PRODUCT INFORMATION



Dexloxiglumide

Item No. 34759

CAS Registry No.: Formal Name:	119817-90-2 (4R)-4-[(3,4-dichlorobenzoyl)amino]- 5-[(3-methoxypropyl)pentylamino]- 5-oxo-pentanoic acid	
MF:	$C_{21}H_{30}CI_2N_2O_5$	
FW:	461.4	
Purity:	≥98%	
Supplied as:	A solid	
Storage:	-20°C	
Stability:	≥4 years	Cl
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Laboratory Procedures

Dexloxiglumide is supplied as a solid. A stock solution may be made by dissolving the dexloxiglumide in the solvent of choice, which should be purged with an inert gas. Dexloxiglumide is soluble in the organic solvent DMSO (sonicated) at a concentration of approximately 50 mg/ml.

Description

Dexloxiglumide is a cholecystokinin (CCK) receptor antagonist and is the single (R) isomer of loxiglumide (Item No. 25534).^{1,2} It selectively binds to the CCK_1 receptor over CCK_2 in CHO cells expressing the rat receptors (K_is = 0.0234 and 5.88 μ M, respectively).¹ Dexloxiglumide (20 μ mol/kg) inhibits guinea pig gallbladder contractions induced by the CCK agonist CCK octapeptide (sulfated) (CCK-8S; Item No. 23371). It reduces CCK-8-induced gastric emptying in rats ($ID_{50} = 1.14 \text{ mg/kg}$).²

References

- 1. Morton, M.F., Barrett, T.D., Yan, W., et al. 3-[5-(3,4-Dichloro-phenyl)-1-(4-methoxy-phenyl)-1H-pyrazol-3-yl]-2-m-tolyl-propionate (JNJ-17156516), a novel, potent, and selective cholecystokinin 1 receptor antagonist: In vitro and in vivo pharmacological comparison with dexloxiglumide. J. Pharmacol. Exp. Ther. 323(2), 562-569 (2007).
- 2. Scarpignato, C., Kisfalvi, I., D'Amato, M., et al. Effect of dexloxiglumide and spiroglumide, two new CCKreceptor antagonists, on gastric emptying and secretion in the rat: Evaluation of their receptor selectivity in vivo. Aliment. Pharmacol. Ther. 10(3), 411-419 (1996).

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

SAFETY DATA

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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