Dapaconazole

MedChemExpress

Cat. No.:	HY-16719		
CAS No.:	1269726-67	-1	
Molecular Formula:	C ₁₉ H ₁₅ Cl ₂ F ₃ N	1 ₂ 0	
Molecular Weight:	415.24		
Target:	Cytochrome P450		
Pathway:	Metabolic Enzyme/Protease		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month

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SOLVENT & SOLUBILITY

		Solvent Mass Concentration	1 mg	5 mg	10 mg		
	Preparing Stock Solutions	1 mM	2.4082 mL	12.0412 mL	24.0825 ml		
		5 mM	0.4816 mL	2.4082 mL	4.8165 mL		
		10 mM	0.2408 mL	1.2041 mL	2.4082 mL		
	Please refer to the so	lubility information to select the app	propriate solvent.				
vo	1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.5 mg/mL (6.02 mM); Clear solution						
		2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: 2.5 mg/mL (6.02 mM); Suspended solution; Need ultrasonic					
		one by one: 10% DMSO >> 90% corn oil ng/mL (6.02 mM); Clear solution					

BIOLOGICAL ACTIVITY			
Description	Dapaconazole, as an antifungal agent, inhibits sterol 14α -demethylase cytochrome P450 activity with an IC ₅₀ of 1.4μ M ^[1] .		
IC₅₀ & Target	Cytochrome P450 (CYP26) 1.4 μM (IC ₅₀)		
In Vitro	Dapaconazole inhibits sterol 14 α -demethylase cytochrome P450 activity with an IC ₅₀ of 1.4 ± 0.3 μ M ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.		

Product Data Sheet

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r=N N∍ In Vivo

Dapaconazole (20 mg/kg; p.o.) shows that the bioavailability is 97.3 $\%^{[1]}$.

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Animal Model:	Male beagle dogs ^[1]
Dosage:	20 mg/kg (Pharmacokinetic Analysis)
Administration:	P.o.
Result:	Showed that the bioavailability was 97.3 %.

REFERENCES

[1]. Juliana SP, et al. Pharmacokinetics of Dapaconazole, A Novel Antifungal Agent, in Beagle Dogs and Inhibition of Cytochrome P450 Family 51. J Eur Acad Dermatol Venereol. 2018 Jun 10.

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