Mechanical: (a) Picking. (b) Chipping. (c) Scratching.
7. Wash in distilled water to remove soluble chlorides.
PART III

SILICEOUS AND RELATED MATERIALS
CHAPTER XIV

STONE

Since early in the present century the preservation of stone has engaged the attention of scientists the world over. The phenomena of decay are commonly to be seen on public buildings and outdoor monuments: weathering, which weakens the stone though it may sometimes enhance its appearance; staining, which may result in permanent disfigurement; and the crystallization of salts, which in some cases may cause powdering of the surface, cracking, and even complete disintegration. Deterioration may be traced to bad technique such as carelessness, or lack of knowledge or experience in the handling of stone by architects or builders, but it is more often due to natural causes. In this case, the most that can be done is to try to find a way of mitigating the damage, by washing, or periodic steam cleaning, or, it may be, by some form of impregnation.

While some reference will be made to the treatment of outdoor stone monuments, it is principally the preservation of stone objects indoors that will be dealt with here. This is the simpler task, and although it introduces problems, it may fairly be claimed that most of them are capable of solution.

MINERALS

The earth's crust is composed of minerals associated together to form rocks, and, while the rocks have been used as such for building and sculpture, certain of the minerals have attracted the artist-craftsman from earliest times because of their special qualities for fine work, e.g. hardness, texture, and colour. In the Far East and in the Americas, minerals such as jade, rock crystal, and malachite were used for fine carvings; in the Middle East, lapis lazuli, turquoise, and carnelian were employed extensively for inlaying furniture and for
the decoration of metal objects; and alabaster provided the Egyptian craftsman with an attractive material from which to fashion the lamps and unguent pots that were an essential furnishing both of the temple and of the tomb.

Minerals, as a class, may be regarded as inert, and objects fashioned from them are unlikely to need attention other than periodic cleaning, which can be done in some cases by washing with warm soapy water, or by brushing with spirit.

There is, however, one exception, and this is the mineral called marcasite which is sometimes used for inlays. Marcasite is a sulphide of iron, being a variety of iron pyrites, and it has a similar brassy appearance. It is very liable to decomposition which takes place as a result of oxidation in the presence of moisture, the sulphide being converted to ferrous sulphate which forms a white feathery incrustation on the surface, sulphuric acid being produced at the same time. Treatment may take the form of washing, or, in bad cases, soaking in dilute ammonia prior to washing; and since the acid is, of course, the major factor in causing decomposition, washing is continued until acid can no longer be detected on testing the wash water with litmus.

When fossils contain decomposing marcasite and calcium carbonate they are often beyond recovery, because the acid resulting from the decomposition of the marcasite reacts with the calcium carbonate causing disintegration. Such fossils are said to be 'pyritized', and the best method of treating them is to expose the fossil to the fumes of strong ammonia in a closed vessel. This neutralizes any free acid, but the treatment may take many months and even then can never be guaranteed to be completely successful as it may not be possible for the ammonia fumes to reach all of the acid. After washing in water, drying is conducted quickly through baths of alcohol and ether, and the fossil is then impregnated with a lacquer to exclude air and moisture. For this purpose a silicone lacquer\footnote{Midland Silicones Ltd., 19 Upper Brook Street, London, W. 1.} is very suitable, as it is a water repellent; good results have also been obtained by using polyvinyl acetate in a mixed solvent of nine volumes of toluene to one volume of acetone.
IGNEOUS ROCKS—GRANITE AND BASALT

Of the igneous rocks, granite and basalt are most widely distributed. These are generally hard, non-porous, and very stable under a wide variety of conditions. Granite contains over 66 per cent. of silica, and is an acidic rock, whereas basalt is a basic rock containing less than 52 per cent. of silica. Between these two extremes there are many well-defined rocks of an intermediate silica content. Many have been used for monuments in antiquity and have survived weathering extremely well.

A problem is presented, however, when such stones are brought away from their natural surroundings and exposed in the open to weathering of a nature that is foreign to them. Cleopatra’s Needle, a granite obelisk from Heliopolis, now on the Embankment in London, was observed to have suffered a marked deterioration in its condition on being taken from the dry atmosphere of Egypt to the humid atmosphere of England. A like deterioration was observed in a similar granite monolith that was taken from Egypt to New York, and it was found necessary in each case to treat the stone so as to prevent access of moisture.

A study of the London monolith made in 1952\(^1\) showed it to be a typical granite composed of crystals of quartz, felspars, hornblende, and some white mica. While the stone was originally of a pink colour, it had become badly stained with a black film. This proved to be mostly carbonaceous in nature, but it contained an inorganic residue which consisted mainly of iron oxide and silica. Light coloured patches marked places where flakes of the stone had become detached by the action of frost which was regarded as the prime cause of the deterioration. Conservation involved cleaning, drying, and waterproofing the stone. Cleaning proved to be difficult; the roughened surface of the granite necessitated the use of scrubbing brushes and even wire brushes, and it was found that the application of an organic solvent was essential to soften the black film before it could be removed. The solvent was composed of nine volumes of carbon teta-

chloride and one volume of benzene emulsified by the addition of a surface active agent, e.g. 1 per cent. of Lissapol N. When the cleaning was completed and the natural colour restored, the stone was hosed down and allowed to dry; it was then given two applications of a 10 per cent. solution of paraffin wax (M.P. 50° C.) dissolved in white spirit. After the solvent had evaporated, the stone was warmed gently, using a blow-lamp, with the object of getting the wax into the cracks and porous sections, and thus excluding moisture. This worked well, and the stone, inasmuch as it has been waterproofed, is now protected from the action of frost which was the main object of treatment. The staining by soot, however, cannot be prevented, especially with a surface as rough as this Egyptian granite, and within three years the stone had become as black as ever.

Basalts are more homogeneous than granites, and they seem to be less liable to deterioration. Certain monuments of lava from British Honduras were observed to be very crumbly on the surface after having been exposed in the portico of the British Museum for a few years and had to be brought indoors. These stones are porous and therefore prone to damage by frost, but how far the industrial atmosphere may have contributed to their deterioration is hard to assess.

Such incidents serve to illustrate the general principle that when an object is exposed out of doors it is liable to suffer deterioration from causes that are beyond control, and although the object may never have been intended for a museum, it may be necessary to keep it indoors so that it will be protected from dirt and the rigours of a climate that will hasten its destruction.

SEDIMENTARY ROCKS—SANDSTONE AND LIMESTONE

The problem of preserving sedimentary rocks is essentially one of overcoming surface frailty. The particles of grit of which they are composed are cemented together in a matrix which, if disrupted, leaves the particles free to fall away in powder. On the other hand, some stones develop a natural skin or patina which may afford a certain degree of protection. If this skin is damaged, it exposes a powdery under-surface, and experiments made in an effort to repair such a skin,
or to apply an artificial protective skin, have proved to be unsuccessful as a means of conservation. Indeed, studies in stone preservation have brought out the fact that, as often as not, the application of impervious surface films to stone (as distinct from deep impregnation) seems to intensify deterioration instead of affording protection.

The deterioration of the surface of sandstones and limestones which are comparatively porous is caused mainly by the crystallization of soluble salts. When these are present in the stone, or, indeed, in any porous substance, they will work their way towards that part of the surface where there is greatest evaporation; they will become concentrated there, sometimes as a hard deposit but often in the form of filamentous crystals which appear as if extruded from the pores (Pls. 50a and b). The growth of the crystals and the accompanying pressure may impose so much strain on the stone that the surface is disrupted, ornament is defaced, and inscriptions gradually become illegible. This can be prevented if the salts are extracted either by washing or by using a process involving the use of moist paper pulp.

1. Removal of soluble salts by washing

In specimens where the salt has not yet started to crystallize on the surface, the stone should be immersed in running water and finally in changes of distilled water to remove the salt before it can do any damage. When, however, crystallization has occurred on the surface, the salts should be removed as far as possible by dusting them off gently with a small soft brush prior to washing. When once the stone is wet, it should not be allowed to dry until it has been ascertained by means of the silver nitrate test that all but traces of chloride have been eliminated (see p. 199).

Unfortunately, it is not always possible to treat specimens in this way. Statuary may be large and already weakened by salt action, and in the case of heavy stone the element of risk involved in moving it may not be justified. In the case of outdoor statuary it may be possible to get rid of salts and dirt by using a hose, but for stones indoors that are bulky, the most convenient way of removing soluble salts is to employ the paper pulp method.
2. Removal of soluble salts by the paper pulp method

Paper pulp is available commercially, but it can easily be prepared by boiling soft paper with distilled water and beating it until completely disintegrated and the cellulose fibres mat together forming a pulp. When cold, the paper pulp is thrown against the stone so that it adheres, and a kind of papier mâché garment is gradually built up until the object is concealed completely (Pl. 51). The water in the pulp will be absorbed by the stone and the pulp will shrink slightly as it dries so that, eventually, the stone will be covered with a porous layer of material about ¼ to ½ in. thick. The water absorbed by the stone will dissolve the soluble salts, and, at first, carry them farther in; but, because salts in solution tend to move towards a surface where evaporation is taking place, they will in time change direction and eventually leave the stone, forming an incrustation in the pulp. The pulp is left in position for about three weeks, when it may easily be removed; it is then replaced with fresh pulp and if the stone is very salty a third application may be necessary.

In cases where the surface of the stone has been loosened by incrustations of crystals, or where there is painted decoration, as in Egyptian tomb paintings, consolidation is necessary before applying the pulp. This is done by painting the stone freely with a 2 per cent. solution of celluloid in a solvent consisting of equal volumes of amyl acetate and acetone. By this means the powdery grains are held together and the paint fixed, and when quite dry the stone is ready for the pulp treatment. The film of nitro-cellulose does not prevent the extraction of the salts, but as the diffusión takes place more slowly it may be necessary to continue the treatment for several months, renewing the pulp coating from time to time during this period. If the painted decoration appears to be shiny after treatment, the gloss can be removed by dabbing the areas in question with cotton-wool moistened with a little acetone, but this should be done with care as it is liable to dissolve too much of the nitrocellulose and possibly

1 While the paint may become glossy by such treatment, stones that contain or have contained much salt are generally very porous and can absorb large quantities of celluloid solution without change in appearance.
loosen the consolidated surface. It is usual for the texture and appearance of the painted surface to be improved after treatment by the paper pulp method and sometimes they may even be enhanced by a final coating of a 2 per cent. solution of celluloid. It should be noted that although the paper pulp method can be extremely effective, it does not remove all the soluble salt from an incrusted stone; a certain residue is bound to remain, but if the work has been carried out properly this will have no material effect upon the stability of the specimen when it is preserved under museum conditions.

As an example of what may be achieved through the successful application of this treatment we may instance the case of an Egyptian Mastabah tomb of limestone which was frail and encrusted with salts. As a preliminary step it was consolidated with a solution of celluloid. This was followed by three applications of paper pulp spread over a period of four months. By this time the incrustation had disappeared and what remained of the natural surface of the limestone was exposed, bringing to light delicate carvings in bas relief and painted decoration which had long been completely concealed from view.

3. Removal of insoluble salts

Sometimes the prevailing incrustation is insoluble in water so that washing or treatment of the stone by the paper pulp method are of no avail and alternative methods must then be employed. These will depend on the chemical nature of the incrustation. If the salt is carbonate, it can be decomposed by any acid, but a warning is necessary since even sandstones may contain calcite (calcium carbonate) as an essential part of their structure; before using acid, therefore, it must be established that the rock is neither a limestone nor a sandstone containing calcium carbonate. The rock, when tested, should not itself effervesce with acid. Even when all seems well, acids should only be applied locally and sparingly, not with the object of dissolving salts wholesale, but rather with the object of loosening the crystals so that they may be dislodged mechanically. For this
purpose, dilute hydrochloric acid (5 per cent. or even weaker) may be used, followed by thorough washing with water.

When the incrustation is selenitic, i.e. when it consists of gypsum or calcium sulphate that has been laid down in glassy crystals, it may be very disfiguring. Such incrustations have formed very slowly and are probably harder in the mass than the surface of the stone which will certainly have been weakened during their growth. It is unwise, therefore, to try any chipping or grinding process until preliminary steps have been taken to try to soften the incrustation. Unfortunately, no solvents are available that will dissolve gypsum readily. If feasible, the best treatment is to keep the stone immersed in lukewarm water for a long time, changing the water every 24 hours; this has been found to be more effective than immersion in either cold or hot water. Solutions of sodium thiosulphate (10 per cent.), or ammonium carbonate (10 per cent.), may be applied locally, but these chemicals act slowly and they must be washed out very thoroughly afterwards or the cure may be worse than the disease.

A novel method of treatment that at times has proved to be very effective depends on the fact that selenite is a hydrated mineral, and that it goes to powder when it is dehydrated by heating. An incrustation may sometimes be heated with an electric soldering iron, and if the temperature is strictly under control it is possible to disintegrate the gypsum so that it can be brushed away without damage to the stone. The method is obviously not without danger, but in cases where a fine object is suffering from a gross disfigurement of selenite which would otherwise be difficult to remove, it may be worthy of consideration. Needless to say, the stone must be quite dry when heat is applied, and it should be borne in mind that overheating a limestone converts it into quicklime.

4. Consolidation

After salts have been removed from stone, it is often necessary to strengthen the surface. In the British Museum Laboratory hardening tests were carried out upon a number of Coptic stele of very porous limestone that had been desalted by the pulp method, using such
consolidating liquids as white beeswax in turpentine, white shellac in alcohol, and solutions of cellulose nitrate and polyvinyl acetate. It was found that, under museum conditions, success was related directly to the degree of penetration of the consolidating agent. After a lapse of twenty years a further series of observations was made on the same stele after they had been exposed to damp in an unheated war-damaged building. This confirmed the earlier observations that the most satisfactory consolidating agents are those which can be made to penetrate well into the material and not merely remain on the surface, especially if the stones are liable to be exposed to damp and varying temperature conditions.

It is clear from the tests that it is not possible to preserve a powdery stone surface by merely giving it a coat of varnish. Porous stone contains air that will expand and contract with variations in temperature, even under museum conditions, and no thin surface film could long withstand the consequent movement without fracture. This is the reason why the most successful results have been achieved either by securing deep penetration of the consolidating agent, or by employing a form of surface treatment that leaves the stone free to ‘breathe’. These treatments will now be described in detail.

(1) Impregnation with wax. If properly carried out, wax treatment has been shown to stand the test of time satisfactorily. Where possible the stone should be immersed in a tank of molten wax—a method of treatment that has already been described in the case of wooden objects (see p. 127). Special plant is required for such treatment, however, and the size and weight of the stone might be such as to make this method impracticable. When wax has to be applied to the surface of a large object that cannot be immersed, the only way to ensure penetration into the pores of the stone is to heat the stone for a time before applying the wax. The stone must, of course, be thoroughly dry before it is heated.

A convenient method of heating is to use an electric radiator placed about three feet from the stone. By this means, a wide area can be dealt with at once. When the stone is hot, the radiator is switched off and white beeswax applied in the form of a thin salve having the con-
sistency of vaseline. This salve is prepared by stirring molten beeswax at about 85° C. into petroleum ether (B.P. 80–100° C.), care being taken not to do so in the presence of naked lights. When the salve is applied to the warm stone, the wax is absorbed into the pores and the inflammable solvent evaporates. After all the solvent has evaporated heating may be repeated, and further coats applied in the same manner as long as the wax is being absorbed.

Perhaps it should be mentioned, in the interests of aesthetics, that waxing invariably causes a certain dullness and lowering of tone particularly noticeable on light stones, and also that waxed stone is easily stained by dirt. This latter problem is not so serious when microcrystalline waxes are employed (see Appendix XII).

The waxing method may be used also for statuary that is kept out of doors, but it is absolutely essential that the stones should be quite dry before they are waxed and that the wax should penetrate well. Where stones are set in a wall, wax should never be used as a consolidating agent as there is always the possibility of water getting into the foundation or behind the wall and causing future trouble. An examination of many frescoes on the walls of public buildings and churches has shown, without doubt, that treatment with wax has only been successful where the fabric of the building has been kept continuously dry. Where water has had access to the walls by the chance blocking of a gutter, or perhaps through inadequate drainage on the outside, the moisture cannot escape through the waterproofed surface; it is held back by the wax barrier where it may cause deterioration at the interface, with the result that large flakes become detached, or, in some cases, even the foundation layers of the fresco.

(2) Impregnation with lacquers. In attempting to impregnate a stone with film-forming solutions (lacquers), one has to face the difficulty that any solvent that goes into the stone must eventually evaporate and come out again, at the same time bringing much of the solid content of the lacquer with it to the surface, or the immediate subsurface. This is easily demonstrated by cutting sections of a white porous stone that has been impregnated with a coloured lacquer: it will be seen that the colour has penetrated well into the stone, but
that the solids remaining after the solvent has evaporated are distributed near the surface. Lacquering methods might, therefore, seem to be of limited use, but, where applicable, as when stones are kept indoors, they can be regarded as satisfactory. It must be emphasized, however, that this method of treatment affords no protection to stones that are exposed in the open to the undermining action of rain and frost.

Lacquer solutions must be applied with a brush in a series of coats, the first very dilute, and the succeeding ones progressively more concentrated, allowing time for each coat to dry before applying the next. As has already been mentioned, white shellac gives good results. This is obtainable as French polishers’ ‘white polish’, and is diluted with methylated spirit for use on stone. Other impregnating media that have proved satisfactory for indoor work are polyvinyl acetate in toluene/acetone solvent, and Bedacryl 122 X¹ in xylene/toluene solvent.

(3) *Vacuum impregnation.* For small stone objects that are in a powdery condition, the best treatment is to impregnate them *in vacuo* with one or other of the above media well diluted with solvent. This is carried out in a vessel that can be made air-tight and is at the same time strong enough to withstand the strain of being evacuated. For small work a thick glass desiccator² serves well (see Fig. 5, p. 152).

The stone is immersed in the impregnating solution, air is sucked from the vessel, and bubbles are seen rising from the stone through the solution as the pressure falls. After most of the air has been removed and the bubbling has subsided, air may then be readmitted to the vessel. The pressure of the incoming air will force the solution into the pores of the stone. It is most important that the air should be admitted gently, as a sudden inrush might cause the stone to rupture. After impregnation, the object is allowed to drain above the solution in the atmosphere of the solvent to reduce the rate of drying. Hasty drying is liable to bring too much of the lacquer to the surface,

¹ Imperial Chemical Industries Ltd.
43. SWORD HILT (SUTTON HOO)
Surviving elements displayed in a Perspex mount
A. As excavated

B. After restoration

44. SHIELD BOSS (SUTTON HOO)
leaving the stone with a shiny appearance. The apparatus has been described in greater detail under the impregnation of wooden objects (see p. 130).

