

Product Data Sheet

Posovolone

Cat. No.: HY-141795

CAS No.: 256955-84-7

Molecular Formula: $C_{26}H_{40}N_2O_3$ Molecular Weight: 428.61

Target: Others
Pathway: Others

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 (233.31 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.3331 mL	11.6656 mL	23.3312 mL
	5 mM	0.4666 mL	2.3331 mL	4.6662 mL
	10 mM	0.2333 mL	1.1666 mL	2.3331 mL

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

In Vivo

1. Add each solvent one by one: 10% DMSO >> 90% (20% SBE- β -CD in saline) Solubility: \geq 2.5 mg/mL (5.83 mM); Clear solution

BIOLOGICAL ACTIVITY

Description	Posovolone (Co 134444) is an orally active, neuroactive steroid. Posovolone has anticonvulsant and anxiolytic-like activity as well as ataxic effects ^[1] .
In Vitro	In mice, Co 134444 (0.5-10 mg/kg, IP, 15 min; 1-60 mg/kg, PO; 30 min) demonstrates efficacy against pentylenetetrazol (PTZ)-induced seizures with an ED $_{50}$ of 3.3 mg/kg after IP administration and 9.8 mg/kg orally. Similarly, protection against maximal electroshock (MES) is observed with an ED $_{50}$ of 4.8 mg/kg IP and 20.6 mg/kg PO. In rats, Co134444 (10-40 mg/kg; PO; 30 min) protects against PTZ-induced seizures and MES-induced seizures with an ED $_{50}$ of 23.6 mg/kg and 25.3 mg/kg, respectively ^[1] .

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

REFERENCES



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