# **KPL Silver Enhancer Kit**

for Microscopy Applications



CAT. NO.

<u>Size</u> 50 mL

<u>Catalog No.</u> 5520-0021 (50-22-01)

# DESCRIPTION

KPL Silver Enhancer Kit for Microscopy Applications is a sensitive and simple method for the enhancement of gold labeled samples when viewed using a light microscope. The resulting brown/black stain is permanent and offers sharp resolution and contrast with colored counterstains.

# CONTENTS

This kit contains: KPL Silver Enhancer Solution A (red cap) KPL Silver Enhancer Solution B (blue cap)

## STORAGE/STABILITY

Store at 2 - 8°C. Stable for a minimum of 4 months from date of receipt when stored at 2 - 8°C. **DO NOT FREEZE.** Do not expose to extreme heat or light.

## **APPLICATIONS**

The KPL Silver Enhancer Kit for Microscopy Applications is suitable for use in enhancing the sensitivity of gold conjugates used in light microscopy applications.

#### PROCEDURE

- 1. If using paraffin embedded tissue sections, deparaffinize according to standard protocols.
- 2. Apply the gold conjugated primary antibody or primary antibody followed by a gold conjugated secondary antibody, and incubate as instructed.
- 3. Wash as instructed.
- 4. Mix equal volumes of KPL Solution A and KPL Solution B into a plastic tube. The recommended amount per slide is  $500 \ \mu L 1.0 \ m L/slide$ .
- 5. Incubate the slide with KPL Silver Enhancer.

**NOTE:** The incubation time may need to be optimized depending upon the assay system. As soon as a brownish color is seen, just before the color turns black, the reaction should be stopped.

6. After suitable color intensity is observed, stop the reaction by rinsing the slide in deionized water using a squirt bottle three times for 5 minutes each.

**NOTE:** Direct the stream of water on the slide and not the tissue.

 Counterstain, if desired, with Eosin, KPL Contrast Green or other appropriate counterstain for 30 - 60 seconds.

- 8. Rinse with deionized water for 1 minute.
- 9. Dehydrate through graded ethanol for 3 minutes each in 20%, 40%, 80% and 100% EtOH.

**NOTE:** Floating sections or whole mounts may be fixed to slides by drying under low heat followed by a 1 minute rinse in 95% ethanol.

- 10. Place the sample into two washes of xylene or a xylene substitute, for 1 minute each.
- 11. Air dry thoroughly.
- 12. Mount slides in an organic mounting media.

## PRODUCT SAFETY AND HANDLING

See SDS (Material Safety Data Sheet) for this product.

## RELATED PRODUCTS

KPL Silver Enhancer Kit for	5450-0012 (50-22-02)
Membrane Applications	
KPL Contrast Green	5540-0003 (71-00-11)

# TROUBLESHOOTING GUIDE

Problem	Possible Cause	Corrective Measure
Excessive	Silver enhancer	Shorten/optimize silver
Development	incubation time	enhancer incubation
and /or	too long.	time.
Background		
Floating	Excess antibody;	Dilute primary antibody
Precipitate	reaction too fast.	and/or gold conjugate.
	Silver enhancer incubation time too long.	Optimize silver enhancer time.
Purple or	Excess	Shorten counterstain
Other Color	counterstain.	incubation time.

#### REFERENCES

1. Danscher, G., Hacker, G., et. al., *J. Histotechnology*, 16(3):201-207, 1993.

2. Hacker, G., Grimelius, L., et. al., *J. Histotechnology*, 11(4):213-221, 1988.

3. Holgate, C., Jackson, P., Cowen, P., and Bird, C., *J. Histo/Cytochemistry*, 31(7):938-944, 1983.

4. Danscher, G., *Histochemistry*, 71:1-16, 1981.

The product listed herein is for research use only and is not intended for use in human or clinical diagnosis.