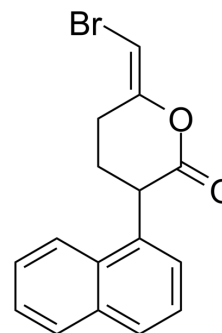


Bromo-enol lactone

| | | | |
|---------------------------|--|-------|----------|
| Cat. No.: | HY-107411 | | |
| CAS No.: | 88070-98-8 | | |
| Molecular Formula: | C ₁₆ H ₁₃ BrO ₂ | | |
| Molecular Weight: | 317.18 | | |
| Target: | Phospholipase | | |
| Pathway: | Metabolic Enzyme/Protease | | |
| Storage: | Powder | -20°C | 3 years |
| | In solvent | -80°C | 6 months |
| | | -20°C | 1 month |



SOLVENT & SOLUBILITY

In Vitro

DMSO : 50 mg/mL (157.64 mM; Need ultrasonic)

| Preparing Stock Solutions | Solvent Concentration | Mass | | |
|---------------------------|-----------------------|-----------|------------|------------|
| | | 1 mg | 5 mg | 10 mg |
| | 1 mM | 3.1528 mL | 15.7639 mL | 31.5278 mL |
| | 5 mM | 0.6306 mL | 3.1528 mL | 6.3056 mL |
| | 10 mM | 0.3153 mL | 1.5764 mL | 3.1528 mL |

Please refer to the solubility information to select the appropriate solvent.

In Vivo

- Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline
Solubility: 2.5 mg/mL (7.88 mM); Suspended solution; Need ultrasonic
- Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline)
Solubility: 2.5 mg/mL (7.88 mM); Clear solution; Need ultrasonic
- Add each solvent one by one: 10% DMSO >> 90% corn oil
Solubility: ≥ 2.5 mg/mL (7.88 mM); Clear solution

BIOLOGICAL ACTIVITY

Description

Bromo-enol lactone ((6E)-Bromo-enol lactone) is a suicide-based irreversible, selective, potent inhibitor of calcium-independent phospholipase A₂ (iPLA₂β) with an IC₅₀ value of approximately 7 μM, which inhibits antigen-stimulated mast cell exocytosis without blocking Ca²⁺ influx^{[1][2]}.

IC₅₀ & Target

PLA2
7 μM (IC₅₀)

In Vitro

In RBL 2H3 and bone marrow-derived mast cells (BMMCs), Ca²⁺ entry is critical for exocytosis. Bromo-enol lactone inhibits

exocytosis when stimulated using a Ca^{2+} ionophore A23187, which passively transports Ca^{2+} down a concentration gradient and also in permeabilised mast cells where Ca^{2+} entry is no longer relevant. Moreover, Bromoenol lactone has only a minor effect on antigen- or thapsigargin-stimulated Ca^{2+} signalling, both the release from internal stores and sustained elevation due to Ca^{2+} influx^[1].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

CUSTOMER VALIDATION

- Genomics. 2022 Sep 5;114(5):110479.

See more customer validations on www.MedChemExpress.com

REFERENCES

[1]. Fensome-Green A, et al. Bromoenol lactone, an inhibitor of Group V1A calcium-independent phospholipase A2 inhibits antigen-stimulated mast cell exocytosis without blocking Ca^{2+} influx. *Cell Calcium*. 2007 Feb;41(2):145-53.

[2]. Takuma T, et al. Role of Ca^{2+} -independent phospholipase A2 in exocytosis of amylase from parotid acinar cells. *J Biochem*. 1997 Jun;121(6):1018-24.

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

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