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Product Data Sheet

Piperaquine tetraphosphate tetrahydrate

Cat. No.:	HY-B1896B	
CAS No.:	915967-82-7	4 HO-P-OH
Molecular Formula:	$C_{29}H_{32}Cl_{2}N_{6} \cdot 4H_{3}O_{4}P \cdot 4H_{2}O$	N OH
Molecular Weight:	1000	↓ 4 H ₂ O
Target:	Parasite	5
Pathway:	Anti-infection	
Storage:	4°C, sealed storage, away from moisture	
	* In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture)	N

SOLVENT & SOLUBILITY

In Vitro	H ₂ O : 6 mg/mL (6.00 DMSO : < 1 mg/mL (i	mM; Need ultrasonic) nsoluble or slightly soluble)		
		Solvent Mass Concentration	1 mg	5 mg
	Preparing Stock Solutions	1 mM	1.0000 mL	5.0000 mL
	Stock Solutions	5 mM	0.2000 mL	1.0000 mL
		10 mM		

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

BIOLOGICAL ACTIV			
Description	Piperaquine tetraphosphate tetrahydrate is a bisquinoline antimalarial agent. Piperaquine tetraphosphate tetrahydrate can be used in antimalarial research in combination with Artemisinin ^{[1][2]} .		
In Vivo	Piperaquine (10-90 mg/l Piperaquine (90 mg/kg; 33.5 mg•h/L, 1.55 L/h/kg malaria-infected mice ^[1] MCE has not independer	kg; a single i.p.) decreases parasitemia at all of the doses tested in mice ^[1] . a single i.p.) exhibits the t _{1/2} , apparent clearance, and apparent volume of distribution 17.8 days, g, and 956 L/kg, respectively, in healthy mice and 16.1 days, 27.3 mg•h/L, 1.9 L/h/kg, and 1,059 L/kg in ¹ . ntly confirmed the accuracy of these methods. They are for reference only.	
	Animal Model:	Male BALB/c mice (7 to 8 weeks) are inoculated with Plasmodium berghei parasites $^{[1]}$	
	Dosage:	0, 10, 30, 90 mg/kg	
	Administration:	A single i.p. administration	

CI

10 mg

10.0000 mL

2.0000 mL

Result:	The median survival time was 10 days at dose of 10 mg/kg.
	The median survival time was 54 days at dose of 30 mg/kg.
	All mice were active and alert and had stable body weights throughout the course of
	study at dose of 90 mg/kg.
Animal Model:	Male Swiss mice (6 weeks old) ^[1]
Dosage:	90 mg/kg (Pharmacokinetic Analysis)
Administration:	A single i.p. administration
Result:	t _{1/2} =17.8 d; AUC=33.5 mg•h/L; apparent clearance=1.55 L/h/kg; apparent volume of

CUSTOMER VALIDATION

• Cell Rep. 2021 Apr 6;35(1):108959.

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REFERENCES

[1]. Moore BR, et, al. Pharmacokinetics and pharmacodynamics of piperaquine in a murine malaria model. Antimicrob Agents Chemother. 2008 Jan; 52(1): 306-11.

[2]. Davis TME, et, al. Piperaquine: a resurgent antimalarial drug. Drugs. 2005; 65(1): 75-87.

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

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