

# medilines

## Perls' Iron Colour

### Perls' Prussian Blue

#### DESIGN TO DEMONSTRATE

Deposits of Hemosiderin

#### REAGENTS & PREPARATIONS

##### **Perl Solution A**

##### **Perl Solution B**

##### **Perl Solution 1**

Mix

Perl Solution A            1 volume

Perl Solution B            1 volume

Solution should be made immediately before use.

**Perl Solution 2**            Ready to use

**Perl Solution 3**            Ready to use

#### TISSUE SAMPLE

4-5 $\mu$  paraffin sections of neutral buffered formalin fixed tissue are suitable. Avoid iron containing materials and jars while fixing as these may contaminate the tissue. Acid containing fixatives may remove some of the iron deposits, but apart from that most are satisfactory.

#### STAINING PROCEDURES

1. Remove Paraffin and bring sections to distilled water.
2. Place into the **Perl Solution 1** for 30 minutes at 60°C or 1 hour at room temperature.
3. Rinse in distilled water.
4. Counterstain with **Perl Solution 2** for 2-3 minute.
5. Wash in **Perl Solution 3** or Rinse well with distilled water.
6. Dehydrate with absolute alcohol
7. Clear in xylene. Mount in synthetic resin

#### EXPECTED RESULTS

Ferric iron	– blue or Green
Nuclei	– red
Background	– pink

#### Notes

1. Avoid washing with tap water before placing into the **Perl Solution 1**, as rust in the water or tap fixtures could cause false positive staining.

#### Reference

*Culling, C.F.A., Allison, R.T. and Barr, W.T. Cellular Pathology Technique, Ed.4. Butterworth, London, UK.*

*Susan Budavari, Editor, (1996) The Merck Index, Ed. 12 Merck & Co., Inc., Whitehouse Station, NJ, USA*

medilines  
medilines@medilines.com