Data Sheet (Cat.No.T12375)

C13H17NaO2



Fezagepras sodium

Chemical Properties

Formula:

CAS No.: 1254472-97-3

Molecular Weight: 228.26

Appearance: no data available

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

Biological Description

Description	Fezagepras sodium (Setogepram sodium salt) is an orally active GPR40 agonist and is an antagonist or inverse agonist for GPR84, with anti-fibrotic, anti-inflammatory and anti-proliferative actions. GPR			
Targets(IC50)				
In vitro	PBI-4050 inhibited kidney macrophage infiltration, oxidative stress, and TGF-β-mediated fibrotic signaling pathways, and it also protected against the development of tubulointerstitial fibrosis. To confirm a direct antiinflammatory/antifibrotic effect in the kidney, further studies with a nondiabetic model of EGFR-mediated proximal tubule activation confirmed that PBI-4050 dramatically decreased the development of the associated tubulointerstitial injury and macrophage infiltration[1].			
In vivo	eNOS-/- db/db mice were treated with PBI-4050 from 8-20 weeks of age (early treatment) or from 16-24 weeks of age (late treatment). PBI-4050 treatment ameliorated the fasting hyperglycemia and abnormal glucose tolerance tests seen in vehicle-treated mice. In addition, PBI-4050 preserved (early treatment) or restored (late treatment) blood insulin levels and increased autophagy in islets. PBI-4050 treatment led to significant improvements in lifespan in the diabetic mice. Both early and late PBI-4050 treatment protected against progression of DN, as indicated by reduced histological glomerular injury and albuminuria, slow decline of glomerular filtration rate, and loss of podocytes[1].			

Solubility Information

Solubility	H2O: 90.0 mg/mL (394.3 mM),		
	DMSO: 60 mg/mL (262.86 mM),		
	(< 1 mg/ml refers to the product slightly soluble or insoluble)		

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Preparing Stock Solutions

	1mg	5mg	10mg
1 mM	4.381 mL	21.9048 mL	43.8097 mL
5 mM	0.8762 mL	4.381 mL	8.7619 mL
10 mM	0.4381 mL	2.1905 mL	4.381 mL
50 mM	0.0876 mL	0.4381 mL	0.8762 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.

Reference

Li Y, et al. Fatty acid receptor modulator PBI-4050 inhibits kidney fibrosis and improves glycemic control. JCI Insight. 2018 May 17;3(10). pii: 120365.

Inhibitor · Natural Compounds · Compound Libraries · Recombinant Proteins

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