

UNC2250

Catalog No: tcsc2318



Available Sizes

Size: 5mg

Size: 10mg

Size: 50mg

Size: 100mg



Specifications

CAS No:

1493694-70-4

Formula:

$C_{24}H_{36}N_6O_2$

Pathway:

Protein Tyrosine Kinase/RTK

Target:

TAM Receptor

Purity / Grade:

>98%

Solubility:

DMSO : ≥ 46 mg/mL (104.41 mM)

Observed Molecular Weight:

440.58

Product Description

UNC2250 is a phosphorylation of endogenous Mer inhibitor with an IC50 of 9.8 nM and blocked ligand-stimulated activation of a

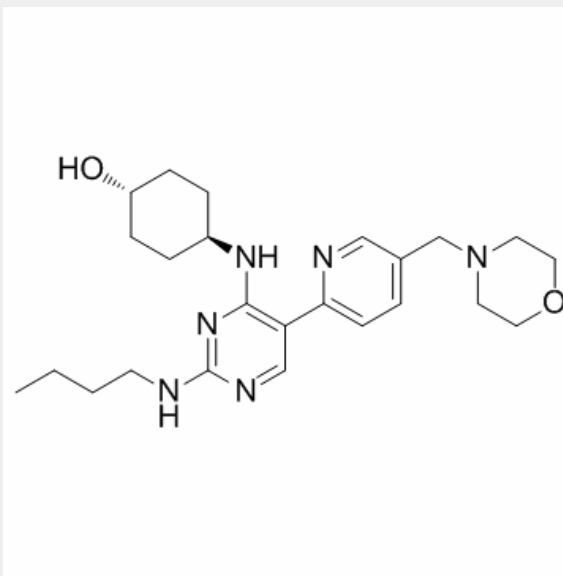
chimeric EGFR-Mer protein.

IC50 Value: 9.8 nM [1]

Target: Others

in vitro: UNC2250 is 160-fold more active for Mer versus Axl and 60-fold versus Tyro3. UNC2250 had a moderate half-life, clearance, and volume of distribution as well as reasonable oral bioavailability and good solubility and was thus chosen for characterization of kinase selectivity and further evaluation in cell-based studies of Mer activity. UNC2250 efficiently inhibited ligand-dependent phosphorylation of a chimeric protein consisting of the extracellular and transmembrane domains of the epidermal growth factor (EGF) receptor and the intracellular tyrosine kinase domain of Mer. Moreover, UNC2250 incubation inhibited colony formation in soft agar cultures of the BT-12 rhabdoid tumor and the Colo699 NSCLC cell lines. In the Colo699 NSCLC cell line, the concentrations of UNC2250 required to inhibit colony formation and Mer phosphorylation were similar. These data suggest that the functional antiproliferative activity mediated by UNC2250 resulted from Mer inhibition rather than a consequence of off-target inhibition of other kinases [1,2].

In vivo:



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