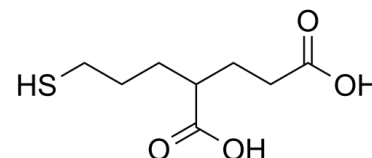


Data Sheet

| | |
|---------------------------|---|
| Product Name: | 2-MPPA |
| Cat. No.: | CS-0027694 |
| CAS No.: | 254737-29-6 |
| Molecular Formula: | C ₈ H ₁₄ O ₄ S |
| Molecular Weight: | 206.26 |
| Target: | Carboxypeptidase |
| Pathway: | Metabolic Enzyme/Protease |
| Solubility: | DMSO : 100 mg/mL (484.82 mM; Need ultrasonic); Methanol : 250 mg/mL (1212.06 mM; Need ultrasonic) |



BIOLOGICAL ACTIVITY:

2-MPPA (GPI-5693) is an orally active and selective **glutamate carboxypeptidase II (GCP II; PSMA)** inhibitor with an **IC₅₀** of 90 nM [1][2]. **IC₅₀ & Target:** IC₅₀: 90 nM (GCP II)[2] **In Vivo:** 2-MPPA (10, 30 or 100 mg/kg) significantly attenuates Dizocilpine (HY-15084B) (0.1 mg/kg)-induced prepulse inhibition (PPI) deficits in mice, in a dose dependent manner. The efficacy of 2-MPPA on dizocilpine-induced PPI deficits is significantly antagonized by pretreatment with the selective group II metabotropic glutamate receptor (mGluR) antagonist LY341495 (HY-70059) (1.0 mg/kg)[1].

2-MPPA (30 mg/kg) significantly prevents the deficit in SNCV induced by both 5 and 25 mg/kg taxol by 96.3 ± 4.4% and 98.3 ± 11.6% respectively[2].

2-MPPA (10 mg/kg) inhibits tumor growth in the high expressor PSMA model CWR22RS by 70%[2].

References:

[1]. Takatsu Y, et al. Orally active glutamate carboxypeptidase II inhibitor 2-MPPA attenuates dizocilpine-induced prepulse inhibition deficits in mice. *Brain Res.* 2011 Jan 31;1371:82-6.

[2]. She Y, et al. 2-MPPA, a selective inhibitor of PSMA, delays prostate cancer growth and attenuates taxol-induced neuropathy in mice. *Journal of Clinical Oncology*, 2005, 23(16_suppl): 8054-8054.

CAIndexNames:

Pentanedioic acid, 2-(3-mercaptopropyl)-

SMILES:

O=C(O)C(CCCS)CCC(O)=O

Caution: Product has not been fully validated for medical applications. For research use only.

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