Proteins

Product Data Sheet

Zabofloxacin hydrochloride

Cat. No.: HY-106410A CAS No.: 623574-00-5 Molecular Formula: $C_{19}H_{21}ClFN_5O_4$

Molecular Weight: 437.85

Target: Bacterial; Topoisomerase; Antibiotic Pathway: Anti-infection; Cell Cycle/DNA Damage Storage: -20°C, sealed storage, away from moisture

* In solvent: -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture)

SOLVENT & SOLUBILITY

In Vitro

DMSO: 6.67 mg/mL (15.23 mM; ultrasonic and warming and heat to 60°C)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.2839 mL	11.4194 mL	22.8389 mL
	5 mM	0.4568 mL	2.2839 mL	4.5678 mL
	10 mM	0.2284 mL	1.1419 mL	2.2839 mL

Please refer to the solubility information to select the appropriate solvent.

BIOLOGICAL ACTIVITY

Description Zabofloxacin hydrochloride (DW-224a) is a potent and seletive inhibitor of the bacterial type II and IV topoisomerases. Zabofloxacin hydrochloride has excellent activity against gram-positive pathogens including Steptococcus aureus, Streptococcus pyogenes and S.pneumonia. Zabofloxacin hydrochloride is a novel fluoronaphthyridone quinolone that is considered as an alternative antibiotic for treatment of quinolone-susceptible (QSSP) and quinolone-resistant gonorrhea

	(QRSP) ^[1] .					
IC ₅₀ & Target	Topoisomerase I	Quinolone	Topoisomerase II	Quinolone		
In Vitro	Zabofloxacin shows a highly potent in vitro activity against clinical isolates of penicillin-sensitive S. pneumoniae (minimum inhibitory concentration, MIC_{90} : 0.03 mg/L) and penicillin-resistant S. pneumoniae (MIC_{90} : 0.03 mg/L). Against quinolone-resistant S. pneumoniae, zabofloxacin (MIC_{90} : 1 mg/L) is more active than ciprofloxacin, sparfloxacin, and moxifloxacin ^[1] .					

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

1]. Park HS, et al. Antimicrobia	al Activity of Zabofloxacin again	st Clinically Isolated Streptococ	cus pneumoniae. Molecules. 2016 Nov 17;21(11). pii: E1562.
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