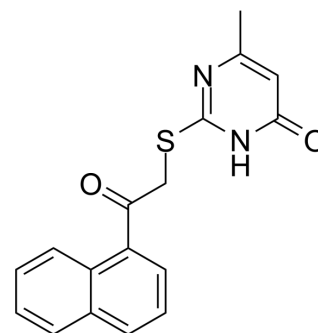


## I3MT-3

Cat. No.:	HY-128206	
CAS No.:	459420-09-8	
Molecular Formula:	C <sub>17</sub> H <sub>14</sub> N <sub>2</sub> O <sub>2</sub> S	
Molecular Weight:	310.37	
Target:	Hippo (MST)	
Pathway:	Stem Cell/Wnt	
Storage:	Powder	-20°C 3 years
	In solvent	-80°C 6 months
		-20°C 1 month



### SOLVENT & SOLUBILITY

In Vitro	DMSO : 125 mg/mL (402.75 mM; Need ultrasonic)					
	Preparing Stock Solutions	Solvent Concentration	Mass	1 mg	5 mg	10 mg
			1 mM	3.2220 mL	16.1098 mL	32.2196 mL
			5 mM	0.6444 mL	3.2220 mL	6.4439 mL
			10 mM	0.3222 mL	1.6110 mL	3.2220 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: 2.08 mg/mL (6.70 mM); Suspended solution; Need ultrasonic					
	2. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.08 mg/mL (6.70 mM); Clear solution					

### BIOLOGICAL ACTIVITY

Description	I3MT-3 (HMPSNE) is a potent, selective, and cell-membrane permeable inhibitor of 3-Mercaptopyruvate sulfurtransferase (3MST) (IC <sub>50</sub> =2.7 μM). I3MT-3 is inactive for other H <sub>2</sub> S/sulfane sulfur-producing enzymes. I3MT-3 targets a persulfurated cysteine residue located in the active site of 3MST <sup>[1]</sup> .
IC <sub>50</sub> & Target	IC <sub>50</sub> : 2.7 μM (3-Mercaptopyruvate sulfurtransferase (3MST)) <sup>[1]</sup>
In Vitro	I3MT-3 (1 μM) is selective for 3MST and shows a high inhibitory activity (80–90%) even at 10 μM in cell lysate of 3MST-overexpressing HEK293 cells. Besides, it is almost inactive towards the other two H <sub>2</sub> S-producing enzymes even at 100 μM <sup>[1]</sup> . I3MT-3 (1 μM) shows a high selectivity for 3MST, it completely suppresses the 3MST activity in COS7 cells living cells <sup>[1]</sup> . I3MT-3 produces a concentration-dependent inhibition of the AzMC (the fluorescent H <sub>2</sub> S probe) signal when incubated with purified human recombinant enzyme, the inhibition of the catalytic activity of 3-MST produces a concentration-dependent

inhibition of H<sub>2</sub>S production with an IC<sub>50</sub> of 13.6 μM<sup>[1]</sup>.

I3MT-3 shows a dose-dependent inhibition of 3-MST activity from CT26 homogenates, which contain the murine form of the enzyme. The IC<sub>50</sub> of HMPSNE for murine 3-MST is calculated as 2.3 μM with a concentration-dependent decrease of AzMC fluorescence<sup>[1]</sup>.

I3MT-3 (10 μM-100 μM; after 3 h probe AzMC) causes a partial inhibition of the signal, while at 100 μM, HMPSNE causes a complete inhibition of the AzMC-guided H<sub>2</sub>S fluorescence at 100 μM. Additionally, HMPSNE is capable of inhibiting its target in situ in CT26 cells (with an IC<sub>50</sub> of approximately 30 μM)<sup>[2]</sup>.

I3MT-3 (0-300 μM; 5-50 hours) does not enhance MTT conversion at 10 μM, while at 100 and 300 μM it produces an inhibitory response, without increasing the LDH signal, i.e., without inducing any detectable degree of cell necrosis. It also produces a decreased oxygen consumption rate (OCR) profiles in CT26 cells<sup>[2]</sup>.

I3MT-3 (0-300 μM; 48 hours) inhibits CT26 cells proliferate with increasing concentrations of I3MT-3. Confluence of cells treated with HMPSNE is recorded each hour for 48 h by the IncuCyte method<sup>[2]</sup>.

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

Cell Proliferation Assay<sup>[2]</sup>

Cell Line:	CT26 cells
Concentration:	0 μM; 10 μM; 30 μM; 100 μM; 300 μM
Incubation Time:	48 hours
Result:	Slowed down proliferation of CT26 cells.

## CUSTOMER VALIDATION

- Cell Death Dis. 2022 Oct 30;13(10):913.
- Biochim Biophys Acta Mol Basis Dis. 2023 Dec 7:166983.
- J Biol Chem. 2023 Apr 13;104710.

See more customer validations on [www.MedChemExpress.com](http://www.MedChemExpress.com)

## REFERENCES

[1]. Kenjiro Hanaoka, et al. Discovery and Mechanistic Characterization of Selective Inhibitors of H<sub>2</sub>S-producing Enzyme: 3-Mercaptopyruvate Sulfurtransferase (3MST) Targeting Active-site Cysteine Persulfide. Sci Rep. 2017 Jan 12;7:40227

[2]. Fiona Augsburger, et al. Role of 3-Mercaptopyruvate Sulfurtransferase in the Regulation of Proliferation, Migration, and Bioenergetics in Murine Colon Cancer Cells. Biomolecules. 2020 Mar 13;10(3):447.

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

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: [tech@MedChemExpress.com](mailto:tech@MedChemExpress.com)

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA