

LY2119620

Chemical Properties

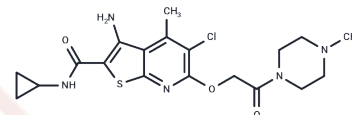
CAS No. : 886047-22-9

Formula: C₁₉H₂₄ClN₅O₃S

Molecular Weight: 437.94

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

Actual storage temperature shall be subject to the COA.



Biological Description

Description	LY2119620 is a specific, and allosteric agonist of human M2 and M4 muscarinic acetylcholine receptors.
Targets(IC50)	AChR
In vitro	LY2119620 displays modest allosteric agonism and positively modulates the functional G protein–signaling ability of an agonist at the M2/M4 receptor subtypes by placing the M2 and M4 receptors into an active G protein–bound state. LY2119620 enhances the potency of three muscarinic acetylcholine receptor agonists, ACh, Oxo-M and iperoxo. [1] [3H]LY2119620 can be used as a probe for the human M(2) and M(4) muscarinic receptor allosteric binding sites. [2]
Kinase Assay	Biochemical Assay: For IC ₅₀ determination, EI1 is serial diluted threefold in DMSO for a total of 12 concentrations, with the starting concentration at 1 μM. The reaction is incubated at room temperature for 120 min, and stopped by adding quench solution (2.5% TFA with 320 nM d4-SAH). SAH production is quantitated using an API 4000 triple quadrupole mass spectrometry with Turbulon Spray coupled with Prominence UFLC. The percentage of inhibition is normalized using positive (no inhibitor) and negative (no enzyme) controls, and IC ₅₀ calculated using PRISM. Enzymology studies of S-Adenosyl methionine (SAM) competition are performed with slight modification of reaction condition: 10 μM EI1 is used as the starting dose for serial dilution. SAM is titrated over a range between 1 μM and 50 μM (corresponding to 1 × K _m and 50 × K _m), and substrate peptide is present in the final reaction mixture at its saturated condition (10 μM). For histone methyltransferase (HMT) profiling in Table 1, all HMTs are purified recombinant proteins from either Escherichia coli or Baculovirus system. The catalytic domain of G9a, SuV39H2, Set7/9, CARM1, SETD8, NSD3, SETD2, and Dot1L, and the full-length SmyD2 protein were used in the biochemical assays. HMT biochemical reactions are carefully characterized with enzymology studies and the SAM and substrate K _m determined. The SAM and substrate concentrations are kept at their respective K _m for most of the HMTs, except the ones (SmyD2 and Set7/9) with low SAM-K _m value, for which 0.5 μM SAM is used. All HMT reactions are performed using the same assay format where the production of SAH from the biochemical reaction is quantitated by LC-MS.

Solubility Information

A DRUG SCREENING EXPERT

Solubility	DMSO: 50 mg/mL (114.17 mM),Sonication is recommended. Ethanol: 10 mg/mL (22.83 mM),Heating is recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble)
In vivo Formulation	10% DMSO+40% PEG300+5% Tween 80+45% Saline: 2 mg/mL (4.57 mM),Sonication is recommended. <i>Please add the solvents sequentially, clarifying the solution as much as possible before adding the next one. Dissolve by heating and/or sonication if necessary. Working solution is recommended to be prepared and used immediately. The formulation provided above is for reference purposes only. In vivo formulations may vary and should be modified based on specific experimental conditions.</i>

Preparing Stock Solutions

	1mg	5mg	10mg
1 mM	2.2834 mL	11.4171 mL	22.8342 mL
5 mM	0.4567 mL	2.2834 mL	4.5668 mL
10 mM	0.2283 mL	1.1417 mL	2.2834 mL
50 mM	0.0457 mL	0.2283 mL	0.4567 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

Croy CH, et al. Mol Pharmacol. 2014, 86(1), 106-115.

Schober DA, et al. Mol Pharmacol. 2014, 86(1), 116-123.

Inhibitor · Natural Compounds · Compound Libraries · Recombinant Proteins

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