



## **Gatifloxacin**

Catalog No: tcsc1841



## **Available Sizes**

Size: 1g

Size: 5g



## **Specifications**

CAS No:

112811-59-3

Formula:

 $C_{19}^{H_{22}}FN_{3}^{O_{4}}$ 

**Pathway:** 

Anti-infection

**Target:** 

Bacterial

**Purity / Grade:** 

>98%

**Solubility:** 

H2O: 1 mg/mL (2.66 mM; Need ultrasonic)

**Alternative Names:** 

BMS 206584-01;PD 135432;AM-1155

**Observed Molecular Weight:** 

375.39

## **Product Description**

Gatifloxacin is an antibiotic of the fourth-generation fluoroquinolone family, it inhibits the bacterial enzymes DNA gyrase and topoisomerase IV.





Target: Antibacterial

Gatifloxacin is an antibiotic of the fourth-generation fluoroquinolone family, that like other members of that family, inhibits the bacterial enzymes DNA gyrase and topoisomerase IV. Gatifloxacin had activity equal to that of tosufloxacin and activity more potent than those of norfloxacin, ofloxacin, ciprofloxacin, and sparfloxacin against the second-step mutants (grlA gyrA; gatifloxacin MIC range, 1.56 to 3.13 microg/ml) and had the most potent activity against the third-step mutants (grlA gyrA grlA; gatifloxacin MIC range, 1.56 to 6.25 microg/ml), suggesting that gatifloxacin possesses the most potent inhibitory activity against singly mutated topo IV and singly mutated DNA gyrase among the quinolones tested [1].

Ophthalmic gatifloxacin 0.3% is at least as effective as ciprofloxacin at healing corneal ulcers infected with Pseudomonas aeruginosa when gatifloxacin is administered less frequently than ciprofloxacin. Trends favored gatifloxacin in fluorescein retention scores [2].

Clinical indications: Bacterial infection

FDA Approved Date:

Toxicity: Hepatotoxicity; Acute pancreatitis [3]; Torsades de pointes [4]

All products are for RESEARCH USE ONLY. Not for diagnostic & therapeutic purposes!