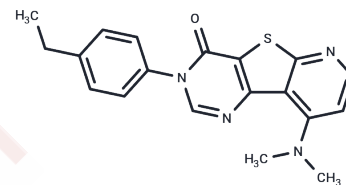


A-794282

Chemical Properties

CAS No. :	869802-44-8
Formula:	C ₁₉ H ₁₈ N ₄ O ₅
Molecular Weight:	350.44
Storage:	Store at low temperature Powder: -20°C for 3 years In solvent: -80°C for 1 year <small>Actual storage temperature shall be subject to the COA.</small>



Biological Description

Description	A-794282 is a selective and potent mGlu1 receptor antagonist with analgesic activity and may be useful in the study of neurologic disorders.
Targets(IC50)	GluR
In vivo	METHODS: In this study, the effects of A-794282, a potent and selective mGlu1 receptor antagonist (ip) were evaluated in a rat skin incision-induced postoperative pain model. Postoperative pain was examined 2 h after surgery, using the difference in weight bearing between the injured and uninjured paws as a measure of spontaneous pain. RESULTS: In this model, A-794282 significantly attenuated spontaneous postoperative pain behaviors induced by A-794282, with an ED50 of 50 µmol/kg, respectively. [2]

Preparing Stock Solutions

	1mg	5mg	10mg
1 mM	2.8536 mL	14.2678 mL	28.5356 mL
5 mM	0.5707 mL	2.8536 mL	5.7071 mL
10 mM	0.2854 mL	1.4268 mL	2.8536 mL
50 mM	0.0571 mL	0.2854 mL	0.5707 mL

Please select the appropriate solvent to prepare the stock solution, according to the solubility of the product in different solvents. Please use it as soon as possible.

Note: The dilution table applies only to solid products. For liquid products, please calculate the stock solution based on the stated concentration and/or density.

Reference

Yang YL,et al. [Research progress of selective mGluR1 antagonists]. Yao Xue Xue Bao. 2011 Oct;46(10):1167-72. Review. Chinese.

Zhu CZ,et al. Analgesic activity of metabotropic glutamate receptor 1 antagonists on spontaneous post-operative pain in rats. Eur J Pharmacol. 2008 Feb 12;580(3):314-21.

Zheng GZ,et al. Structure-activity relationship of triazafluorenone derivatives as potent and selective mGluR1 antagonists. J Med Chem. 2005 Nov 17;48(23):7374-88.

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