Resazurin (Sodium Salt)

Dyes and **Stains**

Cell proliferation and viability dye

Catalog # 75005 5 q



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Product Description

Resazurin (7-hydroxy-10-oxidophenoxazin-10-ium-3-one, sodium) is a blue fluorogenic dye used as a redox indicator in cell viability and proliferation assays for bacteria, yeast, or mammalian cells. The blue form of the dye is irreversibly reduced by enzymes in viable cells to generate a highly red-fluorescent product, resorufin, which exhibits an emission maximum at ~595 nm and can be detected by flow cytometry, fluorescence microscopy, and high-throughput screening methods. Resazurin is minimally toxic to living cells, making it suitable for use in long-term cell culture. The dye has also been used to assay L-glutamate and to measure the metabolic activity of mitochondria.

Chemical Name: sodium; 10-oxido-7-oxophenoxazin-10-ium-3-olate

Alternative Names: 7-hydroxy-10-oxidophenoxazin-10-ium-3-one, sodium; Diazoresorcinol, sodium salt

CAS Number: 62758-13-8 Chemical Formula: C₁₂H₆NO₄ · Na Molecular Weight: 251.2 g/mol

Excitation Wavelength: 530 - 545 (resorufin product) Emission Wavelength: 585 - 595 (resorufin product)

Properties

Storage: Store at 15 - 25°C.

Product stable until expiry date (EXP) on label. Protect product from prolonged exposure to light. Shelf Life:

Format/Formulation: A crystalline solid

Applications

Verified: FΑ

Reported: Enzyme assay, FA (Cytotoxicity, Proliferation, Viability), FC, Fluorescence microscopy, Fluorometry,

Spectroscopy

Special Applications: This product has been verified for analyzing hematopoietic cells cultured in StemSpan™ SFEM

(Catalog #09650) and StemSpan™ SFEM II (Catalog #09655).

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

PREPARATION

A stock solution may be made by dissolving Resazurin in the diluent of choice. Resazurin is soluble in aqueous buffers and in organic solvents. Guidelines for the solubility of Resazurin are as follows:

- · Phosphate-buffered saline (PBS), pH 7.2 ≤ 5 mg/mL
- · Ethanol ≤ 0.5 mg/mL
- \cdot DMSO \leq 0.5 mg/mL
- · Dimethyl formamide ≤ 0.5 mg/mL

NOTE: If making a stock solution using an organic solvent, further dilutions into aqueous buffers or isotonic saline should be made prior to performing biological experiments. Ensure that the residual amount of organic solvent is insignificant, as organic solvents may have physiological effects at low concentrations.

Wherever possible, prepare and use the stock solutions on the same day. Protect stock solutions from prolonged exposure to light. If stock solutions must be made in advance, it is recommended that they are stored in aliquots in tightly sealed vials at -20°C, protected from prolonged exposure to light. Generally these will be stable for up to 1 month.

The dye should be titrated for optimal performance for each cell type and application.

Notes and Tips

Data may be collected using either fluorescence-based or absorbance-based instrumentation. Absorbance can be measured using a spectrophotometer at 570 nm, or 600 nm if wavelength correction is available. It is recommended that the dye be assayed at pH 6.8 - 7.4.

Data/Structure

Chemical structure of Resazurin (Sodium Salt).

Dyes and Stains Resazurin (Sodium Salt)



References

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- 2. Diril MK et al. (2012) Cyclin-dependent kinase 1 (Cdk1) is essential for cell division and suppression of DNA re-replication but not for liver regeneration. Proc Natl Acad Sci USA 109(10): 3826–31. (Cell proliferation assay)
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- 7. Wilson BAP et al. (2011) High-throughput screen identifies novel inhibitors of cancer biomarker α-methylacyl coenzyme A racemase (AMACR/P504S). Mol Cancer Ther 10(5): 825–38. (Cell growth, viability assays)
- 8. Xu S et al. (2011) Marek's disease virus type 1 microRNA miR-M3 suppresses cisplatin-induced apoptosis by targeting Smad2 of the transforming growth factor beta signal pathway. J Virol 85(1): 276–85. (Cell viability assay)
- 9. Hamid R et al. (2004) Comparison of alamar blue and MTT assays for high through-put screening. Toxicol In Vitro 18(5): 703–10. (Cell viability assay, Fluorescence microscopy)
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- 11. Ahmed SA et al. (1994) A new rapid and simple non-radioactive assay to monitor and determine the proliferation of lymphocytes: an alternative to [3H]thymidine incorporation assay. J Immunol Methods 170(2): 211–24. (Cell proliferation assay)

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