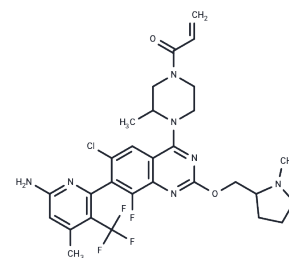


Divarasib

Chemical Properties

CAS No. :	2417987-45-0
Formula:	C ₂₉ H ₃₂ ClF ₄ N ₇ O ₂
Molecular Weight:	622.06
Appearance:	no data available
Storage:	Powder: -20°C for 3 years In solvent: -80°C for 1 year



Biological Description

Description	Divarasib (GDC-6036) (GDC-6036) is an investigational, oral, high-potency and selective KRAS G12C inhibitor with an IC ₅₀ of < 0.01 μM. Divarasib irreversibly locks KRAS G12C oncoprotein in an inactive state and inhibits tumor cell growth. It is being investigated for use in solid tumors, including non-small cell lung and colorectal cancers and other types of cancer.
Targets(IC ₅₀)	Ras
In vitro	Divarasib exhibits a potency with an EC ₅₀ value of 2 nM in K-Ras G12C-alkylation HCC1171 cells.[2]
In vivo	Administration of Divarasib (10-100 mg/kg/day; PO for 7 days) results in a reduction in the ratio of free KRAS G12C.[1]

Solubility Information

Solubility	DMSO: 90.0 mg/mL (144.7 mM), Sonication is recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble)
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Preparing Stock Solutions

	1mg	5mg	10mg
1 mM	1.6076 mL	8.0378 mL	16.0756 mL
5 mM	0.3215 mL	1.6076 mL	3.2151 mL
10 mM	0.1608 mL	0.8038 mL	1.6076 mL
50 mM	0.0322 mL	0.1608 mL	0.3215 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

Meng L, et al. Assessment of KRAS G12C Target Engagement by a Covalent Inhibitor in Tumor Biopsies Using an Ultra-Sensitive Immunoaffinity 2D-LC-MS/MS Approach. Anal Chem. 2022;94(37):12927-12933.

Inhibitor · Natural Compounds · Compound Libraries · Recombinant Proteins

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