Product Data Sheet

STM2457

Cat. No.:HY-134836CAS No.:2499663-01-1Molecular Formula: $C_{25}H_{28}N_6O_2$ Molecular Weight:444.53Target:ApoptosisPathway:Apoptosis

Storage: Powder -20°C

20°C 3 years 4°C 2 years

In solvent -80°C 6 months

-20°C 1 month

SOLVENT & SOLUBILITY

In Vitro

DMSO: 50 mg/mL (112.48 mM; ultrasonic and warming and heat to 60°C)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.2496 mL	11.2478 mL	22.4957 mL
	5 mM	0.4499 mL	2.2496 mL	4.4991 mL
	10 mM	0.2250 mL	1.1248 mL	2.2496 mL

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

In Vivo

- 1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.08 mg/mL (4.68 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE- β -CD in saline) Solubility: \ge 2.08 mg/mL (4.68 mM); Clear solution
- 3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.08 mg/mL (4.68 mM); Clear solution

BIOLOGICAL ACTIVITY

Description	STM2457 is a first-in-class, highly potent, selective and orally active METTL3 inhibitor with an IC ₅₀ of 16.9 nM. STM2457 can be used for the research of acute myeloid leukaemia (AML) ^{[1][2]} .
In Vitro	STM2457 (Compound 72) inhibits MOLM13 cells proliferation with an IC $_{50}$ of 8.699 μ M $^{[1]}$. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

CUSTOMER VALIDATION

- Nat Biotechnol. 2023 Mar;41(3):355-366.
- Gastroenterology. 2022 Jun 11;S0016-5085(22)00629-1.
- Cancer Commun (Lond). 2022 Mar 9.
- Cell Rep Med. 2023 Aug 15;4(8):101144.
- J Hazard Mater. 2023 Jun 19;458:131891.

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REFERENCES

[1]. Wesley Peter Blackaby, et al. Mettl3 inhibitory compounds. WO2020201773A1.

[2]. Eliza Yankova, et al. Small molecule inhibition of METTL3 as a strategy against myeloid leukaemia. Nature. 2021 Apr 26.

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

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