# PRODUCT INFORMATION



## **Ningetinib**

Item No. 25233

CAS Registry No.: 1394820-69-9

Formal Name: N-[3-fluoro-4-[[7-(2-hydroxy-2-methylpropoxy)-4-

quinolinyl]oxy]phenyl]-2,3-dihydro-1,5-dimethyl-3-

oxo-2-phenyl-1H-pyrazole-4-carboxamide

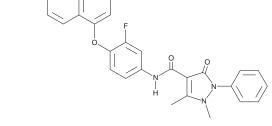
MF:  $C_{31}H_{29}FN_4O_5$ 

556.6 FW: ≥98% **Purity:** 

UV/Vis.:  $\lambda_{max}$ : 234, 294 nm A crystalline solid Supplied as:

Storage: -20°C Stability: ≥2 years

Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis.



## **Laboratory Procedures**

Ningetinib is supplied as a crystalline solid. A stock solution may be made by dissolving the ningetinib in the solvent of choice. Ningetinib is soluble in organic solvents such as DMSO and dimethyl formamide (DMF), which should be purged with an inert gas. The solubility of ningetinib in these solvents is approximately 2 and 15 mg/ml, respectively.

Ningetinib is sparingly soluble in aqueous buffers. For maximum solubility in aqueous buffers, ningetinib should first be dissolved in DMF and then diluted with the aqueous buffer of choice. Ningetinib has a solubility of approximately 0.16 mg/ml in a 1:5 solution of DMF:PBS (pH 7.2) using this method. We do not recommend storing the aqueous solution for more than one day.

### Description

Ningetinib is a multi-kinase inhibitor that inhibits c-Met, VEGFR2/KDR, and AxI ( $IC_{50}$ s = 19, 37, and 11 nM, respectively). In vivo, ningetinib (20 mg/kg) inhibits tumor growth in MKN45, Čaki-1, NCI-H441, Huh-7, U87MG, and MDA-MB-231 mouse xenograft models by 53-97%.

### Reference

1. Xi, N. Substituted quinoline compounds and methods of use. Sunshine Lake Pharma Co. Ltd. US20120219522A1, (2012).

WARNING
THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

#### WARRANTY AND LIMITATION OF REMEDY

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