Web: www.taiclone.com
Tel: +886-2-2735-9682
Email: order@taiclone.com

AMG-208

Catalog No: tcsc0185

Available Sizes

Size: 5mg

Size: 10mg

Size: 50mg

Size: 100mg

Specifications

CAS No:
1002304-34-8

Formula:
$C_{22}H_{17}N_5O_2$

Pathway:
Protein Tyrosine Kinase/RTK

Target:
c-Met/HGFR

Purity / Grade:
>98%

Solubility:
DMSO : 7.8 mg/mL (20.34 mM; Need ultrasonic and warming)

Observed Molecular Weight:
383.4

Product Description

AMG-208 is a potent small molecular c-Met inhibitor with an IC50 of 9.3 nM.

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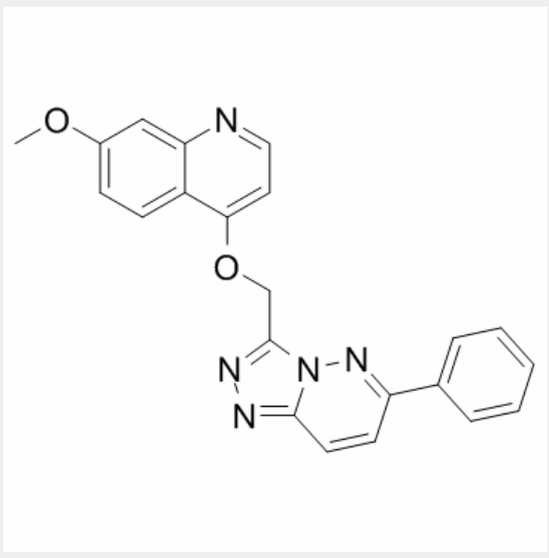
IC50 value: 9.3 nM

Target: c-Met

in vitro: AMG-208 shows the potent inhibition of kinase c-Met activity with IC50 of 9 nM in a cell-free assay. Besides, AMG-208 treatment also leads to the inhibition of HGF-mediated c-Met phosphorylation in PC3 cells with IC50 of 46 nM [1]. Pre-incubation of AMG-208 with human liver microsomes for 30 minutes shows a potent time-dependent inhibition for CYP3A4 metabolic activity with IC50 of 4.1 μM, which is an eightfold decrease relative to the IC50 (32 μM) without preincubation [2]. AMG-208 is identified to be a c-MET and RON dual selective inhibitor [3].

in vivo: In male Sprague Dawley rats, AMG-208 (0.5 mg/kg i.v.) shows a high bioavailability with Cl of 0.37 L/h/kg, Vss of 0.38 L/kg and T1/2 of 1 hour[1].



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