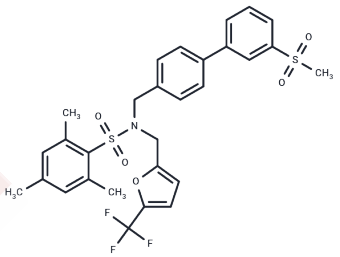
Data Sheet (Cat.No.T15427)

TargetMol

GSK2033

Chemical Properties



CAS No. : 1221277-90-2

Formula: C29H28F3NO5S2

Molecular Weight: 591.66

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	GSK2033 is an antagonist of LXR (pIC50s: 7 and 7.4 for LXRα or LXRβ, respectively).
Targets(IC50)	Liver X Receptor
In vitro	GSK2033 dose-dependently inhibits basal transcription in full-length LXRα or full-length LXRβ cotransfection assays (IC50s: 17 nM and 9 nM, respectively). GSK2033 also effectively inhibits the transcription of an ABCA1 driven luciferase reporter dose-dependently (IC50s: 52 nM for LXRα and 10 nM for LXRβ). GSK2033 also suppresses the expression of both of fatty acid synthase (FASN) and SREBP1[2].
In vivo	GSK2033 (one month) treatment, does not have obviously effects on hepatic triglyceride levels. GSK2033 treatment also have no effects on plasma triglyceride levels[2].

Solubility Information

Solubility	DMSO: 11.9 mg/mL (20.11 mM),Sonication is recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble)

Preparing Stock Solutions

	1mg	5mg	10mg
1 mM	1.6902 mL	8.4508 mL	16.9016 mL
5 mM	0.338 mL	1.6902 mL	3.3803 mL
10 mM	0.169 mL	0.8451 mL	1.6902 mL
50 mM	0.0338 mL	0.169 mL	0.338 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

Zuercher WJ, et al. Discovery of tertiary sulfonamides as potent liver X receptor antagonists. J Med Chem. 2010 Apr 22;53(8):3412-6.
Liu Y, Wang Z, Jin H, et al.Squalene-epoxidase-catalyzed 24 (S), 25-epoxycholesterol synthesis promotes trained-immunity-mediated antitumor activity.Cell Reports.2024, 43(4).
Griffett K, et al. Promiscuous activity of the LXR antagonist GSK2033 in a mouse model of fatty liver disease. Biochem Biophys Res Commun. 2016 Oct 21;479(3):424-428.