

H2DCFDA

#Cat: NB-64-06719-1ml Size: 1 ml #Cat: NB-64-06719-50mg Size: 50 mg #Cat: NB-64-06719-100mg Size: 100 mg

Chemical Properties:

| CAS No: | 4091-99-0 | |
|-------------------|--|--|
| Formula: | C ₂₄ H ₁₆ Cl ₂ O ₇ | |
| Molecular Weight: | 487.29 | |
| Appearance: | no data available | |
| Storage: | keep away from direct sunlight, store at low temperature | |
| | Powder: -20°C for 3 years In solvent: -80°C for 1 year | |



Biological Description:

| Description | H2DCFDA (DCFH-DA) belongs to the class of green fluorescent dyes and is a probe for the detection of intracellular reactive oxygen species (ROS) (Ex/Em=488/525 nm) with cell membrane neuropeitite. | | |
|----------------|---|--|--|
| | membrane permeability. | | |
| Targets (IC50) | Reactive Oxygen Species | | |
| In vitro | METHODS : Flow cytometry was used to detect ROS levels: 1, H2DCFDA was dissolved into 10 mM DMSO stock solution and further diluted with PBS before use. 2. Adherent cells are incubated with 5 μ M H2DCFDA solution for 30 min at 37°C, protected from light, then harvested with 0.05% trypsin-EDTA solution, suspended in fresh medium and immediately analyzed by flow cytometry (488 nm). [1] METHODS: Confocal microscopy was performed to detect ROS levels: 1. H2DCFDA was dissolved into 10 mM DMSO stock solution and further diluted with PBS before use. 2. Coverslips containing cells were placed in 5 μ M H2DCFDA staining solution and incubated for 60 min at 37°C, protected from light, then washed and imaged with a confocal laser scanning microscope Leica TCS SL equipped with an argon laser. | | |
| In vivo | [1] METHODS : Fluorescence microscopy was used to analyze the oxidative activity of LPS induced peritonitis in mice: 1, H2DCFDA was dissolved in 100 μL ethanol and further diluted with PBS before use. 2. C57BL/6J mice were injected intraperitoneally with LPS (0.1-1 mg/mL) to induce peritonitis. 3.5 h later, H2DCFDA (0.1-0.8 mg/ml) was injected intraperitoneally. 3. 30 min after H2DCFDA injection, the animals were killed by cervical dislocation, and the peritoneal cells were recovered by rinsing with 5 mL of ice-cold HBSS solution at pH 7.4. 4. Peritoneal cells were washed and resuspended in PBS. Macrophages were removed by adhesion method after incubation in a polystyrene dish at 37°C for 30 min. The supernatant was recovered and approximately 20,000-25,000 leukocytes were added to microscope slides using a cytocentrifuge. The slides were then analyzed using fluorescence microscopy. [2] | | |

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Solubility Information

| | Ethanol: 14.29 mg/mL (29.33 mM), Sonication is recommended. |
|------------|---|
| Solubility | DMSO: 50 mg/mL (102.61 mM), |
| | (< 1 mg/ml refers to the product slightly soluble or insoluble) |

Preparing Stock Solutions

| | 1mg | 5mg | 10mg |
|-------|-----------|------------|------------|
| 1 mM | 2.0522 mL | 10.2608 mL | 20.5217 mL |
| 5 mM | 0.4104 mL | 2.0522 mL | 4.1043 mL |
| 10 mM | 0.2052 mL | 1.0261 mL | 2.0522 mL |
| 50 mM | 0.041 mL | 0.2052 mL | 0.4104 mL |

Please select the appropriate solvent to prepare the stock solution, according to the solubility of the product in different solvents. Please use it as soon as possible.

Reference

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$\label{eq:Inhibitor} Inhibitor \cdot Natural Compounds \cdot Compound Libraries \cdot Recombinant Proteins \\ This product is for Research Use Only \cdot Not for Human or Veterinary or Therapeutic Use \\ \end{tabular}$