

Certificate of Analysis

Catalog Number	BP10145
Product Name	Sitravatinib

Physical and Chemical Properties

Synonyms	MG516, MGCD516
CAS No.	1123837-84-2
Chemical Formula	C33H29F2N5O4S
Molecular Weight	629.68
Solubility	DMSO: 32 mg/mL
Storage	Powder: -20°C for 2 years In solvent: -80°C for 1 year
Chemical Structure OR Tested Image	HN HN O H

Product Information

Description	Sitravatinib (MGCD516) is an inhibitor targeting multiple RTKs involved in driving sarcoma cell growth, including c-Met, c-Kit, PDGFRα/β, PDGFR, and Axl.
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Targets&IC50	DDR2: 0.5 nM, EPHA3: 1 nM, Mer: 2 nM, VEGFR3 (FLT4): 2 nM, Axl: 1.5 nM
In vitro	Cell lines: DDLS,LS141,and MPNST. Concentrations: 62.5,125,250,500,1000,2000 nM2,000-3,000 cells were plated in 96-well plates in RPMI/DME media with 10% FBS and then treated with the indicated drugs the next day. After 72 hours, media was replaced with 100 µL of media with 10% serum and 10% CCK-8 solution. After 1 hour, the optical density was read at 450 nm to determine viability. Background values from negative control wells without cells were subtracted for final sample quantification. Data was plotted as % cell viability compared to DMSO control.
In vivo	Animal Models: ICR/SCID mice. Formulation: 0.5% hydroxypropyl methylcellulose (HPMC) and 0.1% Tween-80 solution (pH 1.4). Dosages: 15 mg/kg. Administration: p.o.

Analytical Data

HPLC	Shows Min >99% purity
H-NMR	Consistent with structure
Stability and Solubility Advice	Information on product stability, especially in solution, has rarely been reported and in most cases we can only provide a general guideline. We recommend that once the stock solution has been prepared, it be stored in equal quantities in sealed vials and used within 1 month. Avoid repeated freezing and thawing cycles. Storage conditions for some special products should be referred to their storage details.

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