

Instructions For Use

CEA-IFU

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Cresyl Echt Violet Solution (0.1%)

Description and Principle

Cresyl Echt Violet Solution (0.1%) is designed for staining Nissil substance in neurons on formalin fixed, paraffin-embedded tissue. Nissl granules are purple, nuclei of neuroglia and endothelial cells are slightly bluer than Nissl granules.

Expected Results

Nissil Substance: Violet

Neuroglia Nuclei: Violet to Dark Blue

Kit Contents	<u>Storage</u>
Additional Kit Reagents Sold Separately	
Cresyl Echt Violet Solution	2-8° C
Luxol Fast Blue Solution	18-25°C
3. Lithium Carbonate Solution (0.05%)	18-25°C
4. Alcohol, Reagent (70%)	18-25°C

Suggested Controls (not provided)

Cerebral Cortex

Uses/Limitations

Not to be taken internally.
For In-Vitro Diagnostic use only.
Histological applications.
Do not use if reagent become cloudy.
Do not use past expiration date.
Use caution when handling reagent.
Non-Sterile.

Storage

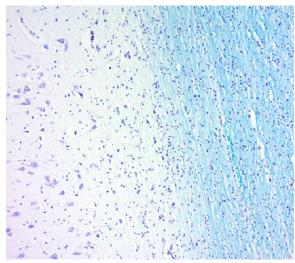
Store at 2-8°C.

Safety and Precautions

Please see current Safety Data Sheets (SDS) for this product and components GHS classification, pictograms, and full hazard/precautionary statements.

Procedure

- 1. Deparaffinize sections if necessary and hydrate to distilled water.
- 2. Pour Luxol Fast Blue Solution into a staining jar and Incubate slide for 24 hours at room temperature or 2 hours at 60° C. Solution is alcoholic and will readily evaporate at smaller volumes.
- 3. Rinse thoroughly in distilled water.
- 4. Differentiate section by dipping in Lithium Carbonate Solution (0.05%) several times (up to 20 seconds).
- 5. If needed, continue differentiation by repeatedly dipping in Alcohol, Reagent (70%) until gray-matter is colorless and white-matter remains blue.
- 6. Rinse slide in 2 changes of distilled water.
- 7. Incubate slide in Cresyl Echt Violet (0.1%) for 2-5 minutes.



White-matter and gray-matter of Human Brain stained with Luxol Fast Blue Stain Kit

- 8. Rinse quickly in 1 change of distilled water.
- 9. Dehydrate quickly in 3 changes of absolute alcohol.
- 10. Clear as desired and mount in synthetic resin.

References

- 1. Sheenan, D.C., Hrapchak, B.B. Theory and Practice of Histotechnology, 2nd Edition. Battelle Press, Columbus, OH. Page 262-264. 1980
- Kluver, H., Barrera, E.A. A Method for the combined staining of cells and fibers in the nervous system. Journal of Neuropathology and Experimental Neurology, 1953, 12: pages 400-403.
- 3. Margaret M. Powers & George Clark (1955) An Evaluation of Cresyl Echt Violet Acetate as a Nissl Stain, Stain Technology, 30:2, 83-88.





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