

# Instructions For Use GMG-1-IFU

**Revision: 2** 

Rev. Date: Feb. 5, 2012

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P.O. Box 3286 - Logan, Utah 84323, U.S.A. - Tel. (800) 729-8350 - Tel. (435) 755-9848 - Fax (435) 755-0015 - www.scytek.com

# Giemsa Stain Kit (May-Grunwald) (For Bone Marrow)

Description:		ald) is intended for use in the visualization of cells present in microorganisms. This kit may be used on formalin-fixed, paraffin-
	Nuclei: Cytoplasm Collagen: Muscle Fibers: Erythrocytes: Rickettsia: Helicobacter Pylori: Mast Cells:	Blue/Violet Light Blue Pale Pink Pale Pink Gray, Yellow or Pink Reddish-Purple Blue Dark Blue with Red Granules
Uses/Limitations:	Not to be taken internally. For In-Vitro Diagnostic use only. Histological applications. Do not use if reagents become clo Do not use past expiration date. Use caution when handling reager Non-Sterile.	
Control Tissue:	Blood Film. Bone Marrow. Spleen. Any well fixed tissue.	and the second
Ordering informatior on back page!	n regarding individual compone	nts

### Avialability/Contents:

Item #	Kit Contents	Volume	Storage
MAY500	May-Grunwald Stock Solution	500 ml	18-25 <i>°</i> C
GGS500	Giemsa Stock Solution	500 ml	18-25 <i>°</i> C
PBM500	Phosphate Buffer Solution, pH 6.8	500 ml	18-25 <i>°</i> C

### **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.

Storage: 18° C







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# **Preparation of Reagents Prior to Beginning:**

- 1. Prepare <u>Working May-Grunwald Solution</u> by mixing 25ml of May-Grunwald Solution (MAY500) with 25ml of Phosphate Buffer Solution, pH 6.8 (PBM500).
- Prepare <u>Working Giemsa Solution</u> by mixing 2.5ml of Giemsa Stock Solution (GGS500) with 50ml of Phosphate Buffer Solution, pH 6.8 (PBM500).

# **Procedure (Standard):**

- 1. Deparaffinize sections if necessary and hydrate to distilled water.
- 2. Place slide in staining tray and flood 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 until stain no longer runs off.
- 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. Dip slide twice in Xylene or Xylene Substitute.
- 9. Mount in synthetic resin.

# **Procedure (Mast Cells):**

- 1. Deparaffinize sections if necessary and hydrate to distilled water.
- 2. Place slide in staining tray and flood 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 until stain no longer runs off.
- 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. Differentiate by dipping slide in Acetic Acid Solution (0.25%) until background is desired intensity.
- 7. Dip slide for 10 seconds in Phosphate Buffer Solution, pH 6.8 while agitating gently.
- 8. Dip slide quickly in distilled water to remove buffer and air dry at room temperature.
- 9. Dip slide twice in Xylene or Xylene Substitute.
- 10. Mount in synthetic resin.

Storage: 18° C



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#### Bulk Reagent Ordering Information and Current Pricing at www.scytek.com

Description:	Catalog #	Volume
May-Grunwald Stock Solution	MAY125 MAY500 MAY999	125 ml 500 ml 1000 ml
Giemsa Stock Solution	GGS125 GGS500 GGS999	125 ml 500 ml 1000 ml
Phosphate Buffer Solution, pH 6.8	PBM500 PBM999	500 ml 1000 ml

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