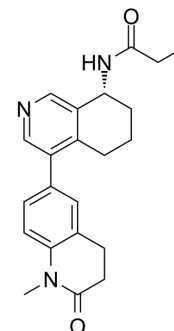


Baxdrostat

Cat. No.:	HY-132809		
CAS No.:	1428652-17-8		
Molecular Formula:	C ₂₂ H ₂₅ N ₃ O ₂		
Molecular Weight:	363.45		
Target:	Mineralocorticoid Receptor		
Pathway:	Metabolic Enzyme/Protease; Vitamin D Related/Nuclear Receptor		
Storage:	Powder	-20°C	3 years
	In solvent	-80°C	6 months
		-20°C	1 month



SOLVENT & SOLUBILITY

In Vitro	DMSO : 100 mg/mL (275.14 mM; Need ultrasonic)				
		Solvent Concentration	Mass		
	Preparing Stock Solutions			1 mg	5 mg
		1 mM		2.7514 mL	13.7570 mL
		5 mM		0.5503 mL	2.7514 mL
	10 mM		0.2751 mL	1.3757 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: ≥ 2.08 mg/mL (5.72 mM); Clear solution 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.08 mg/mL (5.72 mM); Clear solution				

BIOLOGICAL ACTIVITY

Description	Baxdrostat is a aldosterone synthase inhibitor ^[1] .
IC₅₀ & Target	aldosterone synthase ^[1]
In Vitro	Baxdrostat is a aldosterone synthase inhibitor ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

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

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