

Odevixibat

Chemical Properties

CAS No. : 501692-44-0

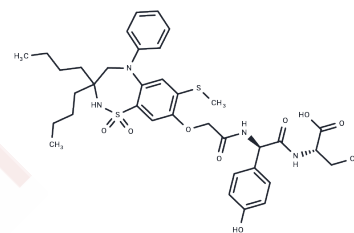
Formula: C37H48N4O8S2

Molecular Weight: 740.93

Keep away from direct sunlight

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	Odevixibat (A4250) is an effective, selective and orally active ileal bile acid transporter inhibitor. Odevixibat has the potential for the treatment of primary biliary cirrhosis. Odevixibat reduces cholestatic liver and bile duct injury in mice model.
Targets(IC50)	Others,ASBT
In vivo	Odevixibat obviously decreases bile flow and biliary BA output, which correlates with reduced bsep transcription, while Ntcp and Cyp7a1 are induced. Odevixibat (0.01% (w/w) in chow diet; 4 weeks) improves sclerosing cholangitis and significantly decreases serum alanine aminotransferase, alkaline phosphatase and BAs levels, hepatic expression of pro-inflammatory and pro-fibrogenic genes and bile duct proliferation in Mdr2-/- mice [1].

Solubility Information

Solubility	DMSO: 150 mg/mL (202.45 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: 5 mg/mL (6.75 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	1.3497 mL	6.7483 mL	13.4966 mL
5 mM	0.2699 mL	1.3497 mL	2.6993 mL
10 mM	0.135 mL	0.6748 mL	1.3497 mL
50 mM	0.027 mL	0.135 mL	0.2699 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

Baghdasaryan A, et al. Inhibition of intestinal bile acid absorption improves cholestatic liver and bile duct injury in a mouse model of sclerosing cholangitis. *J Hepatol.* 2016 Mar;64(3):674-81.

Inhibitor · Natural Compounds · Compound Libraries · Recombinant Proteins

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