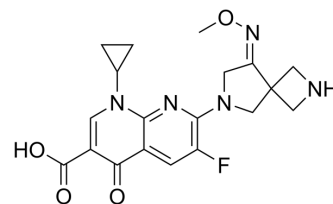


## Zabofloxacin

<b>Cat. No.:</b>	HY-106410		
<b>CAS No.:</b>	219680-11-2		
<b>Molecular Formula:</b>	C <sub>19</sub> H <sub>20</sub> FN <sub>5</sub> O <sub>4</sub>		
<b>Molecular Weight:</b>	401.39		
<b>Target:</b>	Bacterial; Topoisomerase; Antibiotic		
<b>Pathway:</b>	Anti-infection; Cell Cycle/DNA Damage		
<b>Storage:</b>	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



### BIOLOGICAL ACTIVITY

<b>Description</b>	Zabofloxacin (DW-224a Free base) is a potent and selective inhibitor of the bacterial type II and IV topoisomerases. Zabofloxacin has excellent activity against gram-positive pathogens including <i>Staphylococcus aureus</i> , <i>Streptococcus pyogenes</i> and <i>S. pneumoniae</i> . Zabofloxacin is a novel fluoronaphthyridone quinolone that is considered as an alternative antibiotic for treatment of quinolone-susceptible (QSSP) and quinolone-resistant gonorrhea (QRSP) <sup>[1]</sup> .		
<b>IC<sub>50</sub> &amp; Target</b>	Topoisomerase I	Topoisomerase II	Quinolone
<b>In Vitro</b>	Zabofloxacin shows a highly potent in vitro activity against clinical isolates of penicillin-sensitive <i>S. pneumoniae</i> (minimum inhibitory concentration, MIC <sub>90</sub> : 0.03 mg/L) and penicillin-resistant <i>S. pneumoniae</i> (MIC <sub>90</sub> : 0.03 mg/L). Against quinolone-resistant <i>S. pneumoniae</i> , zabofloxacin (MIC <sub>90</sub> : 1 mg/L) is more active than ciprofloxacin, sparfloxacin, and moxifloxacin <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.		

### REFERENCES

[1]. Park HS, et al. Antimicrobial Activity of Zabofloxacin against Clinically Isolated *Streptococcus pneumoniae*. *Molecules*. 2016 Nov 17;21(11). pii: E1562.

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

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