

# Dyes and Stains

## Propidium Iodide

Cell viability dye (DNA-labeling dye)

Catalog # 75002

10 mg



Scientists Helping Scientists™ | [WWW.STEMCELL.COM](http://WWW.STEMCELL.COM)

TOLL FREE PHONE 1 800 667 0322 • PHONE +1 604 877 0713

[INFO@STEMCELL.COM](mailto:INFO@STEMCELL.COM) • [TECHSUPPORT@STEMCELL.COM](mailto:TECHSUPPORT@STEMCELL.COM)

FOR GLOBAL CONTACT DETAILS VISIT OUR WEBSITE

## Product Description

Propidium Iodide (PI) is a red-fluorescent cell viability dye which is excluded from live cells with intact membranes, but penetrates dead or damaged cells and binds to DNA and RNA by intercalating between the bases. It is widely used as a counterstain to differentiate and exclude non-viable cells in flow cytometric analyses, and can be excited using blue (488 nm), green (532 nm), or yellow-green (561 nm) laser lines, with detection in the FL2 or FL3 channels. PI is used in DNA fluorescence imaging applications to discriminate early and late stages of apoptosis, to study cell-mediated cytotoxicity, and for chromosome analysis. It is also commonly used in quantitative DNA assays.

<b>Chemical Name:</b>	3,8-diamino-5-[3-(diethylmethylammonio)propyl]-6-phenylphenanthridinium diiodide
<b>Alternative Names:</b>	3,8-Diamino-5-{3-[diethyl(methyl)ammonio]propyl}-6-phenylphenanthridinium diiodide; PI; Propidium diiodide
<b>CAS Number:</b>	25535-16-4
<b>Chemical Formula:</b>	$C_{27}H_{34}N_4 \cdot 2I$
<b>Molecular Weight:</b>	668.4 g/mol
<b>Excitation Wavelength:</b>	488 - 535 nm (DNA or RNA complex)
<b>Emission Wavelength:</b>	617 nm (DNA or RNA complex)

## Properties

<b>Storage:</b>	Store at -20°C.
<b>Shelf Life:</b>	Product stable until expiry date (EXP) on label. Protect product from prolonged exposure to light.
<b>Format/Formulation:</b>	A crystalline solid

## Applications

<b>Verified:</b>	FC
<b>Reported:</b>	FC, FISH, Fluorescence microscopy, Fluorometry, ICC
<b>Special Applications:</b>	This product has been verified for viability assessments of cells isolated with EasySep™ and RosetteSep™ kits.

Abbreviations: CellSep: Cell separation; ChIP: Chromatin immunoprecipitation; FA: Functional assay; FC: Flow cytometry; ICC: Immunocytochemistry; IF: Immunofluorescence microscopy; IHC: Immunohistochemistry; IP: Immunoprecipitation; RIA: Radioimmunoassay; WB: Western blotting

## Handling/Directions for Use

For preparing a stock solution, PI is soluble in aqueous buffers and organic solvents, as follows:

- Phosphate-buffered saline (PBS), pH 7.2  $\leq$  2 mg/mL
- Ethanol  $\leq$  0.2 mg/mL
- Dimethyl sulfoxide (DMSO)  $\leq$  2.5 mg/mL
- Dimethyl formamide (DMF)  $\leq$  3.3 mg/mL

NOTE: If preparing stock solution using an organic solvent, further dilute into aqueous buffer or isotonic saline before performing biological experiments. Ensure that the residual amount of organic solvent is insignificant, as it may have physiological effects at low concentrations.

Whenever possible, prepare and use stock solution on the same day. Protect stock solution from prolonged exposure to light. If stock solution must be made in advance, aliquot and store in tightly sealed vials at  $-20^{\circ}\text{C}$  and protect from prolonged exposure to light. Generally these will be stable for up to 1 month.

### FLOW CYTOMETRY

1. Prepare a 1 mg/mL (1.5 mM) stock solution by dissolving solid PI in PBS.
2. Add to cells at a final concentration of  $\leq$  1  $\mu\text{g/mL}$ .
3. Incubate for 5 - 10 minutes in the dark, then analyze immediately.

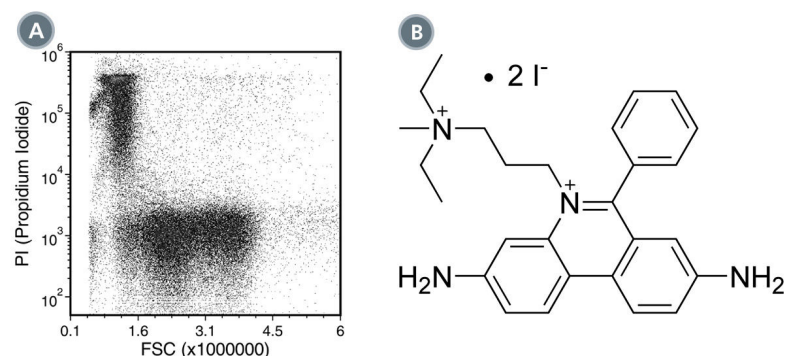
Titrate the dye for optimal performance in each application.

