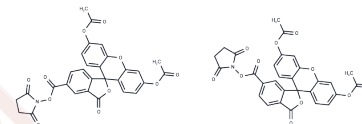


CFSE

Chemical Properties

| | |
|-------------------|---|
| CAS No. : | 150347-59-4 |
| Formula: | C ₂₉ H ₁₉ NO ₁₁ |
| Molecular Weight: | 557.46 |
| 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 | CFSE (CFDA-SE) is a fluorescent dye with cell membrane permeability. CFSE irreversibly binds to intracellular proteins in living cells and is used for the detection of cell proliferation. The labeled cells fluoresce in green color with excitation wavelength of 488 nm and emission wavelength of 518 nm. |
| Targets(IC50) | Others |
| In vitro | <p>METHODS: 1 mL of cells and CFSE (5 μM in 110 μL PBS) were flipped and mixed to label the cells. CFSE-labeled CD8+OT-I T cells were cultured with dendritic cells pulsed with varying amounts of OVA for 3 days, and CFSE profiles were examined using Flow Cytometry.</p> <p>RESULTS: CD8+ T cells divided 1-3 times according to the CFSE dilution peak, and more T cells divided at higher antigen concentrations. [1]</p> <p>METHODS: Human erythroleukemia cell line K562, mouse lymphoma cell line YAC-1, human breast cancer cell line MCF-7, and human melanoma cell line A375 were treated with CFSE (1-10 μM) for 1-6 h. Cell death was detected by Flow Cytometry.</p> <p>RESULTS: CFSE was non-toxic to the cells, as the cell death rate due to CFSE labeling was less than 5%. [2]</p> |
| In vivo | <p>METHODS: CFSE-labeled CD8+OT-I T cells were injected intravenously into the tail of C56BL/6J mice, followed by intravenous injection of OVA (20 μg), and CFSE profiles were measured three days later.</p> <p>RESULTS: Most of the cells fell within 7 CFSE peaks, indicating that the cells had undergone up to 6 divisions. [1]</p> <p>METHODS: To label thymocytes in vivo, CFSE (10 μM) was injected into the thymic lobes of anesthetized C56BL/6 mice.</p> <p>RESULTS: CFSE labeled a representative sample of all thymocyte subpopulations and these cells migrated to peripheral lymphoid organs at a rate of approximately 2-3 x 10⁶ cells/day. They enter the lymph nodes on day 1 post-injection and remain there for at least 21 days, while turnover is faster in the spleen. [3]</p> |
| Cell Research | <p>Instructions:</p> <p>I. Solution preparation</p> <p>1. Preparation of mother solution: Take 1 mg CFDA-SE and dissolve it in 0.1794 mL DMSO to obtain 10 mM CFDA-SE mother solution.</p> <p>Note: The mother solution is recommended to be stored at -20°C or -80°C away from</p> |

light to avoid repeated freezing and thawing.

2. Preparation of working solution: Use pure DMEM to dilute the mother solution, usually with a concentration of 1-10 μ M.

Note: Please adjust the concentration of CFDA-SE working solution according to actual conditions.

II. Cell staining

1. Cell type:

1) Suspended cells: Centrifuge at 4°C, 1000 g for 3-5 minutes, discard the supernatant. Wash with PBS twice, 5 minutes each time.

2) Adherent cells: Discard the cell culture medium, add trypsin to dissociate the cells, and make a single cell suspension. Centrifuge at 4°C, 1000 g for 3-5 minutes, discard the supernatant. Wash with PBS twice, 5 minutes each time.

3. Add 1 mL CFDA-SE working solution and incubate at room temperature for 30 minutes.

4. Centrifuge at 400 g for 3-4 minutes at 4°C

5. Wash twice with PBS, 5 minutes each time.

6. Resuspend cells in serum-free cell culture medium or PBS and detect by fluorescence microscopy or flow cytometry.

Solubility Information

| | |
|------------|--|
| Solubility | Ethanol: < 1 mg/mL (insoluble or slightly soluble), DMSO: 93 mg/mL (166.83 mM), Sonication is recommended. H ₂ O: < 1 mg/mL (insoluble or slightly soluble), (< 1 mg/ml refers to the product slightly soluble or insoluble) |
|------------|--|