(4) Silicon ester spray. For slabs of sandstone and siliceous limestone of large dimensions which are kept indoors, a most successful strengthening agent is silicon ester, but it is essential that it be applied strictly in accordance with the manufacturer’s directions.

Silicon ester is available in the form of a dilute solution in alcohol, and it is necessary to apply it with a spray or an atomizer as it is not effective when merely brushed on the stone.

The atomizer should be held at some distance from the stone, so that the liquid reaches the surface only as a fine mist. Further applications are made not oftener than once a week, thus allowing time between each operation for the ester to decompose and coat the grains of stone with silica. After about three applications the surface may be tested by rubbing lightly with the fingers, and if the dust has been fixed, this will indicate that the process is likely to prove successful. Eventually, after about three to seven applications, a trace of permanent milkiness will be observed—a sign that treatment should be discontinued, as the surface of the stone is now well consolidated without the pores being clogged. Sandstones so treated have been found to remain in good condition in the museum for many years.

The method is not so effective with fine limestones, though in some cases it seems to harden them. Disappointing results were found in the case of lavas and it would seem that the type of stone that responds best to this spray treatment is one having a gritty surface, the silicon ester reinforcing the decomposed matrix and thus consolidating the gritty constituent of the stone.

It should be added that there is a tendency for silicon ester to clog the jet of the spray, and once this happens it is difficult to clear it for use again. The spray should therefore be dismantled between operations and the jet washed with alcohol and kept, preferably, in a bottle of spirit.

1 A suitable brand of silicon ester for the preservation of siliceous stones in museums is that known as Silicaseal No. 1A, obtainable from Silicaseal Ltd., Newcastle upon Tyne.
Silicon ester has a comparatively short shelf life, and it should be used within a few weeks of purchase, as even dilute solutions become viscous and unfit for use after a lapse of a few months.

MARBLE

Both igneous and sedimentary rocks may be changed in form, depending on their composition and history, granitic rocks becoming gneisses, sandstones quartzites, and limestones marbles. It is the marbles that are of most interest in museum work as they have been used to such a large extent for sculpture as well as for building purposes.

Marble is a metamorphic rock arising from the action of heat or pressure, or both, on limestone. By this action the original character of the limestone is lost, and it becomes an aggregate of calcite crystals which may be pure white, coloured, veined, or black, according to the impurities present. As a result of this transformation the pore-space in the stone is reduced and is smaller than that of limestone, and marble is thus capable of taking a higher polish than limestone; but the pores are not so small as to prevent the marble from being easily stained, and special care is necessary to prevent the accumulation of dust in the hollows of white marble sculpture, as ingrained stains may be very difficult, or even impossible, to remove.

Dusting and washing. For the preservation of marble in the museum, periodic dusting is essential, and this should be done with a large feather whisk or soft brush. Cloths should never be used as they tend to rub the dust into the stone.

Washing may be necessary at times, and for this purpose it is important to use water that is free from iron, and preferably distilled. A little good-quality soap may be used, just enough to cause a slight frothing when applied with a soft brush. The solution used in the British Museum for the periodic washing of marbles consists of:

- Soft soap (B.P.) . . . 10 grams.
- Distilled water . . . 100 ml.
- Ammonia (0.88) . . . 1 ml.
This is made up as required, kept in a glass bottle, and used from a glass or pottery vessel. The mixture must not be allowed to come in contact with iron, and should not be used from an iron pail, even if galvanized, as this might lead to eventual staining. After the marble has been dusted with a feather whisk, washing operations are commenced at the top of the statuary. The cleansing solution is worked with a soft brush into a froth on a small area at a time, taking care to prevent dirty water from lying in the hollows or running down in rivulets over areas of unwashed marble. When clean, this area should be dried with a soft towel before proceeding farther. Only when all the marble has been cleaned and mopped dry is the stone washed freely with fresh water to remove the last traces of soap, which would soon collect dust again if allowed to remain.

For marble that has been neglected and allowed to accumulate dirt for years, it may be necessary to use a stronger detergent, such as the froth from Lissapol, Teepol, or one of the proprietary cleansers made from such materials. A convenient quantity is 2 oz. of detergent in one gallon of water, and this concentration should not be exceeded. Such a solution is not for regular use and should only be applied in exceptional circumstances, taking the same precautions as before to prevent rivulets of dirty water from running down the marble.

*Removal of stains.* The common staining ingredients of dust are soot and traces of iron, and these give rise to grey and to rust-coloured marks respectively. White marbles easily pick up colour from contact with packing materials such as paper and straw if these are damp, and even damp clean dust-sheets have been observed to cause staining. In one such case the source of the trouble was a slight mildewing of the textile which caused the appearance on the marble of a brown line following the course of the retaining string. The susceptibility of marble, and particularly of white statuary marble, to such forms of staining makes it essential to take special precautions in packing. Where there is a possibility of exposure to damp, white marble should not be allowed to come in contact with any organic material liable to fungoid attack. This fact was brought home by the
discovery of intensive polychrome staining all over a fine white marble bust of Voltaire by Houdin. The marble had been wrapped in a white silk eiderdown, boxed up, and stored in a garage where conditions were at one time so damp that the wood of the box had rotted. When the box was opened the packing was found to be concealed in a mass of mould growths, and the marble was stained brown, red, and green at all points of contact. Much of the staining was removed by the use of an aqueous 2 per cent. solution of chloramine-T, but certain areas of dark staining resisted all efforts to bleach them.

Another type of staining by contact with organic material was that suffered by the Michelangelo tondo in the Royal Academy of Arts. In this case the trouble was caused by felt padding pressing against the white marble, and this resulted in the appearance of a rust-coloured smudge which was very disfiguring. Fortunately, most of the stain responded to washing and treatment with chloramine-T, but rather than take the risk of using a stronger reagent, the residual yellowing was, for aesthetic reasons, chalked over to conceal the blemish. In the course of time the yellowing became decidedly less intense. The gradual disappearance of the stain was not due in any way to the presence of the chalk but no doubt to the volatility of the residual staining material.

It is not an easy matter to restore the pristine whiteness of Carrara marble that has been stained. Although it may be possible to remove coloured matter from the surface, in most cases the stain will have penetrated into the stone where it can no longer be reached by solvents, and may remain as a permanent disfigurement. The problem of removing stains from marble is further complicated by the fact that all acids disintegrate marble, and for this reason the use of specific acidic solvents for removing stains is out of the question. The common neutral organic solvents are safe to use, and slightly alkaline solvents are permissible, but when the latter are used they should not be left for long in contact with the marble and should be washed off very thoroughly afterwards as there is a possibility of their causing yellow stains by contact with colourless impurities present in the stone.
When the marble is sound and the staining is general, thorough washing, and brushing in the wet condition, may be undertaken, but when the stain is localized and highly coloured, it is usually best to determine the nature of the stain and employ selective solvents without prior washing. Thus, a freshly prepared 2 per cent. aqueous solution of chloramine-T will remove stains caused by red ink and also ink stains of the blue-black type, though in the latter case a yellowish residue is likely to remain. This residual coloration may respond to treatment with hydrogen peroxide (20 vols.) to which a drop of ammonia has been added. After the stain is discharged the marble should be thoroughly washed.

A common type of staining is that due to oil paint. A thick blob of hard oil paint should be carefully scraped off without damaging the stone, and any residue treated with a suitable solvent. Pyridine and morpholine are useful in dealing with bad oil stains, or staining of a bituminous character, as is also a mixture of equal volumes of benzene, ammonia (0.88), and methylated spirit. These liquids are applied locally with a stencil brush, and, after mopping with cotton-wool, the marble is washed down thoroughly with water. It should be noted, however, that the oily ingredient of the paint will certainly have been absorbed to some extent by the marble, and, even if it is possible to eradicate all of the surface pigment, the chances of extracting all the oil from the pores of the stone are very remote indeed, and a grey or brownish residue is almost certain to remain. This is not a serious matter if the area affected is in an inconspicuous part, but if the stain is a bad disfigurement it is much better to cover it over with chalk rather than to torture the surface of the stone with chemicals in the remote hope of improving matters.

A method that is sometimes of value in removing surface stains is to use a stripping film according to the procedure developed for recovering fossil imprints from coal. This consists in applying to the stain a viscous solution of nitrocellulose so compounded that, on drying, it leaves behind a very elastic film; when this is peeled off it may take the stain with it. The method has given good results in the case of stained marble but could be applied equally well to
any other fine-grained stone. The following solvent mixtures are suggested by Duerden:

(1) **Rapid drying:**
- Alcohol 1 vol.
- Ether 1 vol.
- Castor oil 5 per cent. by vol.

(2) **Slow drying:**
- Acetone 2 vols.
- Amyl acetate 1 vol.
- Triacetin 2 per cent. by vol.

A sufficient quantity of nitrocellulose (celluloid cuttings) must be added in each case to give a viscous solution.

A common disfigurement is the formation of organic deposits caused by the growth of lichens, algae, &c., on the surface of marble exposed in the open. These deposits may be softened and removed by treatment with a little dilute ammonia. In cases where the algal growths are extensive, they may be arrested by spraying with formalin, and this treatment facilitates their subsequent removal.

When the surface of a finely polished marble is blemished locally by an insoluble deposit the only cure may be to wet the marble and remove the deposit by rubbing with a small chisel-shaped slip of water-of-Ayr stone or snake-stone. ‘Snaking’, when applied lightly, does not impair the polish of the marble.

**Decomposition and consolidation.** Marble has been described as an aggregate of crystals of calcite, and these crystals have a different coefficient of thermal expansion in two directions. The result is that when marble is heated it tends to become distorted. This distortion is detectable even below 100° C., but when the temperature is as high as 400° C. warping may be considerable. Such deformity is sometimes noticed on old mantelpieces, and, as it is a case of the stone having taken on a permanent set, there is no way of recovering the original shape.

The second type of deterioration resulting from heat causing irregular expansion is the granulated condition sometimes observed on projecting portions of statuary. In this case, expansion at different

---

rates in different directions causes the crystals to lose their power of cohesion and it is possible to release granules of the marble by merely rubbing such surfaces with the finger. The affected portions are a staring white, as they scatter the light more than the surrounding stone. Examples have been found in the Parthenon marbles, and in marbles from many other sources. In one case, an extended limb of a large Canova group was worn down to a shapeless mass of loosely adherent crystals having the appearance of granulated sugar.

It is possible to do something to consolidate museum marbles in this condition, but the same difficulty arises as in the consolidation of sandstones and limestones, namely that impregnating agents requiring the use of solvents that evaporate cannot be persuaded to fill the pores of the stone because the solvents, having gone in, must find their way out again. For some purposes the most satisfactory treatment may be to impregnate the stone with wax, even though this may seem illogical, as it involves applying heat to the stone. The aim, however, is to get the wax to penetrate well into the granular marble, and this can be achieved to a depth of an inch or more, if the stone is warmed by an electric radiator. The radiator should not be placed too near the stone, and sufficient time should be given for the stone to warm through gradually. After removing the radiator, treat the stone with a soft buttery salve prepared by stirring paraffin wax of M.P. 46° C. into petroleum ether (B.P. 40–60° C.). It will penetrate the granulated marble easily, replacing the air and filling the pores so that the marble now has a more or less uniform appearance and the enfeebled stone is consolidated. As petroleum ether is highly inflammable, care should be taken to see that there are no naked lights in the room during operations.

Another method of dealing with granular marble in the museum is to impregnate the porous areas with lime water, giving three coats, and allowing a few days’ interval between each for the marble to dry. This is followed, if necessary, by one or two coats of a 10 per cent. solution of ‘soluble casein’. By this means the granules are consolidated by a film that consists essentially of calcium caseinate and calcium carbonate. In cases where the original surface of the
marble remains to a large extent unimpaired, and there are only isolated granulated patches, there may be justification for stippling the staring white of the granulated areas with a little pastel colour to match the surrounding unaffected marble. This masks deformities and assists in recovering the aesthetic appeal of the work. (For repairs to broken marble see p. 317.)

It may be of interest to reproduce laboratory notes relating to a marble which presented several problems in one—removal of an incrustation, removal of stains, consolidation, retouching, and repair. The marble in question is the head regarded as that of Mithras, from the Walbrook Mithraeum in London, excavated by Mr. W. F. Grimes in 1955 (Pl. 52).

This fine head came from a moist site where it had been in contact with ferruginous clay, and it arrived at the laboratory damp, muddy, and encrusted with iron compounds. The stone, particularly in the region of the hair and peak of the Phrygian cap, was found to be weakened by a form of sugary decomposition, which might have resulted from the action of intense heat. There was no evidence to show that the temple had at any time been burned down, but a carbon residue was found in the incrustation, suggesting that the marble may have been exposed to altar fires.

As the stone was considered to be too fragile for treatment in its damp condition, it was given time to dry out, and was then treated as follows:

1. The bulk of the incrustation was removed with pointed matchsticks, exposing a surface that was porous and deeply stained in parts by iron compounds. Match-sticks were used because they are softer than marble and could not injure the surface.

2. The iron stains were removed by sequestrating reagents, applied to the surface with cotton-wool wrapped round the end of a stick. This was a long and painstaking operation, as it was only possible to work on one small area at a time. The most effective reagent was found to be Versenol,¹ and, although this substance also sequestrates

calcium, it had no apparent effect on the marble in the dilute solutions employed. Care was taken not to carry this treatment too far. The discoloration resulting from the burning could not be obliterated, and it was therefore inadvisable to make the rest of the marble too white, as this would have caused it to have a patchy appearance.

3. The sugary condition of the burnt areas was dealt with by applications of lime water (calcium hydroxide), which, in time, was converted in the pores of the marble to calcium carbonate, a material of the same chemical composition as the marble itself. A small amount of crystalline efflorescence was brushed away, and, finally, two coats of a 10 per cent. solution of soluble casein were applied to the areas of sugary marble to consolidate the surface, and the front of the Phrygian cap was lightly touched out for aesthetic reasons with ground pastel colour.

During the course of this work the neck of Mithras was excavated from the temple. It proved to be in a much better condition than the head and could be cleaned by ordinary washing. The neck and head were then joined together by a stout dowel of Delta metal¹ inserted with plaster of Paris into opposing holes drilled in the fractured surfaces.

The Mithras marble was only slightly burnt and was therefore amenable to treatment, but where marbles have been exposed to temperatures sufficiently high to convert the surface into quicklime, there is nothing that can be done to restore them. Sometimes the heating may not have proceeded so far, and the surface may still be hard and intact though blackened with tarry matter. Some improvement should be possible here, but it will depend on the nature of the specimen. The use of pyridine or morpholine, applied in a coating of paper pulp, or mixed with an absorbent powder like kieselguhr and held in intimate contact with the stone until the solvent evaporates, may help to remove such staining. Ammoniacal benzene (see p. 309) may also help, but the final treatment may have to take the form of rubbing down the surface with water-of-Ayr stone or even with fine emery powder. Such treatment would obviously be

¹ Delta Metal Co. Ltd., 205 Tunnel Avenue, London, S.E. 10.
unsuitable for fine sculpture, but it might be the means of saving a good piece if the stain were restricted to the plinth, or to a small part where the modelling was suggested, rather than executed with precision. But, in general, mechanical methods of cleaning marbles involving the use of abrasives can seldom be justified in museum work, and are only used under exceptional conditions.

The use of copper chisels has been advocated as a means of cleaning marble gravestones and the like. Since copper is of the same order of hardness as fresh marble the procedure is not so drastic as it would appear to be. It is, nevertheless, quite unsuitable for museum work where surfaces are studied at close quarters and patina, an important feature of the specimen, must be preserved at all costs.

The patina of marble. The patina of marble differs from that of metals in that it is essentially of the same chemical composition as the material upon which it has been laid down, whereas, in the case of metals, the patina is non-metallic and generally consists of a variety of chemical compounds, oxides, chlorides, carbonates, &c. Marble patinas are formed in the same way as stalagmite by virtue of the fact that calcium carbonate is slightly soluble in water that contains dissolved carbon dioxide gas. When the solution (calcium bicarbonate) drips steadily from a height, the carbon dioxide gas is released, and the calcium carbonate is redeposited on the ground, where it may accumulate to form pillar-like structures or stalagmites. In the same way, when old marbles have been washed by rain for generations, there is a tendency towards solution and redeposition of calcium carbonate. Even though this be on a comparatively minute scale, a change of surface takes place and may be recognized as a variation in texture, translucency, or even colour, warm tones resulting from the presence of a little ferric iron in the solution, and cooler greenish tones from ferrous iron or copper. Plants cause staining and organic growths such as lichens or algae may also leave a greenish tinge on the marble and the patinated surface is often of such porosity that it retains coloured impurities that add interest to the stone. Where patina exists, it is an essential feature of old marble and must be retained. It will survive cleaning by the general methods described, but
would be ruined by the use of abrasives, sharp tools, or acids of any kind.

Examination of marble by U.V. fluorescence. It is instructive to examine a piece of ancient marble in a darkened room by ultra-violet light, and to compare the fluorescence of the patinated surface with that of a recent scratch or fracture in the same stone. Freshly exposed surfaces usually fluoresce brightly, whereas the ancient surfaces are either deep brown or unchanged in appearance. It is often possible by examining a piece of marble sculpture under ultra-violet light to determine whether it is recent or ancient and to reveal any modern cutting on an old marble, but as there can be no absolute guide as to the time required for a marble to acquire a patina recognizable as ancient, it is impossible to date marbles by the quality of their fluorescence. Discretion is required in making such examinations, and, as the fluorescence of ancient marble is low in tone, it is essential to adapt the eye to a condition of total darkness for a few minutes before making the examination, so that slight differences in fluorescence can be discriminated. The problem is much simpler in the case of a composite piece made up from fragments of marble of different origins. An example is provided by a small Greek marble figure reclining against a tree trunk. This appeared to be homogeneous but in a darkened room under the lamp was found to be heterogeneous; the torso, apparently original, was of one kind of marble, while the head and limbs were cut in two others, and the tree in yet a third. Even when the position of the joints had been discovered by this means, they were still very difficult to discern in daylight.

