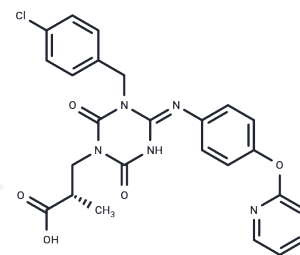


Sivopixant

Chemical Properties

CAS No. : 2414285-40-6
 Formula: C₂₅H₂₂ClN₅O₅
 Molecular Weight: 507.93
 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	Sivopixant (S-600918) (S-600918) is a potent and selective antagonist of P2X3 receptor (P2X3 IC ₅₀ =4.2 nM; P2X2/3 IC ₅₀ =1100 nM). Sivopixant shows strong analgesic effect .
Targets(IC50)	P2X Receptor
In vivo	Sivopixant (S-600918) was identified as a clinical candidate with potent and selective antagonistic activity (P2X3 IC ₅₀ , 4.2 nM; P2X2/3 IC ₅₀ , 1100 nM) and a strong analgesic effect in the rat partial sciatic nerve ligation model (Seltzer model) of allodynia (ED ₅₀ , 0.4 mg/kg).

Solubility Information

Solubility	DMSO: 55 mg/mL (108.28 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: 1.67 mg/mL (3.29 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	1.9688 mL	9.8439 mL	19.6878 mL
5 mM	0.3938 mL	1.9688 mL	3.9376 mL
10 mM	0.1969 mL	0.9844 mL	1.9688 mL
50 mM	0.0394 mL	0.1969 mL	0.3938 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

Kai H, et al. Discovery of clinical candidate Sivopixant (S-600918): Lead optimization of dioxotriazine derivatives as selective P2X3 receptor antagonists [published online ahead of print, 2021 Sep 26]. *Bioorg Med Chem Lett.* 2021; 52:128384.

Wang D P, Zhang M, Li M, et al. Druggable site near the upper vestibule determines the high affinity and P2X3 homotrimer selectivity of Sivopixant/S-600918 and its analogue DDTPA. *British Journal of Pharmacology.* 2023

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