
Product Data Sheet

Product Name: Omeprazole-d3-1

Cat. No.: GC64070

Chemical Properties

Cas. No. 934293-92-2

Formula C₁₇H₁₆D₃N₃O₃S

M.Wt 348.43

Solubility DMSO : ≥ 30 mg/mL (86.10 mM); DMF : ≥ 30 mg/mL (86.10 mM); Ethanol : ≥ 5 mg/mL (14.35 mM)

Store
Storage at -
20°CGeneral
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 **Background**

Omeprazole-d₃ is intended for use as an internal standard for the quantification of omeprazole by GC- or LC-MS. Omeprazole is a selective and irreversible inhibitor of the gastric H⁺/K⁺ ATPase pump (IC₅₀ = 1.1 μM).¹ It is a racemic mixture of two enantiomers, S-omeprazole and R-omeprazole (sodium salt) , which are prodrugs of the active sulfonamide which is formed by acid-stimulated conversion.^{2,3} Both enantiomers are extensively metabolized by the cytochrome P450 (CYP) isomers CYP2C19 and CYP3A4.³

1.Smolka, A.J., Goldenring, J.R., Gupta, S., et al. Inhibition of gastric H,K-ATPase activity and gastric epithelial cell IL-8 secretion by the pyrrolizine derivative ML 3000BMC Gastroenterol.4(4)(2004) 2.Richardson, P., Hawkey, C.J., and Stack, W.A. Proton pump inhibitors. Pharmacology and rationale for use in gastrointestinal disorders Drugs56(3)307-335(1998) 3.Shi, S., and Klotz, U. Proton pump inhibitors: An update of their clinical use and pharmacokinetics Eur. J. Clin. Pharmacol.64(10)935-951(2008)

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

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