In detecting repairs to marble and estimating their extent, an ultra-violet examination is of the greatest value, and this type of investigation is facilitated by the fact that plasters and glues fluoresce vividly. A valuable Italian marble that had been lent to an exhibition in London was alleged to have suffered damage in transit assessed at a high figure. When examined by ultra-violet light, however, the breakages were shown to be merely ancient repairs that had become unstuck on the journey. It was also revealed that on one part of the marble there was a slight abrasion (invisible in
daylight) which fluoresced brightly and was obviously of modern origin.

It is perhaps not irrelevant to add that some of the modern surface-active agents used as detergents have a characteristic bright fluorescence in ultra-violet light, and hence it is not advisable to use them for cleaning marble that might have to be studied under the lamp. Detergents may have to be used for washing very dirty-neglected marbles, but traces of these materials left on or in the powdery surface of the stone would be likely to complicate fluorescence analysis to such an extent as to vitiate this line of inquiry.

Effects of weathering. Numerous as are the problems of conservation of stone objects indoors, they are seldom so profound or so crucial as those presented by stones exposed outside. These have to face extremes of temperature and humidity, and the disintegrating action of frost, to say nothing of atmospheric impurities, and adventitious damage due to animals and plants. Sand or dust storms gradually erode the surface, and rain causes dirt to collect in hollows of the stone where stains are formed that can never be removed.

A most striking example of weathering is provided by comparing photographs of Lord Elgin’s plaster casts of the West Frieze of the Parthenon made in 1802 (which show the condition of the marbles at that time) with photographs of the same marbles taken in situ in 1938, i.e. 136 years later. The photographs show (Pls. 53 a and b) that, in what may be called the industrial age, the modelling has everywhere lost its sharpness, facial expressions have changed, and whole features have gone for ever, due not to the fault of anyone, but to the variety of causes that we know collectively as ‘weathering’.

Marbles of all kinds are liable to be seriously damaged by the acids present in an industrial atmosphere. In the presence of moisture these destroy the surface, particularly at angles and corners, and open the joints to the disintegrating action of frost. It is not surprising, therefore, that the industrial age has already taken a heavy toll of some of the finest marble monuments of antiquity. In the case of exposed buildings, the damage can only be arrested by the application of stringent anti-pollution laws, but where statuary is concerned, a
practical alternative may be offered by the possibility of covering in
the marbles or bringing them indoors, so that they may be maintained
under conditions where conservation is possible.

**Packing marble sculptures.** The art of packing heavy stone objects
may only be learnt by experience, but even the experience gained with
sculptures of another kind may not suffice in the case of white marble
sculpture, which is so susceptible to staining. White marbles should
be packed in strong wooden boxes, made of well-seasoned timber,
assembled with brass screws. As the main requirement is to dispense
with all but the essential packing material, and at the same time
ensure that the sculpture will not move in the case, wooden bars
are screwed firmly in strategic places so that the marble is retained
rigidly in position. The wooden bars must be padded with the best
quality white cotton-wool covered with tissue paper. The final
retaining member should be a wooden wedge similarly padded, and
driven home gently with a mallet to anchor the stone in position.
The wedge should be retained, if possible, with a screw and be
marked as the first member to be removed when the marble is being
unpacked. On no account should the use of newspaper, wood-wool,
or straw be permitted. When a marble has been allowed to remain
in its box over a long period it is desirable to inspect the padding and
verify the tightness of the retaining wedge if removal of the box is
contemplated, as the padding may have lost its resilience and the
object may no longer be held securely in position.

**Repair of stone objects**

Adhesives that are quite satisfactory for the repair of small stone
objects are inadequate where weights of 1 lb. and over have to be
carried. In such cases the fractured surfaces are brought together
with metal dowels or pins, and by this means the strain is no longer
confined to the fractured surfaces but carried by the body of the
stone.

The choice of metal for dowelling is important. Much damage has
been done under the guise of conservation by dowelling stones with
iron, which, on rusting, swells and opens old cracks and gives rise to
new ones. Stainless steel may be used, but generally copper or one of its alloys is a better choice, one of the best and strongest for the purpose being Delta metal. As a rule the metal dowel should be of rectangular cross-section, and not so heavy that its insertion weakens the stone. The ends may be bifurcated or the edges nicked with a chisel to help the metal to key with the cementing material and form a reliable bond. In joints of this kind the dowels are made to do the work and the crack stopped afterwards with some inconspicuous filler. The procedure of fitting a metal dowel is briefly as follows:

Drill or cut a hole in one of the fractured faces of the stone at the appropriate angle, and to a depth of a little more than half the length of the dowel, and of a diameter just large enough to admit it. With the dowel temporarily in position, mark the opposite face, and sink a similar hole at the correct angle, so that the faces can be brought into correct register. (If a second dowel is required, provision is made for it in the same manner.) The dowel has now to be fixed in one-half of the joint, and this is done by wetting the stone, filling the hole to a reasonable depth with a cement such as plaster of Paris, and pushing in the dowel, and, if necessary, wedging it firmly in position. It is allowed to set overnight. Next day the two parts are brought together and any necessary adjustment made to the hole that is to accommodate the projecting dowel. It now remains to wet the stone and apply the cementing material to the second hole, and it may be desirable to coat the surfaces as well with a thin wash of plaster to help to bed the joint in position when the pieces are brought together. When the joint is permanently set, then the crack has to be made good. This is done with some form of plastic stopping which may be a cement or a mastic, i.e. adhesive plus an inert substance to act as a filler.

The following cements may be used—plaster of Paris; Keen’s cement, which is a retarded plaster of a harder nature; Sirapite, a white dolomitic gypsum containing about 1 per cent. of free lime; or Portland cement, the hard strong cement obtained by heating a mixture of clay and chalk. A suitable mastic may be prepared by grinding kieselguhr in a binder such as silicate of soda or glue size. For
use with antiquities, the best filler is undoubtedly plaster of Paris because of the ease with which it can be applied and removed if this should ever be necessary at a future date. It is used for filling, and for repairing, and even for mounting heavy sculpture on stone plinths. When dealing with large plaster mixes the quick rate of setting becomes a problem but this may be overcome by adding a little glue to the plaster when it is made up. This has the effect of making the material workable for a longer time.

For mounting and consolidation work (see p. 323) it sometimes happens that a cement is required which is anhydrous and yet guaranteed not to shrink. This can be made from a 10 per cent. solution of celluloid cuttings dissolved in a solvent consisting of equal volumes of acetone and amyl acetate. If 50 grams of fine white sand are worked up with 12 grams of this nitrocellulose solution, the mixture will be found to be of the consistency of soft putty which can be shaped with a palette knife. As it dries it adheres very firmly. It takes about four days for a block $\frac{1}{2}$ in. thick to dry through, and then it has the appearance of light-coloured sandstone. A test block 4 in. by 2 in. by $\frac{1}{2}$ in. made according to this formula showed negligible contraction on drying.

**CLAY**

Clay is a hydrated aluminium silicate of variable composition which has the fundamental characteristic of being plastic when moist so that it can be modelled or moulded; it hardens on drying, but can be softened again by the action of water. The nature of clay is changed, however, by the action of heat, so that it becomes rigid, stony, and almost imperishable without change of form, and these qualities have been exploited by man in making bricks, terracotta objects, pottery, and porcelain. When baked above a temperature of 600° C. the change is irreversible, but if inadequately baked the material is of uncertain hardness and not necessarily permanent. There are thus three types of material presented for conservation—baked clay, unbaked clay, and clay that has been inadequately baked. In the latter categories are the writing tablets of clay that are too frail to handle until they have been hardened by heating in a kiln.
Clay tablets and their treatment

Clay was widely used in Sumeria, Babylonia, and Assyria in the plastic condition for writing upon, cuneiform characters being impressed in the moist surface with a stylus. The clay soon regained its solidity in the hot sun and there is evidence that, on occasion at least, it was further hardened by firing, but it is not known to what extent kiln firing was general. The hardness of the clay tablets that are excavated varies considerably and it seems likely that many were not kiln-hardened but merely sun-dried. Hundreds of thousands of clay documents have come down to us—the British Museum alone contains upwards of 100,000—and they have survived astonishingly well, on account of the dry climate of Mesopotamia.

But clay objects suffer in the same way as all porous material that has been buried in salty ground—they absorb salts which will tend to crystallize out after excavation and cause disintegration. The usual treatment for the removal of salts is washing, but with clay objects this is not possible as water would simply convert the clay into mud. As it is impossible to judge by inspection what heat treatment a cuneiform tablet may have been given in antiquity, the first requirement is a controlled rebaking of the old tablet in order to convert it into brick. The tablet may then be washed, or indeed soaked for prolonged periods without being any the worse. The laboratory treatment of clay tablets thus consists in baking, washing, repairing, and cleaning the surface by mechanical methods should this be necessary to render the writing legible. Pls. 54a and b illustrate how, by this treatment, encrusted tablets can be strengthened and the writing made legible.

Baking and washing clay tablets. In the British Museum baking is carried out in an oven or furnace in which the temperature is gradually raised to 750° C.¹ and maintained for some hours. This results in the loss of what is known as 'combined water', and the conversion of the clay into brick. The tablets are allowed to cool in the closed furnace

¹ P. Delougaz (Oriental Institute of the University of Chicago, Studies in Ancient Oriental Civilization, No. 7, 1933), in describing 'The Treatment of Clay Tablets in the Field' advocates a final heating at a temperature of 1,500–1,600° C., but such a high temperature would appear to be unnecessary and undesirable.
45. IRON AND GILT BRONZE HELMET (SUTTON HOO)

A reconstruction
Condition as excavated
Radiograph disclosing outline of inlays under rust
Preliminary exploration

Obverse. Mars
Equipment employed for removing the rust. Radiograph (*extreme right*) used as guide
Reverse. Military standards

46. FRAGMENT OF A ROMAN SWORD FROM SOUTH SHIELDS (c. 200 A.D.)
overnight. When the baking has been properly conducted the tablets are converted to a clean hard material of a biscuit or pale ochre colour, depending on the amount of iron present in the clay. Any organic matter is, of course, burnt away, and in cases where there is saline impurity of a selenitic nature, this is dehydrated in the course of heating, leaving a white powdery residue which may be removed by brushing. When, as is generally the case, soluble salts are present, it will be necessary to immerse the baked tablets in running water for about six hours to wash away all but a negligible residue of soluble saline matter.

Unlike metal objects, where it is necessary to wash away the last trace of salt because even a small residue might cause a renewed outbreak of corrosion, stone, whether natural or artificial, is not affected by the presence of a slight amount of salt and may be considered safe as long as the bulk of the salt has been eliminated.

Rendering cuneiform writing legible. To render cuneiform inscriptions legible, the old-fashioned methods of surface cleaning such as picking with a needle or treating the baked tablets with hydrochloric acid are now out of date for routine purposes, although they may still be useful on occasion. These methods have been largely superseded by the sand-blasting technique. The sand or shot-blast method of cleaning\(^1\) is both speedy and effective; while removing debris and adherent particles from the inscription, it need not impair the surface of baked clay tablets in any way. A useful portable form of the apparatus has been devised in the British Museum Laboratory. This consists of a brass cylinder containing the sand or other grit. Air is blown through a tube below sand level, so that the particles form a suspension in the air stream and are ejected with the air from the nozzle which may be made from a curved glass or Polythene tube. Such an apparatus may be improvised from ordinary laboratory equipment, the air supply being obtained from a vacuum compressor at about 5 lb. per sq. in. pressure. The potency of the instrument will clearly depend on the nature and size of the particles, the shape of the jet, and the pressure and volume of the air.

\(^1\) The commercial form of shot-blast apparatus is referred to on p. 204.
The apparatus illustrated in Fig. 11 was designed for use in the field. It requires only a source of compressed air, which may be conveniently obtained either from a cylinder or from a motor-driven compressor. For cleaning tablets the jet should not be held too near the writing, nor should the pressure be allowed to become too strong; this is controlled by a movement of the finger over a hole in the feed pipe. The size of the jet must obviously be related to the coarseness of the sand, which is screened by passing it through a sieve of appropriate mesh before it is fed into the cylinder. This excludes the larger particles that would clog the jet.

It should be added that while sand may safely be used out of doors
STONE

(working down wind), it should never be used indoors for health reasons, as it is the cause of the disease known as silicosis. Bauxite grit, however, may be used indoors with safety: it is an aluminium hydrate and cannot cause silicosis though it must be admitted that the grit is just as unpleasant as all other forms of dust, and the operator is well advised to do the work in a fume cupboard or where there is good ventilation. Garnet powders are also sold for grit spraying.

Repairing cuneiform tablets. When a tablet is broken the fragments are baked separately, stuck together with a nitrocellulose adhesive such as Durofix, and then washed and dried. Occasionally one comes across cases where the clay is so crumbly that it would fall to powder if it were heated, and then the only course of action is to try and make good without baking, by consolidating and sticking the particles together. Tablets so treated may be further strengthened, if necessary, by filling cracks with a mixture of nitrocellulose and baked clay powder applied as a thick paste.

Consolidation of objects of unbaked clay

In cases where clay cannot be hardened by firing, as in certain ethnographical specimens and in wall-paintings, consolidation is only possible by impregnation and for this purpose thin lacquers of polyvinyl acetate, polymethacrylate, or nitrocellulose have given successful results. It is not advisable to use aqueous consolidating agents such as glue or casein. These might have a disintegrating effect on the clay in the first instance and, moreover, the water would be liable to affect colours adversely and to bring any soluble salts to the surface. Durofix may be used for repairs, mixed, if necessary, with fine clay. Such mixtures shrink on setting and the non-shrinking nitrocellulose/sand cement referred to on p. 319 is to be preferred.

This cement was used for the surround and backing of Sir Aurel Stein's large frescoes from Chinese Turkestan, now at the British Museum. The ground on which the frescoes were painted was mud and on arrival at the Museum the slabs were laid face downwards and the thick clay backing was cut away. About $\frac{1}{4}-\frac{1}{2}$ in. of the original ground was left and this was reinforced either by a textile
covering or by a sheet of expanded aluminium, the adhesive being nitrocellulose and sand. The frescoes that were backed with textile were then framed upon a solid wooden backing, and then the surround and lacunae in the frescoes filled with the nitrocellulose cement, which had the colour and general appearance of artificial sandstone.

The pieces that were mounted on expanded aluminium frames all belonged to one large fresco and were hung on the wall by attaching them to an armature that had been prepared for their reception. It was possible to set each slab in its permanent position in relation to its neighbours by moving pairs of screws at right angles. All that remained to be done was to make good the joints, using once again the nitrocellulose/sand cement which proved invaluable.

**KIMMERIDGE SHALE**

The dark bituminous substance known as Kimmeridge shale is easily carved and turned, and on this account it was used for making a variety of objects—spinning whorls, weights, bowls, and even furniture in the Roman period. It has deteriorated considerably with the passage of time, however, so that artifacts of shale are usually found in fragments or in such a condition that fragmentation soon follows excavation, and this is due to the shrinkage that accompanies drying.

The form taken by the disintegrating shale depends upon how the shape and ornament are orientated relative to the bedding planes. When the two are roughly parallel, the shale splits into a series of thin distorted leaves which may break up still further; but where the carving is at an angle to the bedding planes, a series of small crescent-shaped cracks appears, and these open up in the manner of a ripened fir-cone as the shale dries. Disintegration will follow unless the object is immersed in water; if this is done in good time, however, the cracks will close up and the surface may assume its original appearance.

The only certain way of preserving all surface details is to keep an object of Kimmeridge shale continuously wet, and this can be done by sealing it in a jar of water containing about 15 per cent. of
A. As received

B. After cleaning

47. T'SUBA FROM THE CITY MUSEUM AND ART GALLERY, BIRMINGHAM
48. 7th-century Frankish Iron Buckle Inlaid with Silver

Left: Before treatment. Right: After treatment

49. Merovingian Purse Mount

Below: X-radiograph of the mount, showing decorations within a mass of iron rust
glycerine, to which about 1 per cent. of carbolic acid is added to act as a preservative and prevent the growth of moulds.

In cases where it is considered desirable to dry the shale, and where a certain distortion of surface can be tolerated, the most effective procedure is to pass the object through a series of alcohols and then to impregnate in a vacuum with a consolidant such as an alcoholic solution of shellac. This, however, is not enough; the surface must be kept under continuous compression until the shellac hardens, and no further movement of the shale is possible. This is carried out as follows. The surface is first covered with absorbent paper (which can easily be removed at the end of treatment) and the object padded, if necessary, and bound tightly with Polythene tape. If the shape allows, the shale is then placed in a carpenter’s vice provided with thick facings of soft wood or cork and allowed to dry for several days under pressure which is so exerted as to close the cracks; additional protection may be given to any protruding portions by the application of shaped pieces of wood and the use of cramps. While such treatment prevents the loss of major fragments, it will be found, after removing the protective tape, that the surface is covered with a system of micro-cracks which have resulted from shrinkage. They are no great disfigurement as they are eventually filled with shellac during the final cleaning process. Ethyl lactate is recommended as the most effective solvent to use for the removal of excessive shellac at the conclusion of treatment.

Objects of shale differ greatly in their appearance and behaviour after excavation, and a warning should be added that the restoration of important specimens should not be lightly undertaken as the process of drying and consolidation may well extend over several weeks, and the work may require regular attention during this time if the results are to be satisfactory.
CHAPTER XV

CERAMICS

It has already been noted that while clay is plastic in the moist condition, it can be hardened to a stone-like mass on heating to redness. This hard condition persists on cooling, and no amount of soaking in water will soften the material again because a permanent and irreversible change has taken place. Baked clay is also highly resistant to chemical action and in this sense it is one of the most stable materials known, rivalling gold itself in permanence.

Raw clays vary in their chemical composition, and in the nature and quantity of the impurities they contain. It is only to be expected, therefore, that considerable diversity will be found in clay products. Examples of pottery of widely differing physical properties may even be made from one and the same clay, depending on the temperature and duration of firing and whether this has been carried out in the presence or absence of oxygen. Surface texture may also be varied in many ways. A common method is to dip the air-dried clay object in an aqueous suspension (slip) of an entirely different clay, prior to firing. Another method is to polish the surface before the clay is fired; this is possible because, in the normal course of air-drying, wet clay passes through a stage described by potters as being 'leather-hard' and in this condition it can be turned on the lathe and burnished, so that when eventually fired the surface is smooth and has a dull polished appearance. Or again, the fired clay or biscuit ware may be glazed.

UNGLAZED WARE

Though it may be regarded chemically as an inert material, baked clay or pottery varies considerably in its physical properties. In-
adequate baking yields pottery which is soft and porous and may be very frail in the wet condition. When excavated, pots are often fragmentary. The pieces require cleaning in order to remove earthy deposits and salts: they require strengthening, and in due course piecing together—with possibly some reconstruction of missing portions in plaster of Paris.

