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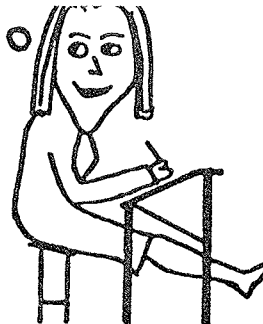
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NOTES FOR
DIPLOMA STUDENTS



Information sheet

Alcohol: Ethyl alcohol C_2H_5OH (ethanol or absolute alcohol) with not more than 1% water and approx. 75° over proof (OP)*. Diluted to 95-90% by volume = "rectified spirit" (approx. 57-66° OP)

Isopropyl alcohol $CH_3.CH(OH)CH_3$ with not more than 3-4% water

Methylated spirit: ethyl + methyl (methanol) alcohols + acetone (some brands) + methyl acetate + water to give 80-90% strengths

- a) mineralised methylated spirit: 90 vols ethanol + 10 vols methanol + mineral oil(s) (paraffin or pyridine) + colouring. Useless for preserving biological specimens
- b) Industrial methylated spirit (IMS): 95 vols ethanol + 5 vols methanol to give 99-92% strength (74-60 OP); some brands contain pyridine. Diluted to 70-80% standard biological preservative.

*Proof alcohol = 57% strength at 60°F; by definition 5 vols 90-95% alcohol SPG .838 diluted with 3 vols water.

<u>dilution by vol.</u>	<u>degrees over proof (approximated)</u>
69%	22
74%	31
80%	40
84%	48
90%	57
95%	66
100	75

Formaldehyde: Commercial grades 40%. A 4% solution adequate for most museum specimens i.e. 10ml formaldehyde (40%) + 90ml water; neutralise with borax or hexamine (saturate stock solution).

Propylene Phenoxetol/Glycol: specimens must be adequately fixed before storing in solution no II

Solution I fixative:	propylene phenoxetol	1ml
	propylene glycol	5ml
	40% formaldehyde	10ml
	distilled water	84ml
Solution II preservative:	propylene phenoxetol	1ml
	propylene glycol	10ml
	distilled water	89ml

Colour preservation: modification of Kaiserling's tripartite method

Solution I	Formaldehyde (40%)	400ml
	Potassium acetate	50g
	Potassium nitrate	30g
	distilled water	1000ml
Solution II	80-90% alcohol until colour returns	
Solution III	Glycerine	3000ml
	Potassium acetate	2000g
	distilled water	9000ml

Some dates in preservation techniques

Dry preservation	Wet preservation	Osteological <u>in situ</u> preparations
<p>484 BC Herodotus Egyptian embalming methods (see Pettigrew 1834)</p> <p>*1490 Leonardo da Vinci casting brain ventri- cles and wax casting of heart (see Dobson 1956)</p> <p>1642 Ole Worm's catalogue of Museum, all dried specimens (see Anon 1642)</p> <p>1656 Tradescants catalogue of museum (probably all dried specimens) (see Allan 1964)</p>	<p>1660 Ashmole showed Charles II specimens "in a solution of Dr Warner's invention" (see Gunther 1927)</p> <p>1662 Boyle recorded use of spirits of wine for preserving tissues (see Birch 1746)</p> <p>*1670 Swammerdam men- tioned spirits of wine in catalogue to museum</p>	

Dry preservation	Wet preservation	Osteological <u>in situ</u> preparations
<p>1890[†] Altmann described method for freeze drying tissues</p>	<p>1681 Grew mentioned liquid preparations in Royal Society catalogue</p> <p>1710 Ruysch mentioned liquid preps. in his catalogue</p> <p>*1768 Hunter collection - large number of liquid preps. mentioned (see Laskey 1813)</p> <p>1786 Seba - many specimens preserved in "kilduivel" (= killing devil or spirits of wine) (see Engel 1937)</p> <p>1859[†] Butlerov discovered formaldehyde</p> <p>1867[†] von Hofmann demonstrated production of formaldehyde</p> <p>1888[†] Loew discussed antiseptic properties of formaldehyde</p> <p>1893[†] Blum tissue preserving qualities of formaldehyde; colour restoration in alcohol</p> <p>colour preservation 1896[†] Melnikow Raswedenkow introduced salts into final preservative</p> <p>colour preservation 1896[†] Jores added salts to first solution</p>	

1920 Hochstetter
displayed wax impreg-
nated specimens

1926 Noble & Jackle
described wax impreg-
nation method based
on Hochstetter's
technique

1927 Hochstetter
published his method

Freeze drying
1932[†] Gersh modified
Altmann's methods

Freeze drying
1948[†] Mercie described
method for fungi

Freeze drying
1954[†] Davies described
method for whole
animals and plants

Freeze drying
1960[†] Meryman
described methods for
whole vertebrates (USA)

1964 Harris developed
methods in UK

1922 Kaiserling
reviewed development
of formaldehyde
preservation/colour
stability

colour preservation
1936 Pulvertaft
described method
using sodium hydro-
sulphate

1956 Owen & Steedman
described experiments
with propylene
phenoxetol as
preservative

colour preservation
1962 Yoshida
described anti-
oxidant sodium
ascorbic method

colour preservation
1965 Waller
experimented with
Butylated Hydro-
xytoluene (BHT)

1894 Schultz published
method for rendering
whole animal trans-
parent using sodium
hydroxide

1904-1912 Lundvall[†]
used alizerine to
stain bones

1911 Spalteholz[†]
improved transparency
methods

1926 Dawson[†] improved
Spalteholz's methods

1953 Williams[†] used
toluidine blue + aliz-
erine to distinguish
cartilage from bone

[†]see Edwards & Edwards 1959 for references

*approximate date

[†]see Harris 1964 for references

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We hope that 'Notes for Diploma Students' will appear as a regular feature in future B.C.G. Newsletters. The idea came about when one of us realised that various Newsletter articles, particularly the 'Biodeterioration' special by Reg Harris in 1978, made extremely useful revision notes for the Museums Association Diploma Curatorial Examination. In addition, many of us regret the demise of the 'old-style' curatorial courses, which enabled students to get to grips with biological and geological practices in some depth over a solid eleven day period. This proposed series of notes will at least go towards making up for this loss.

We wish to thank Dr. Ray Ingle (British Museum, Natural History) for permission to reproduce his information sheet, in this issue; and also to appeal to everyone for further articles in the series. Otherwise some gentle arm-twisting may be necessary!! Seriously though, this is an area where B.C.G. can play an active role in training museum biologists, and even the rest of us should find such articles useful 'refresher' material.

- Editors