PRODUCT INFORMATION



(R)-Ofloxacin

Item No. 29601

CAS Registry No.:	100986-86-5	
Formal Name:	(3R)-9-fluoro-2,3-dihydro-3-methyl-	
	10-(4-methyl-1-piperazinyl)-7-oxo-7H-	
	pyrido[1,2,3-de]-1,4-benzoxazine-6-	
	carboxylic acid	
Synonym:	Dovtroflovacin	
Synonym.	Dextronoxacin	
MF:	$C_{18}H_{20}FN_{3}O_{4}$	
FW:	361.4	
Purity:	≥98%	F
UV/Vis.:	λ _{max} : 227, 299 nm	öö
Supplied as:	A crystalline solid	
Storage:	-20°C	
Stability:	≥4 years	
Information represent	s the product specifications. Batch specific analytic	cal results are provided on each certificate of analysis

Laboratory Procedures

(R)-Ofloxacin is supplied as a crystalline solid. A stock solution may be made by dissolving the (R)-ofloxacin in the solvent of choice, which should be purged with an inert gas. (R)-Ofloxacin is soluble in organic solvents such as ethanol, DMSO, and dimethyl formamide (DMF). The solubility of (R)-ofloxacin in ethanol is approximately 1 mg/ml and approximately 20 mg/ml in DMSO and DMF.

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 (R)-ofloxacin can be prepared by directly dissolving the crystalline solid in aqueous buffers. The solubility of (R)-ofloxacin in PBS, pH 7.2, is approximately 10 mg/ml. We do not recommend storing the aqueous solution for more than one day.

Description

(R)-Ofloxacin is a fluoroquinolone antibiotic and the (R) isomer of the antibiotics ofloxacin (Item No. 22891) and levofloxacin (Item No. 20382).¹ It is active against certain Gram-positive and Gram-negative bacteria, including E. coli, P. aeruginosa strains 32104 and 32122, S. aureus strains 209P and Smith, and S. epidermis strain 56556 (MICs = 0.78, 12.5, 6.25, 25, 12.5, and 25 µg/ml, respectively) but not S. epidermis strain 56500, S. pyogenes, or S. faecalis (MICs = >100 μ g/ml for all).² (R)-Ofloxacin inhibits E. coli DNA gyrase with an IC₅₀ value of 75 μ g/ml, which is approximately 30- and 50-fold lower than inhibition by ofloxacin and levofloxacin, respectively.1

References

- 1. Morrissey, I., Hoshino, K., Sato, K., et al. Mechanism of differential activities of ofloxacin enantiomers. Antimicrob. Agents Chemother. 40(8), 1775-1784 (1996).
- 2. Hayakawa, I., Atarashi, S., Yokohama, S., et al. Synthesis and antibacterial activities of optically active ofloxacin. Antimicrob. Agents Chemother. 29(1), 163-164 (1986).

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

SAFFTY 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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1180 EAST ELLSWORTH RD ANN ARBOR, MI 48108 · USA PHONE: [800] 364-9897 [734] 971-3335 FAX: [734] 971-3640 CUSTSERV@CAYMANCHEM.COM WWW.CAYMANCHEM.COM