

H-1152

Catalog No: tcsc1536



Available Sizes

Size: 5mg

Size: 10mg



Specifications

CAS No:

451462-58-1

Formula:

$C_{16}H_{21}N_3O_2S$

Pathway:

TGF-beta/Smad;Stem Cell/Wnt;Cell Cycle/DNA Damage

Target:

ROCK;ROCK;ROCK

Purity / Grade:

>98%

Solubility:

10 mM in DMSO

Observed Molecular Weight:

319.42

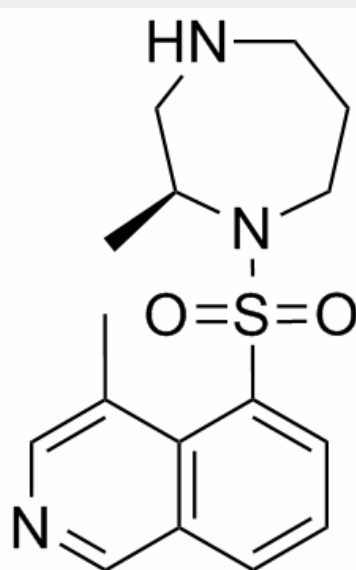
Product Description

H-1152 is a membrane-permeable and selective **ROCK** inhibitor, with a **K_i** value of 1.6 nM, and an **IC₅₀** value of 12 nM for **ROCK2**.

IC50 & Target: IC50: 12 nM (ROCK2)^[1]

Ki: 1.6 nM (ROCK)^[2]

In Vitro: H-1152 is an inhibitor of Rho-kinase, with an IC_{50} of 12 nM for ROCK2. H-1152 (H-1152P) also shows less inhibitory activities against CaMKII, PKG, AuroraA, PKA, Src, PKC, MLCK, Abl, EGFR, MKK4, GSK3 α , AMPK, and P38 α , with IC_{50} s of 0.180, 0.360, 0.745, 3.03, 3.06, 5.68, 28.3, 7.77, 50.0, 16.9, 60.7, 100, and 100 μ M, respectively^[1]. H-1152 potently inhibits Rho kinase, with a K_i of 1.6 nM, and slightly suppresses PKA, PKC and MLCK, with K_i s of 0.63, 9.27, and 10.1 μ M, respectively. H-1152 (0.1-10 μ M) highly inhibits MARCKS phosphorylation, with an IC_{50} value of 2.5 μ M in LPA-treated cells, but shows no such obvious effects in PDBu-treated cells^[2]. H-1152 (0.5-10 μ M) causes no decreased neuronal survival. H-1152 (1, 5 or 10 μ M) also exerts no alterations in the ratios of different neuronal morphologies. Furthermore, H-1152 (10 μ M) increases neurite length in both BMP4 and LIF cultures^[3].



All products are for RESEARCH USE ONLY. Not for diagnostic & therapeutic purposes!