

# medilines

## Best's Carmine Colour

DESIGN TO DEMONSTRATE Glycogen

### REAGENTS & PREPARATIONS

<b>Best Solution A</b>	<b>Best Solution A1</b>	
	<b>Best Solution A2</b>	Mix equal volumes of A1 and A2 prior to use (violet black in colour) and can be use within a weak or discard if it become brown.
<b>Best Solution B</b>	<b>Best Solution B1</b>	1.5 volume
	<b>Best Solution B2</b>	1.0 volume
		This should be made up immediately before use and can be use within 2-3 days
<b>Best Solution C</b>		Ready to use

### TISSUE SAMPLE

4-5 $\mu$  paraffin sections of 10% neutral buffered formalin fixed tissue. Carnoy's or alcoholic-formalin fixed tissues are more suitable

### STAINING PROCEDURES

1. Remove Paraffin and bring sections to distilled water.
2. Place into the **Best Solution A** for 1 minute to stain Nuclei.
3. Rinse in distilled water.
4. Place into the **Best Solution B** for 15 minutes. (In some Ref 30min)
5. Differentiate in **Best Solution C** for a few seconds.
6. Rinse quickly in 70% alcohol.
7. Dehydrate in graded alcohols.
8. Clear in xylene, three or four changes.
9. Mount with synthetic resin.

### EXPECTED RESULTS

Glycogen	Pink to red granules
Nuclei	dark blue

CONTROL Liver.

### References: (Best, 1906)

p 219 Cook H C, Manual of Histological Demonstration Techniques p 190 Bancroft JD & Stevens A, Theory and Practice of Histological Techniques 2nd ed.

Mallory, F.B.: Pathological Technique, New York, Hafner Publishing Co., 1961, pp. 126-128.