

## VG 染色实验报告

### 一、实验器材及试剂

#### 1、实验器材

名称	厂家	型号
脱水机	DIAPATH	DONATELLO
包埋机	武汉俊杰电子有限公司	JB-P5
病理切片机	上海徕卡仪器有限公司	RM2016
冻台	武汉俊杰电子有限公司	JB-L5
组织摊片机	浙江省金华市科迪仪器设备有限公司	KD-P
烤箱	天津市莱玻瑞仪器设备有限公司	GFL-230
载玻片	Wanwu	
正置光学显微镜	日本尼康	NIKON ECLIPSE E100
成像系统	日本尼康	NIKON DS-U3

#### 2、主要实验试剂

试剂名称	厂家	货号
无水乙醇	国药集团化学试剂有限公司	100092683
二甲苯	国药集团化学试剂有限公司	10023418
VG 染液套装	Wanwu	G1046
中性树脂	国药集团化学试剂有限公司	10004160

### 二、实验步骤

1、石蜡切片脱蜡至水：依次将切片放入二甲苯I20min-二甲苯II20min-无水乙醇I5min-无水乙醇II5min-75%酒精 5min，自来水洗。

2、VG 染色：Van Gieson (VG) 染液 B 9ml 加入 Van Gieson (VG) 染液 A 1ml (配制后可反复使用多次) 混合成 VG 染液 (按比例配制用多少配多少)，染 1min，快速水洗，无水乙醇三缸快速脱水。

~~3、切片放入二甲苯透明，中性树脂封片。透明封片：两缸干净的二甲苯透明各 20s、5min (二甲苯专用不与其他二甲苯共用)，中性树脂湿封。~~

4、显微镜镜检，图像采集分析。

### 三、结果判读：

胶原纤维呈红色，其他组织成分呈黄色。

### 四、注意事项：

~~1.4~~水洗要快，防止红色褪掉，无水乙醇也要快脱，防止黄色变暗。

## **VG staining experimental report STAINING- EXPERIMENT REPORT**

### **1. Experimental equipment and reagents**

#### **1.1 Experimental equipment**

<b><u>Name</u></b>	<b><u>Manufacturer</u></b>	<b><u>Model</u></b>
<u>Dehydrator</u>	<u>DIAPATH</u>	<u>Donatello</u>
<u>Embedding machine</u>	<u>Wuhan Junjie Electronics Co., Ltd</u>	<u>JB-P5</u>
<u>Pathology slicer</u>	<u>Shanghai Leica Instrument Co., Ltd</u>	<u>RM2016</u>
<u>Frozen platform</u>	<u>Wuhan Junjie Electronics Co., Ltd</u>	<u>JB-L5</u>
<u>Tissue spreader</u>	<u>Zhejiang Kehua Instrument Co., Ltd</u>	<u>KD-P</u>
<u>Oven</u>	<u>Tianjin Laibo Rui Instrument Equipment Co., Ltd</u>	<u>GFL-230</u>
<u>Slides</u>	<u>Wanwu</u>	
<u>Orthostatic microscope</u>	<u>NIKON, JAPAN</u>	<u>NIKON ECLIPSE E100</u>
<u>Image system</u>	<u>NIKON, JAPAN</u>	<u>NIKON DS-U3</u>

### **1. Experimental equipment and reagents**

experiment-equipment

<b><u>name</u></b>	<b><u>factory</u></b>	<b><u>model</u></b>
<u>Dehydrator</u>	<u>DIAPATH</u>	<u>Donatello</u>
<u>Embedding machine</u>	<u>Wuhan Junjie Electronics Co., Ltd.</u>	<u>JB-P5</u>
<u>Pathology slicer</u>	<u>Shanghai Leica Instruments Co., Ltd.</u>	<u>RM2016</u>
<u>Frozen platform</u>	<u>Wuhan Junjie Electronics Co., Ltd.</u>	<u>JB-L5</u>
<u>Organizer</u>	<u>Jinhua City, Zhejiang Province Cody</u> <u>Instrument-Equipment-Co.,Ltd.</u>	<u>KD-P</u>
<u>oven</u>	<u>Tianjin=Lai=Bo=Rui=Instrument=Equipment=GFL-230</u> <u>Co.,Ltd.</u>	
<u>Glass slide</u>	<u>Wanwu</u>	
<u>UPRIGHT OPTICAL</u> <u>MICROSCOPE</u>	<u>NIKON JAPAN</u>	<u>NIKON ECLIPSE E100</u>

IMAGING SYSTEM

NIKON JAPAN

NIKON DS-U3

## 1.2 Main experimental reagents

### Main experimental reagents

<u>Reagent name</u>	<u>Reagent name</u>	<u>Manufacturer</u>	<u>factory</u>	<u>Article number</u>	<u>Article number</u>
Absolute ethanol		Sinopharm Group Chemical Reagent Co., Ltd.		100092683	
Xylene		Sinopharm Group Chemical Reagent Co., Ltd.		10023418	
VG <u>staining</u>		Wanwu		G1046	
Neutral gum		Sinopharm Group Chemical Reagent Co., Ltd.		10004160	

## 2. Experimental steps

2.1 Paraffin slides dewaxed as follow: Two changes of pure xylene for 20 min; Two changes of pure ethanol for 5min; 75% ethanol for 5min; Keep slides in tap water.

2.2 VG dyeing: Add 9ml of Van Gieson (VG) dye solution B to 1ml of Van Gieson (VG) dye solution A (can be used repeatedly after preparation) and mix into VG dye solution (prepared in proportion to use as much as possible), dye for 1 minute, fast Washed with water, dehydrated quickly in anhydrous ethanol triplex.

2.3 ~~III. The slides were immersed in xylene to transparent and then coverslip with neutral resin. Transparent sealing: two cylinders of clean xylene are transparent for 20s and 5min each (xylene is not shared with other xylenes), neutral gum wet seal.~~

~~IV 2.4. Observed under microscope and took images.~~

~~Microscope inspection, image acquisition and analysis.~~

## 3. The results were as follows

Collagen fibers ~~was~~ red, and other tissue components ~~was~~ yellow.

## 4 Four, Precautions

1. It should be washed quickly to prevent the red from fading, and the absolute ethanol should also be taken off quickly to prevent the yellow from darkening.