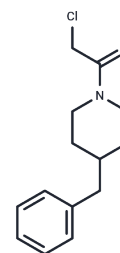


GSTO1-IN-3

Chemical Properties

| | |
|-------------------|---|
| CAS No. : | 158890-32-5 |
| Formula: | C ₁₄ H ₁₈ ClNO |
| Molecular Weight: | 251.75 |
| 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 | GSTO1-IN-3 is a potent inhibitor of GSTO1-1, with an IC ₅₀ value of 0.11 μM, and exhibits selectivity towards GSTO2-2, GSTA1-1, and GSTP1-1 (IC ₅₀ > 100 μM). It enhances the cytotoxic effects of Cisplatin on human breast cancer cells and inhibits IL-1β release in mouse bone marrow-derived macrophages (BMDM). Additionally, GSTO1-IN-3 reduces inflammation in mice and is applicable for research related to breast cancer and inflammation. |
| In vitro | GSTO1-IN-3 (compound 5C-1) demonstrates favorable in vitro safety, as it shows no thiol reactivity in ALARM NMR experiments and lacks mutagenicity in Ames tests with *Salmonella typhimurium* strains TA98 and TA100 at concentrations up to 2 mg/mL. It exhibits greater stability in human liver microsomes compared to mouse microsomes. GSTO1-IN-3 forms a covalent bond with the Cys32 of GSTO1-1, with an inhibition efficiency constant (k _{on} /K) of 3.4 × 10 ⁴ M ⁻¹ s ⁻¹ . At a concentration of 5 μM, GSTO1-IN-3 reduces IL-1β release by 59.2% in mouse bone marrow-derived macrophages (BMDM). Additionally, at 5-50 μM for 24 hours, it exhibits a synergistic effect with Cisplatin, significantly enhancing cytotoxicity in a concentration-dependent manner against human breast cancer MDA-MB-231 cells. |
| In vivo | GSTO1-IN-3, when administered intraperitoneally at a dose of 30 mg/kg, 30 minutes prior to LPS injection, can mitigate the inflammation induced by lipopolysaccharide (LPS) in mice. |

Preparing Stock Solutions

| | 1mg | 5mg | 10mg |
|-------|------------|------------|-------------|
| 1 mM | 3.9722 mL | 19.861 mL | 39.7219 mL |
| 5 mM | 0.7944 mL | 3.9722 mL | 7.9444 mL |
| 10 mM | 0.3972 mL | 1.9861 mL | 3.9722 mL |
| 50 mM | 0.0794 mL | 0.3972 mL | 0.7944 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.

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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