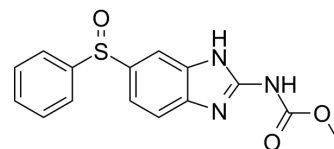


Oxfendazole

Cat. No.:	HY-B0291		
CAS No.:	53716-50-0		
Molecular Formula:	C ₁₅ H ₁₃ N ₃ O ₃ S		
Molecular Weight:	315.35		
Target:	Parasite		
Pathway:	Anti-infection		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	2 years
		-20°C	1 year



SOLVENT & SOLUBILITY

In Vitro

DMSO : 31.25 mg/mL (99.10 mM; ultrasonic and warming and heat to 60°C)
 H₂O : 0.67 mg/mL (2.12 mM; Need ultrasonic)

Concentration	Mass		
	1 mg	5 mg	10 mg
1 mM	3.1711 mL	15.8554 mL	31.7108 mL
5 mM	0.6342 mL	3.1711 mL	6.3422 mL
10 mM	0.3171 mL	1.5855 mL	3.1711 mL

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

In Vivo

- Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline
 Solubility: ≥ 2.5 mg/mL (7.93 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline)
 Solubility: ≥ 2.08 mg/mL (6.60 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% corn oil
 Solubility: ≥ 2.08 mg/mL (6.60 mM); Clear solution

BIOLOGICAL ACTIVITY

Description

Oxfendazole is a sulfoxide form of fenbendazole that is effective when taken orally. Oxfendazole fights parasites and has tumor-promoting activity^{[1][2][3]}.

In Vivo

Oxfendazole (30 mg/kg, oral, single dose) is effective in preventing and controlling natural infections of gastrointestinal nematodes in pigs^[1].
 Oxfendazole (0-500 mg/kg, oral, 8 weeks) has the potential to promote liver tumors in rats^[2].

Pharmacokinetic parameters of Oxfendazole administered in different doses to goats through the rumen or directly into the reticulum^[3]

Route	Dose (mg/kg)	AUC (µg/mL·h)	C _{max} (µg/mL)	T _{max} (h)	βT _{1/2} (h)
Intraruminal	5	12.7	0.35	12	6.5
Intraruminal	10	26.7	0.55	15.5	8.5
Intraruminal	20	35.8	0.61	17.6	9.1
Intra-abomasal	10	18.3	0.9	4.1	/

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

Animal Model: Naturally parasitized pigs^[1]

Dosage: 30 mg/kg; single dose

Administration: Oral

Result: Was effective against natural infections in piglets caused by adult roundworms, esophageal tapeworms, pig tapeworms, and giant roundworms.

Animal Model: Fisher 344 rat^[2]

Dosage: 0, 10, 100, 250, 500 mg/kg; 8 weeks

Administration: Oral

Result: Led to an increase in relative liver weight, with noticeable hypertrophy of centrilobular hepatocytes, a significant increase in smooth endoplasmic reticulum in liver cells, and a marked rise in the number of placental form glutathione S-transferase positive single cells.

CUSTOMER VALIDATION

- Commun Biol. 2024 Jan 24;7(1):123.
- bioRxiv. 2020 Jun.

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REFERENCES

- [1]. Alvarez, L., et al., Efficacy of a single high oxfendazole dose against gastrointestinal nematodes in naturally infected pigs. Veterinary parasitology, 2013. 194(1): p. 70-74.
- [2]. K Mitsumori, et al. Liver Tumour-promoting Effects of oxfendazole in rats. Food Chem Toxicol. 1997 Aug;35(8):799-806.
- [3]. N C Sangster, et al. Disposition of oxfendazole in goats and efficacy compared with sheep. Res Vet Sci. 1991 Nov;51(3):258-63.

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

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA