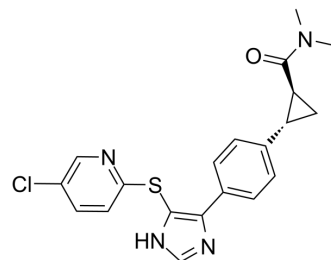


FAAH-IN-1

Cat. No.:	HY-111389		
CAS No.:	1242441-47-9		
Molecular Formula:	C ₂₀ H ₁₉ ClN ₄ OS		
Molecular Weight:	398.91		
Target:	FAAH; Autophagy		
Pathway:	Metabolic Enzyme/Protease; Neuronal Signaling; Autophagy		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



SOLVENT & SOLUBILITY

In Vitro	DMSO : 100 mg/mL (250.68 mM; Need ultrasonic)			
		Solvent Concentration	Mass	
			1 mg	5 mg
			10 mg	
Preparing Stock Solutions	1 mM	2.5068 mL	12.5342 mL	25.0683 mL
	5 mM	0.5014 mL	2.5068 mL	5.0137 mL
	10 mM	0.2507 mL	1.2534 mL	2.5068 mL
Please refer to the solubility information to select the appropriate solvent.				
In Vivo	<ol style="list-style-type: none"> Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.5 mg/mL (6.27 mM); Clear solution Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (6.27 mM); Clear solution Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (6.27 mM); Clear solution 			

BIOLOGICAL ACTIVITY

Description	FAAH-IN-1 is a fatty acid amide hydrolase (FAAH) inhibitor, with IC ₅₀ s of 145 nM and 650 nM for rat and human FAAH, respectively.
IC₅₀ & Target	IC ₅₀ : 145 nM (Rat FAAH), 650 nM (Human FAAH) ^[1]
In Vitro	FAAH-IN-1 (Compound 8) is a fatty acid amide hydrolase (FAAH) inhibitor, with IC ₅₀ s of 145 nM and 650 nM for rat and human FAAH, respectively ^[1] .

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

REFERENCES

[1]. Liu P, et al. Discovery of MK-3168: A PET Tracer for Imaging Brain Fatty Acid Amide Hydrolase. ACS Med Chem Lett. 2013 Apr 20;4(6):509-13.

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