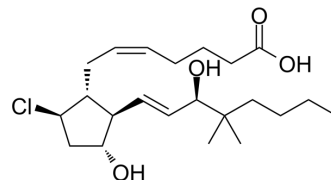


Nocloprost

Cat. No.:	HY-106054
CAS No.:	79360-43-3
Molecular Formula:	C ₂₂ H ₃₇ ClO ₄
Molecular Weight:	400.98
Target:	Prostaglandin Receptor
Pathway:	GPCR/G Protein
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Nocloprost, a prostaglandin E2 (PGE2) analog, is an orally active EP1- and EP3-receptor agonist. Nocloprost inhibits evoked [³ H]ACh release. Nocloprost has gastroprotective and ulcer-healing properties. Nocloprost accelerates the healing of chronic gastric ulcerations and enhances mucosal growth in rats ^{[1][2]} .									
IC₅₀ & Target	EP1	EP3								
In Vivo	<p>Nocloprost (intragastrically; 0.01-10 µg/kg) 30 min before 100% ethanol, acidified Aspirin (HY-14654), acidified Taurocholate (HY-N0545), water immersion, or restraint stress dose dependently prevents the formation of gastric lesions in rats^[1].</p> <p>Nocloprost (i.g.; 0.01-100 µg/kg) fails to affect gastric acid secretion or intestinal secretion (enteropooling) but prevents the increased gastroduodenal alkaline secretion^[1].</p> <p>Nocloprost (s.c.) shows protective activity against ethanol damage but is ineffective when applied intraduodenally^[1].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <table border="1"> <tr> <td>Animal Model:</td> <td>220-250 g rats^[1]</td> </tr> <tr> <td>Dosage:</td> <td>0.01-10 µg/kg</td> </tr> <tr> <td>Administration:</td> <td>Intragastrically; single dose</td> </tr> <tr> <td>Result:</td> <td>30 min before 100% ethanol, acidified Aspirin (ASA), acidified Taurocholate, water immersion, or restraint stress dose dependently prevented the formation of gastric lesions, the ID₅₀ values being 0.25, 0.58, 0.06 and 0.12 µg/kg in rats, respectively.</td> </tr> </table>		Animal Model:	220-250 g rats ^[1]	Dosage:	0.01-10 µg/kg	Administration:	Intragastrically; single dose	Result:	30 min before 100% ethanol, acidified Aspirin (ASA), acidified Taurocholate, water immersion, or restraint stress dose dependently prevented the formation of gastric lesions, the ID ₅₀ values being 0.25, 0.58, 0.06 and 0.12 µg/kg in rats, respectively.
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REFERENCES

- [1]. S J Konturek, et al. Nocloprost, a unique prostaglandin E2 analog with local gastroprotective and ulcer-healing activity. *Eur J Pharmacol.* 1991 Apr 3;195(3):347-57.
- [2]. T Reinheimer, et al. Prostanoid receptors of the EP3 subtype mediate inhibition of evoked [³H]acetylcholine release from isolated human bronchi. *Br J Pharmacol.* 1998 Sep;125(2):271-6.

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

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