Pottery is often found to be damp through and through when freshly excavated,¹ and in this condition it had best be allowed to dry before much is done in the way of cleaning. Pieces should be marked on the inside surface with waterproof Indian ink, and at this stage any interesting matter examined very carefully in case it should provide information that may be used in interpretation. Cleaning in the field should be delayed when the pot bears evidence of food remains or imprints of seeds as these will require to be submitted to expert examination. In other cases surface dirt may be removed by washing, adding, if necessary, a small amount of a peptizing agent, e.g. a surface-active agent such as Teepol² that will tend to convert the mud from its normal condition of a colloidal gel into that of a sol; it then ceases to adhere to the pottery and can be washed away. Deposits of lime and chalk may remain, and before using solvents to remove them it will be necessary to determine the nature of any existing decoration and its ability to survive cleaning. Soaking in dilute hydrochloric acid might be ideal for cleaning one type of sherd but would certainly not do for all. Thus, acid would be likely to ruin any pots that had been fired at a low temperature or even coarse terracotta excavated from chalky soil where the pores of the ware are filled with calcium carbonate. In this case the acid would react with the carbonate and the resulting effervescence might cause disintegration of the pottery.

 Unglazed earthenware is, of course, easily disrupted by salt, and Egyptian amphorae often display a broken external surface due to the cumulative effect of salt crystallizations. Salt must be washed out

¹ The excavation, recording, and repair of pottery is fully dealt with in R. J. C. Atkinson's Field Archaeology (Methuen & Co. Ltd.), 2nd ed., revised 1953.

by soaking in changes of water and, after drying, the ware may be strengthened by impregnation with a solution of nitrocellulose.

In cases where a pot has been inadequately fired, or where painted decoration is not adhering well, washing with water should be avoided unless the object has been previously consolidated with nitrocellulose. Ostraka are in this category. The writing is normally in a carbon ink, but this might easily be lost in the washing process unless it is first fixed by applying coats of a 2 per cent. solution of nitrocellulose in equal parts of acetone and amyl acetate. After this protective coating is quite dry it is then possible to wash the ostraka in the usual way to remove the salts without any fear of loss of legibility.

**GLAZED WARE**

As glazed pottery is covered by a vitreous film which is waterproof, it is much more permanent than unglazed ware. If, however, the glaze layer is incomplete or imperfect, soluble salts may get into the body of the ware; these will tend to crystallize and cause the glaze to flake off. Although glazes exist in infinite variety, they only seem to give serious trouble when the attachment to the body is threatened.

It is usually a lengthy and difficult business to extract the salt from the body if the only means of getting it out is through cracks in the glaze. This may be facilitated by immersing the object in dilute alcohol, which, though not a very good solvent for salts, will penetrate better than water. The most effective treatment seems to be to apply the paper pulp technique,\(^1\) see p. 299. This is not always possible, however, because the glaze may be loose, or fragments may be held to the ware only by the salt crystals. In such cases it is necessary to fix the glaze by applying a dilute solution of nitrocellulose dissolved in equal volumes of amyl acetate and acetone, and, later on, to wash out the salt in changes of distilled water. Some further consolidation will no doubt be necessary after the salts have been removed.

\(^1\) Penetration will be facilitated by the addition of one volume of Lissapol N to 200 volumes of distilled water.
In common with certain kinds of enamel and glass, glaze is subject to devitrification—a loss of translucency caused by crystallization or deposition of one or more of its ingredients. When silica is deposited in quantity, the glaze will become opaque or it may take the form of an opalescent film veiling the colour and ornament. In the case of certain pieces of Persian pottery it was possible to restore the transparency and to reveal underglaze decoration by local applications of a 1 per cent. solution of hydrofluoric acid for a few minutes at a time, washing with water, and rubbing the treated areas afterwards with wet glass-paper of medium grade. When the opalescent deposit is confined to the immediate surface, as is usually the case, this treatment restores transparency. The cloudiness may go deeper, however, or it may be deposited irregularly, and in this event it may be unsafe to attempt to do more than treat the surface. If by such treatment it is not possible to clear the surface in the first instance, it may be that lacquering will restore transparency, and for this purpose any clear lacquer of high refractive index will suffice.

A common feature of some Chinese pottery is a crackled glaze, and this might seem at first sight to be a defect, but this is very far from being the case. Crackle may arise from many accidental causes, but in the long run it is due to differences in the relative contraction and rate of cooling of the glaze and the body. The Chinese potter was able to control conditions so as to produce a definite artistic crackle and thus make capital out of an apparent deficiency.

REPAIR OF POTTERY

Pottery, earthenware, tiles, and suchlike are all porous, and if they require to be strengthened, this is done by impregnation, using dilute synthetic lacquers containing polyvinyl acetate or polymethacrylate. In some cases, vacuum impregnation may be desirable (see p. 304). Repairs may be carried out with Durofix and, if necessary, a joint may be strengthened by dowelling (see p. 317), or by attaching a bandage along the back where it will be inconspicuous.

It is not possible to repair dusty joints. The first step, therefore, consists in getting rid of dust; then, if the ware is not very absorbent,
the inside of the fractured edge is roughened in order to give a key for the adhesive. The next step consists in painting adhesive on both edges and this is allowed to become tacky. A thin film of fresh adhesive is then applied to the edges, and the joints are pressed firmly together. Pressure should be maintained for at least half an hour by an elastic band or by some other means to ensure that satisfactory joints are formed. In this connexion a sand-box is a great convenience as the repaired pieces may be stuck in the sand temporarily at any appropriate angle, so that joints will not be under strain whilst the adhesive is hardening. Small muslin bags of sand are also useful to support larger and heavier pieces of pottery in the course of reconstruction work. When dry, any excess of adhesive that has been squeezed out from the joint may be removed with a sharp chisel, or even by rubbing with a cloth moistened with a little solvent, though, if the latter expedient is adopted, care must be taken that there is not enough solvent present to run into any cracks and soften the joints.

The following adhesives may be used as alternatives to Durofix. For large work, a retarded plaster, e.g. 'white glue' (hot glue to which has been added plaster of Paris); for dark pottery in which repaired joints may have to be softened by heat in the course of making a complicated reconstruction, thick shellac in spirit or even solid flake shellac applied with the aid of a tiny flame.

For making permanent joints that are very strong and waterproof an epoxy resin adhesive, e.g. Araldite 101, is recommended. In using this adhesive, the pieces must be heated, preferably to at least 180°C, and the Araldite then applied to the hot joints like sealing wax or as a powder according to which is the more convenient; the pottery or porcelain is wired, or otherwise held firmly together, whilst the Araldite is polymerized in the oven. The cured adhesive has a brown colour but is very strong and reliable.

Another adhesive, marketed by the same firm, and which can be specially recommended for unglazed ware, is the resorcinol-formaldehyde resin known as Aerodux 185. This is a cold-setting, gap-filling

1 This is marketed by Aero Research Ltd., Duxford, Cambridge, in the form of sticks or as a powder ready for application.
adhesive resistant to boiling water; it is mixed with a hardener before being applied to the fractured surfaces. The fact that it sets in the cold is a great convenience for many types of objects; and, since it is a gap-filling cement, it has the advantage that the joints which are being fitted together need only be lightly clamped. See also p. 131.

In the reconstruction of pots from broken fragments, it often happens that pieces are missing, and it may be desirable to fill the lacunae with plaster that has been coloured to match the pot. The procedure is as follows. If the missing portion is triangular, clean the three sides and, if the pottery is finely grained, score the edges longitudinally in order to give the plaster a key. Soften some Plasticine by warming and rolling it on a glass plate, and apply a slab, a quarter inch thick, to the inside of the pot covering the missing portion, so that, when the space is filled with soft plaster, the Plasticine will form a backing and reproduce the correct curvature. With narrow-necked vessels this is not possible, and the plaster must be built up from outside, little by little, to fill the space without any backing; the repair will bulge slightly on the inside where it is not seen.

Before applying plaster it is necessary to stop the suction, i.e. to treat the pottery with a liquid in order to prevent it absorbing moisture from the plaster and thus reducing its cohesion. This could be done with water, but pottery varies so much in its condition—strength, porosity, &c., particularly when it has been broken in pieces—that water is better avoided in this kind of repair work. A dilute solution of shellac is much more effective. The joint is painted, therefore, with thin bleached shellac or French polish diluted with methylated spirits.

Coarse baked plaster of Paris is used as a rule unless for exceptionally fine work when boiled plaster is the grade preferred.¹

If a coloured plaster is required for repair work, dry powder colours are added to the plaster powder before it is made up with water. These colours should be mixed well with the plaster on a glass plate, using a steel spatula. When water is added the colour will appear much darker, but on evaporation of the water the plaster

¹ Caffarate and Co. Ltd., Newark on Trent.
will gradually recover the original hue that was established before wetting.

To mix the plaster, put a quantity of cold water in a bowl and shake the powdered plaster on to the surface until it stands as a conical heap above the level of the water. When air bubbles cease to rise, stir with a spoon until a thick cream is obtained that shows signs of becoming thicker within half a minute, when it is ready to be applied to the break. If the plaster fails to thicken quickly in the bowl, do not stir violently, but add some more plaster powder until the correct consistency is obtained before using it to repair the pot.

When the pot has been prepared and the plaster mixed, it may be filled into the gap with the aid of a spoon. Avoid imprisoning air-bubbles and continue filling the gap until the fresh plaster is everywhere slightly above the surface. After a few minutes the plaster will have set to a buttery consistency and it is then an easy matter to pare it down with a plaster scraper (a metal spatula having a comb-like edge) until it is level with the surface of the pot. After half an hour or so the surface may be finished with the appropriate grade of glasspaper—0, 1\(\frac{1}{2}\), or 2, as the case may be. The Plasticine is then removed from the back of the repair and the exposed surface of the plaster made good as required.

If, after drying, it is found that the colour of the repaired area does not match the pot, the whole area of new plaster will have to be stopped again with a thin coating of white French polish before retouching with fresh colour. This is essential, because, if colour is applied without previous stopping, the binding medium will be sucked from the fresh colour into the body of the repair, leaving a ‘watermark’ stain which is difficult to remove. To prepare a matt paint for retouching pottery, the pigments—ochres, umbers, black, white, or green—should be mixed and ground either in skimmed milk or in dilute size solution.
A. Before treatment—the shape almost obliterated by salt crystals

B. After treatment—revealed as the model of a left foot

50. EGYPTIAN SCULPTOR'S TRIAL PIECE OF A FOOT IN LIMESTONE
   (PTOLEMAIC PERIOD)
CARVED STONE FIGURE, PROBABLY HUAXTEC CULTURE, FROM MEXICO
Coated with paper pulp to remove soluble salts
CHAPTER XVI

GLASS

Glass results from the fusion of acidic and basic oxides, the chief acidic oxides being silica and boron oxide, and the chief basic oxides soda, potash, lime, alumina, litharge, and magnesia. When selected mixtures of these oxides are fused at a high temperature, a clear liquid is obtained, which, on cooling, becomes a transparent amorphous solid. Thus, soda-lime glass is made by fusing soda, lime, and clean sand (= silica) in a refractory crucible. This is the commonest type of glass.

The physical properties of glass may be varied according to the nature and proportions of its constituent silicates. For example, potash-lime glass (crown glass) has a high fusion point. Potash-lead glass (flint glass)\(^1\) is softer and more easily cut and it has, moreover, a high refractive index, and is sometimes referred to as ‘crystal’.

In the molten condition, glass is an excellent solvent for metallic oxides, and some, such as cobalt oxide, copper oxide, and iron oxide, are used to give the glass characteristic colours while retaining transparency; others, such as tin and antimony oxides, cause opacity and impart to the glass a white appearance. Opacity results also from the use of larger quantities of colouring oxides.

Until the seventeenth century, almost all glass was composed of silica, lime, and soda or potash. The glass-makers themselves thought of their material as being made from only two constituents. Neri\(^2\) defines glass as ‘a concrete of salt and sand or stones’. The salt, at

\(^1\) The presence of lead in glass may be detected by applying a drop of hydrofluoric acid in an inconspicuous place and after a moment absorbing the drop in filter paper that has been previously moistened with ammonium sulphide solution. A black stain on the paper indicates the presence of lead.

least in Europe, was derived from plant ashes, but plant ashes contain much besides the alkali salts; they contain lime, alumina, &c., and because of this a stable glass was produced. This early glass was generally coloured dark green by the presence of iron oxide, and unfortunately, in their desire to produce a pure white glass, some makers used salts leached from the ashes and produced a glass dangerously low in lime. This was noticed by Neri, who wrote: 'furthermore, in the finest Glasses, wherein the salt is most purified, and in a greater proportion of salt to the sand, you shall find that such Glasses standing long in subterraneous and moist places will fall to pieces, the union of the salt and sand decaying'.

If the early glass-makers had been using pure ingredients, their product, potassium or sodium silicate (known as water-glass), would have been soluble in water. Other ingredients are essential to make a stable glass. Of the silicates that are combined to produce soda-lime glass, sodium silicate is soluble in water and readily fusible, calcium silicate is insoluble and fuses with difficulty; but a correctly balanced mixture of the two combines to form a glass that is insoluble and fuses at a moderate temperature.

In an interesting paper on the composition and working properties of ancient glasses, Matson\(^1\) states that 'The eutectic mixture in the soda-lime-silica system is, approximately, silica 73 per cent., lime 5 per cent., soda 22 per cent.', and he advocates using this as a yardstick in studying analyses of ancient glasses, some of which he quotes. By studying the complete analysis, he shows that one can hazard a guess at the working properties of the glass, e.g. excess of silica makes the glass very difficult to work. But what is more important from the point of view of conservation, the analysis reflects also upon the stability of the glass, excess of lime tending to promote devitrification and excess of soda tending to make the glass susceptible to attack by water. It is not only the presence of certain ingredients but the proportions in which they are present that determine the stability of the material.

In considering the stability of glass there is yet another factor to

\(^1\) Matson, F. R., *Journal of Chemical Education*, 1951, 28, p. 82.
be taken into account. From the physico-chemical point of view, glass is regarded as a state of matter rather than a specific material. It is a supercooled liquid and therefore a metastable substance, and it is characteristic of material in this condition that crystallization may take place at any time. Glasses having a balanced formula are remarkably stable substances, but where injudicious changes are made there is a tendency for one or other of the ingredients of glass to crystallize out, and when this happens the condition is known as devitrification. As we have seen in the case of glazes, devitrification causes opacity and opalescence; a partial devitrification may sometimes enhance the appearance of the glass, particularly striking effects of iridescent colour resulting from the diffraction of light by the thin films or flakes of which the surface of devitrified glass is composed.

Deterioration of glass

The decomposition of glass is generally accompanied by the liberation of free alkali, and the form taken by the disintegration will be determined by the nature and amount of the alkali liberated. Free alkali is more or less hygroscopic, and liberation of lime and soda will cause deposition of moisture. Carbon dioxide will be absorbed from the atmosphere by this moist alkali, with the result that an incrustation of alkaline carbonates is gradually laid down, interspersed with silica, and with tiny flakes of semi-decomposed glass, the result being the creation of a surface that is opalescent.

An opalescent surface requires no special treatment as it is stable under dry conditions. Unfortunately such a surface is frail and the glass itself brittle, and as any attempt to fix the flaking particles with lacquer might destroy the colour effect, lacquering should in general be avoided.¹ To remove dirt, such glass may be washed with a detergent and dried, preferably through baths of alcohol and ether.

When much alkali is present, conditions are very different.

¹ Nevertheless, remarkable effects have been achieved by using a special polymethacrylate lacquer under controlled conditions as described by Hedvall, J. A., et al., Chalmers Tekniska Högskolas Handlingar, 1951, Nr. 118.
Potassium salts, for example, are much more hygroscopic than those of lime and when present in excess no incrustation forms, but instead, beads of potassium carbonate solution of a strongly alkaline nature run down the glass and hasten its decomposition. Glass undergoing such decomposition is known as ‘sweating’ or ‘weeping’ glass and, if left untreated, will soon deteriorate in appearance and in time be destroyed completely. So long as the glass is damp it may remain transparent; unfortunately when the hygroscopic salts are removed and the glass is dry, the transparency is lost and can never be restored completely even by lacquering.

‘Weeping’ glass is treated as follows: after washing in running water for a few minutes, place in a bath of sulphuric acid (2 per cent.) for several days to remove free alkali. Wash it once more and then dry, preferably through alcohol and ether. This treatment retards disintegration but can offer no permanent cure. Glass liable to ‘sweat’ must be kept under reasonably dry conditions. Individual pieces may be kept in airtight boxes supplied with enough silica gel to keep the contents dry. Such boxes can be constructed from Perspex and sealed at the joints with Polythene tape. As an alternative, an exhibition case may be adapted for the purpose as shown in Fig. 9.

Much ancient glass must have been lost to us as a result of decomposition arising from an excess of lime in the formula or from imperfect mixing of the ingredients, due to inadequate heating during manufacture. In the case of opaque glass having what appears to be a stable surface, any heterogeneity in the interior will be concealed from view so long as the outer shell is intact. Such an exceptional condition was illustrated in the case of a large Egyptian scarab of the eighteenth dynasty that had been made by casting opaque blue glass in a mould. The scarab had remained intact for 3,000 years; it had been on exhibition in the British Museum for a generation, and there had been no reason to regard it as being in any respect abnormal. It was reported one day to have developed cracks, and was found to be so hygroscopic that it would not remain dry for two minutes on end. Soon afterwards it fell into about a dozen pieces. From an examination of the interior of the scarab it was obvious that the condition of
52. THE MITHRAS HEAD FROM WALBROOK, LONDON

A. *Left*: Before treatment.  B. *Right*: After cleaning and reuniting with the neck, later discovered on the same site
A. Plaster cast made in 1802 which shows the condition of the marble at that time

B. The marble photographed in situ in 1938, indicating the deterioration resulting from 136 years’ exposure in the industrial age

53. SCULPTURE FROM THE WEST FRIEZE OF THE PARTHENON
the inside was different from that of the exterior—a white incrustation formed on parts of the fractured surfaces but no such crystallization took place on the outside. It only required a micro-crack to form in the outer shell for moisture to have access to the hygroscopic material inside, and when this happened disintegration was inevitable.

