

Produktinformation



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Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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PRODUCT INFORMATION



Naloxegol (oxalate)

Item No. 23731

CAS Registry No.: 1354744-91-4

Formal Name: $(5\alpha,6\alpha)-4,5$ -epoxy-6-(3,6,9,12,15,18,21-

heptaoxadocos-1-yloxy)-17-(2-propen-1-yl)-

morphinan-3,14-diol, monoethanedioate

Synonyms: AZ 13337019,

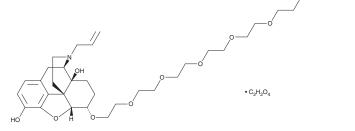
NKTR-118

MF: C₃₄H₅₃NO₁₁ • C₂H₂O₄

FW: 741.8 **Purity:** ≥98% UV/Vis.: λ_{max} : 212 nm A crystalline solid Supplied as:

-20°C Storage: ≥2 years Stability:

Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis.



Laboratory Procedures

Naloxegol (oxalate) is supplied as a crystalline solid. A stock solution may be made by dissolving the naloxegol (oxalate) in the solvent of choice. Naloxegol (oxalate) is soluble in organic solvents such as ethanol, DMSO, and dimethyl formamide, which should be purged with an inert gas. The solubility of naloxegol (oxalate) in these solvents is approximately 33, 20, and 16 mg/ml, respectively.

Further dilutions of the stock solution into aqueous buffers or isotonic saline should be made prior to performing biological experiments. Ensure that the residual amount of organic solvent is insignificant, since organic solvents may have physiological effects at low concentrations. Organic solvent-free aqueous solutions of naloxegol (oxalate) can be prepared by directly dissolving the crystalline solid in aqueous buffers. The solubility of naloxegol (oxalate) in PBS, pH 7.2, is approximately 10 mg/ml. We do not recommend storing the aqueous solution for more than one day.

Description

Naloxegol is a peripherally acting antagonist of the μ -opioid receptor (K_i = 7.42 nM; pA₂ = 7.95). It is selective for the μ -opioid receptor over the δ -opioid receptor ($K_i = 866$ nM). Naloxegol also acts as a partial agonist of κ -opioid receptors in vitro (EC₅₀ = 47 nM for [35 S]GTP γ S binding) but lacks activity ex vivo at concentrations up to 10 µM. In vivo, naloxegol reverses morphine-induced decreases in gastrointestinal motility and analgesia in a hot-plate assay in rats (ED_{50} s = 23.1 and 55.4 mg/kg, respectively), demonstrating a two-fold separation for peripheral versus CNS effects. Naloxegol also exhibits a brain uptake rate comparable to atenolol, a low-permeation standard with no brain uptake, in a rat brain perfusion model.

Reference

1. Floettmann, E., Bui, K., Sostek, M., et al. Pharmacologic profile of naloxegol, a peripherally acting µ-opioid receptor antagonist, for the treatment of opioid-induced constipation. J. Pharmacol. Exp. Ther. 361(2), 280-291 (2017).

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

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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