Proteins

Product Data Sheet

(S)-Sabutoclax

Cat. No.: HY-15191B CAS No.: 1228178-73-1 Molecular Formula: $C_{42}H_{42}N_{2}O_{8}S$ Molecular Weight: 732.84 Target: **Bcl-2 Family**

Pathway: **Apoptosis**

Powder Storage: -20°C 3 years In solvent -80°C 6 months

-20°C 1 month

SOLVENT & SOLUBILITY

In Vitro

DMSO: 30 mg/mL (40.94 mM; Need ultrasonic)

H₂O: < 0.1 mg/mL (insoluble)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	1.3646 mL	6.8228 mL	13.6455 mL
	5 mM	0.2729 mL	1.3646 mL	2.7291 mL
	10 mM	0.1365 mL	0.6823 mL	1.3646 mL

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

In Vivo

1. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: 3 mg/mL (4.09 mM); Suspended solution; Need ultrasonic

BIOLOGICAL ACTIVITY

Description

(S)-Sabutoclax ((S)-BI-97C1), an optically pure apogossypol derivative, is pan-active inhibitor of antiapoptotic B-cell lymphoma/leukemia-2 (Bcl-2) family proteins. (S)-Sabutoclax (Compound II) inhibits the binding of BH3 peptides to Bcl-XL, Bcl-2, Mcl-1, and Bfl-1 with IC₅₀ values of 0.31, 0.32, 0.20, and 0.62 μM, respectively. (S)-Sabutoclax also potently inhibits cell growth of human prostate cancer, lung cancer, and lymphoma cell lines with EC $_{50}$ values of 0.13, 0.56, and 0.049 μ M, respectively. (S)-Sabutoclax can be used for the research of apoptosis-based therapies against cancer^[1].

IC₅₀ & Target

IC50: 0.31 μM (Bcl-XL), 0.32 μM (Bcl-2), 0.20 μM (Mcl-1), 0.62 μM (Bfl-1); EC50: 0.13 (human prostate cancer cell lines), 0.56 (lung cancer cell lines), 0.049 μM (lymphoma cell lines)^[1].

REFERENCES

1]. Wei J, et al. BI-97C1, an opt 2010 May 27;53(10):4166-76.	ically pure Apogossypol deri	ivative as pan-active inhibitor of a	ntiapoptotic B-cell lymphoma/leukemi	a-2 (Bcl-2) family proteins. J Med Chem.
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