Enamels

An enamel is a lead glass to which some metallic oxide has been added to colour it or make it opaque. It is applied thinly to a metal base, usually of gold or silver. When glass is melted on metal, as in the manufacture of enamels, the difference in the degree of contraction on cooling between the glass and the metal is considerable, and sometimes a state of strain is set up between the two. In order to equalize the strains, the enamel is sometimes applied all over the back of the metal as well as the front.

That the strains are not always equalized is evident from a singular occurrence that took place in the Wallace Collection in 1929.¹

An eighteenth-century enamelled snuff-box on exhibition in the museum suddenly disintegrated, scattering glass powder throughout the case. When the powder was examined in the British Museum Laboratory the particles were shown to be of the same shape as those formed by the bursting of Prince Rupert drops, that is, rounded and not splintery. These drops, which have been known for centuries, are pear-shaped blobs of glass made by pouring molten glass into water. Though apparently quite stable, and so strong that they can even be hit at the thick end with a hammer without breaking, they are, in fact, under great internal strain, and have the astonishing property of flying to powder when the smallest fragment is broken from the fine glass filament forming the stem of the pear. When made in the form of a phial it suffices to drop a tiny grain of sand inside for the glass to fall in two, and it is likely that some such simple trigger action caused the enamel in the Wallace Collection to ‘explode’.

Sometimes strain causes an enamel to become fractured and detached from the base, and when this happens the only thing to do is

¹ Camp, S. J., Art Work, 1930, 21, p 47.
to attempt consolidation by flooding the area with dilute lacquer so that it seeps through the cracks. When it has completely set, the excess of lacquer may be removed. In the coarser type of enamel work it is sometimes possible to execute repairs in coloured cellulose paints or varnishes that have been suitably dyed.

A curious effect occasionally to be observed is that of fungus apparently growing on one specific colour—say on a translucent crimson. On closer inspection it is found that the nutrient material is gelatine, which has been used to reinforce the weak areas of the enamelling and possibly to act as a base for retouching with pigment. The treatment in this case is to remove the gelatine with warm water, sterilize the area with Santobrite (2 per cent.), and after drying consolidate with lacquer tinted as required.

The opaque red glass of sealing-wax colour, often referred to as cuprous enamel, which was of universal use in the Bronze Age for the decoration of metal, is often found to be oxidized to a green colour. The earliest examples that have come to the British Museum Laboratory are from a glass furnace excavated at Nimrud (650 B.C.) and are in the form of lenticular cakes from the crucible, some 8 inches in diameter, and up to 2 inches in their greatest thickness. These are invariably green on the surface, and are like corroded bronze in appearance. The green layer is porous and extends to a depth of almost 1/16th inch, the interior being apparently in a perfect state of preservation, rich purplish-red, opaque, and having a characteristic glassy fracture. Unfortunately, in dealing with inlays of this red glass, the only means of revealing the red colour is to grind or scrape away the superficial green layer. This treatment is by no means always practicable, but has been successfully carried out in the case of a number of Saxon hanging bowls decorated with escutcheons in which this red glass was used for enamelling. The removal of the green film enhanced the appearance of the enamel considerably. When the glass is unattached to metal, or where an inlay can be removed for cleaning, it might be worth while trying to dissolve the green surface layer with a dilute solution of hydrofluoric acid. Precautions are necessary, however. Immersion should be for short
periods only; the work is inspected at frequent intervals by washing under a running tap and the acid treatment stopped before all of the green has disappeared. When washing with hydrofluoric acid a plastic photographic dish should be used, and rubber gloves should be worn to protect the skin, particularly under the finger-nails, where it is very sensitive to this acid.

*Repairing glass*

Broken glass is one of the most difficult materials to repair. If thermo-setting adhesives can be used, Araldite 101 gives excellent results and it has actually been used to mend the broken stem of a wine glass. But it is seldom possible to take the risk of heating old glass which, if devitrified, would almost certainly disintegrate. Durofix is probably the most useful general-purpose adhesive, and another good one is made by dissolving chips of Perspex in the following mixed solvent, to give a solution of 10–15 per cent. strength:

\[
\begin{align*}
\text{Ethylene dichloride} & \quad \ldots \quad 195 \text{ ml.} \\
\text{Glacial acetic acid} & \quad \ldots \quad 5 \text{ ml.}
\end{align*}
\]

Acid solutions of glue may also be recommended, and a reliable adhesive can be made by dissolving Scotch glue (20 grams) in 30 ml. of hot glacial acetic acid to which 1 gram of ammonium bichromate is added. The glue is warmed in the usual fashion before application.

Adhesives are almost useless for the repair of glass that is very thin or much fractured. Where fragments of a glass vessel are in such a condition that its shape can be recovered, a transparent Polythene tape may be used, applying it to the inside, and beginning the reconstruction preferably at the mouth of the vessel. This technique, however, requires dexterity and leaves much to be desired. It must be admitted that in this category of repair work there are cases that are apparently beyond recovery.
APPENDIXES
I. STRENGTH OF SOLUTIONS

1. Conventions

Liquids dissolved in liquids are measured in fluid ounces per pint or in millilitres per litre: i.e. volume/volume, (v/v.).
Solids dissolved in liquids are measured in ounces avoirdupois per pint or in grams per litre: i.e. weight/volume, (w/v).
10 per cent. v/v solutions contain 2 fluid oz./pint or 100 mls./litre.
10 ,, w/v ,, ,, 2 avoird. oz./pint or 100 gms./litre.

Acids and alkalis. Concentrated acids and alkalis in liquid form are sold by specific gravity as they are not necessarily of 100 per cent. strength, thus:

<table>
<thead>
<tr>
<th>Acid</th>
<th>Sp. Gr.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sulphuric acid</td>
<td>1.84</td>
</tr>
<tr>
<td>Nitric acid</td>
<td>1.42</td>
</tr>
<tr>
<td>Hydrochloric acid</td>
<td>1.18</td>
</tr>
<tr>
<td>Ammonia</td>
<td>0.88</td>
</tr>
</tbody>
</table>

The convention adopted in this book, in preparing dilute solutions, is to assume that the commercial chemical is 100 per cent. and dilute accordingly. Thus,

1 per cent. nitric acid is made by dissolving 1 fluid oz. of nitric acid (1.42) in 5 pints distilled water.
10 per cent. sulphuric acid is made by dissolving 2 fluid oz. of sulphuric acid (1.84) in 1 pint of distilled water.
50 per cent. ammonia is made by mixing equal volumes of ammonia (0.88) and distilled water.

2. Hydrogen peroxide

This is available, commercially, as follows:

'100 volumes' = approx. 30 per cent. w/v hydrogen peroxide, sometimes called Perhydr. Perhydr. and 100 per cent. w/v hydrogen peroxide.

'20 volumes' = approx. 6 per cent. w/v hydrogen peroxide.

These solutions are supplied in coloured bottles and should be stored in a cool place protected from light.
3. Enzymes

Enzymes are naturally-occurring organic substances that have a specific action in catalysing many organic reactions, including the breakdown of sugars, fats, and proteins. Mixtures of enzymes are obtainable, called 'digestor powder', and this may be used for rendering certain organic stains soluble, particularly those occurring on textiles (see p. 105), so that they may be removed subsequently by washing. Enzymes act best in weak solutions at temperatures between 37 and 50° C.—they are soon destroyed at temperatures in excess of 60° C. The hydrolytic action of enzymes takes time to complete, perhaps an hour or longer, and the process is known as digestion.
II. DANGEROUS CHEMICALS

Although many chemicals may be regarded as potentially dangerous, special care is needed when using the following:

(1) Concentrated sulphuric acid
This can cause a serious burn if brought in contact with the skin. In preparing dilute solutions of sulphuric acid the concentrated acid should always be added slowly to the water with continuous stirring to dissipate the large amount of heat evolved. Never add water to the concentrated acid.

(2) Caustic soda
The solid should not be allowed to come in contact with the skin. In preparing solutions of caustic soda there is a considerable evolution of heat which may cause a thick glass vessel to crack. Such solutions are therefore best made by slowly adding the solid (with care to prevent splashing) to water contained in a porcelain dish or an iron or stainless steel vessel standing in a sink.

(3) Concentrated nitric acid
Strongly corrosive and should not be allowed to come in contact with the skin, which it stains a deep yellow.

(4) Concentrated hydrochloric acid
Care should be taken when opening a bottle of this acid, as fumes are immediately evolved which will cause pain if they come in contact with the eyes.

(5) Hydrofluoric acid
Strongly corrosive. Usually handled in Polythene bottles. The fumes are very unpleasant if inhaled and will damage any glass with which they come in contact.

(6) Formic acid. Carbolic acid (Phenol)
Powerful skin irritants.
(7) Concentrated ammonia
Care should be taken when opening a bottle of concentrated ammonia; the stopper may fly off due to pressure. The bottle should be kept in a cool place and must never be allowed to remain in the sun.

(8) Hydrogen peroxide '100 vols.' Perhydrol
This must be kept in a cool place as great pressure may be set up in the bottle if left in the sun or near a source of heat.

(9) Diethyl ether (B.P. 34·6° C.)
Highly volatile and inflammable. Forms explosive mixtures with air.

(10) Carbon disulphide (B.P. 46° C.)
Highly volatile and inflammable. Poisonous. Forms explosive mixtures with air.
N.B. A foam fire extinguisher should be at hand in any room where ether or carbon disulphide are used.

FIRST AID

(1) Acid burns. Flood the affected area with water and then wash with a dilute solution of sodium bicarbonate.

(2) Caustic soda burns. Flood the affected area with water and then wash with very dilute acid, e.g. vinegar.
III. PREPARATION AND USE OF FLOUR PASTE

Adhesives for paper should be free from staining material and harmful preservatives, and while the best photographic mountants may satisfy these conditions, cheap office pastes and adhesives supplied for hanging wall-paper should be avoided as well as all preparations of doubtful or unknown composition.

For general work there is nothing to surpass freshly prepared flour paste (bookbinder's paste), as it is easy to make and to apply and has excellent adhesive properties. It is made from ordinary (not self-raising) flour and water in the following proportions—

<table>
<thead>
<tr>
<th>Ingredient</th>
<th>Quantity</th>
</tr>
</thead>
<tbody>
<tr>
<td>White flour (wheat)</td>
<td>500 grams</td>
</tr>
<tr>
<td>Water</td>
<td>2.5 litres</td>
</tr>
</tbody>
</table>

Mix the flour with a little of the water in an aluminium or enamel pan, the lumps being broken up with the hand to form a smooth cream. Boil the remainder of the water separately and add it to the cream, stirring continuously. The paste is now heated, not directly, but by standing the pan in a container of water kept boiling (double pan. See Pl. 55). It should be stirred meantime and will soon thicken. After about ten minutes it may be decanted into a suitable vessel, and to prevent a crust forming as it cools place a sheet of paper on the surface of the paste and pour a little water above the paper. The mixture as prepared is too thick for general use: small quantities should be removed to a pasting dish and thinned with water as required. The paste keeps well for a few days in cool surroundings but should be discarded on the first sign of souring.

If it is desired to keep the paste for as long as a week, about 10 ml. of formalin may be stirred into the mixture while fresh, but for ordinary use no other form of preservative can be recommended. Alum is often present as an ingredient in formulas for paste; it hardens the adhesive so that it tends to become insoluble in water, but alum has been shown to be harmful as it makes the paper acid (see p. 52). For use in the tropics, however, the paste must contain a substance that will inhibit the growth of mould and render it unattractive to insects, and for this purpose poisonous material is introduced. Suitable formulas have been suggested
containing small quantities of phenyl mercuric acetate, or phenyl mercuric borate, say, about half a gram for every 2½ litres of paste.

In applying paste it should be spread evenly and thinly so that it will dry quickly, and there should never be so much water present as to cause the paper to stretch appreciably. The paste may be applied with a brush or with an ivory or bone paper-knife. After pasting, the paper should be dried in the press or under a sheet of heavy plate glass.
IV. PEARSON’S SQUARE FOR SIMPLIFYING DILUTION CALCULATIONS

\[ x = B - C \]
\[ y = A - B \]

Examples:
(1) How much water has to be added to a 12 per cent. solution to make a 5 per cent. solution?
Let \( A = 12 \) per cent.
\( B = 5 \) per cent.
\( C = 0 \) per cent. (water)
Then \( x = B - C = 5 - 0 = 5 \) parts of the 12 per cent. solution \((A)\)
\( y = A - B = 12 - 5 = 7 \) parts of water \((C)\)
Answer: Mix 5 parts of the 12 per cent. solution with 7 parts of water.

(2) How much of a 45 per cent. solution has to be added to a 17 per cent. solution to obtain a 30 per cent. solution?
Let \( A = 45 \) per cent.
\( B = 30 \) per cent.
\( C = 17 \) per cent.
Then \( x = B - C = 30 - 17 = 13 \) parts of the 45 per cent. solution \((A)\)
\( y = A - B = 45 - 30 = 15 \) parts of the 17 per cent. solution \((C)\)
Answer: Mix 13 parts of the 45 per cent. solution with 15 parts of the 17 per cent. solution.
V. ACIDITY, NEUTRALITY, ALKALINITY

USE OF INDICATORS

An acid may be defined as a substance which forms hydrogen ions when dissolved in water, and an alkali as a substance which forms hydroxyl ions when dissolved in water. Acids and alkalis are capable of neutralizing one another to form salts.

When dealing with small concentrations of acids and alkalis it is cumbersome to express the actual concentrations of hydrogen and hydroxyl ions in terms of weight per volume. A convenient method was proposed in 1909 by Sörensen who introduced the hydrogen ion exponent, commonly known as pH. The relationship between pH and hydrogen ion concentration is shown in the following nomograph:

![Diagram showing the pH scale and hydrogen ion concentrations]

The advantage of the system is that very small concentrations of acid or alkali can be expressed as whole numbers. A solution in which the hydrogen and hydroxyl ion concentrations are equal is an exactly neutral solution and the pH = 7. In an acid solution the hydrogen ion concentration exceeds that of the hydroxyl ion and the pH is less than 7, whilst in an alkaline solution the reverse is the case and the pH is greater than 7.

Certain substances known as indicators show characteristic colour changes depending upon the pH of the solution to which they are added. Litmus, for example, is red in acid and blue in presence of alkalis. Its pH range is 5.0–8.0. A more precise estimation of the acidity or alkalinity of a solution may be made by adding to it a few drops of Universal Indicator\(^1\) which shows graded colour changes from red to violet throughout the pH range as follows:

\(^1\) Indicators of this type are B.D.H. Universal Indicator obtainable from British Drug Houses, Poole, Dorset, and B.T.L. Universal Indicator obtainable from Messrs. Hopkin & Williams Ltd., Freshwater Road, Chadwell Heath, Essex.
<table>
<thead>
<tr>
<th>Colour of indicator</th>
<th>Nature of solution</th>
<th>pH value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Red</td>
<td>Very acid</td>
<td>3-4</td>
</tr>
<tr>
<td>Orange</td>
<td>Moderately acid</td>
<td>5-5.5</td>
</tr>
<tr>
<td>Yellow</td>
<td>Slightly acid</td>
<td>6-6.5</td>
</tr>
<tr>
<td>Greenish-yellow</td>
<td>Neutral</td>
<td>7-7.5</td>
</tr>
<tr>
<td>Green</td>
<td>Slightly alkaline</td>
<td>8</td>
</tr>
<tr>
<td>Bluish-green</td>
<td>Moderately alkaline</td>
<td>8.5</td>
</tr>
<tr>
<td>Blue</td>
<td>Very alkaline</td>
<td>9.5</td>
</tr>
<tr>
<td>Violet</td>
<td>Intensely alkaline</td>
<td>10-12</td>
</tr>
</tbody>
</table>
## VI. THERMOMETRIC EQUIVALENTS

<table>
<thead>
<tr>
<th>Centigrade</th>
<th>Fahrenheit</th>
<th>Centigrade</th>
<th>Fahrenheit</th>
<th>Centigrade</th>
<th>Fahrenheit</th>
</tr>
</thead>
<tbody>
<tr>
<td>100° C.</td>
<td>212° F.</td>
<td>66° C.</td>
<td>150°8° F.</td>
<td>32° C.</td>
<td>89°6° F.</td>
</tr>
<tr>
<td>99</td>
<td>210.2</td>
<td>65</td>
<td>149</td>
<td>31</td>
<td>87.8</td>
</tr>
<tr>
<td>98</td>
<td>208.4</td>
<td>64</td>
<td>147.2</td>
<td>30</td>
<td>86</td>
</tr>
<tr>
<td>97</td>
<td>206.6</td>
<td>63</td>
<td>145.4</td>
<td>29</td>
<td>84.2</td>
</tr>
<tr>
<td>96</td>
<td>204.8</td>
<td>62</td>
<td>143.6</td>
<td>28</td>
<td>82.4</td>
</tr>
<tr>
<td>95</td>
<td>203</td>
<td>61</td>
<td>141.8</td>
<td>27</td>
<td>80.6</td>
</tr>
<tr>
<td>94</td>
<td>201.2</td>
<td>60</td>
<td>140</td>
<td>26</td>
<td>78.8</td>
</tr>
<tr>
<td>93</td>
<td>199.4</td>
<td>59</td>
<td>138.2</td>
<td>25</td>
<td>77</td>
</tr>
<tr>
<td>92</td>
<td>197.6</td>
<td>58</td>
<td>136.4</td>
<td>24</td>
<td>75.2</td>
</tr>
<tr>
<td>91</td>
<td>195.8</td>
<td>57</td>
<td>134.6</td>
<td>23</td>
<td>73.4</td>
</tr>
<tr>
<td>90</td>
<td>194</td>
<td>56</td>
<td>132.8</td>
<td>22</td>
<td>71.6</td>
</tr>
<tr>
<td>89</td>
<td>192.2</td>
<td>55</td>
<td>131</td>
<td>21</td>
<td>69.8</td>
</tr>
<tr>
<td>88</td>
<td>190.4</td>
<td>54</td>
<td>129.2</td>
<td>20</td>
<td>68</td>
</tr>
<tr>
<td>87</td>
<td>188.6</td>
<td>53</td>
<td>127.4</td>
<td>19</td>
<td>66.2</td>
</tr>
<tr>
<td>86</td>
<td>186.8</td>
<td>52</td>
<td>125.6</td>
<td>18</td>
<td>64.4</td>
</tr>
<tr>
<td>85</td>
<td>185</td>
<td>51</td>
<td>123.8</td>
<td>17</td>
<td>62.6</td>
</tr>
<tr>
<td>84</td>
<td>183.2</td>
<td>50</td>
<td>122</td>
<td>16</td>
<td>60.8</td>
</tr>
<tr>
<td>83</td>
<td>181.4</td>
<td>49</td>
<td>120.2</td>
<td>15</td>
<td>59</td>
</tr>
<tr>
<td>82</td>
<td>179.6</td>
<td>48</td>
<td>118.4</td>
<td>14</td>
<td>57.2</td>
</tr>
<tr>
<td>81</td>
<td>177.8</td>
<td>47</td>
<td>116.6</td>
<td>13</td>
<td>55.4</td>
</tr>
<tr>
<td>80</td>
<td>176</td>
<td>46</td>
<td>114.8</td>
<td>12</td>
<td>53.6</td>
</tr>
<tr>
<td>79</td>
<td>174.2</td>
<td>45</td>
<td>113</td>
<td>11</td>
<td>51.8</td>
</tr>
<tr>
<td>78</td>
<td>172.4</td>
<td>44</td>
<td>111.2</td>
<td>10</td>
<td>50</td>
</tr>
<tr>
<td>77</td>
<td>170.6</td>
<td>43</td>
<td>109.4</td>
<td>9</td>
<td>48.2</td>
</tr>
<tr>
<td>76</td>
<td>168.8</td>
<td>42</td>
<td>107.6</td>
<td>8</td>
<td>46.4</td>
</tr>
<tr>
<td>75</td>
<td>167</td>
<td>41</td>
<td>105.8</td>
<td>7</td>
<td>44.6</td>
</tr>
<tr>
<td>74</td>
<td>165.2</td>
<td>40</td>
<td>104</td>
<td>6</td>
<td>42.8</td>
</tr>
<tr>
<td>73</td>
<td>163.4</td>
<td>39</td>
<td>102.2</td>
<td>5</td>
<td>41</td>
</tr>
<tr>
<td>72</td>
<td>161.6</td>
<td>38</td>
<td>100.4</td>
<td>4</td>
<td>39.2</td>
</tr>
<tr>
<td>71</td>
<td>159.8</td>
<td>37</td>
<td>98.6</td>
<td>3</td>
<td>37.4</td>
</tr>
<tr>
<td>70</td>
<td>158</td>
<td>36</td>
<td>96.8</td>
<td>2</td>
<td>35.6</td>
</tr>
<tr>
<td>69</td>
<td>156.2</td>
<td>35</td>
<td>95</td>
<td>1</td>
<td>33.8</td>
</tr>
<tr>
<td>68</td>
<td>154.4</td>
<td>34</td>
<td>93.2</td>
<td>0</td>
<td>32</td>
</tr>
<tr>
<td>67</td>
<td>152.6</td>
<td>33</td>
<td>91.4</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
A. Before treatment—the tablets covered with a salt deposit making them illegible

