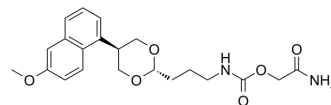


SSR411298

Cat. No.:	HY-123863		
CAS No.:	666860-59-9		
Molecular Formula:	C ₂₁ H ₂₆ N ₂ O ₆		
Molecular Weight:	402.44		
Target:	FAAH		
Pathway:	Metabolic Enzyme/Protease; Neuronal Signaling		
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 (248.48 mM; Need ultrasonic)

Concentration	Mass		
	1 mg	5 mg	10 mg
1 mM	2.4848 mL	12.4242 mL	24.8484 mL
5 mM	0.4970 mL	2.4848 mL	4.9697 mL
10 mM	0.2485 mL	1.2424 mL	2.4848 mL

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

BIOLOGICAL ACTIVITY

Description

SSR411298 is an orally active, selective and reversible fatty acid amide hydrolase (FAAH) inhibitor. SSR411298 has the potential for post-traumatic stress disorder research^[1].

In Vitro

SSR411298 (0.1, 1, 10, 100, 1000, 10000 nM) inhibits mouse brain FAAH in a concentration-dependent manner^[1]. SSR411298 produces maximal inhibition of FAAH between 0.3 and 30 mg/kg in the ex-vivo experiment^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

In Vivo

SSR411298 (0.3, 1, 3 mg/kg; p.o.; twice a day during 2 days; 60 minutes before pretest, immediately after, and 60 minutes before the second session of testing) produces robust antidepressant-like activity in the rat forced-swimming test and in the mouse chronic mild stress model, restoring notably the development of inadequate coping responses to chronic stress in male Wistar rats weighing 235-290 g^[1]. SSR411298 (0.3, 1, 3, 10 mg/kg; po; single dose; two hours prior to sacrifice) produces a significant global increase of tissular AEA, PEA and OEA levels in the hippocampus, but not of 2-AG levels^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Guy Griebel, et al. The selective reversible FAAH inhibitor, SSR411298, restores the development of maladaptive behaviors to acute and chronic stress in rodents. Sci Rep. 2018 Feb 5;8(1):2416.

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