## Notes and Tips

For flow cytometric analysis, PI can be detected in the FL2 (DNA content) or FL3 (viability) channels. Use FL2 to analyze PI staining if it is being used as a counterstain with fluorescein-conjugated Annexin V.

For microscopy analysis, PI can be viewed using a rhodamine (red) filter. Cells will be stained with PI if the membrane has been permeated, e.g. as a result of natural cell death or detergent treatment.

## Data/Structure



(A) Flow cytometry analysis of human peripheral blood mononuclear cells (PBMCs) labeled with PI.

(B) Chemical structure of PI.

## References

1. Beggs KM et al. (2014) Molecular mechanisms of hepatocellular apoptosis induced by trovafloxacin-tumor necrosis factor- $\alpha$  interaction. *Toxicol Sci* 137(1): 91–101. (FC)
2. Jeong SM et al. (2014) SIRT4 protein suppresses tumor formation in genetic models of Myc-induced B cell lymphoma. *J Biol Chem* 289(7): 4135–44. (FC)
3. Kong S et al. (2014) DBC1 is a suppressor of B cell activation by negatively regulating alternative NF- $\kappa$ B transcriptional activity. *J Immunol* 193(11): 5515–24. (FC)

4. Sonnemann J et al. (2014) p53-dependent and p53-independent anticancer effects of different histone deacetylase inhibitors. *Br J Cancer* 110(3): 656–67. (FC)
5. Yun B et al. (2014) Serine hydrolase inhibitors block necrotic cell death by preventing calcium overload of the mitochondria and permeability transition pore formation. *J Biol Chem* 289(3): 1491–504. (ICC, IF)
6. Zhao W et al. (2013) The peroxisome-proliferator activated receptor- $\gamma$  agonist pioglitazone modulates aberrant T cell responses in systemic lupus erythematosus. *Clin Immunol* 149(1): 119–32. (FC)
7. Riccardi C & Nicoletti I. (2006) Analysis of apoptosis by propidium iodide staining and flow cytometry. *Nat Protoc* 1(3): 1458–61. (FC)
8. Sakimoto I et al. (2006) Alpha-sulfoquinovosylmonoacylglycerol is a novel potent radiosensitizer targeting tumor angiogenesis. *Cancer Res* 66(4): 2287–95. (FC)
9. Kral T et al. (2005) Propidium iodide and PicoGreen as dyes for the DNA fluorescence correlation spectroscopy measurements. *J Fluoresc* 15(2): 179–83. (Fluorometry)
10. Lecoer H. (2002) Nuclear apoptosis detection by flow cytometry: influence of endogenous endonucleases. *Exp Cell Res* 277(1): 1–14. (FC)
11. Coder DM. (2001) Assessment of cell viability. *Curr Protoc Cytom* Chapter 9: Unit 9.2. (FC)
12. Douglas RS et al. (1995) A simplified method for the coordinate examination of apoptosis and surface phenotype of murine lymphocytes. *J Immunol Methods* 188(2): 219–28. (FC)
13. Martin SJ et al. (1995) Early redistribution of plasma membrane phosphatidylserine is a general feature of apoptosis regardless of the initiating stimulus: inhibition by overexpression of Bcl-2 and Abl. *J Exp Med* 182(5): 1545–56. (FC)
14. Crompton T et al. (1992) Propidium iodide staining correlates with the extent of DNA degradation in isolated nuclei. *Biochem Biophys Res Commun* 183(2): 532–7. (FC)
15. De Caestecker MP et al. (1992) The detection of intracytoplasmic interleukin-1 alpha, interleukin-1 beta and tumour necrosis factor alpha expression in human monocytes using two colour immunofluorescence flow cytometry. *J Immunol Methods* 154(1): 11–20. (FC)

## Related Products

For a complete list of related products available from STEMCELL Technologies, visit [www.stemcell.com/dyesandstains](http://www.stemcell.com/dyesandstains) or contact us at [techsupport@stemcell.com](mailto:techsupport@stemcell.com).

STEMCELL TECHNOLOGIES INC.'S QUALITY MANAGEMENT SYSTEM IS CERTIFIED TO ISO 13485. PRODUCTS ARE FOR RESEARCH USE ONLY AND NOT INTENDED FOR HUMAN OR ANIMAL DIAGNOSTIC OR THERAPEUTIC USES UNLESS OTHERWISE STATED.

Copyright © 2018 by STEMCELL Technologies Inc. All rights reserved including graphics and images. STEMCELL Technologies & Design, STEMCELL Shield Design, Scientists Helping Scientists, EasySep, and RosetteSep are trademarks of STEMCELL Technologies Canada Inc. All other trademarks are the property of their respective holders. While STEMCELL has made all reasonable efforts to ensure that the information provided by STEMCELL and its suppliers is correct, it makes no warranties or representations as to the accuracy or completeness of such information.