B. After treatment—the clay stabilized, the salts removed, and legibility restored

54. A COLLECTION OF CLAY TABLETS INSCRIBED IN SUMERIAN, BABYLONIAN, AND ASSYRIAN CUNEIFORM SCRIPTS
55. APPARATUS FOR MOUNTING PRINTS

A Glass-topped Table; B Over-throw Mount; C Guard Papers; D Paste-pot and Brush; E Sponge; F Spatulas for spreading Paste; G Guillotine; H Slotted Card with Chamfer Template; I Print-trimming Knives; J Weighing Flour to make Paste; K Paste-making; L Covered Paste; M Improvised Press
VII. METHODS OF RECORDING RELATIVE HUMIDITY

Temperature and relative humidity readings are taken with a thermometer and hygrometer respectively. Several types of hygrometer are available,¹ the most convenient being the sling or whirling hygrometer or psychrometer which consists of wet- and dry-bulb thermometers fixed side by side on a frame for swinging, the wet bulb being kept damp by a wick dipping into a reservoir containing distilled water. To use the instrument it is held well away from the body and is whirled vigorously for at least a minute, readings on the wet-bulb and dry-bulb thermometers being taken immediately. Further observations should be taken till constant readings are obtained. The wet bulb will generally record a temperature lower than the dry one. This difference represents the depression in degrees resulting from evaporation of water, and from this depression, taken in conjunction with the temperature of the dry bulb, it is possible to determine the relative humidity by consulting hygrometric or psychrometric tables.²

The sling hygrometer has the advantage that it may be used in different parts of the room, and it will reveal the existence of any humidity gradient in the atmosphere, so that the effect of ventilation near doors and windows can be readily assessed. When daily readings are to be taken in a room, the instrument must be used in the same places, under the same conditions, and at the same time of day if results are to be comparable. When mildew is detected in a cupboard or among books, it is obvious that readings taken at the door or near a window, where the results would be influenced by ventilation, would not be a true indication of the atmospheric conditions in the danger area. Readings should be taken at the point where damp is suspected. In making a thorough assessment, conditions at night should not be neglected because in a closed room the relative humidity of the atmosphere increases with fall of temperature. The effect is most striking when heating appliances are of a type that become suddenly cold, such as steam radiators or electric fires.

² Hygrometric Tables, on Ivorine Cards, Negretti & Zambra Ltd., London; Psychrometric Tables, Marvin, C. F., U.S. Weather Bureau, 1941.
Self-recording types of hygrometer are essential when it is necessary to obtain a continuous record. In the usual recording hygrometer an ink line is drawn by a pen on a paper cylinder which is revolved by clockwork. The up-and-down movements of the pen are controlled by the contraction and expansion of some moisture-sensitive substance, e.g. strands of hair, stretched within the instrument. By this means it is possible to obtain a record of the relative humidity as a graph on a time scale. Readings from self-recording hygrometers may be misleading, however, unless such instruments are calibrated by a weekly check with the whirling hygrometer (the more reliable instrument), and the humidity graph should be annotated at the time of observation. In this way it is possible to observe the amount of any slight error on the chart which may arise from slowness of response due to friction, &c., and the correction is taken into account in interpreting the records.

Dial forms of non-recording paper hygrometer have the merit of compactness and are convenient for use in restricted spaces, but suffer like the self-recording types from a tendency towards slow response to rapid changes in the moisture content of the air.
VIII. THE PEROXIDE TEST FOR VEGETABLE-TANNED LEATHER

This is an accelerated decay test, and its value lies in that it will reveal in the course of a week whether a vegetable-tanned leather is suitable for bookbinding, i.e. whether it will survive exposure to a polluted atmosphere without deterioration. It may be applied to all vegetable-tanned leathers available today, but the test would not be applicable in the case of leathers coming on the market in which the iron has been sequestrated (see p. 36).

The test is carried out as follows:

A sample of leather, 2½ in. square and weighing from 2 to 6 grams, is laid on a glass plate, flesh side upwards, and evenly moistened with sulphuric acid (5 per cent.) in the proportion of 1 ml. per gram of air-dried leather. The acid can conveniently be applied with a capillary pipette, and smoothed out on the leather with a glass rod. After remaining at room temperature overnight, hydrogen peroxide (10 vol. strength) is added evenly to the leather, dropwise, in the proportion of 0·6 ml. per gram of leather. It is left for twenty-four hours, and then given five further daily doses of hydrogen peroxide; this treatment will cause unsatisfactory leather to be blackened and gelatinized, but durable leather will survive, except for possible discoloration of the edges. Changes in the colour of the dye-stuffs are immaterial.
IX. POTASSIUM LACTATE SOLUTION

Dissolve 2 oz. of potassium lactate in 1 pint of water. Vegetable-tanned leather that is free from any sign of chemical deterioration may be protected against the deleterious action of sulphur dioxide by spraying or sponging it with this solution on both sides of the skin. This ensures, as far as possible, that freshly tanned leather will be a durable material.

The solution should be applied to the outside of leather bindings that have been washed, in order to replace protective salts that may have been washed from the surface of the leather.

Chemical decay, when once started, cannot be arrested by applying potassium lactate; the treatment is a form of protection, not a cure.

Do not store potassium lactate solution for long periods of time; it may be preserved for a limited period in a stoppered bottle by adding a little chloroform, but it is better to use the solution fresh and discard any residue.

It is unnecessary to apply lactate solution to parchment or vellum, or to alum-tawed bindings, as these are not subject to chemical deterioration by sulphur dioxide.
X. BRITISH MUSEUM LEATHER DRESSING

The ingredients are:

- Lanolin (anhydrous) 7 oz. or 200 g.
- Cedarwood Oil 1 fluid oz. or 30 ml.
- Beeswax 1/2 oz. or 15 g.
- Hexane (or Petroleum Ether B.P. 60-80° C.) 11 fluid oz. or 330 ml.

These are compounded to form a yellow cream. It is highly inflammable, and no naked light must be allowed in the room during application of the preservative and for some time afterwards. It should be applied sparingly and rubbed into the leather, and two days later the surface may be polished with a soft cloth or brush.

This dressing is obtainable from Messrs. Baird & Tatlock (London) Ltd., 14 St. Cross Street, Hatton Garden, E.C. 1.
XI. SILICA GEL

Silica gel is an effective drying agent which has the considerable advantage that it does not become moist to the touch in use, nor does it cause staining; in fact the appearance of the material remains unchanged. Since it is essential to have some means of telling when it is 'spent' and no longer able to absorb moisture, it is tinted during manufacture with cobaltous salts which are deep blue in the dry condition, but become pink when silica gel has absorbed moisture. When silica gel has assumed a pink colour, it can be readily reconditioned by heating in an oven until it regains its original deep-blue colour. This cycle of reheating can be repeated indefinitely.
XII. MICROCRYSTALLINE POLISHING WAXES

There are two types, the aqueous emulsion type and the conventional solvent type, and the latter is generally preferable for application as a surface coating to antiquities and works of art.

A microcrystalline wax salve may be applied with the object of removing surface dirt, adjusting the optical quality, enhancing the appearance, or excluding moisture, and when compounded with a Polythene wax as described in the following recipe it has been found to stabilize painted or varnished surfaces that are liable to bloom.

Several varieties of microcrystalline and Polythene waxes are available and the gloss of the synthetic wax film can be varied by altering the grades and proportions of the waxes used.

The following basic recipe has been found to be generally satisfactory: Cosmolloid 80 Hard¹ (100 g.) and BASF Wax A² (25 g.) are cut into small pieces and melted together, care being taken to ensure that the Polythene wax is thoroughly dispersed. The molten mixture is poured quickly into white spirit (300 ml.), taking precautions against fire risk, and, while cooling, is constantly stirred so that a paste of agreeable consistency is obtained. This is stored in screw-top cosmetic jars.

Matt waxes can be produced by using Cosmolloid Soft in the proportion of 10 parts to 1 of Polythene wax.

[It should be noted that although the white spirit evaporates, it may have a softening action on varnish of the polycyclohexanone type (AW2). As this is not the case with petroleum ether (boiling range 80-100° C.), the latter should be used as the diluent instead of white spirit where varnishes based on Resin AW2 are to be waxed.]

The basic formula can be modified for use in the Tropics as a leather dressing and it will afford protection against the attack of insects if lauryl pentachlorophenate is incorporated in a concentration of about 10 per cent. When a wax polish is required for preserving bright steel, about 10 per cent. of sodium benzoate should be incorporated in the standard formula.

¹ Cosmolloid microcrystalline paraffin wax, marketed by Astor, Boisellier, & Laurence Ltd., 9 Savoy Street, London, W.C. 2.
² BASF Wax A (Polythene wax), marketed by Bush, Beach, & Gent Ltd., Marlow House, Lloyd's Avenue, London, E.C. 3, is now obtainable in the form of pellets.
XIII. METHOD OF ADAPTING MAINS ELECTRIC SUPPLY FOR ELECTROLYTIC REDUCTION

**Apparatus Required:**
- One Transformer 18 v. 12 a.
- One Bridge-Connected Funnel-Cooled Selenium Rectifier 17 v. 12 a.
- One Variable Resistance 3.5 ohm 12 amp. max.
- One Ammeter 0–15 a, M.C. Flush Mount.
- One Pair of Large Terminals: One Red, One Black (Belling Lee).
- One Pair of Slydlock (or other) Fuses.
INDEX

Abrasives: for polishing metals, 205-6, 213, 216, 231, 237, 268, 291; for rubbing down stonework, 313-14.
Acclimatization of waterlogged objects, 141.
ACP Deoxidine No. 125, 277, n. 3.
Adhesives, used in repairing: baked clay, 324; bronze, 255; canvas paintings, 167-8, 169; clay tablets, 323; frail textiles, 106, 108; glass, 339; ivory and bone, 152; leather, 32; marble, 313, 317; painted panels, 160, 164; pottery, 320-2; seals, 91-92; steel, 285; stone objects, 317, 318-19; tears in paper, 86, 87, 347-8; wood, 131.
Aerodux No. 185, 330-1.
Aktolite No. 300, 132.
Agora, lead objects from, 261.
'Ain Feshkha, Palestine, finds at, 40.
Air conditioning, 4 ff. & n.
Alabaster, 295.
Alcho-Re solder paste, 288.
Alcohol-ether-resin, and waterlogged wood, 136-8.
Alginate, use for moulding, 153.
Alloys, 188-9, 189 n.; decay and, 190; drying of, 199; cleaning of, 205; of gold, 207, 208; of silver, 220, 221, 224, 230; of copper, 232 ff., 248, 256; gun-metal and bell-metal, 247; speculum, 248-9, 288; of lead and tin, 258, 270; and soldering, 286.
Alum, 23, 24, 25, 52, 135-6, 139, 347.
Amber, 156.
Amberlite IR No. 120, 263 n.
Antimony, 258.
Antisun glass, 112 n.
Antler, 155-6.
Antrobus, Mrs. Guy, 109.
Armitage, F. D., 52 n. 2, 54 n. 2.
Athenaeum Club, 19.
Atkinson, R. J. C., 327 n. 1.

Atmospheric pollution, 4-5, 11 ff., 213, 259, 277, 316; dust, 95, 171, 210, 307; paint fumes, 259; sea-air, 11; soot, 4, 11, 95, 171, 210, 297, 307; sulphurous gases, 11-12, 19, 34, 35, 36, 37, 52-53, 112, 166, 170-1, 188, 213-15, 277.
Atomizers, 30, 115.
Australia, Hartogs plate from, 266.
Bacteria: and leather degradation, 22, 26; and textiles, 93; sulphate-reducing, 138-9, 272-3, 282; and water paints, 213; and iron corrosion, 272-3.
Bakelite, 130 & n. 3.
Baker's Soldering Fluid, 287.
Bark-cloth, 133-4.
Barker, H., 244 n.
Basalt, 296, 297.
BASF Wax A, 359 & n. 2.
Basketry, 133.
Bastille medal, 268.
Bedacryl 122 X, 130 & n. 1, 257, 304.
—L, 151.
Beetles, wood-boring, 121-3.
Belgian Museums, Central Laboratory of, Brussels, 128 & n. 2.
Bellinger, Louisa, 105.
Bengtson, Bengt, 215.
Bennister, H. L., 279 n. 2.
Bick, L., 279 n. 2 & 3.
Binding mediums in painting, 158, 169-70.
Birch bark, 44.
Black Prince relics, 107 & n.
Bleaching: of paper, 59-60, 64-65, 75-76; of stained prints, 74 ff.; of old textiles, 100; of waterlogged wood, 136, 138.
—agents, 75 ff.
Blistering, paint, 164-6.
Bloom, 10, 172-3.
INDEX

