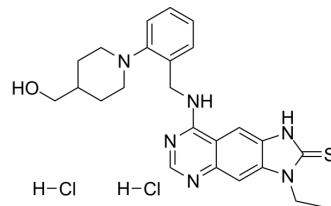


Thioquinapiperifil dihydrochloride

Cat. No.:	HY-119611A
CAS No.:	204077-66-7
Molecular Formula:	C ₂₄ H ₃₀ Cl ₂ N ₆ OS
Molecular Weight:	521.51
Target:	Phosphodiesterase (PDE)
Pathway:	Metabolic Enzyme/Protease
Storage:	4°C, sealed storage, away from moisture and light * In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture and light)



SOLVENT & SOLUBILITY

In Vitro	DMSO : 125 mg/mL (239.69 mM; Need ultrasonic)																					
	<table border="1"> <thead> <tr> <th rowspan="2">Solvent</th> <th rowspan="2">Mass</th> <th colspan="3">Concentration</th> </tr> <tr> <th>1 mg</th> <th>5 mg</th> <th>10 mg</th> </tr> </thead> <tbody> <tr> <td rowspan="3">Preparing Stock Solutions</td> <td>1 mM</td> <td>1.9175 mL</td> <td>9.5875 mL</td> <td>19.1751 mL</td> </tr> <tr> <td>5 mM</td> <td>0.3835 mL</td> <td>1.9175 mL</td> <td>3.8350 mL</td> </tr> <tr> <td>10 mM</td> <td>0.1918 mL</td> <td>0.9588 mL</td> <td>1.9175 mL</td> </tr> </tbody> </table>	Solvent	Mass	Concentration			1 mg	5 mg	10 mg	Preparing Stock Solutions	1 mM	1.9175 mL	9.5875 mL	19.1751 mL	5 mM	0.3835 mL	1.9175 mL	3.8350 mL	10 mM	0.1918 mL	0.9588 mL	1.9175 mL
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	Please refer to the solubility information to select the appropriate solvent.																					
In Vivo	<ol style="list-style-type: none"> Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.08 mg/mL (3.99 mM); Clear solution Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.08 mg/mL (3.99 mM); Clear solution 																					

BIOLOGICAL ACTIVITY

Description	Thioquinapiperifil dihydrochloride (KF31327), a potent, selective and non-competitive phosphodiesterase-5 (PDE-5, IC ₅₀ of 0.074 nM) inhibitor, is used for sexual enhancement study ^{[1][2]} .			
IC₅₀ & Target	PDE5 0.074 nM (IC ₅₀)	PDE1 380 nM (IC ₅₀)	PDE2 670 nM (IC ₅₀)	PDE3 38 nM (IC ₅₀)
	PDE4 800 nM (IC ₅₀)			
In Vitro	Thioquinapiperifil can be found in dietary supplements ^[1] . Thioquinapiperifil dihydrochloride (KF31327) (0.1-10 μM) concentration dependently inhibits platelet aggregation. In the			

absence of nitroglycerin, higher concentrations 1 and 10 μM of Thioquinapiperifil dihydrochloride (KF31327) are required to inhibit platelet aggregation^[2].

Thioquinapiperifil dihydrochloride (KF31327) and shows significant increase in cyclic GMP at 10 μM . After 5 min incubation, the mean cyclic GMP levels of Thioquinapiperifil dihydrochloride (KF31327)-treated cells is 0.95 ± 0.17 pmol/ 10^8 cells^[2].

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

REFERENCES

[1]. Nahoko Uchiyama, et al. Determination of a new type of phosphodiesterase-5 inhibitor, thioquinapiperifil, in a dietary supplement promoted for sexual enhancement. Chem Pharm Bull (Tokyo). 2008 Sep;56(9):1331-4.

[2]. R Hirose, et al. KF31327, a new potent and selective inhibitor of cyclic nucleotide phosphodiesterase 5. Eur J Pharmacol. 2001 Nov 9;431(1):17-24.

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

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