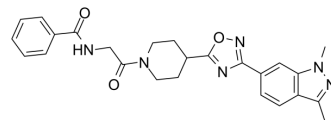


YTX-465

Cat. No.:	HY-124751
CAS No.:	2225824-53-1
Molecular Formula:	C ₂₅ H ₂₆ N ₆ O ₃
Molecular Weight:	458.51
Target:	Stearoyl-CoA Desaturase (SCD)
Pathway:	Metabolic Enzyme/Protease
Storage:	4°C, protect from light * In solvent : -80°C, 6 months; -20°C, 1 month (protect from light)



SOLVENT & SOLUBILITY

In Vitro	DMSO : 100 mg/mL (218.10 mM; Need ultrasonic)				
		Solvent Concentration	Mass		
	Preparing Stock Solutions		1 mg	5 mg	10 mg
		1 mM	2.1810 mL	10.9049 mL	21.8098 mL
		5 mM	0.4362 mL	2.1810 mL	4.3620 mL
	10 mM	0.2181 mL	1.0905 mL	2.1810 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: ≥ 2.5 mg/mL (5.45 mM); Clear solution				
	2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (5.45 mM); Clear solution				
	3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (5.45 mM); Clear solution				

BIOLOGICAL ACTIVITY

Description	YTX-465 is a stearoyl-CoA desaturase (Ole1/SCD) inhibitor. YTX-465 inhibits Ole1 and SCD1 with IC ₅₀ s of 0.039 μM and 30.4 μM, respectively. YTX-465 can be used in the research of Parkinson's disease and other synucleinopathies ^[1] .	
IC ₅₀ & Target	Ole1 0.039 μM (IC ₅₀)	SCD1 30.4 μM (IC ₅₀)
In Vitro	YTX-465 (1-10000 nM) has an EC ₅₀ value of 0.013 μM for α-synuclein (α-syn) toxicity rescue ^[1] . YTX-465 (0.05 μM; 0-2 days) rescue the growth of α-Syn-expressing yeast ^[1] . YTX-465 (0, 10, 40, 160, 640, 2500 nM; 4h) increases the level of Ole1 protein in wild-type yeast expressing OLE1 in a	

concentration-dependent manner, that indicate YTX-465 induce a negative feedback loop^[1].

YTX-465 (0, 0.03, 0.09, 0.27, 0.81 μM ; 6 hours) reduces fatty desaturation in a concentration-dependent manner in wild-type yeast, with a 50% reduction in desaturation at 0.03 μM ^[1].

YTX-465 (0.25 μM ; 8 hours) decreases the desaturation index (DI) for all major classes of membrane phospholipids in wild-type yeast^[1].

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

REFERENCES

[1]. Vincent BM, et al. Inhibiting Stearoyl-CoA Desaturase Ameliorates α -Synuclein Cytotoxicity. Cell Rep. 2018 Dec 4;25(10):2742-2754.e31.

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

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