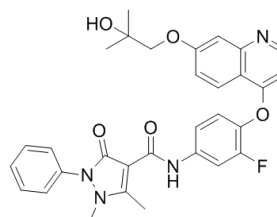


**Product Name** : Ningetinib  
**Synonyms** : CT-053, CT053PTSA  
**Cat No.** : M22115  
**CAS Number** : 1394820-69-9  
**Molecular Formula** : C<sub>31</sub>H<sub>29</sub>FN<sub>4</sub>O<sub>5</sub>  
**Formula Weight** : 556.58  
**Chemical Name** : —



**Description** : Ningetinib (CT053PTSA) is an orally bioavailable tyrosine kinase inhibitor with IC<sub>50</sub>s of <1.0, 1.9 and 6.7 nM for Axl, VEGFR2, and c-Met, respectively. In cell-based functional assays, Ningetinib inhibits VEGF and HGF-stimulated HUVEC proliferation and microvascular angiogenesis in rat aortic rings with IC<sub>50</sub> values of 6.3 and 8.6 nM, respectively. In the orthotopic U87MG human glioblastoma xenograft model, Ningetinib prolongs the median survival time and yields a significant increase in life-span value (ILS=32%) at an oral dose of 20 mg/kg/day (dosed 21 days) versus the vehicle-treated group. When single dosed orally (3 mg/kg) to U87MG tumor-bearing nude mice, Ningetinib potently inhibits the phosphorylation of c-Met and its downstream signaling kinases AKT and ERK1/2 for up to 6 hours in tumor tissues.

**Pathway** : Angiogenesis  
**Target** : c-Met/HGFR  
**Receptor** : c-Met; VEGFR2; AXL  
**Solubility** : DMSO: 15 mg/mL (26.95 mM; Need ultrasonic)  
**SMILES** : Cc1c(C(=O)Nc2ccc(Oc3ccnc4cc(OCC(C)(C)O)ccc34)c(F)c2)c(=O)n(-c2ccccc2)n1C  
**Storage** : (-20°C)  
**Stability** : ≥ 2 years  
**Reference** :

1. Ning Xi, et al. Abstract 1755: CT053PTSA, a novel c-MET and VEGFR2 inhibitor, potently suppresses angiogenesis and tumor growth. Cancer Res 2014;74(19 Suppl):Abstract nr 1755.