Thioquinapiperifil

Molecular Weight:

Cat. No.: HY-119611 CAS No.: 220060-39-9 Molecular Formula: $C_{24}H_{28}N_6OS$

Target: Phosphodiesterase (PDE) Pathway: Metabolic Enzyme/Protease

448.58

Storage: Please store the product under the recommended conditions in the Certificate of

Analysis.

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Product Data Sheet

BIOLOGICAL ACTIVITY

Description	Thioquinapiperifil (KF31327 free base), a potent, selective and non-competitive phosphodiesterase-5 (PDE-5, IC ₅₀ of 0.074 nM) inhibitor, is used for sexual enhancement study $^{[1][2]}$.				
IC ₅₀ & Target	PDE5 0.074 nM (IC ₅₀) PDE4 800 nM (IC ₅₀)	PDE1 380 nM (IC ₅₀)	PDE2 670 nM (IC ₅₀)	PDE3 38 nM (IC ₅₀)	
In Vitro	Thioquinapiperifil can be found in dietary supplements ^[1] . Thioquinapiperifil (KF31327 free base) (0.1-10 μ M) concentration dependently inhibits platelet aggregation. In the absence of nitroglycerin, higher concentrations 1 and 10 μ M of Thioquinapiperifil (KF31327 free base) are required to inhibit platelet aggregation ^[2] .				

Thioquinapiperifil (KF31327 free base) and shows significant increase in cyclic GMP at 10 μM. After 5 min incubation, the

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.

mean cyclic GMP levels of Thioquinapiperifil (KF31327)-treated cells is 0.95±0.17 pmol/10⁸ cells^[2]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

[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.

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