Product Data Sheet

Picoprazole

 $\begin{array}{lll} \textbf{Cat. No.:} & \text{HY-15384} \\ \textbf{CAS No.:} & 78090\text{-}11\text{-}6 \\ \\ \textbf{Molecular Formula:} & \textbf{C}_{17}\textbf{H}_{17}\textbf{N}_3\textbf{O}_3\textbf{S} \\ \end{array}$

Molecular Weight: 343.4

Target: Proton Pump

Pathway: Membrane Transporter/Ion Channel

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

Analysis.

BIOLOGICAL ACTIVITY

Description	Picoprazole is a specific inhibitor of H $^+$ /K $^+$ -ATPase with IC $_{50}$ of 3.1 \pm 0.4 μ M.
IC ₅₀ & Target	IC50: 3.1±0.4 μM (H ⁺ /K ⁺ -ATPase) ^[1]
In Vitro	Picoprazole inhibits the H $^+$ /K $^+$ -ATPase activity in a concentration-dependent manner. The IC $_{50}$ value is $3.1\pm0.4~\mu\text{M}^{[1]}$. Picoprazole is a specific inhibitor of H $^+$ /K $^+$ -ATPase and binds to 100 -kDa polypeptides of the enzyme, dose dependently inhibited opening of the Cl $^-$ conductance by Cu $^{2+}$ -o-phenanthroline, indicating that the Cl $^-$ conductance is part of the function of the H $^+$ /K $^+$ -ATPase $^{[2]}$. The inhibitory effect of the three benzimidazole derivatives Timoprazole, Picoprazole, and Omeprazole on histamine and dbcAMP stimulated 14 C-aminopyrine accumulation (H $^+$ secretion) has been studied in isolated and enriched guinea-pig parietal cells. All compounds tested inhibit H $^+$ secretion in a concentration dependent manner with IC $_{50}$ values of $8.5\pm1.9~\mu\text{M}$ for Timoprazole, $3.9\pm0.7~\mu\text{M}$ for Picoprazole, and $0.13\pm0.03~\mu\text{M}$ for Omeprazole $^{[3]}$. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Beil W, et al. Inhibition of partially purified H⁺/K⁺-ATPase from guinea-pig isolated and enriched parietal cells by substituted benzimidazoles. Br J Pharmacol. 1984 Jul;82(3):651-7.

[2]. Takeguchi N, et al. Disulfide cross-linking of H,K-ATPase opens Cl- conductance, triggering proton uptake in gastric vesicles. Studies with specific inhibitors. J Biol Chem. 1986 Feb 25;261(6):2560-6.

[3]. Sewing KF, et al. Effect of substituted benzimidazoles on acid secretion in isolated and enriched guinea pig parietal cells. Gut. 1983 Jun;24(6):557-60.

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

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