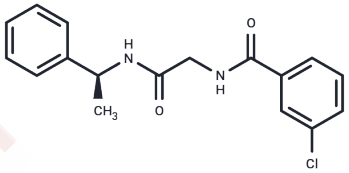
Data Sheet (Cat.No.T3119)

TargetMol

JNJ-63533054

Chemical Properties



CAS No. : 1802326-66-4

Formula: C17H17ClN2O2

Molecular Weight: 316.78

Appearance: no data available

Storage: Powder: -20°C for 3 years | In solvent: -80°C for 1 year

Biological Description

Description	JNJ-63533054 is a potent and selective agonist of hGPR139 with an EC50 = 16 nM.
Targets(IC50)	GPCR
In vitro	JNJ-63533054 specifically activates human GPR139 in the calcium mobilization (EC50 = 16 ± 6 nM) and GTPγS binding (EC50 = 17 ± 4 nM) assays. JNJ-63533054 is found to be clean of any cross reactivity as judged by an external selectivity panel of 50 known GPCRs, ion channels, and transporters as well as our own internal whole cell lead generation biology selectivity panel.
In vivo	JNJ-63533054 is found to cross the blood-brain barrier and have good drug-like properties amenable for oral dosing in rat. JNJ-63533054 exhibits good stability in both human and rat microsomes and high solubility in aqueous media, and no DDI potential was found. JNJ-63533054 also activates the rat and mouse GPR139 receptor with similar potency (rat EC50 = 63 ± 24 nM, mouse EC50 = 28 ± 7 nM).

Solubility Information

Solubility	DMSO: 45 mg/mL (142.05 mM),Sonication is recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble)

Preparing Stock Solutions

	1mg	5mg	10mg
1 mM	3.1568 mL	15.7838 mL	31.5676 mL
5 mM	0.6314 mL	3.1568 mL	6.3135 mL
10 mM	0.3157 mL	1.5784 mL	3.1568 mL
50 mM	0.0631 mL	0.3157 mL	0.6314 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.

Reference

Dvorak CA, et al. ACS Med Chem Lett. 2015 Jul 20;6(9):1015-8.
Liu C, et al. Mol Pharmacol. 2015 Nov;88(5):911-25.

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