Blotting-paper, use of, for: relaxing papyrus, 43; print cleaning, 70 ff., 83; crease-removal, 85; mounting, 88; absorbing grease stains, 101, 108; applying reagents, 148, 149, 154; flattening wooden panels, 163.
Boats, recovery of, 140-3.
Bone, composition of, 144-5; preservation and restoration of, 145 ff.
— objects, 2, 144, 146; conservation of, 147-8, 151.
Bookbinder’s paste, 54, 347.
Bookbindings, 9, 10, 12, 19, 24, 25-26; insecticides and, 30; deterioration of, 34-35; ‘peroxide’ test and, 35-36, 37, 355; preservation of, 36-39; ivory, 144.
Book of the Dead, 42.
Books, 9; mould growths on, 10, 27, 55; sea air and, 11; coal gas and, 19; binding of, 25-26; attacks by insects on, 30; causes of deterioration, 38; dressings for, 38; washing of, 38-39; use of parchment for, 48; paper for, 52; sterilization of, 57, 58-60; repair of, 61, 72, 88; removal of pages from, 84; ivory covers for, 146.
Bradley, M. C., 178 n. 2.
Brass, 189, 283 n. 5.
British Honduras, lava monuments from, 297.
— Leather Manufacturers’ Research Association, 19 & n. 2, 35 & n., 37 n.
— Museum, 4, 54, 123, 147; objects referred to: leather bindings, 34-35, 36-37; King’s Library, 46; Sutton Hoo finds, 116; clay tablets, 320; Chinese frescoes, 323.
— Leather Dressing, 38, 357.
— Records Association, 58.
Brommelle, N., 172 n., 176 n. 2.
Bronze, 139, 188-9, 233; decay and, 190; ‘disease’, 234-6, 246, 252, 256, 272, 273; decorated with tin, 265.
— Age enamel, 338.
— objects: treatment of incrustation on, 192, 193, 197, 198, 204, 205, 210, 234 ff.; reconstruction of, 286.
Bronzes, Chinese, 94, 188, 247-9, 250, 251.
Brushing, of metallic objects, 192, 196, 197, 203-4, 221, 222, 224, 227, 253; and the application of wax, 280-1; of stone objects, 298; of marble, 309.
Buddhist documents, 44.
Burgess, S. G., 273 n. 2, 296 n.
Burial, effect of, on: alloys of gold, 207; bone and ivory, 145; clay objects, 320; copper, 233, 234; iron and steel, 272-3, 278; metals, 187-8; silver coins, 224; tin and lead, 258, 259, 265.
Burnishing, 205-6.
Butlin, K. R., 273 n. 1.
Caley, E. R., 197 & n., 261 & n.
Calgon, 247.
Calligraphy, 65.
Canoes, recovery of, 140-3.
Canova, 311.
Canvas, see Paintings, easel.
Carbon, in ink, 63-65, 328; and steel, 189.
— disulphide, 124-5, 346.
Carnelian, 153, 154, 294.
Carpets, 98-99.
Carrara marble, 308.
Carruthers, R. H., 58 n. 1.
Carson, F. T., 53 n. 3.
Cartouches, 253.
Carving, ivory and bone, 144, 152-4.
Casein, 311.
Catechol group of vegetable tannins, 25, 37.
Cellophane, 29.
Cellulose acetate, 61-62, 133.
Cements, for repairing: stone objects, 318-19; clay objects, 323-4; glass and porcelain, 339.
Cennini, Cennino, 158 n. 1.
Ceramics, see pottery.
Chakravorti, S. K., 59 n. 1.
Chalk, 158, 193.
— drawings, cleaning of, 83, 84.
Chatterton’s Compound, 132.
Chemicals, dangerous, 345-6.
Cheshire, A., 36 n. 1.
Chinese: brass, 189; bronzes, 94, 188, 247-9, 250, 251; frescoes, 323; inks, 63; ivories, 131; lacquer, 21; paper-making, 50-51; pottery, 329; silk, 93.
Chipping metallic incrustation, 202-3.
Chloride, test for, 198, 199, 237-8, 241, 243, 298; and rust, 272, 273-4.
Chlorine dioxide, 60, 75, 77.
Christensen, B. B., 137 and n., 138.
Ciba Experimental Dye-house Laboratories 99.
Clay, 205, 273, 319 ff.; for writing upon, 320; methods of firing, 326; see also pottery.
— objects, 319; treatment of, 320 ff.
Cleavage in easel paintings, 158, 159, 164, 166.
Cleopatra’s Needle, 296.
Cockerell, D., 86 & n.
Cody’s Tree, 130.
Coins: embedding of, 267-8; electrum, 207; gold, 207, 209, 212; lead, 264; Roman, 189; silver, 212, 224-5; Sutton Hoo, 26; tin, 265, 266.
Coleoptera, 28, 121.
Consolidation, see reinforcement.
Cooper, B. S., 111 n. 1.
Copper, 188-9, 195, 199, 267; in gold alloys, 207, 208; gilding of, 209; incrusted on silver, 219-22, 223-4, 231; oxidation of, 232-4, 239; conservation of, 234, 249; treatment of, 239 ff., 279; decoration of, 249 ff.; alloys of, 256; in pewter, 258; and soldering, 289.
Coptic stele, 301-2.
Coral, 156.
Coraline solder paste, 288.
Cored Ersin Solder, 288.
Coremans, P., 128 n. 2.
Corrosion, 93, 138, 139, 185; treatment of, 186, 191 ff.; in buried metals, 187-8, 248; process of, 189-91, 271; mechanical removal of, 200 ff.; of silver coins, 224-5; of copper and bronze, 233 ff.; of lead and tin, 258-9, 265-6; of pewter, 268; of iron, 271-4; soldering and, 287.
Cosmoloid microcrystalline waxes, 359 & n. 1.
Cracking of wooden panels, 159, 160, 161, 163, 166.
Craquelure, 141, 142, 158-9, 171.
Crepe, 61.
Cripps, E. S., 279 n. 3.
Croffles, 196.
Cupping of painted panels, 118, 163.
Curtains, cleaning, 98-99.
Damascene work, 281.
Damask, cleaning, 98.
Damp: staining by, 5, 21, 47, 55; degradation of leather by, 26, 31; and insect pests, 58; and textiles, 93-94, 114; effect of, on wood, 116-17, 120; effect of, on bone and ivory, 144-6; and metallic oxidation, 188, 233-4, 245, 274; and stone waxing, 303; and marble, 307-8.
Da Vinci, Leonardo, 66.
Daylight, see sunlight.
DDT, 29, 59, 115, 125.
Dead Sea Scrolls, 40.
De Bruyne, N. A., 132 n. 1.
Decay, electro-chemical, 189-90.
Decaying matter, and textiles, 93, 95.
Decoration, concealing, 14; cleaning of, 104, 299, 327, 328; with ivory, 144, 153; with gold, 209-10, 249-50; with silver and tin, 249, 250 ff.
Delougaiz, P., 320 n.
Delta metal, 313, 318.
Desch, H. E., 118 & n. 2.
Desiccation, damage by, 9, 21; and skin products, 31; by silica gel, 31, 200, 245, 246, 267, 336, 358; and parchment, 47; its effect on wood, 116, 118-19, 132-3, 161; of metals, 199-200, 223, 246.
Detergents, for use with: bone and ivory, 147, 154; leather, 335; marble, 307, 316; silver, 229; textiles, 97.
Devitrification, 329, 335.
Diastase, 105 & n. 2.
Dorchester, silver coins from, 225.
Drawings: sterilization of, 57; bleaching of, 60, 75-81; study of, by photography, 66; cleaning and repair of, 68 ff.; mounting of, 88-91.
Dry-cleaning, of prints &c., 72-73; of
textiles, 98–100; of corroded metals, 204–
5, 252.
Dry rot, 120 & n. 3.
Duering, H., 310 & n.
Dura-glitz, 216, 227, 257.
Dye-stuffs: soap and, 33 n.; and leather, 35;

desiccation and, 47; in inks, 65; removal of,
75, 79; in cleaning, 81, 82; mould
growths and, 83–84; in textiles, 95, 96, 97,
98, 108; fixing of, 96, 97; organic solvents and,
99, 100; and Terylene, 106–7;
light and, 111; and ivory, 146–7.

Edwards of Halifax, 46.
Egg-tempera, 158, 169, 170, 175.
Egypt, ancient: skin products in, 21; tann;
ing in, 25; tomb paintings from, 25, 299; leather scrolls from, 39–40; use of
‘paper reed’ (papyrus) in, 41–42; writing
fluids in, 63; mummy wrappings from,
94; wood from tombs of, 116; ivory
objects from, 144, 148, 149–50; bronzes
from, 193, 244, 246–7, 252–4; gold
objects from, 208; niello decoration in, 221;
and alabaster, 295; vessels from, 327;
glass scarabs, 336–7.
Electrolytes, and corrosion, 189–91; reduc;
tion by, 194–7, 222, 223.
Electro-plating, 289.
Electrum, 207 ff.
El Greco, 170.
Embedding in plastic materials, 267–8.
Embroidery, 14, 93; cleaning and mount;
ing of, 107–10.
Emesa Helmet, restoration of, 226–9, 288.
Enamel, 337–9.
Enkomi Cup, 220.
Engravings: bleaching of, 60, 75–81;
cleaning and repair of, 68 ff.; mounting
of, 88–91.
Environment: influence of, 1 ff.; and
identification of objects, 13; and painted
panels, 160–1; and canvas paintings, 166;
and corroded bronze, 235.
Enzymes, 344.

Erclaine lacquer, 229, 244, 253, 257.
Etchings: bleaching of, 60, 75–81; cleaning
and repair of, 68 ff.; mounting of, 88–91.
Ethnographical material, 10, 20, 27, 100,
115, 116, 121, 323; sterilization of, 123.
Excavated objects, 1, 2–3; bone and ivory,
151; bronze, 235, 251–2; clay, 320, 327;
lead, 259; silver, 216–17; textile, 103.

Faraday, Michael, 19 & n. 1.
Farnsworth, Miss M., 247 n.
Fibres: animal and vegetable, 93, 117; sun;
light and, 110; synthetic, 107.
Fillers: for repairing—wood, 132; ivory,
150; silver, 228; by dowelling, 318, 319.
Fire, precautions against, 50; deciphering of
writing after, 67; effect of, on bone and
ivory, 145, 146; gilding by, 209; effect of,
on marble, 312, 313.
Fisher, R. C., 122 n.
Flags, mounting of, 106–7.
Flaking paint, 164–5.
Flints, 156.
Fluxes, soldering, 287–8.
Fly-marks on papers, 73, 78.
Forest Products Research Laboratory, 120
& n. 3, 122 n., 125 n. 2, 126 & n.
Formalin, 57, 60.
Fossilization, of bone and ivory, 145, 148.
Fossils: ‘pyritized’, treatment of, 295; im;
prints of, 309.
Fowler, J. N., 102.
Footing of papers, 53, 54, 60, 82.
Franks Casket, 147.
French polish, 331, 332.
Frescoes, 303, 323.
Frigilene lacquer, 229.
Frost, effects of, 296–7, 304, 316.
Fryolux solder paste, 288.
Fuels, and air pollution, 12, 19, 37, 112.
Fumigation, 28–29, 30, 49, 115; and mould
growths, 53 ff.; and insect pests, 123–5, 180.
Fungicides, 27–28, 49, 56, 58, 120, 121, 181.
Fungoid growths, 10; and skin products, 27, 49; and paper, 54; and textiles, 113; and
wood, 116, 118, 120, 121; and canvas
paintings, 180–1; and marble, 307–8; and
enamel, 338.
INDEX

Horn, 155.
Horridge, G. A., 130.
Houdin, bust of Voltaire, 308.
Houwink, R., 132 n. 1.
Hughes, A. W. McKenny, 58 & n. 1, 115 n.
Humber, the Iron Age boats from, 141.
Humidity: changes and effects of, 1 ff.,
27, 117, 158, 164; on skin products, 26,
27, 30–31; parchment, 47, 49; paper
storage, 53–54; textiles, 110, 111; sea-
soning of wood, 118, 129; easel paintings,
158 n. 2, 159; painted panels, 163, 164;
canvas paintings, 166; rusting, 277,
stone monuments, 296; measurement of,
353.
Hunter, Dard, 51 & n. 1.
Huntington Library, San Marino, 58.
Hydrogen peroxide test, 35–36, 37, 355.
Hygrometers, 353–4.
Hygroscopic materials, 8, 10, 47, 49, 53,
117, 335, 336.
Hyphae, 10.
Igepal CA Extra, detergent, 97, 105.
liams, T. M., 58 n. 2.
Illumination, in MSS., 44, 45, 47–48.
Illumination, ultra-violet and infra-red,
66–67, 176; examination of marble by,
315–16.
impatto, 169, 171, 174, 177.
Impregnation, 56, 57–58, 123, 125, 126, 127–
31; methods used for: horn and antler,
&c., 155; incrusted ivory, 150; lead and
tin alloys, 270; marbles, 311; pottery,
328; 'pyritized' fossils, 295; silver coins,
224; stone objects, 302–6; unbaked clay,
323; water-logged wood, 135–8.
Incrustation, 13; as evidence, 13–14, 188,
203, 249, 284; of ivory, 148–50, 153; of
metallic objects, 185 ff., 206; of silver,
216 ff.; of copper, 233 ff.; of lead & tin,
258 ff.; of pewter, 268–9; of iron, 273 ff.;
of minerals, 295, 298, 299 ff.; of glass,
336, 337.
— removal of, by: brushing, 203–4, 216,
222, 224, 227, 240, 241, 253; chemicals,
207, 210, 219 ff., 238 ff., 250, 261 ff.;
electro-chemical processes, 191–7, 214,
217, 221 ff., 235, 237, 239, 241, 253, 265,
INDEX


India proof papers, 78; cleaning of, 82–83.

Indicators, 260, 350–1.

Inhibitors, rust, 277–9.

Inks, 41, 42, 47, 63 ff.; puncturing by, 53; preserving of, 53; carbon, 63–65, 328; iron, 64–65; cleaning and, 70, 75, 76, 86; stain removal, 79.

Inlay, 9, 132, 275, 294; marcasite, 295; niello, 221; red glass, 338; silver, 226.

Innes, R. Faraday, 35, 36 n. 2, 37 n., 53.

Inpainting, 163, 167, 175–6.

Inscriptions, treatment of, 224, 253, 266, 269.

Insecticides, 28–30, 114, 115 & n., 121–2, 124, 125, 126, 180.

Insects, attacks by, on: basketry and bark cloth, 133–4; panel paintings, 180; parchment and paper, 58; skin products, 28 ff.; storage materials, 50; textiles, 113–15; wooden objects, 121–6; control of, 28–30, 114.

International Institute for the Conservation of Museum Objects, 111.

Ion-exchange principles, 262–4, 268.

Iron, 188, 190, 195–6, 203, 205, 271; in gold alloys, 208; in juxtaposition to silver, 226; to brass, 283 No. 8; rusting of, 271–3; sulphides of, 295.


Islamic bronze, 242.

Istituto di Patologia del Libro, Rome, 67.

Italian marbles, 315.

Ivory: composition of, 144–5; preservation and restoration of, 145–55.

— objects, 2, 144, 146; conservation of, 147–55; moulding of, 152–3.

Jade, 294.

Japanese: ivories, 146; mulberry tissue, 108; prints, 82, 87; swords, 206, 280, 283; vellum, 82.

Jet, 156.

Jewellery, 156, 205, 207, 265.

Jones, G. A., 67 n. 1.

Kantrowitz, M. S., 51 n. 2.

Keeley, T. R., 4 n.

Kimberley, A. E., 53 & n. 2.

Kramer, W., 136, 138.

Lace washing, 95, 96.

Lacquer objects, 2–3.

Lacquers, used in protecting: clay objects, 323, 329; copper and its alloys, 244, 245, 253, 257; enamel, 338; fossils, 295; glass, 335 & n.; iron and steel, 279; lead, 265; silver, 215, 229, 231; stone objects, 303–5.

Lamination, of paper, 60–63, 70, 89.

Langwell, W. H., 53 & n.

Lanolin, 277.

Lapis lazuli, 153, 154–5, 211, 204.

Lassoband adhesive tapes, 154 & n.

Lavas, 305.

Lead, 188–9, 197, 259; in Chinese bronze, 248; in solder, 286–7; in glass, 333 & n. 1, 337.

Leadens objects: treatment of incrustation on, 198, 258 ff.; oxidation of, 258–9; washing of, 260–1; acid treatment of, 261–2; ion-exchange resins and, 262–4; storage of, 264–5, 270.


Libraries, mould growths in, 10, 27, 38; care of books in, 38.


Lighting: in museums, 111–12; fluorescent and tungsten, 111–12.

Lime, 193, 207; deposits of, on bronze, 247.

Limestones, 298, 300, 301, 305.

Liquid Erzin soldering flux, 288.

Lissapol detergent, 60, 74, 97, 103 n., 105, 133, 207, 221, 229, 297, 307.

INDEX

Lumarith, 104 n.
Lumite, 104 n.