Preparing Stock Solutions

| | 1mg | 5mg | 10mg |
|-------|-----------|-----------|------------|
| 1 mM | 1.7939 mL | 8.9693 mL | 17.9385 mL |
| 5 mM | 0.3588 mL | 1.7939 mL | 3.5877 mL |
| 10 mM | 0.1794 mL | 0.8969 mL | 1.7939 mL |
| 50 mM | 0.0359 mL | 0.1794 mL | 0.3588 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

- Quah BJ, et al. The use of carboxyfluorescein diacetate succinimidyl ester (CFSE) to monitor lymphocyte proliferation. *J Vis Exp*. 2010 Oct 12;(44):2259.
- Guo E, Xiao R, Wu Y, et al. WEE1 inhibition induces anti-tumor immunity by activating ERV and the dsRNA pathway. *Journal of Experimental Medicine*. 2021, 219(1): e20210789.
- Li P, Xie Y, Wang J, et al. Gene engineered exosome reverses T cell exhaustion in cancer immunotherapy. *Bioactive Materials*. 2024, 34: 466-481.
- Qiu C, Peng B, Xiao C, et al. A novel method for identifying SARS-CoV-2 infection mutants via an epitope-specific CD8+ T cell test. *Biosafety and Health*. 2024
- Zhang Y, Zhang J X, Xiao L X, et al. The Synergistic Effect of Huangqi Gegen Decoction on Thrombosis relates to the Astragalus Polysaccharide-improved Oral Delivery of Puerarin. *Journal of Ethnopharmacology*. 2024: 118622.
- Chen L, Tang J, Chang Y, et al. SMURF1 leads to the β -catenin signaling-mediated progression of esophageal squamous carcinoma by losing PATZ1-induced CCNG2 transcription. *Biochemical Pharmacology*. 2024: 116688.
- Lan T, Gao F, Cai Y, et al. The protein circPETH-147aa regulates metabolic reprogramming in hepatocellular carcinoma cells to remodel immunosuppressive microenvironment. *Nature Communications*. 2025, 16(1): 333.
- Liao Y, Huang J, Liu P, et al. Downregulation of LNMAS orchestrates partial EMT and immune escape from macrophage phagocytosis to promote lymph node metastasis of cervical cancer. *Oncogene*. 2022: 1-13.
- Wang XQ, et al. Carboxyfluorescein diacetate succinimidyl ester fluorescent dye for cell labeling. *Acta Biochim Biophys Sin (Shanghai)*. 2005 Jun;37(6):379-85.
- Graziano M, et al. The fate of thymocytes labeled in vivo with CFSE. *Exp Cell Res*. 1998 Apr 10;240(1):75-85.
- Xiao C, Su J, Zhang C, et al. Effectiveness of Booster Doses of the SARS-CoV-2 Inactivated Vaccine KCONVAC against the Mutant Strains. *Viruses*. 2022, 14(9): 2016.
- Li M, Qin M, Song G, et al. A biomimetic antitumor nanovaccine based on biocompatible calcium pyrophosphate and tumor cell membrane antigens[J]. *Asian Journal of Pharmaceutical Sciences*. 2020
- Xiao C, Mao L, Wang Z, et al. SARS-CoV-2 variant B. 1.1. 7 caused HLA-A2+ CD8+ T cell epitope mutations for impaired cellular immune response. *Iscience*. 2022 Mar 18;25(3):103934. doi: 10.1016/j.isci.2022.103934. Epub 2022 Feb 17.
- Deng J, Pan J, Qiu M, et al. Identification of HLA-A2 restricted CD8+ T cell epitopes in SARS-CoV-2 structural proteins. *Journal of Leukocyte Biology*. 2021
- Jablonski S, Mou H, Otsuka Y, et al. Identification of Potent Small Molecule Inhibitors of SARS-CoV-2 Entry. *SLAS Discovery*. 2021
- Zhao G, Tong Y, Xu J, et al. Jing-Fang powder ethyl acetate extracts attenuate atopic dermatitis by modulating T-cell activity. *Molecular Immunology*. 2023, 160: 133-149.
- Xiao C, Ren Z, Zhang B, et al. Insufficient epitope-specific T cell clones are responsible for impaired cellular immunity to inactivated SARS-CoV-2 vaccine in older adults. *Nature Aging*. 2023: 1-18.
- Yang L, Wu J, Mei G, et al. Corydalis Saxicola Bunting Total Alkaloid Eliminates Porphyromonas gingivalis strain 33277 Internalized into Macrophages by Inhibition of TLR2. *Microbes and Infection*. 2023: 105244.

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