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Lieferung & Zahlungsart

siehe unsere [Liefer- und Versandbedingungen](#)

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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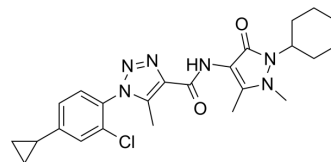
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Smurf1-IN-1

Cat. No.:	HY-149316		
CAS No.:	1824708-03-3		
Molecular Formula:	C ₂₄ H ₂₉ ClN ₆ O ₂		
Molecular Weight:	468.98		
Target:	E1/E2/E3 Enzyme		
Pathway:	Metabolic Enzyme/Protease		
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 (213.23 mM; Need ultrasonic)			
		Solvent Concentration	Mass	
			1 mg	5 mg
			10 mg	
Preparing Stock Solutions	1 mM	2.1323 mL	10.6614 mL	21.3229 mL
	5 mM	0.4265 mL	2.1323 mL	4.2646 mL
	10 mM	0.2132 mL	1.0661 mL	2.1323 mL
Please refer to the solubility information to select the appropriate solvent.				
In Vivo	<ol style="list-style-type: none"> Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: 2.5 mg/mL (5.33 mM); Clear solution; Need ultrasonic Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: 2.5 mg/mL (5.33 mM); Clear solution; Need ultrasonic Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: 2.5 mg/mL (5.33 mM); Clear solution; Need ultrasonic 			

BIOLOGICAL ACTIVITY

Description	Smurf1-IN-1 is an orally active and selective inhibitor of specific E3 ubiquitin protein ligase 1 (SMURF1) with an IC ₅₀ of 92 nM. Smurf1-IN-1 has significant efficacy in rats model of pulmonary hypertension ^[1] .
IC₅₀ & Target	IC ₅₀ : 92 nM Specific E3ubiquitin protein ligase 1 (SMURF1) ^[1]
In Vivo	Smurf1-IN-1 (Compound 38) (1, 3, 10 mg/kg for p.o.) has significant efficacy in rats model of pulmonary hypertension ^[1] . Smurf1-IN-1 (1 mg/kg for i.v., 3 mg/kg for p.o.) shows a T _{1/2} of 7.9, and oral bioavailability of 82% ^[1] .

Pharmacokinetic parameters for Smurf1-IN-1 (Compound 38) in rats ^[1]

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

dose and route	oral bioavailability (%)	p.o. AUC _{last} (nM • h/mL)	p.o. C _{max} (nM/L)	half-life (h)	clearance (mL/min/kg)	V _{ss} (l/kg)
3 mpg p.o./1 mpg i.v.	82	7383	2007	7.9	17.1	2.3

Animal Model:	Hypoxia-Sugen rat model of PAH ^[1]
Dosage:	1, 3, 10 mg/kg
Administration:	Oral gavage (p.o.)
Result:	Caused a 10% increase in muscularization at day 14 and progression to around 18% muscularization was observed in the vehicle group by the end of the study at a dose of 3 mg/kg.

REFERENCES

[1]. Shaw DE, et.al. Design and Synthesis of Inhibitors of the E3 Ligase SMAD Specific E3 Ubiquitin Protein Ligase 1 as a Treatment for Lung Remodeling in Pulmonary Arterial Hypertension. J Med Chem. 2023 Jun 22;66(12):8130-8139.

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

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