Dipyrithione

Cat. No.: HY-N8432 CAS No.: 3696-28-4 Molecular Formula: $C_{10}H_8N_2O_2S_2$ Molecular Weight: 252.31

Target: Apoptosis; Bacterial; Fungal Pathway: Apoptosis; Anti-infection Powder -20°C Storage: 3 years

> 4°C 2 years -80°C In solvent 6 months

-20°C 1 month

Product Data Sheet

SOLVENT & SOLUBILITY

In Vitro

DMSO: 10 mg/mL (39.63 mM; ultrasonic and warming and heat to 60°C)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	3.9634 mL	19.8169 mL	39.6338 mL
	5 mM	0.7927 mL	3.9634 mL	7.9268 mL
	10 mM	0.3963 mL	1.9817 mL	3.9634 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: ≥ 1 mg/mL (3.96 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 1 mg/mL (3.96 mM); Clear solution

BIOLOGICAL ACTIVITY

Description Dipyrithione is a potent antimicrobial agent. Dipyrithione shows antifungal activity and antiproliferative activity. Dipyrithione induces apoptosis and cycle arrest at G1 phase. Dipyrithione shows anti-inflammatory activity in vivo. $\label{limited-potential} \mbox{Dipyrithione shows anti-tumor activity. Dipyrithione has the potential for the research of dermatophytosis $^{[1][2][3]}$.}$

In Vitro Dipyrithione (20 μg/mL) shows antifungal activity with MIC values of 6.03 μM for Trichophyton rubrum^[1].

Dipyrithione (72 h) shows cytotoxic activity against 293 T cells with an IC₅₀ value of 0.22 μ M^[1].

Dipyrithione (1-5 μM; 8.5 h) inhibits LPS (100 ng/ml)-induced up-regulation of iNOS and COX-2 in RAW264.7 cells in a dosedependent manner^[2].

Dipyrithione (1 μM; 8.5 h) suppresses LPS-induced increase of iNOS but not COX-2 mRNA level, inhibits LPS-increased NO production^[2].

Dipyrithione (3 μ M; 2, 5 h) decreases phosphorylation of STAT1 induced by LPS and does not influence LPS-induced MAPK and NF- κ B activation in RAW 246.7 cells^[2].

Dipyrithione (0-5 μ g/mL; 48 h) shows antiproliferative activity for KB, 231, U937 and K562 cells in a dose dependent manner [3].

Dipyrithione (2.5 μ g/ml) induces apoptosis and cycle arrest at G1 phase^[3].

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

Western Blot Analysis^[2]

Cell Line:	RAW264.7 cells	
Concentration:	1-5 μΜ	
Incubation Time:	8.5 h	
Result:	Inhibited the expression of LPS (100 ng/ml)-induced up-regulation of iNOS and COX-2 in a dose-dependent manner.	
Cell Proliferation Assay [[]	2]	
Cell Line:	KB, 231, U937, K562 cells	
Concentration:	2.5 μg/ml	
Incubation Time:	24 h	
Result:	Induced cell cycle arrest at G1 phase with induced p21 accumulation, CyclinD1 and CyclinE1 expressions were downregulated.	
Apoptosis Analysis ^[3]		
Cell Line:	KB, 231, U937, K562 cells	
Concentration:	2.5 μg/ml	
Incubation Time:	36 h	
Result:	Induced apoptosis by induced cleavage of caspase-9, caspase-3 and PARP.	
Western Blot Analysis ^[3]		
Cell Line:	RAW264.7 cells	
Concentration:	1-5 μΜ	
Incubation Time:	8.5 h	
Result:	Inhibited the expression of LPS (100 ng/ml)-induced up-regulation of iNOS and COX-2 in a dose-dependent manner.	

In Vivo

Dipyrithione (0.2 mg/cm²; externally once daily for 10 days) shows great anti-dermatophyte activity effects in guinea pig^[1]. Dipyrithione (1, 2.5, 5 mg/kg; i.p.; daily for 10 days) shows anti-tumor acyivity in mouse^[3].

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Animal Model:	Guinea pig (infected with Trichophyton rubrum) $^{[1]}$
Dosage:	0.2 mg/cm ²

Administration:	Externally once daily for 10 days	
Result:	Showed normal hair growth, with no scaly skin.	
Animal Model:	18-22g male ICR mice ²	
Dosage:	1, 2.5, 5 mg/kg	
Administration:	l.p.	
Result:	Raised the survival rate from 10% to 30%, 60% and 90%, respectively.	
Animal Model:	6 weeks, 18-20 g male ICR mice (hepatoma 22 (H22) cells) ^[3]	
Dosage:	2.5 mg/kg	
Administration:	I.p.; daily for 10 days	
Result:	Inhibited the growth of tumor.	

REFERENCES

[1]. Song X, et al. In vivo antifungal activity of dipyrithione against Trichophyton rubrum on guinea pig dermatophytosis models. Biomed Pharmacother. 2018 Dec;108:558-564.

[2]. Liu Z, et al. Dipyrithione inhibits lipopolysaccharide-induced iNOS and COX-2 up-regulation in macrophages and protects against endotoxic shock in mice. FEBS Lett. 2008 May 28;582(12):1643-50.

[3]. Fan Y, et al. Dipyrithione induces cell-cycle arrest and apoptosis in four cancer cell lines in vitro and inhibits tumor growth in a mouse model. BMC Pharmacol Toxicol. 2013 Oct 21;14:54.

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

Tel: 609-228-6898

Fax: 609-228-5909

 $\hbox{E-mail: tech@MedChemExpress.com}$

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