

Produktinformation



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Zellkultur & Verbrauchsmaterial
Diagnostik & molekulare Diagnostik
Laborgeräte & Service

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SZABO-SCANDIC HandelsgmbH

Quellenstraße 110, A-1100 Wien T. +43(0)1 489 3961-0 F. +43(0)1 489 3961-7 <u>mail@szabo-scandic.com</u> www.szabo-scandic.com

(-)-Blebbistatin (GMP)

MedChemExpress

Cat. No.:	HY-13441G	0
CAS No.:	856925-71-8	∬ о́н
Molecular Formula:	C ₁₈ H ₁₆ N ₂ O ₂	
Molecular Weight:	292.33	
Target:	Myosin	\sim N N
Pathway:	Cytoskeleton	
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.	
	CAS No.: Molecular Formula: Molecular Weight: Target: Pathway:	CAS No.:856925-71-8Molecular Formula:C18H16N2O2Molecular Weight:292.33Target:MyosinPathway:CytoskeletonStorage:Please store the product under the recommended conditions in the Certificate of

BIOLOGICAL ACTIVITY		
Description	(-)-Blebbistatin (GMP) is (-)-Blebbistatin (HY-13441) produced by using GMP guidelines. (-)-Blebbistatin (GMP) is a selective inhibitor of the ATPase activity of non-muscle myosin II ^{[1][2]} .	
In Vitro	Blebbistatin potently inhibits several striated muscle myosins as well as vertebrate nonmuscle myosin IIA and IIB with IC ₅₀ values ranging from 0.5 to 5 μM. Smooth muscle myosin is only poorly inhibited (IC ₅₀ =80 μM) ^[1] . Blebbistatin does not compete with nucleotide binding to the skeletal muscle myosin subfragment-1. The inhibitor preferentially binds to the ATPase intermediate with ADP and phosphate bound at the active site, and it slows down phosphate release. It blocks the myosin heads in a products complex with low actin affinity ^[2] . In culture-activated hepatic stellate cells, blebbistatin is found to change both cell morphology and function. Stellate cells become smaller, acquire a dendritic morphology and have less myosin IIA-containing stress fibres and vinculin-containing focal adhesions. Blebbistatin impairs silicone wrinkle formation, reduces collagen gel contraction and blocks endothelin-1-induced intracellular Ca ²⁺ release. It promotes wound-induced cell migration ^[3] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.	
In Vivo	Blebbistatin dose-dependently and completely relax both KCl- and carbachol-induced rat detrusor and endothelin-1- induced human bladder contraction. Pre-incubation with 10 μM blebbistatin attenuates carbachol responsiveness by 65% while blocking electrical field stimulation-induced bladder contraction reaching 50% inhibition at 32 Hz ^[4] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.	

CUSTOMER VALIDATION

- Cell. 2024 May 9:S0092-8674(24)00448-3.
- Circulation. 2023 Nov 14.
- Adv Sci (Weinh). 2022 Mar 3;e2104682.
- Adv Sci (Weinh). 2020 Jun 17;7(15):1903583.
- Theranostics. 2019 Apr 13;9(9):2555-2571.

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REFERENCES

[1]. Cristina Lucas-Lopez, et al. Absolute Stereochemical Assignment and Fluorescence Tuning of the Small Molecule Tool, (–)-Blebbistatin.

[2]. Ponsaerts R, et al. The myosin II ATPase inhibitor blebbistatin prevents thrombin-induced inhibition of intercellularcalcium wave propagation in corneal endothelial cells. Invest Ophthalmol Vis Sci. 2008 Nov;49(11):4816-27.

Caution: Product has not been fully validated for medical applications. For research use only.

 Tel: 609-228-6898
 Fax: 609-228-5909
 E-mail: tech@MedChemExpress.com

 Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA