PRODUCT INFORMATION



Tegoprazan

Item No. 33243

Synonym: CJ-12420 MF: $C_{20}H_{19}F_2N_3O_3$ FW: 387.4 Purify: >98%	CAS Registry No.: Formal Name:	942195-55-3 7-[[(4S)-5,7-difluoro-3,4-dihydro-2H-1- benzopyran-4-yl]oxy]-N,N,2-trimethyl- 1H-benzimidazole-5-carboxamide	H-N
VUN/Vis.: λ_{max} : 220 nmSupplied as:A solidStorage: -20° C	MF: FW: Purity: UV/Vis.: Supplied as:	$C_{20}H_{19}F_2N_3O_3$ 387.4 ≥98% λ _{max} : 220 nm A solid	

Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis.

Laboratory Procedures

Tegoprazan is supplied as a solid. A stock solution may be made by dissolving the tegoprazan in the solvent of choice, which should be purged with an inert gas. Tegoprazan is soluble in DMSO.

Description

Tegoprazan is a proton pump inhibitor that inhibits H⁺/K⁺-ATPase activity in porcine gastric membrane vesicles (IC₅₀ = 0.53 μ M).¹ It completely inhibits basal gastric acid secretion in pylorus-ligated rats when administered at a dose of 10 mg/kg and reduces esophageal mucosal tissue levels of myeloperoxidase (MPO), a marker of tissue damage, in a rat model of reflux esophagitis (ED₅₀ = 5.9 mg/kg). Tegoprazan reduces ulcer area in rat models of gastric ulcers induced by naproxen, water-immersion restraint stress (WIRS), or ethanol ($ED_{50}s = 0.1, 0.1$, and 1.4 mg/kg, respectively).

Reference

1. Kim, D.K., Lee, K.-H., Kim, S.-J., et al. Effects of tegoprazan, a novel potassium-competitive acid blocker, on rat models of gastric acid-related disease. J. Pharmacol. Exp. Ther. 369(3), 318-327 (2019).

WARNING THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

SAFETY DATA

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

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