

Clindamycin Hydrochloride PRODUCT DATA SHEET

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Product Name: Clindamycin Hydrochloride

Product Number: C035

CAS Number: 21462-39-5

Molecular Formula: $C_{18}H_{33}CIN_2O_5S \cdot HCI$

Molecular Weight: 461.44 g/mol

Form: Powder

Appearance: White or almost white crystalline powder

Solubility: soluble in aqueous solution

Source: Semi-synthetic

Water Content (Karl

Fischer):

3.0-6.0%

Potency (on a dry basis): $\geq 800 \mu g/mg (C_{18}H_{33}CIN_2O_5S)$

pH: 3.0-5.5

Storage Conditions: 2-8 °C. Store in airtight container.

Description: Clindamycin Hydrochloride is a broad-spectrum antibiotic and antiparasitic

agent. It is a semi-synthetic derivative of Lincomycin, a natural lincosamide isolated from *Streptomyces lincolnensis* in 1966. Clindamycin Hydrochloride

is freely soluble in water.

We also offer:

Clindamycin (C233)

Clindamycin Phosphate (<u>C036</u>)

Mechanism of Action: Lincosamides inhibit bacterial protein synthesis by binding the 50S ribosomal

subunit and interfering with tRNA activity during translation.

Spectrum: Clindamycin is a broad-spectrum antibiotic targeting primarily Gram-positive

and Gram-negative bacteria such as Clostridium and Bacteroides species.

Microbiology Applications Clindamycin is commonly used in clinical in vitromicrobiological antimicrobial

susceptibility tests (panels, discs, and MIC strips) against gram positive and gram negative anaerobes. Medical microbiologists use AST results to recommend antibiotic treatment options. Representative ranges include:

• Clostridium difficile 0.25 μg/mL - 32 μg/mL

Bacteroides fragilis 0.25 μg/mL - 4 μg/mL

For a complete list of Clindamycin MIC values, click here.

References:

Dhawan VK and <u>Thadepalli H</u>. (1982) Clindamycin: A review of fifteen years of experience. Clin. Infect. Dis. 4(6):1133-1153 PMID 6818656

Li LH, Kuentzel K L, Shugars KD and Bhuyan BK (1977) Cytotoxicity of several marketed antibiotics on mammalian cells in culture. J. Antibiot (Tokyo) 30(6):506-512 PMID 560364

Lovmar, M and Tanel T (2003) The Mechanism of action of macrolides, lincosamides and streptogramin B reveals the nascent peptide exit path in the ribosome. *J. Molec. Microbiol.* 330(5): 1005-014 PMID 12860123

Magerlein BJ et al (1966) Chemical modification of lincomycin. Antimicrob Agents Chemother. 6:727 PMID 5985307

Wijsman JA, Dekaban GA and Rieder MJ (2013) Differential toxicity of reactive metabolites of clindamycin and sulfonamides in HIV-infected cells: Influence of HIV infection on clindamycin toxicity in vitro. J. Clin. Pharmacol. 45(3):346-351 PMID 15703369

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