

NBD-Pen

Chemical Properties

CAS No. : 1955505-54-0

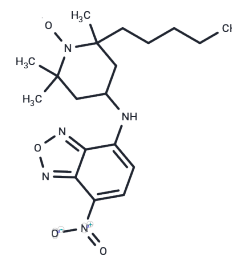
Formula: C₁₉H₂₈N₅O₄

Molecular Weight: 390.46

Store at low temperature

Storage: Powder: -20°C for 3 years | In solvent: -80°C for 1 year

Actual storage temperature shall be subject to the COA.



Biological Description

Description	NBD-Pen is a fluorescent probe that detects lipid free radicals in living cells.
Targets(IC50)	Others
Cell Research	<p>Instructions</p> <p>I. Reagent preparation</p> <ol style="list-style-type: none"> 1. Solvent: NBD-Pen is usually provided in powder form and needs to be dissolved with an appropriate solvent (such as dimethyl sulfoxide (DMSO) or ethanol) before use. 2. Concentration: To prepare the working solution, the common concentration range is 10-50 μM. The specific concentration can be adjusted according to the experimental requirements. <p>II. Operation steps</p> <ol style="list-style-type: none"> 1. Cell treatment: <ol style="list-style-type: none"> 1) Cell culture: Plant the cells to be treated in a suitable culture medium, such as DMEM or RPMI-1640, and culture for 24 hours to adapt the cells. 2) NBD-Pen labeling: Add the dissolved NBD-Pen solution to the cell culture medium to ensure that its concentration is within the working range. Typically, cells are incubated at 37°C for 30 minutes to 1 hour to ensure that NBD-Pen is absorbed by the cells. 3) Washing: After incubation, gently wash the cells with PBS buffer or cell culture medium to remove the probe that has not entered the cells. 2. Fluorescence detection: <ol style="list-style-type: none"> 1) Fluorescence microscopy: When using a fluorescence microscope for detection, the excitation wavelength of NBD-Pen is about 460 nm and the emission wavelength is about 530 nm. The generation of lipid free radicals will lead to a strong enhancement of the fluorescence signal. 2) Flow cytometry: Flow cytometry can also be used to perform fluorescence detection on the treated cells to observe the changes in intracellular fluorescence. 3. Data analysis: <ol style="list-style-type: none"> 1) Fluorescence intensity: By analyzing the fluorescence intensity in the cells, the degree of generation of intracellular lipid free radicals can be evaluated. The higher the fluorescence intensity, the more significant the lipid peroxidation reaction. 2) Standard curve: In some quantitative experiments, the concentration of lipid free radicals can be further quantified by establishing a standard curve.

A DRUG SCREENING EXPERT

Cell Research	<p>Notes:</p> <ol style="list-style-type: none">1. Solubility: NBD-Pen has good solubility, but it should be ensured that the solvent used does not have a negative impact on the activity of the cells.2. Fluorescence bleaching: When performing fluorescence microscopy observation, long-term strong light exposure should be avoided to prevent bleaching of the probe fluorescence signal.3. Experimental control: A blank group and a control group need to be set up in the experiment to ensure the accuracy of the experimental data. <p>The above information is based on published literature. Experimental procedures should be appropriately modified to meet specific research demands.</p>
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Preparing Stock Solutions

	1mg	5mg	10mg
1 mM	2.5611 mL	12.8054 mL	25.6108 mL
5 mM	0.5122 mL	2.5611 mL	5.1222 mL
10 mM	0.2561 mL	1.2805 mL	2.5611 mL
50 mM	0.0512 mL	0.2561 mL	0.5122 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.

Note: The dilution table applies only to solid products. For liquid products, please calculate the stock solution based on the stated concentration and/or density.

Reference

- Takajo T, et al. Mechanism of lipid peroxidation of liposomes by cold atmospheric pressure plasma jet irradiation. *J Clin Biochem Nutr.* 2024 Nov;75(3):183-189.
- Mishima E, et al. Drugs Repurposed as Antiferroptosis Agents Suppress Organ Damage, Including AKI, by Functioning as Lipid Peroxyl Radical Scavengers. *J Am Soc Nephrol.* 2020 Feb;31(2):280-296.
- Ishida Y, et al. Detection and inhibition of lipid-derived radicals in low-density lipoprotein. *Free Radic Biol Med.* 2017 Dec;113:487-493.

Inhibitor · Natural Compounds · Compound Libraries · Recombinant Proteins

This product is for Research Use Only · Not for Human or Veterinary or Therapeutic Use

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