



Avibactam (free acid)

Catalog No: tcsc0593

Z	Available Sizes
Size:	5mg
Size:	10mg
Size:	50mg
	Specifications
CAS N 11925	lo: 00-31-4
Formu	ula: N ₃ O ₆ S
Pathw Anti-in	ray: fection
Targe Bacter	
Purity >98%	/ Grade:
Solub 10 mM	ility: 1 in DMSO
	native Names: 04 (free acid)
Obser 265.24	ved Molecular Weight:
Drod	uct Description





Avibactam (NXL-104) free acid is a covalent, reversible β -lactamase inhibitor, inhibits β -lactamase **TEM-1** and **CTX-M-15** with **IC**₅₀ of 8 nM and 5 nM, respectively.

IC50 & Target: IC50: 8 nM (TEM-1), 5 nM (CTX-M-15)[1]

In Vitro: Avibactam (NXL104) is a molecule with little antibacterial activity, that inhibits class A and C β -lactamases. Avibactam (NXL104) inactivates most important β -lactamases except metallo types and Acinetobacter OXA carbapenemases^[2].

In Vivo: Avibactam (NXL104) sodium displays a slow return of activity with an off-rate of $0.045\pm0.022~\text{min}^{-1}$, which converts to a residence time half-life ($t_{t1/2}$) of $16\pm8~\text{min}$. The measured off-rate for Avibactam (NXL104) suggests that slow deacylation through hydrolysis or reversibility is occurring, and it is in contrast to previously reported extremely long $t_{1/2}$ values of >1 or >7 d for Avibactam (NXL104) inhibition of TEM-1^[1]. Avibactam is a new promising β-lactamase inhibitor, to overcome resistance caused by β-lactamases. Mice are infected with ca.10⁶ CFU of *Pseudomonas aeruginosa* intramuscularly into the thigh or intranasally to cause pneumonia and are given 8 different (single) subcutaneous doses of Ceftazidime and Avibactam (NXL104) in various combined concentrations, ranging from 1 to 128 mg/kg of body weight in 2-fold increases. The mean estimated half-life in plasma of Ceftazidime in the terminal phase is 0.28 h (SD, 0.02 h), and that of Avibactam is 0.24 h (SD, 0.04 h). Volumes of distribution are 0.80 liters/kg (SD, 0.14 liters/kg) and 1.18 liters/kg (SD, 0.34 liters/kg), respectively^[3].

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