

Goldner 三色染色试剂盒

货号: G3550

规格: 6×50mL/6×100mL

保存: 室温, 避光保存, 有效期1年。

产品组成:

| 名称 | | 6×50mL | 6×100mL | 保存 |
|--|---------------|--------|---------|--------|
| 试剂(A): Weigert 铁苏木素染色液 | A1:Weigert染液A | 25mL | 50mL | 室温, 避光 |
| | A2:Weigert染液B | 25mL | 50mL | 室温, 避光 |
| 临用前, 取A1、A2等量混合即为Weigert铁苏木素染色液, 不宜提前配制。 | | | | |
| 试剂(B): 酸性分化液 | | 50mL | 100mL | 室温 |
| 试剂(C): Acid Ponceau染色液 | | 50mL | 100mL | 室温, 避光 |
| 试剂(D): 弱酸溶液 | | 50mL | 100mL | 室温 |
| 试剂(E): Orange G染色液 | | 50mL | 100mL | 室温, 避光 |
| 试剂(F): 亮绿染色液 | | 50mL | 100mL | 室温, 避光 |

产品介绍:

骨染色方法有很多种, 例如甲苯胺蓝法、阿利新蓝法、番红 O 法、Goldner 三色染色法等。Goldner 三色染色又称戈德纳三色染色, 与 Masson 三色染色类似, 三色染色通常是指染细胞核和能选择性的显示胶原纤维和肌纤维。该试剂盒多用于骨类物质的染色, 对细胞染色效果较好, 尤其适用于代谢性疾病(Paget 病、骨性营养不良等)的研究, 以评估成骨细胞和破骨细胞的活性, 并容易辨认骨髓中的转移性肿瘤细胞。

自备材料:

系列乙醇、蒸馏水

操作步骤: (仅供参考)

1. 切片二甲苯脱蜡或脱塑料, 梯度乙醇复水。
2. 入配制好的Weigert铁苏木素染色15-20 min, 流水冲洗1min。
3. 用酸性分化液迅速分化(一般<5s), 流水冲洗5min, 蒸馏水稍洗。
4. 入Acid Ponceau染色液染色5min。
5. 在上述过程中按蒸馏水:弱酸溶液=4:1 比例配制弱酸工作液, 用弱酸工作液洗 15~30s。
6. 入 Orange G 染色液染色, 直至胶原纤维红色脱色, 一般需要 3~10min。
7. 用配制好的弱酸工作液冲洗 15~30s。
8. 直接入亮绿染色液中染色 5min, 用配制好的弱酸工作液冲洗 3 次, 每次 15s。
9. 蒸馏水冲洗, 吸干或晾干, 无水乙醇快速脱水, 中性树胶封固。

染色结果:

| | |
|-----|-------|
| 矿化骨 | 绿色 |
| 类骨质 | 橙色-红色 |
| 软骨 | 紫色 |
| 细胞核 | 蓝色-灰色 |

注意事项:

1. 切片脱蜡应尽量干净。固定起着重要的作用, 使用不同的固定液可延长或缩短染色时间。
2. 取A1、A2等量混合即为 Weigert铁苏木素染液, 一般24h失去染色力。
3. 酸性分化时间应根据切片厚薄、组织的类别和新旧而定。
4. 弱酸溶液可使色彩更清晰鲜艳, 如使用量大可自行配制0.1~1%乙酸溶液予以替代。
5. Orange G染色液染色时在镜下控制, 以丽春红脱去为准。

Goldner Trichrome Stain Kit

Cat: G3550

Size: 6×50mL/6×100mL

Storage: RT, avoid light, valid for 1 year.

Kit Components

| Reagent | | 6×50mL | 6×100mL | Storage |
|---|------------------------|--------|---------|-----------------|
| Reagent(A): Weigert Iron Hematoxylin Solution | A1: Weigert Solution A | 25mL | 50mL | RT, avoid light |
| | A2: Weigert Solution B | 25mL | 50mL | RT |
| Mix equal parts of A1 and A2 to form Weigert Iron Hematoxylin Solution before use. This working solution is not suitable to prepare in advance. | | | | |
| Reagent(B): Acid Differentiation Solution | | 50mL | 100mL | RT |
| Reagent(C): Acid Ponceau Solution | | 50mL | 100mL | RT, avoid light |
| Reagent(D): Weak Acid Solution | | 50mL | 100mL | RT |
| Reagent(E): Orange G Solution | | 50mL | 100mL | RT, avoid light |
| Reagent(F): Light Green Solution | | 50mL | 100mL | RT, avoid light |

Introduction

There are many methods of bone staining, such as Toluidine Blue method, Alcian Blue method, Saffron O method, Goldner Trichrome method and so on. Goldner Trichrome Staining is similar to Masson Trichrome Staining. Trichrome staining usually refers to staining the nucleus and selectively displaying collagen and muscle fibers. This kit is mainly used for staining bone substances and has good effect on cell staining, especially for the study of metabolic diseases (Paget disease, osteodystrophy, etc.), so as to evaluate the activity of osteoblasts and osteoclasts, and easily identify the metastasis tumor cells in bone marrow.

Self Prepared Materials

Series of ethanol, Distilled water

Protocols(for reference only)

1. Dewax or deplasticize in xylene, rinsing in ethanol and soak in distilled water.
2. Stain with prepared Weigert Iron Hematoxylin Solution for 15-20 min, rinse in running water for 1min.
3. Quickly differentiate by Acid Differentiation Solution, rinse in running water for 5min.
4. Stain with Acid Ponceau Solution for 5min.
5. In the above process, mix distilled water and Weak Acid Solution as the ratio of 4:1 to prepare Weak Acid Working Solution, then wash in Weak Acid Working Solution for 15-30s.
6. Stain with Orange G Solution until the red color of collagen fiber decolorizes, generally about for 3-10min.
7. Wash in prepared Weak Acid Working Solution for 15-30s.
8. Directly stain with Light Green Solution for 5min.
9. Wash in prepared Weak Acid Working Solution for three times and each time for 15s.
10. Wash in distilled water and suck dry or air dry. Quickly dehydrate in absolute ethanol and seal with resinene.

Result

| | |
|------------------|------------|
| Mineralized Bone | Green |
| Osteoid | Orange-Red |
| Cartilage | Purple |
| Nucleus | Blue-Grey |

Note

1. Section dewaxing should be as clean as possible. Fixation plays an important role. According to different fixatives, can prolong or shorten the dyeing time.
2. The Weigert Iron Hematoxylin Solution is ready to use, which generally loses its dyeing power for 24 h.
3. The differentiation time of Acid Differentiation Solution should be determined according to the thickness of slice, the type of tissue and the old and new.
4. Weak Acid Solution can make the color more clear and bright. If the use amount is large, can replace with 0.1-1% acetic acid solution.
5. Orange G Solution staining is controlled under the microscope, subject to the removal of Acid Ponceau Solution.