McBurney, C. B. M., 151 & n. 5.
McIntyre, J., 8 n. 1, 54 n. 1.
Malachite, 294.
Malaya, coins from, 266; kris from, 281.
Mallowan, Prof. Max, 153.
Manuscripts: mould growths on, 55; sterilization of, 56–57, 58–60; repair of, 61, 68; mounting of, 89–92; use of gold in, 209.
Maps, mounting of, 63.
Marcasite, 295.
Marco Resin S.B. 26C, 130 & n. 2 267–8.
Marquetry, 9, 133.
Martin, R. S. J., 151 & n. 4, 267 & n. 2.
Matson, F. R., 334 & n.
Maxwell, S., 94 n.
Mayer, R., 181.
Meeuse, Dr. A. D. J., 115 n.
Mercury gilding, 209, 230.
Merovingian objects, 13 n., 283.
Metals: characteristics of, 185 ff.; effect of burial on, 187–8; of antiquity, 188–9, 234–5, 284, 286–9; cathodic protection of, 189–90, 230, 259; incrustation and, 198; drying of, 199; gilt, cleaning of, 210; soldering of, 286–9; patina of, 314; used for dowelling, 317; as a base for enamels, 337.
Meteorites, 271.
<table>
<thead>
<tr>
<th>INDEX</th>
</tr>
</thead>
<tbody>
<tr>
<td>National Library of Wales, Aberystwyth, 36-37.</td>
</tr>
<tr>
<td>— Maritime Museum, 141.</td>
</tr>
<tr>
<td>— Museum of Antiquities of Scotland, Edinburgh, 94 n.</td>
</tr>
<tr>
<td>Neri, A., 333 &amp; n. 2, 334.</td>
</tr>
<tr>
<td>Newspapers, 52, 53.</td>
</tr>
<tr>
<td>Niello, 220-2.</td>
</tr>
<tr>
<td>Nimrud, ivories, 146, 153-5; cuprous enamel, 338.</td>
</tr>
<tr>
<td>Nital, 280.</td>
</tr>
<tr>
<td>Nitro-cellulose treatment, 299-300, 309-10, 328.</td>
</tr>
<tr>
<td>Nonex detergent, 74.</td>
</tr>
<tr>
<td>'Non-tans', 35, 37.</td>
</tr>
<tr>
<td>North Ferriby, 141.</td>
</tr>
<tr>
<td>Nylon bags, use of, 195, 196.</td>
</tr>
<tr>
<td>Old Masters, 168, 170, 172.</td>
</tr>
<tr>
<td>Optical quality, 129, 157, 171, 172, 179.</td>
</tr>
<tr>
<td>Organ, R. M., 198 &amp; n., 199, 244 n., 262 n., 266 n.</td>
</tr>
<tr>
<td>Organic matter, staining by, 310, 314.</td>
</tr>
<tr>
<td>Ochralcum, 283 n. 5.</td>
</tr>
<tr>
<td>Ornamentation: of metallic objects, 186, 192, 216, 220, 241, 242, 244, 266, 275; niello, 220-2; reconstruction of, 285.</td>
</tr>
<tr>
<td>Oseberg ship, 143.</td>
</tr>
<tr>
<td>Oslo, Viking ships at, 142-3.</td>
</tr>
<tr>
<td>Ostraka, 328.</td>
</tr>
<tr>
<td>Paraffin oil, 277.</td>
</tr>
<tr>
<td>— wax, 32, 179, 297.</td>
</tr>
<tr>
<td>Parazyme digestive enzyme mixture, 105.</td>
</tr>
<tr>
<td>Parchment, 9, 10, 21, 24, 40, 41, 55 n. 1; origin of, 44-45; manufacture of, 45-46; alkalinity of, 46-47; humidity and, 47-48, 49; relaxing of, 48-49; storage of, 49-50; sterilization of, 57-58; insect pests and, 58-59; removal of creases from, 85; repair of tears in, 86; resizing and toning of, 87-88; mounting of, 89.</td>
</tr>
<tr>
<td>Paper, 9, 10, 11, 12, 41, 44; advent of, 42, 50-51; raw materials for making, 51-52; tests for permanence of, 52; acidity of, 52-53, 62; storage of, 53-54; sterilization of, against mould, 55-58; insect pests and, 58-59; resizing and bleaching of, 59-60; lamination of, 60-63, 70; examination of before cleaning, 69; removal of creases from, 85-86; repair of tears in, 86-87; resizing and toning of, 87-88; mounting of, 89.</td>
</tr>
<tr>
<td>— pulp, 246; and the treatment of stone objects, 299-300, 313, 328.</td>
</tr>
<tr>
<td>Papyrus, 39, 48; making of, 41-42; from Egyptian tombs, 42-43; method of unwrapping, 43-44; sterilization of, 58; ink used on, 63; bleaching of, 75.</td>
</tr>
<tr>
<td>Parthenon, the, 311, 316.</td>
</tr>
<tr>
<td>Partington, J. R., 212 &amp; n.</td>
</tr>
<tr>
<td>Pastels, sterilization of, 57; cleaning of, 83-84; mounting of, 89.</td>
</tr>
<tr>
<td>Patching, of canvas, 167; of ancient metals, 289; of reconstructed pots, 331.</td>
</tr>
<tr>
<td>Patina, 187, 188, 205, 235; on bronze, 235, 236, 238, 239, 242 ff.; gold, 209; lead, 258-9; marble, 314-15; pewter, 268; silver, 215-16; stones, 297.</td>
</tr>
<tr>
<td>Patterns in sword blades, restoration of, 280-1.</td>
</tr>
<tr>
<td>Pearson's Square, 349.</td>
</tr>
<tr>
<td>Pease, M., 255.</td>
</tr>
<tr>
<td>Peat, objects excavated from, 94 &amp; n., 116, 151.</td>
</tr>
<tr>
<td>Pergamon, Asia Minor, 21, 44.</td>
</tr>
<tr>
<td>'Peroxide' (or 'P.I.R.A.') test, 35-36, 37, 355.</td>
</tr>
<tr>
<td>Index</td>
</tr>
<tr>
<td>-------</td>
</tr>
<tr>
<td>Persian: bronzes, 243; kermes, 65; miniatures, 81; 'parcel gilt' decoration, 209; pottery, 329.</td>
</tr>
<tr>
<td>Petrie, Sir Flinders, 42.</td>
</tr>
<tr>
<td>Pewter, 258; cleaning of, 268-9.</td>
</tr>
<tr>
<td>Photography: and reading faded writing, 66-67; and the repair of textiles, 104; and detecting restoration, 176.</td>
</tr>
<tr>
<td>Picking, 201-2, 203, 241, 250, 264.</td>
</tr>
<tr>
<td>Pigments, 158, 169, 174, 176; tinting plaster for repairing pottery, 331-2.</td>
</tr>
<tr>
<td>Plaster: on bronze, 246; casts, 286.</td>
</tr>
<tr>
<td>Plasticine, 253, 331.</td>
</tr>
<tr>
<td>Plastics, for embedding coins, 267.</td>
</tr>
<tr>
<td>Plesters, Joyce, 267 &amp; n. 3.</td>
</tr>
<tr>
<td>Plenderleith, H. J., 37 n., 54 n. 3, 59 n. 2, 150 n., 151 n. 1, 266 n.</td>
</tr>
<tr>
<td>Plus-Gas Fluid A rust softener, 277 &amp; n. 1.</td>
</tr>
<tr>
<td>Polishing of metals, 205-6.</td>
</tr>
<tr>
<td>— wax, 179, 359.</td>
</tr>
<tr>
<td>Potassium lactate treatment, 35, 39, 356.</td>
</tr>
<tr>
<td>Pottery, 326 ff.; cleaning of, 327-9; repair of, 329-32.</td>
</tr>
<tr>
<td>Prince Rupert drops, 337.</td>
</tr>
<tr>
<td>Prints, sterilization of, 57; bleaching of, 60, 64-65, 75-81; cleaning and repair of, 68 ff.; mounting of, 88-91; see also Japanese.</td>
</tr>
<tr>
<td>Purves, P. E., 151 &amp; n. 4, 267 &amp; n. 2.</td>
</tr>
<tr>
<td>Pyrogaloll group of vegetable tannins, 25, 37.</td>
</tr>
<tr>
<td>Pyx Cloth, 107.</td>
</tr>
<tr>
<td>Quartz, 207.</td>
</tr>
<tr>
<td>Radiation, ultra-violet, 176.</td>
</tr>
<tr>
<td>Ragging for local electroplating, 228, 289.</td>
</tr>
<tr>
<td>Rags, and paper-making, 50-51.</td>
</tr>
<tr>
<td>Rawlins, F. I. G. 4 n.</td>
</tr>
<tr>
<td>Reagents, for removing stains, 102-3.</td>
</tr>
<tr>
<td>Refrigeration, 40.</td>
</tr>
<tr>
<td>Reinforcement, methods used with: clay, 323; copper alloys, 245, 257; enamel, 337-8; marbles, 311-12; oxidized tin, 266; pottery, 328, 329; rusted iron, 281; silver, 227-8, 231; stone objects, 299-300, 301-6.</td>
</tr>
<tr>
<td>Relining easel paintings, 166, 168-9.</td>
</tr>
<tr>
<td>Resin AW 2 and MS 2, 179 n.</td>
</tr>
<tr>
<td>— varnishes, 176-9.</td>
</tr>
<tr>
<td>Resins, Epoxy, 131, 168.</td>
</tr>
<tr>
<td>— used for: canvas repairs, 167, 168, 169; consolidating wood, 129-30; embedding coins, 267-8; flaking paint, 164; fluxes, 287; ion-exchange, 262-4; ivory repair, 150; pottery repair, 330; wood repair, 131, 136.</td>
</tr>
<tr>
<td>Restrainers in controlling solvent activity, 174.</td>
</tr>
<tr>
<td>Rhind Mathematical Papyrus, 39.</td>
</tr>
<tr>
<td>Rijksmuseum, Amsterdam, 266.</td>
</tr>
<tr>
<td>Rochelle salt process, 210, 239, 224, 225, 239-42, 250, 251, 252.</td>
</tr>
<tr>
<td>Rock crystal, 294.</td>
</tr>
<tr>
<td>Rodents, attacks by, on parchment, 50.</td>
</tr>
<tr>
<td>Rolls, leather, unrolling of, 39-40.</td>
</tr>
<tr>
<td>—, papyrus and parchment, sterilization of, 58.</td>
</tr>
<tr>
<td>Roman scabbard of brass, 283 n. 5.</td>
</tr>
<tr>
<td>Romano-British steelyard, 282.</td>
</tr>
<tr>
<td>Rosen, D., 128 &amp; n. 1.</td>
</tr>
<tr>
<td>Rosenberg, G. A., 135 &amp; n.</td>
</tr>
<tr>
<td>Royal Academy of Arts, 308.</td>
</tr>
<tr>
<td>— Institution, 19 &amp; n. 1.</td>
</tr>
<tr>
<td>— School of Needlework, 107 n.</td>
</tr>
<tr>
<td>— Society of Arts, 19.</td>
</tr>
<tr>
<td>Rubber-paste cement, 167.</td>
</tr>
<tr>
<td>Ruhemann, H., 169 n.</td>
</tr>
<tr>
<td>Rust: removal of, 200, 202, 204, 205, 206, 227, 274-8, 291; production of, 271-2; examination of, 274, 284; inhibitors, 279-9; preservation of, 281, 291; as evidence, 284.</td>
</tr>
<tr>
<td>Salty ground, contamination in, 3, 145, 148, 190, 216, 233, 259, 320, 327, 328.</td>
</tr>
<tr>
<td>Sand, 116, 207.</td>
</tr>
<tr>
<td>— blasting, 321.</td>
</tr>
<tr>
<td>Sandstones, 298, 300, 305, 306.</td>
</tr>
</tbody>
</table>
Santobrite, 28, 58, 84, 181, 338.
Santucci, L., 67 n. 4.
Saxon bronzes, 251, 254, 338.
Schaefler, C. F. A., 220 n.
Schafer, R. J., 296 n.
Scraping, 202–3.
Scribner, B. W., 52 n. 1, 53 n. 2, 61 & n. 1.
Sea air, effects of, 11.
Seals, mounting of, 91–92.
Seasoning of wood, 117–18.
Selenite, 301.
Sellotape, 30, 167, 227.
Sequins, 208.
Seyrig, H., 226 & n.
Shale, Kimmersidge, 324–5.
Sheepskin, see parchment.
Sheffield plate, 213.
Shell, 156.
Shellac, 142, 264, 304, 325, 330, 331.
Shot-blasting, 204–5, 321–3.
Sika deer, 155.
Silica gel: desiccation by, 31, 200, 245, 246, 267, 336, 358.
Silicaseal No. 1A, 305 n.
Silicates, in glass, 133–4.
Siliceous materials, 198, 205, 207 246, 294 ff.
Silicones, 295.
Silicon ester, spraying, 305.
Silk, used in lamination, 60–61; and textile mounting, 106–7; in old embroidery, 109; see also textiles.
   — Dip, 214, 231.
Silver-plating, 289.
Simmons, R. H., 51 n. 2.
Sizes, used for: canvas, 165; inks, 63; paper, 51, 52, 54, 69, 83; papyrus, 42; resizing after repair, 87.
Skin and skin products, 19 ff.; used in ancient Egypt, 21; processing of, 21 ff.; fur skins, 23, 28, 29, 114; tawing of, 24, 25; oil dressing of, 24; tanning of, 24–26, 36–37; effect of moisture and moulds on, 26–28, 31; insect attacks on, 28–30; preservation of, 30–32, 34–39; repairing of, 32–33; unrolling of, 39–40; see also parchment.
Slate, 161.
Snaking, 310.
Soap, used with: bone and ivory, 147; leather, 33 n., 39; marble washing, 306; print cleaning, 73–74; textile cleaning, 97.
Sodium chloride, 59, 60, 77.
Solderine, 288.
Soldering, 286–9, 301.
Solutions, preparation of, 343, 349.
Solvent Naphtha, 105.
Solvents, used for cleaning: copper corrosion on silver, 219–22, 231; corroded metals, 186, 216, 257; cuprous enamel, 338–9; lead and tin alloys, 270; pottery, 327, 328, 330; prints, 78–81; rust, 277–9, 291; shellac, 325; stained marble, 308–10; stone objects, 296–7, 300–1, 303; textiles, 98 ff., 105, 108; varnishes, 72, 73, 174, 178.
South Shields sword, 283.
Spencer, E. W., 51 n. 2.
Spraying, 123, 125, 133–4; of varnish, 177–8; of silicon ester on stone objects, 305–6.
Stanwick scabbard, restoration of, 138–40; sword, 282, No. 4.
Statuary: treatment of, 298, 303; weathering and, 316–17; packing of, 317.
Steel, 189, 195, 206, 271; stainless, 189 n., 196.
   — objects, 206, 271, 274; treatment of,
INDEX

277, 282-3, 290-1; protection of, 279; preservation of pattern in, 280-1.
Stein, Sir Aurel, 50, 323.
Sterilization: by fumigation, 55-57; against insects in wood, 123.
Stoddard Solvent, 105.
Stone objects, 295 ff.; cleaning and preservation of, 296 ff.; the paper pulp method and, 299-300; removal of insoluble salts from, 300-1; consolidation of, 301-6; effects of weathering on, 316-17; repair of, 317-19.
Stones, building-, 12; deterioration of, 294.
Stoppings, 163, 167, 175, 176, 226, 227.
Storage of: copper alloys, 257; lead objects, 264-5, 270; marble, 307-8; paper, 53-55; parchment, 49-50; silver, 214-15, 231.
Stout, G. L., 181.
Strasbourg Museum, 13 n.
Strip-lining, 166, 168.
Stromberg, Elizabeth, 110 & n., 111.
Sulphuric acid: and leather, 36, 53, 112; and paper, 53, 112; in ink, 64; and canvas tendering, 166; and copper incrustation, 239 ff.; and bronze, 243-4, 252; preparation of, 343; care of, 345.
Sulphurous gases, 11-12, 19; and leather, 34 ff., 53, 356; and paper, 52-53; and textiles, 112; and easel paintings, 166, 170-1; and metals, 188.
Sunlight, 9; and paper deterioration, 53; a bleaching agent, 75, 79; and textiles, 94, 110-12; and waxed objects, 129; and bone and ivory, 145; and oil paintings, 173.
Surface enrichment, 207.
Sutton Hoo finds, 26, 116, 209, 218, 251, 282, 283; reconstruction of, 284-6.
Swiss National Museum, 136 n. 2.
Tablets, cuneiform: treatment of, 320-1; rendering them legible, 321-3; repair of, 323.
Tandberg, Prof. J., 215 n.
Tannins, 24-25, 64.

Tarnish: on silver, 213-15, 221, 222, 223; on copper, 232.
Taylor, W. D., 67 & n. 3.
Tears, repair of: in paper and parchment, 86-88; in canvas, 167-8.
Teepol detergent, 307, 327.
Temperature changes: effects of, 1 ff., 55, 129, 214; and textile deterioration, 110; and painted panels, 160-1; and canvas paintings, 166; and stone objects, 302; reading of, 353.
Tempering steel, 201 & n.
Tendering of canvas, 166.
Termites, 125-6.
Terracotta, 327.
Terylene, 104; and the mounting of frail textiles, 106-7, 110, 133.
Texibond V4N polyvinyl acetate emulsion, 168 & n. 1, 169.
Textiles, 12, 14, 29, 93 ff.; for storage, 49-50; examination of, 94; cleaning of, 95 ff., 98-100; washing of, 95-98; removal of stains from, 100-3, 105, 344; preservation of, 103-5; mould growths and, 93, 95, 106, 113; repair and mounting of, 106-10; light and, 110-12; atmospheric pollution and, 112-13; insect attacks on, 113-15; isolation of, 114; insecticides and, 115.
Thacker, D. M. D., 279 n. 3.
Thermometric equivalents, 352.
Thymol, 56, 57-58, 73, 109, 113.
Tiles, 329.
Timber, see wood and wooden objects.
Times, The, 12 n.; binding of, 34-35.
Tin, 187, 188-9, 258, 265; in bronze, 233 & n., 236, 247-8; as decoration, 249, 250, 265; soldering of, 286-7.
— objects, 262; cleaning of, 265; oxidation of, 266.
— pest, 265-6.
Todd, W., 151 n. & 2.
Tools, used for: removing corrosion, 200-4; restoring gold objects, 211; repairing crushed silver, 218; restoring bronze objects, 254; cleaning marble, 314.
Tortoise-shell, 155.
INDEX

Tower of London Armouries, 107 n.
Transoptic resin for embedding objects, 267–8.
Tropics, the, 58, 70, 118, 126 n., 347.
Tumbler tank, 225.
Tune ship, 143.
Turquoise, 204.
Tut-ankhamun’s tomb, 21, 208, 212.

Upholstery, 19, 25; leather, preservation of, 33, 37; cleaning of, 98; sterilization of, 123.
Ur of the Chaldees, 188–9, 211, 212, 216, 217, 236.

Vacuum treatment: against insects, 123; for consolidating wood, 130–1, 136, 137; for incrusted ivory, 150, 151; for incrusted metals, 200, 223; for impregnating stone objects, 304–5, 329.
Vapour Phase Inhibitors, 279.

Varnish, dammar and mastic, 172, 173, 175–6.
Vellum, 44–45, 55 n. 1; transparent, 46; cleaning of, 48–49; 83; sterilization of, 57; Japanese, cleaning of, 82; removal of creases from, 85; repair of tears in, 87; mounting of, 88–89.

Veneer, 132–3.
Ventilation, 5, 8, 9, 27, 49, 333; and moulds, 55, 93, 113, 121.
Versenes, for inhibiting rust, 278–9.
Versenol, 312.
Viking ships, 142–3.
Vogt, E., 136 n. 2.

Walbrook Mithraeum, 312.
Walker, J. F., 55 & n. 2.
Wallace Collection, 337.
Wall-paintings, 323.
Walls, dangers of waxing, 303.
Walls, H. J., 67 & n. 3.
Warping: of wood, 118–19, 137, 140; of bone and ivory, 145, 146; in panel paintings, 160, 161; in canvas paintings, 165; of marble, 310–11.
Washing, methods used with: clay tablets, 320–1; excavated pottery, 327–8; glass, 335, 336, 338–9; marble, 306–7, 309; metallic objects, 192, 196, 197–9, 222, 231, 260, 270, 291; stones, 295, 298, 301.
Waterlogging: and wood, 134 ff.; and bone and ivory, 145, 151.
Waterproofing, 120, 143, 296–7, 303.
Waters, C. E., 63 n.
Wax, used for: removing bloom, 173; canvas repairs, 169; impregnation, 127–30, 133, 302–3, 311; iron and steel, 277, 279, 280, 359; ivory repairs, 150; lead preservation, 259 ff.; leather dressing for tropical use, 359; pewter, 269; polishing, 206; re-varnishing, 179–80; wood sterilization, 123, 125; preparation of microcrystalline polishing wax, 359.
Weathering, 13; of stone, 204, 296; effect of, on stone objects, 316–17.
Weeping glass, 336.
Weiss, H. B., 58 n. 1.
Welwyn fire dogs, 282.
Werner, A. E. A., 37 n., 176 n. 1, 179.
Westrosol dry-cleaning solvent, 99.
Windsor Castle, Royal Collection at, 66.

Wood, and paper-making, 51–52, 116; conditions under which it decays, 116–17; directional properties of, 117; seasoning of, 117–18; warping of, 118–19; fungus attacks on, 120–1; insect attack and, 121–3, 180; sterilization of, 123–5; proofing of, 125–6; strengthening of, 126–31, 161–2; repair and maintenance of, 131–3; waterlogging and, 134–43; emitting tannic acid, 264–5.

Wood, R. W., 208 & n.

Wooden objects, 2, 8–9, 120; sterilization of, 124–5; proofing of, 126; consolidation of, 127–9, 302; cleaning of, 133; waterlogged, preservation of, 134–43.
Wood-worm, 121 ff., 130.
Wool, ancient, cleaning of, 104–5; see also textiles.
INDEX

Woolley, Sir Leonard, 211.

Workshop Notes, on the handling of ancient textiles, 104-5.

Wright, E. V. and C. W., 141 n. 2.

Writing: ground materials for, 41 ff.; preserving of, 53; faded, reading of, 66-67; clay tablets for, 319, 320; rendering it legible, 320-3; preservation of ostraka, 328.

X-ray examination: of ivory, 153; of paintings, 176; for hidden ornament, 186; to penetrate incrustation, 220; of rust, 274, 275.

York Minster, 121.

Zinc, 189, 190; and electro-chemical reduction, 191 ff.
Central Archaeological Library, NEW DELHI 18006

Call No. 913.09/Ple
Author H. J. Ploenderleith
Title The Conservation of Antiquities and Works of Art

“A book that is shut is but a block”

CENTRAL ARCHAEOLOGICAL LIBRARY
GOVT. OF INDIA
Department of Archaeology
NEW DELHI.

Please help us to keep the book clean and moving.