

Product Name: Y-27632 dihydrochloride
Revision Date: 10/23/2023

Product Data Sheet

Y-27632 dihydrochloride

Cat. No.: A3008

CAS No.: 146986-50-7; 129830-38-2

Formula: C14H21N3O·2HCI

M.Wt: 320.26

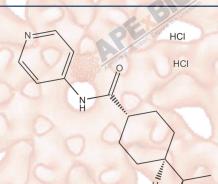
Synonyms: y-27632, Y27632, Y-27632 dihydrochloride, Y

27632

Target: TGF-β / Smad Signaling

Pathway: ROCK

Storage: Desiccate at 4°C or below



Solvent & Solubility

≥52.9mg/ml in H2O; ≥16.013mg/mL in DMSO

Mass 1mg 10mg Solvent 5mg Preparing Concentration In Vitro Stock Solutions 15.6123 mL 1 mM 3.1225 mL 31.2246 mL 5 mM 0.6245 mL 3.1225 mL 6.2449 mL 10 mM 1.5612 mL 3.1225 mL 0.3122 mL

Please refer to the solubility information to select the appropriate solvent.

Biological Activity

Shortsummary	ROCK1 inhibitor		
IC ₅₀ & Target	140 nM (Ki) (ROCK1), 300	140 nM (Ki) (ROCK1), 300 nM (Ki) (ROCK2)	
	Cell Viability Assay		
	Cell Line:	Human (hu) and rat(r) prostatic smooth mus-cle cells (PSM)	
	Preparation method:	The solubility of this compound in DMSO is >10 mM. General tips for obtaining	
In Vitro		a higher concentration: Please warm the tube at 37 °C for 10 minutes and/or	
		shake it in the ultrasonic bath for a while. Stock solution can be stored below	
		-20°C for several months.	
	Reacting conditions:	24h and 48 h; 100 μM	

	Applications:	After identifying prostatic smooth muscle cells and confirming the expression of
		Rho-kinase in these cells we investigated whether the Rho-kinase inhibitor
		Y-27632 affected the viability and proliferation of these cells. In serum-free
		medium huPSM and rPSM were made quiescent for 24 hours. Cell viability
		using neutral red and MTT assays was assessed 24 and 48 hours after
	Carlotte Carlotte	stimulating the cells with 1% serum in the absence and presence of Y-27632
	Expose the limit	(0.01 to 100 μ M). The results of these assays showed that the number of the
	Ju Particular.	cells increased between the 24- and 48-hour incubation periods after
	Konta	re-stimulation with 1% serum. However, in the presence of Y-27632 the
		increase in the number of live cells was less than in the control group. This
		effect was concentration dependent.
In Vivo	Animal experiment	
	Animal models:	Adult Swiss male albino mice
	Dosage form:	0.1 mg/kg/day; intrapertoneal injection
	Applications:	The drug was tested by histopathological examination showed that Y-27632
	the Unknown	administration to EAC-bearing mice diminished pathological structure, to 60-
	Collect Expore	70% degree, toward to normal intact histological structure especially in
	Activie Peru	pre-carcinoma inoculation regime. Respect to this, ROCK inhibition by Y-27632
		decreased significantly tumor invasion and metastasis. Our
		immunohistochemistry results showed that ROCK2 was mainly inhibited by
		Y-27632 in pre-carcinoma, but not in post-carcinoma, groups.
	Other notes:	Please test the solubility of all compounds indoor, and the actual solubility may
		slightly differ with the theoretical value. This is caused by an experimental
		system error and it is normal.

Product Citations

- 1. Ni P, Noh H, et al. "Large-Scale Generation and Characterization of Homogeneous Populations of Migratory Cortical Interneurons from Human Pluripotent Stem Cells." Mol Ther Methods Clin Dev. 2019 Apr 8;13:414-430.PMID:31061832
- 2. Speer JE, Gunasekara DB, et al. "Molecular transport through primary human small intestinal monolayers by culture on a collagen scaffold with a gradient of chemical cross-linking." J Biol Eng. 2019 Apr 27;13:36.PMID:31061676
- 3. Ni P, Noh H, et al. "iPSC-derived homogeneous populations of developing schizophrenia cortical interneurons have compromised mitochondrial function." Mol Psychiatry. 2019 Apr 24.PMID:31019265
- 4. Dakhama A, Al Mubarak R, et al. "Tollip Inhibits ST2 Signaling in Airway Epithelial Cells Exposed to Type 2 Cytokines and Rhinovirus." J Innate Immun. 2019 Mar 29:1-13.PMID:30928973
- 5. Grun D, Adhikary G, et al. "NRP-1 interacts with GIPC1 and SYX to activate p38 MAPK signaling and cancer stem cell survival." Mol Carcinog. 2018 Nov 19.PMID:30456845

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References

[1] Rees R W, Foxwell N A, Ralph D J, et al. Y-27632, a Rho-kinase inhibitor, inhibits proliferation and adrenergic contraction of prostatic smooth muscle cells[J]. The Journal of urology, 2003, 170(6): 2517-2522.

[2] Isler D, Ozaslan M, Karagoz I D, et al. Antitumoral effect of a selective Rho-kinase inhibitor Y-27632 against Ehrlich ascites carcinoma in mice[J]. Pharmacological Reports, 2014, 66(1): 114-120.

Caution

FOR RESEARCH PURPOSES ONLY.

NOT FOR HUMAN, VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

Specific storage and handling information for each product is indicated on the product datasheet. Most APExBIO products are stable under the recommended conditions. Products are sometimes shipped at a temperature that differs from the recommended storage temperature. Shortterm storage of many products are stable in the short-term at temperatures that differ from that required for long-term storage. We ensure that the product is shipped under conditions that will maintain the quality of the reagents. Upon receipt of the product, follow the storage recommendations on the product data sheet.

APExBIO Technology

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