## CHURUKIAN SCHENK METHOD FOR ARGYROPHIL GRANULES

PURPOSE:	For In Vitro Diagnostic Use: Intended for the qualitative demonstration of argyrophil granules.
PRINCIPLE:	Argyrophil cells are capable of absorbing silver but cannot reduce it on their own. Hydroquinone acts as the reducing agent. This procedure uses a low pH to better control staining.
CONTROL:	Pancreas
	Control Slides can be purchased from Histology Control Systems. See inside back cover, Item# cs028.
SPECIMEN PREPARATION:	Formalin fixed, paraffin embedded sections cut at 4-6 micrometers
SOLUTIONS:	<ol> <li>Acidified Water pH 4.2 Item# s2023</li> <li>Silver Nitrate 0.5% for Churukian Schenk Item# s2021</li> <li>Nuclear Fast Red Kernechtrot 0.1% Item# s248</li> <li>Components for Developer Solution (solution is stable for 2 weeks):         <ul> <li>Sodium Sulfite Powder 5 g</li> <li>Hydroquinone Powder 1 g</li> <li>Distilled Water 100 mL</li> </ul> </li> <li>Solutions can be purchased separately from Poly Scientific.</li> </ol>
NOTES:	
REFERENCE:	Churukian, Schenk. "A Modification of Pascual's Argyrophil Method." <u>J. Histotechnology</u> . 1979; 2: 102.

## STAINING PROCEDURE:

- 1. Deparaffinize and hydrate to Acidified Water pH 4.2 for 1 minute.
- Place in Silver Nitrate 0.5% for Churukian Schenk and place in 58°C water bath for 2 hours.
- 3. Rinse in 3 changes of distilled water.
- 4. Transfer to Developer Solution, preheated to 58°C in a water bath, for 5 minutes.
- 5. Rinse in 3 changes of distilled water.
- 6. Place in the same Silver Nitrate 0.5% for Churukian Schenk in the 58°C water bath for 10 minutes.
- 7. Rinse in 3 changes of distilled water.
- 8. Place in the same Developer Solution in the 58°C water bath for 5 minutes.
- 9. Rinse in 3 changes of distilled water.
- 10. Rinse in distilled water.
- 11. Dehydrate in 95% Alcohol, Absolute Alcohol and clear in Xylene.
- 12. Counterstain with Nuclear Fast Red Kernechtrot 0.1% for 3 minutes.
- 13. Mount with Poly Mount (Item# s2153) or any other acceptable mounting medium.

## **RESULTS:**

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