

Product Name : Y-33075 dihydrochloride

Synonyms : —

**Cat No.** : M22457

**CAS Number** : 173897-44-4

Molecular Formula : C16H18Cl2N4O

Formula Weight : 353.25

Chemical Name : ----

Description

Y-33075 dihydrochloride is a selective inhibitor of ROCK(IC50 of 3.6 nM). Y-33075 (Y-39983) is a potent inhibitor of ROCK(IC50 of 3.6 nM). Y-33075 also inhibits PKC and CaMKII more potently than Y-27632(IC50s of Y-27632 and Y-33075 for PKC are 9.0  $\mu$ M and 0.42  $\mu$ M, respectively), whereas the IC50s of Y-27632 and Y-33075 for CaMKII are 26  $\mu$ M and 0.81  $\mu$ M, respectively. The IC50s of Y-27632 and Y-33075 for PKC is 82 and 117 times those for ROCK, respectively, whereas the IC50s of Y-27632 and Y-33075 for CaMKII is 236 and 225 times those for ROCK, respectively. Y-33075 (Y-39983, 10

µM) extends neurites in the retinal ganglion cells (RGCs) compared with those in RGCs treated without Y-39983, Y-33075 (Y-39983, 1 µM) inhibits the contraction of rabbit ciliary artery segments evoked by histamine in Ca2+-free solutions[3].Y-39983 (≥0.01%) significantly lowers intraocular pressure (IOP) at 2 hours after topical administration in rabbits. Y-39983 (0.05%)-treated eyes show significant reduction of IOP between 2 and 7 hours after topical administration in monkeys. Y-39983 (100 µM) increases the regenerating axons of retinal ganglion cells (RGCs) in the eyes of the rats.

Pathway : Cell Cycle/DNA Damage

Target : ROCK

Receptor : ROCK;PKC;CaMKII

Solubility : DMSO:100 mg/mL (283.09 mM; Need ultrasonic);H2O:50 mg/mL (141.54 mM; Need ultrasonic)

**SMILES** : Cl.Cl.C[C@@H](N)c1ccc(cc1)C(=O)Nc1ccnc2[nH]ccc12

Storage : (-20℃)

Stability : ≥ 2 years

Reference :

1. Hideki Tokushige, et al. Effects of Topical Administration of Y-39983, a Selective Rho-Associated Protein Kinase Inhibitor, on Ocular Tissues in Rabbits and Monkeys Invest. Ophthalmol. Vis. Sci. July 2007 vol. 48no. 7 3216-3222