Efinaconazole

Cat. No.:	HY-15660			
CAS No.:	164650-44-6			
Molecular Formula:	$C_{18}H_{22}F_{2}N_{4}O$			
Molecular Weight:	348.39			
Target:	Fungal			
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 : ≥ 100 mg/mL (287.03 mM) * "≥" means soluble, but saturation unknown.						
	Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg		
		1 mM	2.8703 mL	14.3517 mL	28.7035 mL		
		5 mM	0.5741 mL	2.8703 mL	5.7407 mL		
		10 mM	0.2870 mL	1.4352 mL	2.8703 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: ≥ 2.5 mg/mL (7.18 mM); Clear solution						
	2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (7.18 mM); Clear solution						
	3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (7.18 mM); Clear solution						

DIOLOGICAL ACTIV	
Description	Efinaconazole (KP-103) is a triazole antifungal agent and againsts <i>T. mentagrophytes</i> SM-110 and <i>C. albicans</i> ATCC 10231 with MICs of 0.0039 μg/mL and 0.00098 μg/mL, respectively ^[1] . Efinaconazole has a potent in vitro activity against fungal pathogens including dermatophytes, Candida and Malassezia species ^[1] .
IC ₅₀ & Target	Fungal ^[1]





In Vivo

Topical Efinaconazole solution (0.25 to 1%) is dose-dependently effective in treating both dermatophytoses, for 10 guinea pigs with interdigital tinea pedis or tinea corporis is investigated. The follow up study performs on day-30 and day-9 posttreatment demonstrated that the relapse rates for 1% Efinaconazole-treated animals with tinea pedis and for those with tinea corporis are 20 and 30%, respectively. When a single dose of 1% Efinaconazole is applied to the back skin 48 hours before fungal inoculation, 9 of the 10 animals are protected from the dermatophytosis, suggesting that active Efinaconazole is retained in skin tissue for at least 48 hours after dosing^[2].

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

CUSTOMER VALIDATION

• Antimicrob Agents Chemother. 2019 Sep 23;63(10):e00442-19.

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REFERENCES

[1]. Tatsumi Y, et al. Mechanism of action of efinaconazole, a novel triazole antifungal agent. Antimicrob Agents Chemother. 2013 May;57(5):2405-9.

[2]. Tatsum Y, et al. KP-103, a novel triazole derivative, is effective in preventing relapse and successfully treating experimental interdigital tinea pedis and tinea corporis in guinea pigs. Microbiol Immunol. 2002;46(7):425-32.

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

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