Y-27632 (Dihydrochloride)

Small Molecules

RHO/ROCK pathway inhibitor; Inhibits ROCK1 and ROCK2



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Catalog # 72302 1 mg 72304 5 mg

72307 5 x 10 mg
72308 50 mg
100-1044 500 mg

Product Description

Y-27632 (Dihydrochloride) is a cell-permeable, highly potent and selective inhibitor of Rho-associated, coiled-coil containing protein kinase (ROCK). Y-27632 inhibits both ROCKI (Ki = 220 nM) and ROCKII (Ki = 300 nM) by competing with ATP for binding to the catalytic site (Davies et al.; Ishizaki et al.).

Alternative Names: ROCK inhibitor

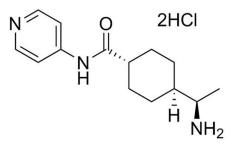
CAS Number: 129830-38-2

Chemical Formula: $C_{14}H_{21}N_3O \cdot 2HCI$ Molecular Weight: 320.3 g/mol

Purity: \geq 98%

Chemical Name: 4-[(1R)-1-aminoethyl]-N-4-pyridinyl-trans-cyclohexanecarboxamide, dihydrochloride

Structure:



Properties

Physical Appearance: A crystalline solid

Storage: Product stable at -20°C as supplied. Protect from prolonged exposure to light. For long-term storage, store with

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a desiccant.

Stable as supplied for 12 months from date of receipt.

Solubility: · PBS (pH 7.2) ≤ 30 mM

· DMSO ≤ 90 mM

· Absolute ethanol ≤ 15 mM

For example, to prepare a 5 mM stock solution in PBS or water, resuspend 1 mg in 624 µL of PBS (pH 7.2) or

water.

Prepare stock solution fresh before use. Stock solutions in PBS or water are stable at -20°C for up to 6 months. Aliquot into working volumes to avoid repeated freeze-thaw cycles. The effect of storage of stock solution on compound performance should be tested for each application.

For use as a cell culture supplement, stock solution should be diluted into culture medium immediately before use. This product has been shown to be effective at a final concentration of 10 μ M (Ungrin et al.; Watanabe et al.). Avoid final DMSO concentration above 0.1% due to potential cell toxicity.

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Published Applications

MAINTENANCE AND SELF-RENEWAL

- · Enhances survival of human embryonic stem (ES) cells when they are dissociated to single cells by preventing dissociation-induced apoptosis (anoikis), thus increasing their cloning efficiency (Watanabe et al.).
- · Improves embryoid body formation using forced-aggregation protocols (Ungrin et al.).
- · Increases the survival of cryopreserved single human ES cells after thawing (Li et al.).
- · Blocks apoptosis of mouse ES-derived neural precursors after dissociation and transplantation (Koyanagi et al.). REPROGRAMMING
- · Direct lineage reprogramming of fibroblasts to mature neurons, in combination with CHIR99021 (Catalog #72052), RepSox (Catalog #73792), Forskolin (Catalog #72112), SP600125 (Catalog #72642), Gö6983 (Catalog #72462), and Valproic Acid (Catalog #72292) (Hu et al.).

DIFFERENTIATION

· Improves survival of human ES cell monolayers at the initiation of differentiation protocols (Rezania et al.)

References

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Ishizaki T et al. (2000) Pharmacological properties of Y-27632, a specific inhibitor of rho-associated kinases. Mol Pharmacol 57(5): 976–83. Koyanagi M et al. (2008) Inhibition of the Rho/ROCK pathway reduces apoptosis during transplantation of embryonic stem cell-derived neural precursors. J Neurosci Res 86(2): 270–80.

Li X et al. (2009) ROCK inhibitor improves survival of cryopreserved serum/feeder-free single human embryonic stem cells. Hum Reprod 24(3): 580–9.

Rezania A et al. (2014) Reversal of diabetes with insulin-producing cells derived in vitro from human pluripotent stem cells. Nat Biotechnol 32(11): 1121–33.

Ungrin MD et al. (2008) Reproducible, ultra high-throughput formation of multicellular organization from single cell suspension-derived human embryonic stem cell aggregates. PLoS One 3(2): e1565.

Watanabe K et al. (2007) A ROCK inhibitor permits survival of dissociated human embryonic stem cells. Nat Biotechnol 25(6): 681-6.

Related Small Molecules

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