

Instructions For Use GMG-IFU

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Giemsa Stain Kit (May-Grunwald)

(For Bone Marrow)

Description and Principle

The Giemsa Stain Kit (May-Grunwald) is intended for use in the visualization and differentiation of cells present in hematopoietic tissues. Giemsa Stain Kit is also used to demonstrate certain microorganisms. Giemsa Stain Kit uses a combination of basic and acid dyes to give a Romanowsky-type of staining. Tissue elements carrying a negative charge are stained predominantly with the basic dyes, methylene blue and azure, whereas tissues carrying a positive charge are stained with the acid dye eosin.

Expected Results

Nuclei:Blue/VioletCytoplasm:Light BlueCollagen:Pale PinkMuscle Fibers:Pale Pink

Erythrocytes: Gray, Yellow or Pink *Rickettsia*: Reddish-Purple

Helicobacter pylori: Blue

Mast Cells: Dark Blue with Red Granules

Kit Contents	Storage
1. May-Grunwald Stock Solution	18-25°C
Giemsa Stock Solution	18-25°C
3 Phosphate Buffer Solution, pH 6.8	18-25°C

Suggested Controls (not provided)

Blood Film, Bone Marrow, Spleen, Any well fixed tissue.

Uses/Limitations

For In-Vitro Diagnostic use only.
Do not use past expiration date.
Use caution when handling reagents.

Non-Sterile

This procedure has not been optimized for frozen sections.

Frozen sections may require protocol modification.

Storage

Store kit and all components at room temperature (18-25°C).

Safety and Precautions

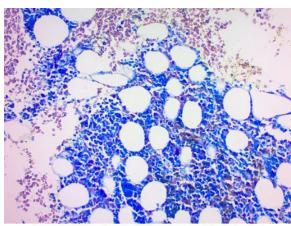
Please see current Safety Data Sheets (SDS) for this product and components GHS classification, pictograms, and full hazard/precautionary statements.

Procedure (Standard):

 Deparaffinize sections if necessary and hydrate to distilled water. For blood smears fix in methanol for 5 minutes before staining.

Prepare Working May-Grunwald Solution by mixing equal parts (1:1) May-Grunwald Stock Solution and Phosphate Buffer Solution, pH 6.8.

- 2. Flood slide with Working May-Grunwald Solution for 5-7 minutes. Note: Agitate slide occasionally to insure proper staining.
- 3. Carefully flood slide with Phosphate Buffer Solution, pH $6.8\,\mathrm{until}$ stain no longer runs off.



Bone Marrow stained with the Giemsa Stain Kit (May-Grunwald) (For Bone Marrow)

When staining tissue samples prepare Working Giemsa Solution by mixing 60μ l (~2 drops) of Giemsa Stock Solution per 1ml of Phosphate Buffer Solution, pH 6.8.

If staining a peripheral blood smear, instead mix 200µl (~6 drops) of Giemsa Stock Solution per 1ml of Phosphate Buffer Solution, pH 6.8.

- 4. Flood slide with Working Giemsa Solution for 10-15 minutes. Note: Agitate slide occasionally to insure proper staining.
- 5. Carefully flood slide with Phosphate Buffer Solution, pH 6.8 until stain no longer runs off.
- 6. Allow slide to remain in Phosphate Buffer Solution, pH 6.8 for an additional 3 minutes.
- 7. Dip slide quickly in distilled water to remove buffer and air dry at room temperature.
- 8. Clear slide in Xylene or Xylene Substitute.
- 9. Mount in synthetic resin.

Notes:

- Background in tissue sections may be differentiated by dipping slide in a solution of 0.25% Acetic Acid (not provided). This may allow for better visualization of mast cells.
- 2. The Working Solutions will immediately begin to precipitate once mixed, use immediately and do not re-use or store for later use.

- References

 1. Sheehan, D., Hrapchak, B., Theory and Practice of Histotechnology: 2nd Edition, 1980, pages 155-156.
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 De Brauwer, E., Jacobs, J., Nieman, F., Bruggeman, C., Drent, M. Test Characterisics of Acridine Orange, Gram, and May-Grunwald-Giemsa Stains for Enumeration of Intracellular Organisms in Bronchoalveolar Lavage Fluid. Journal of Clinical Microbiology, 1999, 37(2): pages 427-429.
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- Lorand-Metze, I., Metze, K. Fractal Characteristics of May-Grunwald-Giemsa Stained Chromatin Are Independent Prognostic Factors for Survival in Multiple Myeloma. 2011, Plos ONE 6(6): e20706. Doi:10.1371/journal.pone.0020706.

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