
Product Data Sheet

Product Name: LY487379

Cat. No.: GC64319

Chemical Properties

Cas. No. 353231-17-1

Formula C₂₁H₁₉F₃N₂O₄S

M.Wt 452.45

Solubility DMSO : 100 mg/mL (221.02 mM; Need ultrasonic) Storage Store at -20°C

General tips For obtaining a higher solubility , please warm the tube at 37 °C and shake it in the ultrasonic bath for a while. Stock solution can be stored below -20°C for several months.

Shipping Condition Evaluation sample solution : ship with blue ice All other available size: ship with RT , or blue ice upon request.

Structure **Background**

LY487379 is a selective human mGluR2 positive allosteric modulator (PAM). LY487379 potentiates glutamate-stimulated [³⁵S]GTPγS binding with EC₅₀ values of 1.7 μM and >10 μM for mGlu2 and mGlu3 receptors respectively. LY487379 promotes cognitive flexibility and facilitates behavioral inhibition in a rat model. LY487379 can be used for schizophrenia research[2].

LY487379 (intraperitoneal injection; 30 mg/kg; injected 30 min before the test) requires significantly fewer trials to criterion during the ED phase of the ASST in attentional set-shifting task in male Sprague-Dawley rats. But there has no significant drug effect during any other discrimination stage[1]. LY487379 hydrochloride (intraperitoneal injection; 10-30 mg/kg) induces an increase in microdialysate norepinephrine levels; the dose-effect resembled a bell-shape relationship increased. And it dose-dependently increases extracellular serotonin levels in the medial prefrontal cortex in male Sprague-Dawley rats[1].

[1]. Nikiforuk A, et al. Effects of a positive allosteric modulator of group II metabotropic glutamate receptors, LY487379, on cognitive flexibility and impulsive-like responding in rats. *J Pharmacol Exp Ther.* 2010;335(3):665-673.

[2]. Schaffhauser H, et al. Pharmacological characterization and identification of amino

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

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acids involved in the positive modulation of metabotropic glutamate receptor subtype 2.
Mol Pharmacol. 2003;64(4):798-810.

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