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**Product Data Sheet**

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Product Name: FRG8701

Cat. No.: GC32050

**Chemical Properties**

Cas. No. 108498-50-6

SMILES O=C(NCCCCOC1=CC=CC(CN2CCCCC2)=C1)CS(CC3=CC=CO3)=OFormula C<sub>22</sub>H<sub>30</sub>N<sub>2</sub>O<sub>4</sub>S M.Wt 418.55

Solubility Soluble in DMSO Storage Store at -20°C

General tips For obtaining a higher solubility , please warm the tube at 37 °C and shake it in the ultrasonic bath for a while. Stock solution can be stored below -20°C for several months.

Shipping Condition Evaluation sample solution : ship with blue ice All other available size: ship with RT , or blue ice upon request.

Structure **Protocol****Cell experiment:**

In this study, male guinea pigs weighing 250-350 g are used. The hearts are excised from animals and right atria are dissected carefully. Atria are suspended in Krebs Henseleit (KH) solution, maintained at 32°C and aerated with 95% O<sub>2</sub> and 5% CO<sub>2</sub>. The tissue is attached with an initial load of 1 g to an isometric transducer and allowed to stabilize for 60 min before histamine application. The histamine response is repeated 10 min after the tissue is incubated with various concentrations of FRG8701 or reference H<sub>2</sub>-antagonist[1].

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

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### Animal experiment:

Gastric mucosal lesions are produced using various kinds of necrotizing agents. One hour after necrotizing agent treatment, the animals are sacrificed. The stomach is removed and inflated by injecting 10 mL of 2% formalin. Subsequently, the stomach is incised along the curvature and examined for macroscopic hemorrhagic damages under a dissecting microscope ( $\times 10$ ). In addition, FRG8701 is given intraperitoneally before HCl-ethanol treatment[1].

### References:

[1]. Shibata M, et al.  
Effects of FRG-8701 on gastric acid secretion, gastric mucosal lesions by necrotizing agents and experimental gastric or duodenal ulcer in rats. Jpn J Pharmacol. 1990 Nov;54(3):277-85.

### Background

FRG-8701 is a new Histamine H<sub>2</sub>-receptor antagonist with an IC<sub>50</sub> of ranging from 0.25 to 0.43  $\mu$ M.

Positive chronotropic response to histamine at 10<sup>-5</sup> M is dose dependently inhibited by FRG8701 (FRG-8701) famotidine or cimetidine; and the IC<sub>50</sub> values of FRG8701, famotidine and cimetidine are 3.3, 3.0 and 108.6 ( $\times 10^{-7}$ M), respectively. The inhibitory potency of FRG8701 is almost the same as that of famotidine and approximately 33 times greater than that of cimetidine[1].

In the pylorus-ligated (4 hr) rats, each drug, given intraduodenally, dose-dependently

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inhibits the total acid output. FRG8701 at 10 or 30 mg/kg, given orally or intraperitoneally, significantly prevent the formation of the gastric mucosal lesions induced by 0.4 N HCl+50% ethanol (HCl•ethanol). Other necrotizing agents-induced gastric lesions are also inhibited by treatment of FRG8701. The oral ED50 values against the lesions range from 1.1 to 9.4 mg/kg. FRG8701, given orally, dose-dependently prevents the development of gastric lesions induced by stress and indomethacin. Duodenal ulcer induced by mepirizole is also inhibited with FRG8701. The ED50 values of FRG8701 for each ulcer model range from 1.7 to 6.9 mg/kg[1].

[1]. Shibata M, et al. Effects of FRG-8701 on gastric acid secretion, gastric mucosal lesions by necrotizing agents and experimental gastric or duodenal ulcer in rats. Jpn J Pharmacol. 1990 Nov;54(3):277-85.

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