PRODUCT INFORMATION



Indolmycin

Item No. 15385

CAS Registry No.: 21200-24-8

5S-[(1R)-1-(1H-indol-3-yl)ethyl]-2-Formal Name:

(methylamino)-4(5H)-oxazolone

Synonyms: PA 155A, TAK-083

MF: $C_{14}H_{15}N_3O_2$ 257.3 FW: **Purity:** ≥95% Supplied as: A powder

Storage: -20°C Stability: ≥4 years

Item Origin: Bacterium/Streptomyces sp.

Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis.

Laboratory Procedures

Ilndolmycin is supplied as a powder. A stock solution may be made by dissolving the indolmycin in the solvent of choice, which should be purged with an inert gas. Indolmycin is soluble in organic solvents such as ethanol, methanol, DMSO, and dimethyl formamide.

Description

Indolmycin is an antibiotic isolated from various Streptomyces strains that functions by competitively inhibiting the prokaryotic tryptophanyl-tRNA synthetase (IC₅₀ = 30 μ M) without significantly affecting the corresponding eukaryotic synthetase. 1,2 It exhibits antibacterial activity against mupirocin- and fusidic acid-resistant strains of Staphylococci.3

References

- 1. Werner, R.G., Thorpe, L.F., Reuter, W., et al. Indolmycin inhibits prokaryotic tryptophanyl-tRNA ligase. Eur. J. Biochem. 68(1), 1-3 (1976).
- 2. Bogosian, G., Haydock, P.V., and Somerville, R.L. Indolmycin-mediated inhibition and stimulation of transcription at the trp promoter of Escherichia coli. J. Bacteriol. 153(2), 1120-1123 (1983).
- Hurdle, J.G., O'Neill, A.J., and Chopra, I. Anti-staphylococcal activity of indolmycin, a potential topical agent for control of staphylococcal infections. J. Antimicrob. Chemother. 52(4), 549-552 (2004).

WARNING
THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

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