

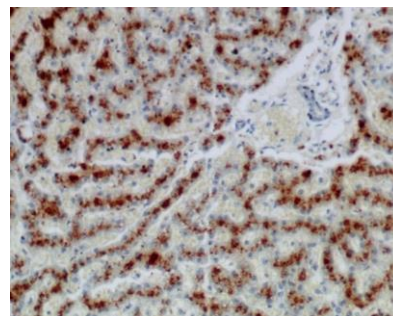
## Copper Stain Kit (For Microwave)

**Description:** The Copper Stain Kit (For Microwave) is intended for the demonstration of copper deposits in tissue sections.

**Uses/Limitations:** Not to be taken internally. For In-Vitro Diagnostic use only. Histological applications. Do not use past expiration date. Use caution when handling reagents. Non-Sterile.

**Control Tissue:** Fetal Liver or a known positive.

**Results:** Copper Deposits: Light Brown to Red  
 Nuclei: Blue



**Availability/Contents:**

<u>Item #</u>	<u>Kit Contents</u>	<u>Volume</u>	<u>Storage</u>
SSC-RSS030	Rhodanine Solution (Stock)	30 ml	2-8°C
SSC-SAB500	Acetate Buffer Solution, pH 8.0	2 x 500 ml	18-25°C
SSC-HMM125	Hematoxylin, Mayer's (Lillie's Mod.)	125 ml	18-25°C

**Mixed Storage Conditions. Separate Contents.**

**For information regarding ordering individual components, please contact us at: 800-442-3573.**

**Precautions:** Keep away from open flame.  
 Avoid contact with skin and eyes.  
 Harmful if swallowed.  
 Follow all Federal, State, and local regulations regarding disposal.  
 Use in chemical fume hood whenever possible.

**Procedure (Standard):****Prepare Working Rhodanine Solution:**

Combine:

4 ml Rhodanine Solution (Stock). Shake Stock Solution immediately before adding to Acetate Buffer.

46 ml Acetate Buffer Solution, pH 8.0

1. Deparaffinize sections if necessary and hydrate to distilled water.
2. Place loosely capped staining jar containing Working Rhodanine in microwave and heat solution until warm but not hot.
3. Place slide in warmed Working Rhodanine Solution and microwave at full power until solution is hot. Do not allow solution to boil.
4. Cap container, gently agitate to mix evenly, and allow solution to cool on countertop to room temperature with occasional agitation.
5. Examine slide microscopically and repeat heating/cooling cycle (steps 3 & 4) until desired staining intensity is achieved.
6. Rinse slide in 2 changes of Acetate Buffer Solution, pH 8.0 for 1 minute each.
7. Stain tissue section with Hematoxylin, Mayer's (Lillie's Modification) for 5-10 seconds.
8. Rinse slide in 3 changes of Acetate Buffer Solution, pH 8.0 for 1 minute each.
9. Dehydrate slide in 3 changes of absolute alcohol.
10. Clear in 2 changes of xylene or xylene substitute, and mount in synthetic resin.

**References:**

1. Sheehan, DC., Hrapchak, BB. Theory and Practice of Histotechnology; 1980, page 230.
2. Lindquist, RR. Studies on the Pathogenesis of Hepatolenticular II: Cytochemical methods for the location of copper. Arch Pathol; 1969, Volume 87: page 370.



Lot-to-Lot Validation Form  
Copper Stain Kit Catalog: SSK-COPPER

Kit Lot Number: \_\_\_\_\_  
Kit Expiration Date: \_\_\_\_\_  
Date Tested: \_\_\_\_\_  
Control Tissue (#) \_\_\_\_\_  
Approved for Use: Y/N \_\_\_\_\_  
Date put into use: \_\_\_\_\_  
If not approved,  
corrective actions  
taken: \_\_\_\_\_  
Approved by: \_\_\_\_\_

Kit Component	Lot #
Rhodanine Solution (stock)	_____
Acetate Buffer, pH 8.0	_____
Hematoxylin, Mayer's (Lillie's Mod.)	_____

Replacement Component if used	Replacement Date	Lot #	Accepted Y/N	Comments
Rhodanine Solution (Stock)				
Acetate Buffer, pH 8.0 Sol.				
Hematoxylin, Mayer's (Lillie's Mod.)				
Approved by: _____				

StatLab is providing this form to assist with reagent lot validation as stated in CLIA'88 Standard 493.1256-For reagent(s), the laboratory must do the following: Check each batch (prepared in-house), lot number (commercially prepared) and shipment of reagents, stains, and identification systems (systems using two or more substrates or two or more reagents, or a combination) when prepared or opened for positive and negative reactivity, if applicable.