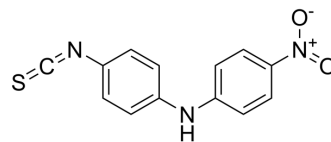


Amoscanate

Cat. No.:	HY-129051
CAS No.:	26328-53-0
Molecular Formula:	C ₁₃ H ₉ N ₃ O ₂ S
Molecular Weight:	271
Target:	Parasite; Oxidative Phosphorylation
Pathway:	Anti-infection; Others
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 : 50 mg/mL (184.50 mM; Need ultrasonic)

Concentration	Solvent	Mass		
		1 mg	5 mg	10 mg
Preparing Stock Solutions	1 mM	3.6900 mL	18.4502 mL	36.9004 mL
	5 mM	0.7380 mL	3.6900 mL	7.3801 mL
	10 mM	0.3690 mL	1.8450 mL	3.6900 mL

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

BIOLOGICAL ACTIVITY

Description

Amoscanate (cgp4540) is phenyl isothiocyanate in which the hydrogen at the para-position has been replaced by a 4-nitroaniliny group. Amoscanate is an anti-schistosomal agent. Amoscanate, as an isothiocyanate compound and uncoupler of oxidative phosphorylation, potently injures rodent ependyma^[1].

IC₅₀ & Target

Schistosome

In Vivo

Amoscanate (500 mg/kg; p.o.; 10 days) destructs ependyma and periventricular brain^[1].
 Amoscanate (250 and 500 mg/kg; p.o.; 28 days) elicits necrosis, Ca⁺⁺-positive microgranules, pyknosis and edema localized in ependyma/subependyma in the medial striatum^[1].
 Amoscanate (25-500 mg/kg; p.o.; 20 days) elicits progressive necrosis of ependyma^[1].
 Amoscanate elicits massive ultrastructural damage in ependymal cells^[1].
 MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model:	Sprague-Dawley rats ^[1]
Dosage:	500 mg/kg

Administration:	P.o.; 10 days
Result:	Destructed ependyma and periventricular brain.
Animal Model:	Sprague-Dawley rats ^[1]
Dosage:	250 and 500 mg/kg
Administration:	P.o.; 28 days
Result:	Elicited necrosis, Ca ⁺⁺ -positive microgranules, pyknosis and edema localized in ependyma/subependyma in the medial striatum.
Animal Model:	Sprague-Dawley rats ^[1]
Dosage:	25~500 mg/kg
Administration:	P.o.; 20 days
Result:	Elicited progressive necrosis of ependymal.

REFERENCES

[1]. Johanson C, et al. The distributional nexus of choroid plexus to cerebrospinal fluid, ependyma and brain: toxicologic/pathologic phenomena, periventricular destabilization, and lesion spread. Toxicol Pathol. 2011;39(1):186-212.

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

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