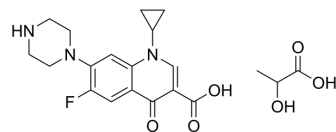


## Ciprofloxacin lactate

<b>Cat. No.:</b>	HY-W040298
<b>CAS No.:</b>	97867-33-9
<b>Molecular Formula:</b>	C <sub>20</sub> H <sub>24</sub> FN <sub>3</sub> O <sub>6</sub>
<b>Molecular Weight:</b>	421.42
<b>Target:</b>	Topoisomerase; Apoptosis; Antibiotic; Bacterial; Mitochondrial Metabolism; Reactive Oxygen Species
<b>Pathway:</b>	Cell Cycle/DNA Damage; Apoptosis; Anti-infection; Metabolic Enzyme/Protease; Immunology/Inflammation; NF-κB
<b>Storage:</b>	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	Ciprofloxacin (Bay-09867) lactate is a potent, orally active topoisomerase IV inhibitor. Ciprofloxacin lactate induces mitochondrial DNA and nuclear DNA damage and lead to mitochondrial dysfunction, ROS production. Ciprofloxacin lactate has anti-proliferative activity and induces apoptosis. Ciprofloxacin lactate is a fluoroquinolone antibiotic, exhibiting potent antibacterial activity <sup>[1][2][3][4]</sup> .																
<b>IC<sub>50</sub> &amp; Target</b>	Quinolone																
<b>In Vitro</b>	<p>Ciprofloxacin (Bay-09867) lactate (5-50 µg/mL; 0-24 h; tendon cells) inhibits cell proliferation and causes cell cycle arrest at the G2/M phase<sup>[1]</sup>.</p> <p>Ciprofloxacin (Bay-09867) lactate shows potent activity against <i>Y. pestis</i> and <i>B. anthracis</i> with MIC<sub>90</sub> of 0.03 µg/mL and 0.12 µg/mL, respectively<sup>[2]</sup>.</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Cycle Analysis<sup>[1]</sup></p> <table border="1"> <tr> <td>Cell Line:</td> <td>Tendon cells</td> </tr> <tr> <td>Concentration:</td> <td>5, 10, 20 and 50 µg/mL</td> </tr> <tr> <td>Incubation Time:</td> <td>24 hours</td> </tr> <tr> <td>Result:</td> <td>Decreased the cellularity of tendon cells.</td> </tr> </table> <p>Apoptosis Analysis<sup>[1]</sup></p> <table border="1"> <tr> <td>Cell Line:</td> <td>Tendon cells</td> </tr> <tr> <td>Concentration:</td> <td>50 µg/mL</td> </tr> <tr> <td>Incubation Time:</td> <td>24 hours</td> </tr> <tr> <td>Result:</td> <td>Arrested cell cycle at the G2/M phase and inhibited cell division in tendon cells.</td> </tr> </table> <p>Western Blot Analysis<sup>[1]</sup></p>	Cell Line:	Tendon cells	Concentration:	5, 10, 20 and 50 µg/mL	Incubation Time:	24 hours	Result:	Decreased the cellularity of tendon cells.	Cell Line:	Tendon cells	Concentration:	50 µg/mL	Incubation Time:	24 hours	Result:	Arrested cell cycle at the G2/M phase and inhibited cell division in tendon cells.
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Cell Line:	Tendon cells
Concentration:	50 µg/mL
Incubation Time:	0, 6, 12, 17 and 24 hours
Result:	Down-regulated the expression of CDK-1 and cyclin B protein and mRNA. Up-regulated the expression of PLK-1 protein.

#### In Vivo

Ciprofloxacin (Bay-09867) lactate (30 mg/kg; i.p.; for 24 hours; BALB/c mice) has protection against *Y. pestis* in murine model of pneumonic plague<sup>[3]</sup>.

Ciprofloxacin (Bay-09867) lactate (100 mg/kg; i.g.; daily, for 4 weeks; C57BL/6J mice) accelerates aortic root enlargement and increases the incidence of aortic dissection and rupture by decreases LOX level and increases MMP levels and activity in the aortic wall<sup>[4]</sup>.

Ciprofloxacin (Bay-09867) lactate (100 mg/kg; i.g.; daily, for 4 weeks; C57BL/6J mice) induces DNA damage and release of DNA to the cytosol, mitochondrial dysfunction, and activation of cytosolic DNA sensor signaling. Ciprofloxacin lactate increases apoptosis and necroptosis in the aortic wall<sup>[4]</sup>.

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

Animal Model:	BALB/c mice <sup>[3]</sup>
Dosage:	30 mg/kg
Administration:	Intraperitoneal injection; for 24 hours
Result:	Reduced the lung bacterial load in murine model of pneumonic plague.

Animal Model:	C57BL/6J mice <sup>[4]</sup>
Dosage:	100 mg/kg
Administration:	Oral gavage; daily, for 4 weeks
Result:	Had aortic destruction that was accompanied by decreased LOX expression and increased MMP expression and activity.

Animal Model:	C57BL/6J mice <sup>[4]</sup>
Dosage:	100 mg/kg
Administration:	Oral gavage; daily, for 4 weeks
Result:	Caused mitochondrial DNA and nuclear DNA damage, leading to mitochondrial dysfunction and ROS production. Increased apoptosis and necroptosis in the aortic wall.

#### CUSTOMER VALIDATION

- Adv Sci (Weinh). 2020 Jul 21;7(17):2001374.
- Water Res. 2023 May 21, 120110.
- Genome Biol. 2023 Apr 30;24(1):98.

- EBioMedicine. 2022 Apr;78:103943.
- Int J Antimicrob Agents. 2018 Aug;52(2):269-271.

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## REFERENCES

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- [1]. Tsai WC, et, al. Ciprofloxacin-mediated cell proliferation inhibition and G2/M cell cycle arrest in rat tendon cells. *Arthritis Rheum*. 2008 Jun;58(6):1657-63.
- [2]. Steenbergen J, et, al. In Vitro and In Vivo Activity of Omadacycline against Two Biothreat Pathogens, *Bacillus anthracis* and *Yersinia pestis*. *Antimicrob Agents Chemother*. 2017 Apr 24;61(5):e02434-16.
- [3]. Hamblin KA, et, al. Inhaled Liposomal Ciprofloxacin Protects against a Lethal Infection in a Murine Model of Pneumonic Plague. *Front Microbiol*. 2017 Feb 6;8:91.
- [4]. LeMaire SA, et, al. Effect of Ciprofloxacin on Susceptibility to Aortic Dissection and Rupture in Mice. *JAMA Surg*. 2018 Sep 1;153(9):e181804.
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**Caution: Product has not been fully validated for medical applications. For research use only.**

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