## **Product** Data Sheet

# **Pyrantel tartrate**

Cat. No.: HY-12641 CAS No.: 33401-94-4 Molecular Formula:  $C_{15}H_{20}N_2O_6S$  Molecular Weight: 356.39

Target: Parasite; nAChR; Antibiotic

Pathway: Anti-infection; Membrane Transporter/Ion Channel; Neuronal Signaling

**Storage:** 4°C, sealed storage, away from moisture

\* In solvent: -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture)

### **SOLVENT & SOLUBILITY**

In Vitro

DMSO : ≥ 34 mg/mL (95.40 mM)

\* "≥" means soluble, but saturation unknown.

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.8059 mL	14.0296 mL	28.0591 mL
	5 mM	0.5612 mL	2.8059 mL	5.6118 mL
	10 mM	0.2806 mL	1.4030 mL	2.8059 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 (5.84 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE- $\beta$ -CD in saline) Solubility:  $\ge$  2.08 mg/mL (5.84 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.08 mg/mL (5.84 mM); Clear solution

## **BIOLOGICAL ACTIVITY**

Description	Pyrantel tartrate is an orally active anthelmintic and an agonist of the nicotinic acetylcholine receptor (nAChR). Pyrantel tartrate can cause spasmodic muscle paralysis in parasites. Pyrantel tartrate can be used in the study of parasitic infections such as ascariasis, hookworm infections, intestinal worms (pinworm infections), trichinosis and trichinosis <sup>[1][2]</sup> .
IC <sub>50</sub> & Target	Parasites $^{[1][2]}$ .
In Vitro	Pyrantel tartrate (10 nM-10 $\mu$ M; 72 h) shows good anti-A. suum and (0-168.2 M; 72 h) anti-N. americanus activity <sup>[1][2]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

A. suum	
10 nM-10 μM	
72 h	
Inhibited A. suum with a pEC <sub>50</sub> value of 7.24.	
N. americanus	
0-168.2 M (0-100 μg/mL)	
72 h	
Inhibited third-stage larvae and adult of N. americanus with IC <sub>50</sub> values of 2.0 and 7.6 mg/mL, respectively.	

#### In Vivo

Pyrantel tartrate (10 mg/kg; p.o.; single) reduces the worms in A. ceylanicum-infected hamsters, with the worm burden reduction of 87.2% and worm expulsion rate of 63.4%  $^{[2]}$ .

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

Animal Model:	Male Syrian Golden hamsters (3-week-old; A. ceylanicum-infected) <sup>[2]</sup> .
Dosage:	10 mg/kg
Administration:	Oral administration; single.
Result:	Exhibited worm burden reduction and worm expulsion rates of 87.2% and 63.4%, respectively.

#### **REFERENCES**

[1]. Martin RJ, et, al. Oxantel is an N-type (methyridine and nicotine) agonist not an L-type (levamisole and pyrantel) agonist: classification of cholinergic anthelmintics in Ascaris. Int J Parasitol. 2004 Aug;34(9):1083-90.

[2]. Tritten L, et, al. In vitro and in vivo efficacy of Monepantel (AAD 1566) against laboratory models of human intestinal nematode infections. PLoS Negl Trop Dis. 2011 Dec;5(12):e1457.

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

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