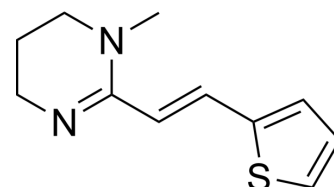


Pyrantel

Cat. No.:	HY-12641A
CAS No.:	15686-83-6
Molecular Formula:	C ₁₁ H ₁₄ N ₂ S
Molecular Weight:	206.31
Target:	nAChR; Parasite; Antibiotic
Pathway:	Membrane Transporter/Ion Channel; Neuronal Signaling; Anti-infection
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Pyrantel is an orally active anthelmintic and an agonist of the nicotinic acetylcholine receptor (nAChR). Pyrantel can cause spasmodic muscle paralysis in parasites. Pyrantel can be used in the study of parasitic infections such as ascariasis, hookworm infections, intestinal worms (pinworm infections), trichinosis and trichinosis ^{[1][2]} .																
IC₅₀ & Target	Parasites ^{[1][2]} .																
In Vitro	<p>Pyrantel (10 nM-10 μM; 72 h) shows good anti-A. suum and (0-168.2 M; 72 h) anti-N. americanus activity^{[1][2]}. MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Viability Assay^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>A. suum</td> </tr> <tr> <td>Concentration:</td> <td>10 nM-10 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>72 h</td> </tr> <tr> <td>Result:</td> <td>Inhibited A. suum with a pEC₅₀ value of 7.24.</td> </tr> </table> <p>Cell Viability Assay^[2]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>N. americanus</td> </tr> <tr> <td>Concentration:</td> <td>0-168.2 M (0-100 μg/mL)</td> </tr> <tr> <td>Incubation Time:</td> <td>72 h</td> </tr> <tr> <td>Result:</td> <td>Inhibited third-stage larvae and adult of N. americanus with IC₅₀ values of 2.0 and 7.6 mg/mL, respectively.</td> </tr> </table>	Cell Line:	A. suum	Concentration:	10 nM-10 μM	Incubation Time:	72 h	Result:	Inhibited A. suum with a pEC ₅₀ value of 7.24.	Cell Line:	N. americanus	Concentration:	0-168.2 M (0-100 μg/mL)	Incubation Time:	72 h	Result:	Inhibited third-stage larvae and adult of N. americanus with IC ₅₀ values of 2.0 and 7.6 mg/mL, respectively.
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In Vivo	<p>Pyrantel (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].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p>																

Animal Model:	Male Syrian Golden hamsters (3-week-old; <i>A. ceylanicum</i> -infected) ^[2] .
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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