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Lieferung & Zahlungsart

siehe unsere [Liefer- und Versandbedingungen](#)

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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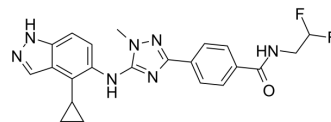
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Zelasudil

Cat. No.:	HY-148808		
CAS No.:	2365193-22-0		
Molecular Formula:	C ₂₂ H ₂₁ F ₂ N ₇ O		
Molecular Weight:	437.45		
Target:	ROCK		
Pathway:	Cell Cycle/DNA Damage; Cytoskeleton; Stem Cell/Wnt; TGF-beta/Smad		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



SOLVENT & SOLUBILITY

In Vitro	DMSO : 100 mg/mL (228.60 mM; Need ultrasonic)				
		Solvent Concentration	Mass 1 mg	5 mg	10 mg
	Preparing Stock Solutions	1 mM	2.2860 mL	11.4299 mL	22.8598 mL
		5 mM	0.4572 mL	2.2860 mL	4.5720 mL
10 mM		0.2286 mL	1.1430 mL	2.2860 mL	
Please refer to the solubility information to select the appropriate solvent.					
In Vivo	1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: 2.5 mg/mL (5.71 mM); Clear solution; Need ultrasonic 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: 2.5 mg/mL (5.71 mM); Clear solution; Need ultrasonic				

BIOLOGICAL ACTIVITY

Description	Zelasudil is a Rho-associated (ROCK) kinase inhibitor. Zelasudil has a ROCK2 binding affinity ^{[1][2]} .
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REFERENCES

- [1]. Lee E, et al. Selective ROCK2 inhibition for treatment of edema and associated conditions: World Intellectual Property Organization, WO2022169946. 2022-08-11.
- [2]. WHO Drug Information-World Health Organization (WHO).

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

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