

<b>Product Name:</b>	Ciprofloxacin hydrochloride
<b>Product Number:</b>	C032
<b>CAS Number:</b>	86393-32-0
<b>Molecular Formula:</b>	$C_{17}H_{18}FN_3O_3 \cdot HCl \cdot H_2O$
<b>Molecular Weight:</b>	385.82 g/mol
<b>Form:</b>	Powder
<b>Appearance:</b>	White or off-white crystalline powder
<b>Solubility:</b>	sparingly soluble in aqueous solution (1.1 µg/mL).
<b>Source:</b>	Synthetic
<b>Water Content (Karl Fischer):</b>	4.7-6.7%
<b>pH:</b>	3.0-4.5
<b>Storage Conditions:</b>	2-8 °C, protected from light.
<b>Description:</b>	<p>Ciprofloxacin hydrochloride is a broad-spectrum second generation 4-fluoroquinolone antibiotic that targets the bacterial enzyme DNA gyrase and is effective against Gram-positive and Gram-negative bacteria and <i>Mycoplasmas</i>. Ciprofloxacin HCl is freely soluble in aqueous solution (35 mg/mL).</p> <p>TOKU-E offers two forms of Ciprofloxacin:</p> <ul style="list-style-type: none"> <li>• Ciprofloxacin hydrochloride (C032)</li> <li>• <u>Ciprofloxacin</u> (C031)</li> </ul>
<b>Mechanism of Action:</b>	Fluoroquinolone antibiotics target bacterial DNA gyrase (topoisomerase type II), an enzyme which reduces DNA strain during replication and is essential for DNA packaging, transcription, and replication. Thus, DNA synthesis and cell division is inhibited.
<b>Spectrum:</b>	Ciprofloxacin is a broad spectrum antibiotic targeting a wide variety of Gram-positive and Gram-negative bacteria. It is also effective against <i>Mycoplasmas</i> .
<b>Microbiology Applications</b>	<p>Ciprofloxacin HCl is commonly used in clinical <i>in vitro</i> microbiological antimicrobial susceptibility tests (panels, discs, and MIC strips) against gram positive and gram negative microbial isolates. Medical microbiologists use AST results to recommend antibiotic treatment options for infected patients.</p> <ul style="list-style-type: none"> <li>• For a complete list of ciprofloxacin MIC values, <a href="#">click here</a>.</li> </ul>

## Plant Biology Applications

Garlic protoplasts were cultured in medium containing ciprofloxacin, and the compound was successful at preventing microbial growth and was not toxic to the protoplasts. (Fellner, 1995)

## Cancer Applications

Two transitional cell carcinoma cell lines, MBT12 and T24, were used in an *in vitro* study of the effects of ciprofloxacin on cell proliferation, and the compound was found to inhibit cell proliferation in a dose-dependent manner (Ebisuno et al, 1997).

## References:

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Gignac SM, Brauer S, Hane B, Quentmeier H, and Drexler HG (1991) Elimination of mycoplasma from infected leukemia cell lines. Leukemia 5(2):162-165 PMID 2020199

Ridgway GL, Mumtaz G, Gabriel FG, and Oriel JD (1986) The Activity of Ciprofloxacin and Other 4-Quinolones Against *Chlamydia trachomatis* and *Mycoplasmas In Vitro*. In: Neu HC, Reeves DS. (eds) Ciprofloxacin. Current Topics in Infectious Diseases and Clinical Microbiology, vol 1. Vieweg+Teubner Verlag, Wiesbaden

Schmitt K, Däubener W, Bitter-Suermann D, Hadding U.(1988) A safe and efficient method for elimination of cell culture mycoplasmas using ciprofloxacin. 109(1):17-25 PMID 3282011

Wolfson, JS and Hooper DC (1985) The Fluoroquinolones: Structures, Mechanisms of Action and Resistance, and Spectra of Activity *in Vitro*. Antimicrob Agents Chemother. 28(4):581-586 PMID 3000292

Zeiler HJ and Grohe K. (1986) The *In Vitro* and *In Vivo* Activity of Ciprofloxacin. In: Neu HC, Reeves DS (eds) Ciprofloxacin. Current Topics in Infectious Diseases and Clinical Microbiology, vol 1. Vieweg+Teubner Verlag, Wiesbaden