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APPENDICES

APPENDICES

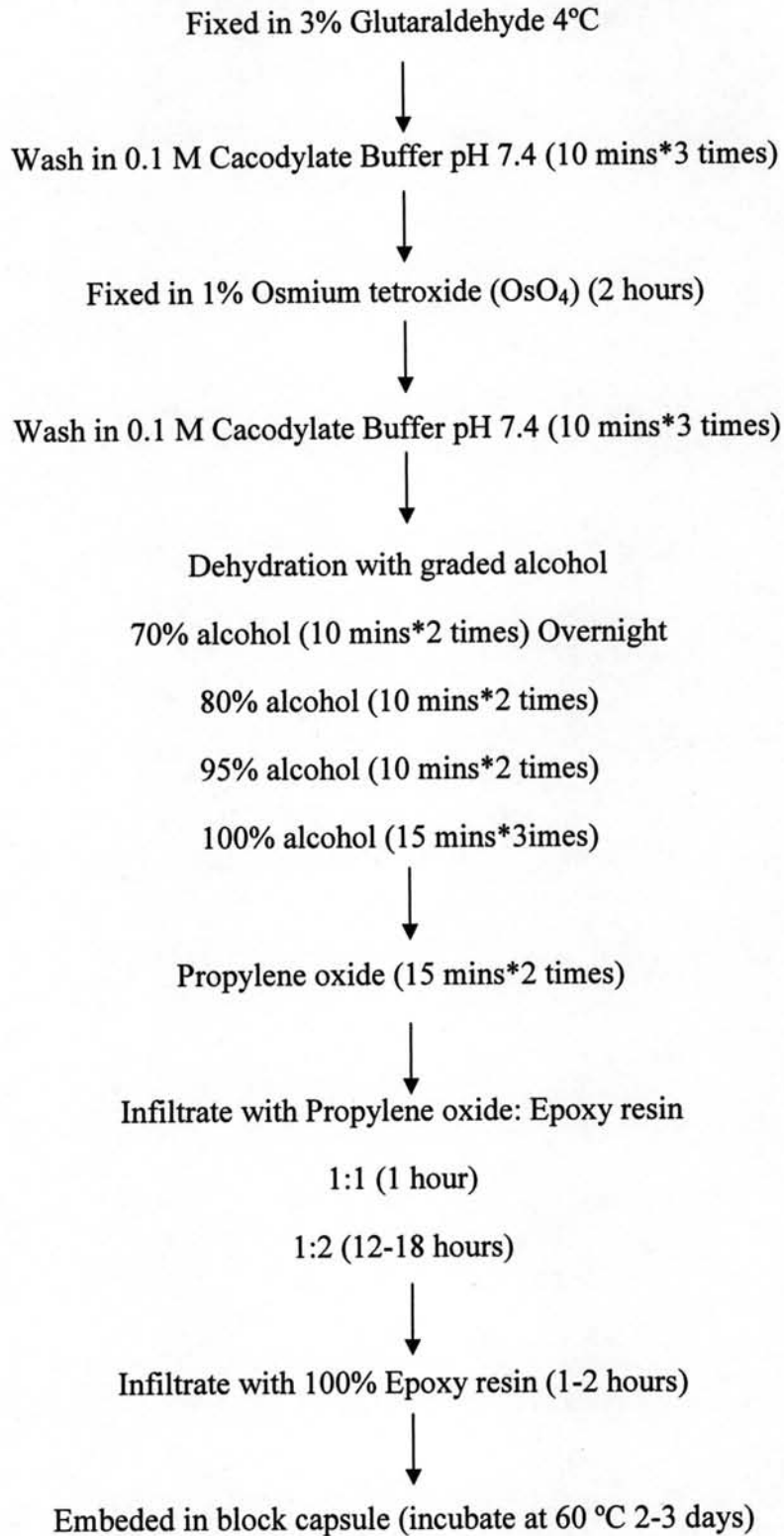
Appendix 1

Preparation of stacking gel and separating (running) gel

*Running gel buffer : 1 M Trizma pre-set crystal , pH8.8

Composition	Running gel (10%)		Stacking gel	
	1 plate	2 plate	1plate	2 plate
30%Acrylamide mix	3.3ml	9.9ml	0.65ml	1.95ml
1M Running gel buffer	3.75ml	11.25ml		
1M Stacking gel buffer			0.625ml	1.875ml
Distilled water	2.75ml	8.25ml	3.28ml	9.84ml
10%SDS	100 μ l	300 μ l	50 μ l	150 μ l
10%APS	100 μ l	300 μ l	37.5 μ l	112.5 μ l
TEMED	10 μ l	30 μ l	5 μ l	15 μ l
Total volume	10ml	30ml	4.6ml	13.9ml

Stacking gel buffer : 1 M Tris-base , pH 6.8

Appendix 2 Tissue processing for ultramicrotome section

Appendix 3 Solutions and dyes used in the tissue processing**0.1 M Cacodylate buffer**

Sodium cacodylate 21.4 g., Trihydrate MW. = 214.02
Cacodylate acid
dH₂O 986 ml.
0.2 M HCl 14 ml.

3% Glutaraldehyde

30% glutaraldehyde 20 ml.
0.1 M cacodylate buffer 230 ml.

1% Osmium

Osmium tetroxide 1 g.
0.1 M cacodylate buffer 100 ml.

1% para-phenylenediamine

Sodium borate 5 g.
Distilled water 500 ml.: 1% sodium borate
Methylene blue 5 g.
1% sodium borate 500 ml.

Lead citrate

1 g Lead Citrate
100 ml freshly distilled water
0.4g NaOH per 1 ml

Toluidine blue

Sodium borate 5 g.
Distilled water 500 ml
Toluidine blue 5g
Azure II 5 g

Appendix 4 Substrate solution for immunological detection in Western blot analysis

To 20 ml of distilled water, add 8 drops of buffer stock (pH 7.5) solution and mix well



Add 16 drops of DAB Stock Solution and mix well



Add 8 drops of the Hydrogen Peroxide Solution and mix well



If a gray-black stain is desired, add 8 drops of the Nickel Solution and mix well

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