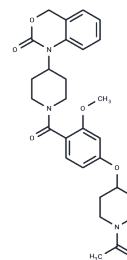


L-371,257

Chemical Properties

CAS No. : 162042-44-6
 Formula: C₂₈H₃₃N₃O₆
 Molecular Weight: 507.58
 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 | L-371,257 is a competitive antagonist of oxytocin receptor with pA ₂ of 8.4 and K _i of 19 nM. L-371,257 shows a K _i of 3.7 nM for vasopressin receptor 1a. |
| Targets(IC ₅₀) | Vasopressin Receptor,Oxytocin Receptor |
| In vivo | L-371,257 (0.5 and 1.0 mg/kg; i.p.) stimulates weight gain (10.5 g) relative to vehicle treatment (4.7 g) in rats[3]. |

Solubility Information

| | |
|---------------------|--|
| Solubility | DMSO: 8.1 mg/mL (15.96 mM),Sonication 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: 0.5 mg/mL (0.99 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 | 1.9701 mL | 9.8507 mL | 19.7013 mL |
| 5 mM | 0.394 mL | 1.9701 mL | 3.9403 mL |
| 10 mM | 0.197 mL | 0.9851 mL | 1.9701 mL |
| 50 mM | 0.0394 mL | 0.197 mL | 0.394 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

Williams PD, et al. 1-(1-[4-[(N-acetyl-4-piperidinyl)oxy]-2-methoxybenzoyl]piperidin-4-yl)-4H-3,1-benzoxazin-2(1H)-one (L-371,257): a new, orally bioavailable, non-peptide oxytocin antagonist. *J Med Chem.* 1995 Nov 10;38(23):4634-6.

Tunstall BJ, et al. Oxytocin blocks enhanced motivation for alcohol in alcohol dependence and blocks alcohol effects on GABAergic transmission in the central amygdala. *PLoS Biol.* 2019 Apr 16;17(4):e2006421.

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