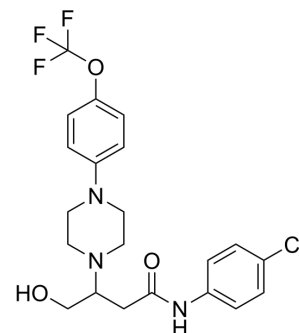


VBIT-4

| | | | |
|---------------------------|--|-------|----------|
| Cat. No.: | HY-129122 | | |
| CAS No.: | 2086257-77-2 | | |
| Molecular Formula: | C ₂₁ H ₂₃ ClF ₃ N ₃ O ₃ | | |
| Molecular Weight: | 457.87 | | |
| Target: | VDAC | | |
| Pathway: | Membrane Transporter/Ion Channel | | |
| Storage: | Powder | -20°C | 3 years |
| | In solvent | -80°C | 6 months |
| | | -20°C | 1 month |



SOLVENT & SOLUBILITY

In Vitro

DMSO : 125 mg/mL (273.00 mM; Need ultrasonic)
 H₂O : 1 mg/mL (2.18 mM; ultrasonic and adjust pH to 3 with HCl)

| Concentration | Solvent | Mass | 1 mg | 5 mg | 10 mg |
|---------------------------|---------|------|-----------|------------|------------|
| | | | 1 mg | 5 mg | 10 mg |
| Preparing Stock Solutions | 1 mM | | 2.1840 mL | 10.9201 mL | 21.8403 mL |
| | 5 mM | | 0.4368 mL | 2.1840 mL | 4.3681 mL |
| | 10 mM | | 0.2184 mL | 1.0920 mL | 2.1840 mL |

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

In Vivo

- Add each solvent one by one: 10% DMSO >> 90% saline
 Solubility: 5 mg/mL (10.92 mM); Suspended solution; Need ultrasonic
- Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline)
 Solubility: ≥ 2.08 mg/mL (4.54 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% corn oil
 Solubility: ≥ 2.08 mg/mL (4.54 mM); Clear solution

BIOLOGICAL ACTIVITY

Description

VBIT-4 is an inhibitor of voltage-dependent anion channel 1 (VDAC1) oligomerization with a binding affinity (K_d) of 17 μM. VBIT-4, as an apoptosis inhibitor, can be used for therapeutic purposes in apoptosis-associated disorders, such as neurodegenerative and cardiovascular diseases^[1].

IC₅₀ & Target

Kd: 17 μM (VDAC1)^[1]
 Apoptosis^[1]

In Vitro

VBIT-4 targets the mitochondrial protein VDAC1, inhibiting apoptosis and protecting against mitochondria dysfunction. VBIT-4 (0.1-10 μM) inhibits VDAC1 oligomerization, Cyto c release from mitochondria and apoptosis in HEK-293 cells with IC_{50} s of $1.9\pm 0.08 \mu\text{M}$, $1.8\pm 0.24 \mu\text{M}$, and $2.9\pm 0.12 \mu\text{M}$, respectively^[1].

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

Apoptosis Analysis^[1]

| | |
|------------------|---|
| Cell Line: | HEK-293 cells |
| Concentration: | 0.1, 1, and 10 μM |
| Incubation Time: | |
| Result: | Inhibited apoptosis with an IC_{50} of $2.9\pm 0.12 \mu\text{M}$. |

CUSTOMER VALIDATION

- Cell. 2020 Oct 29;183(3):636-649.e18.
- Immunity. 2022 Aug 9;55(8):1370-1385.e8.
- Nat Commun. 2023 Feb 16;14(1):872.
- Autophagy. 2021 Jan 19;1-17.
- Sci Rep. 2023 Nov 17;13(1):20126.

See more customer validations on www.MedChemExpress.com

REFERENCES

[1]. Ben-Hail D, et al. Novel Compounds Targeting the Mitochondrial Protein VDAC1 Inhibit Apoptosis and Protect against Mitochondrial Dysfunction. J Biol Chem. 2016 Nov 25;291(48):24986-25003.

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

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