



Instructions For Use FMS-2-IFU

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Revision: 2

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Fontana-Masson Stain Kit

(For Argentaffin Cells and Melanin)

Description:

The Fontana-Masson Stain Kit is intended for use in the histological visualization of Argentaffin cells and Melanin in paraffin or frozen sections. In addition, the Fontana-Masson stain has been reported to be useful in identifying Capsule-Deficient Cryptococcus Neoformans and typical Cryptococcus Neoformans.

| | |
|------------------------|------------|
| Argentaffin Cells: | Black |
| Melanin: | Black |
| Cryptococci Cell Wall: | Black |
| Nuclei: | Red |
| Cytoplasm: | Light Pink |

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:

Any paraffin embedded tissue that contains Hair Follicles or Skin for Melanin.
Small Intestine for Argentaffin.

Availability/Contents:

| <u>Item #</u> | <u>Kit Contents</u> | <u>Volume</u> | <u>Storage</u> |
|---------------|----------------------------------|---------------|------------------|
| GCB030 | Gold Chloride Solution (0.2%) | 30 ml | 2-8° Centigrade |
| SNX009 | Silver Nitrate Solution (10%) | 2 x 9ml | 2-8° Centigrade |
| STB030 | Sodium Thiosulfate Solution (5%) | 30 ml | Room Temperature |
| NFS030 | Nuclear Fast Red Solution | 30 ml | Room Temperature |

Precautions:

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.

Storage: 2° C




25° C



Mixed Storage Conditions.
Separate Contents.

Doc: IFU-TemplateMixedStoragerev2

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Preparation of Reagent Prior to Beginning:

Prepare **Ammoniacal Silver Solution** immediately prior to use.

In new or chemically cleaned glassware, mix 27ml Distilled/Deionized water with one vial of Silver Nitrate Solution (10%) and blend completely. Carefully add Concentrated Ammonium Hydroxide (Not included) one drop at a time, swirling gently after each drop. Initially the mixture will turn dark brown and then gradually become transparent with a fine layer of sediment. The solution is ready for immediate use when all sediment dissolves.

Procedure (Standard):

1. Deparaffinize sections if necessary and hydrate to distilled water.
2. Place freshly mixed Ammoniacal Silver Solution in a 58-60 °C waterbath and allow adequate time for temperature to equalize.
3. Incubate slide in warmed Ammoniacal Silver Solution for 30-60 minutes or until tissue section becomes yellowish/brown in color. (**NOTE:** Melanin typically stains in 30 minutes while Argentaffin stains in 50-60 minutes)
4. Rinse in several changes of distilled water.
5. Incubate slide in Gold Chloride Solution (0.2%) for 30 seconds.
6. Rinse in several changes of distilled water.
7. Incubate slide in Sodium Thiosulfate Solution (5%) for 1-2 minutes.
8. Rinse for 2 minutes in running tap water followed by 2 changes of distilled water.
9. Incubate slide in Nuclear Fast Red Solution for 5 minutes.
10. Rinse for 2 minutes in running tap water followed by 2 changes of distilled water.
11. Dehydrate very quickly in 3 changes of fresh Absolute Alcohol.
12. Clear, and mount in synthetic resin.

References:

1. Sheenan, D.C., Hrapchak, B.B. Theory and Practice of Histotechnology, 2nd Edition. Battelle Press, Columbus, OH.
2. Gaitanis, G., et al. Novel application of the Masson-Fontana Stain for Demonstrating Malassezia Species Melanin-Like Pigment Production In Vitro and in Clinical Specimens. Journal of Clinical Microbiology. 2005, August; 43(8): pages 4147-4151.
3. Kimura, M., et al. Fontana-Masson – stained tissue from culture-proven mycoses. Archives of Pathology & Laboratory Medicine. 1998, December; 122(12): page 11.
4. Lazcano, O., et al. Combined Fontana-Masson-Mucin staining of Cryptococcus neoformans. Archives of Pathology & Laboratory Medicine. 1991, November; 115(11): pages 1145-1149.
5. Ro, J.Y., et al. Advantage of Fontana-Masson stain in capsule-deficient cryptococcal infection. Archives of Pathology & Laboratory Medicine. 1987, January; 111(1): pages 53-57.

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