



北京富百科生物技术有限公司  
www.fluorescence.cn

北京富百科生物技术有限公司  
富百科---用荧光丰富百家学科, 用荧光点亮生命科学  
Beijing Fluorescence Biotechnology Co. Ltd  
Fluorescence---Light up the life sciences

## Propidium diiodide 或 PI 染色液说明书

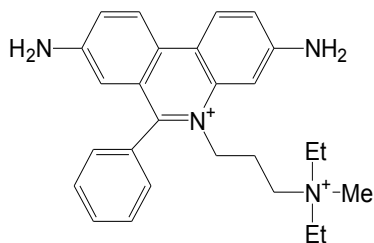
中文名: 3,8-二氨基-5-[3-(二乙基甲氨基)丙基]-6-苯基吩啉二碘; 碘化丙啉

英文名: 3,8-Diamino-5-[3-(diethylmethylammonio)propyl]-6-phenylphenanthridini-um diiodide,

Propidium iodide; Propidium diiodide; PI

分子式: C<sub>27</sub>H<sub>34</sub>I<sub>2</sub>N<sub>4</sub> 分子量: 668.39; CAS#: 25535-16-4

结构式:



性质:

1. 外观: 碘化丙啉固体为红棕色粉末; PI 染色液为紫红色溶液 浓度: 10mM solution in DMSO
2. 纯度: ≥98%超纯级 UltraPure
3. 产品描述:

PI 是一种可对 DNA 染色的细胞核染色试剂。它在嵌入双链 DNA 后释放红色荧光。常用于细胞凋亡 (apoptosis) 或细胞坏死 (necrosis) 的检测, 常用于流式细胞仪分析。尽管 PI 不能通过活细胞膜, 但却能穿过破损的细胞膜而对核染色。PI 经常被用来与 Calcein, AM、Hoechst 33258 或 Hoechst 33342 等一起使用, 能同时对活细胞和死细胞染色。PI-DNA 复合物的激发和发射波长分别为 535nm 和 615nm。

4. 染色过程

- (1) 用 PBS 或适当的缓冲液制备 10~50μM 的 PI 溶液。
- (2) 将 1/10 培养基体积的 PI 溶液加入到细胞培养基中 (也可以用 1/10 浓度的 PI 缓冲液代替培养基)。
- (3) 在 37°C 培养细胞 10~20 分钟。
- (4) 用 PBS 或合适的缓冲液洗涤细胞两次。
- (5) 用 535nm 激发波长, 615nm 发射波长的滤光器的荧光显微镜观察细胞。

储存条件: -20°C 干燥避光保存, 有效期一年。

注意事项:

Sigma 的 MSDS 显示碘化丙啉 PI 与 EB 一样对生殖细胞致突变性, 可能致癌, 请必须注意防护。



北京富百科生物技术有限公司  
www.fluorescence.cn

北京富百科生物技术有限公司  
富百科---用荧光丰富百家学科，用荧光点亮生命科学  
Beijing Fluorescence Biotechnology Co. Ltd  
Fluorescence---Light up the life sciences

## Propidium diiodide and 1.5 mM PI aqueous solution

### Description

Chemical Name: 3,8-Diamino-5-[3-(diethylmethylammonio)propyl]-6-phenylphenanthridini-um diiodide;

Propidium iodide; Propidium diiodide; PI

Appearance: reddish-brown powder

Purity: ≥98% UltraPure; MW: C<sub>27</sub>H<sub>34</sub>I<sub>2</sub>N<sub>4</sub>=668.39; CAS: 25535-16-4

Storage & Shipping Condition : Store at 4°C protected from light. Product is shipped at ambient temperature

### Product Description

Propidium iodide (PI) is an ethidium bromide analog that emits red fluorescence upon intercalation with double-stranded DNA. Though PI does not permeate viable cell membranes, it passes through disturbed cell membranes and stains the nuclei. PI is often used in combination with a fluorescein compound, such as Calcein-AM or FDA, for simultaneous staining of viable and dead cells. The excitation and emission wavelengths of PI-DNA complex are 535 nm and 615 nm, respectively.

### Application---Staining Procedure

1. Prepare 10-50 μM PI solution with PBS or an appropriate buffer
2. Add PI solution with 1/10 of the volume of cell culture medium to the cell culture
3. Incubate the cell at 37°C for 10-20 min.
4. Wash cells twice with PBS or an appropriate buffer.
5. Observe the cells under a fluorescence microscope with 535 nm excitation and 615 nm emission filters.

**Note:** Since PI may be carcinogenic, extreme care is necessary during handling.

### Reference

- T. Crompton, et al., Biochem Biophys. Res. Commun., 183, 532 (1992)  
N. M. Poulin, et al., J. Histochem. Cytochem., 42, 1149 (1994);  
F. Belloc, et al., Cytometry, 17, 59 (1994);  
C. A. Van Hooijdonk, et al., Cytometry, 17, 185 (1994);  
D. L. Garner, et al., J. Androl., 15, 620 (1994);  
C. Souckler, et al., Cytometry, 20, 203 (1995);  
D. L. Garner, et al., Biol. Reprod., 53, 276 (1995);  
M. Wulf, et al., Biotechniques, 19, 368 (1995);  
K. Wrobel, et al., J. Immunol. Methods, 189, 243 (1996);  
W. E. Corver, et al., Cytometry, 28, 329 (1997).