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## Product Data Sheet

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Product Name: Dazopride (AHR-5531)

Cat. No.: GC33555

### Chemical Properties

Cas. No. 70181-03-2

SMILES O=C(NC1CN(CC)N(CC)C1)C2=CC(Cl)=C(N)C=C2OC

Formula  $C_{15}H_{23}ClN_4O_2$  M.Wt 326.82

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

### Background

Dazopride is an antiemetic agent.

Dazopride (0.3 mg/kg) produces significant enhancement of gastric evacuation and is approximately six times more potent than metoclopramide in gastric evacuation assay. Dazopride (0.3-10.0 mg/kg, i.v.) produces a dose-related increase in antral motility primarily by increasing the amplitude of antral contractions in three conscious dogs. Dazopride significantly reduces the emetic frequency from that of the control group[1]. Dazopride (5 mg/kg, i.p.) antagonises the tetralin-induced emesis in all animals, but fails to antagonise the response at 0.25-2.5 mg/kg. Dazopride fails to modify cisplatin-induced emesis at 0.1 mg/kg (i.v.) although a larger dose of 1.0 mg/kg abolishes or attenuates the response and 5.0 mg/kg of dazopride antagonises the development of emesis in all animals[2].

[1]. Alphin RS, et al. Antagonism of cisplatin-induced emesis by metoclopramide and dazopride through enhancement of gastric motility. Dig Dis Sci. 1986 May;31(5):524-9.

[2]. Costall B, et al. The action of dazopride to enhance gastric emptying and block emesis. Neuropharmacology. 1987 Jul;26(7A):669-77.

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

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