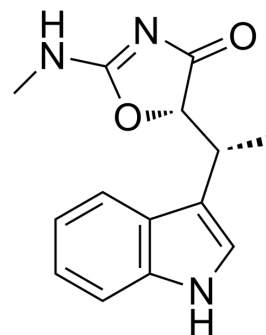


Indolmycin

| | |
|--------------------|---|
| Cat. No.: | HY-117319 |
| CAS No.: | 21200-24-8 |
| Molecular Formula: | C ₁₄ H ₁₅ N ₃ O ₂ |
| Molecular Weight: | 257.29 |
| Target: | Antibiotic; Bacterial |
| Pathway: | Anti-infection |
| Storage: | Please store the product under the recommended conditions in the Certificate of Analysis. |



BIOLOGICAL ACTIVITY

| | |
|--------------------|---|
| Description | Indolmycin (TAK-083), an antibiotic, is a competitive inhibitor of prokaryotic tryptophanyl-tRNA synthetase (TrpS). Indolmycin (TAK-083) possesses both anti-viral and anti-bacterial activity ^{[1][2][3]} . |
| In Vitro | Indolmycin was bacteriostatic and demonstrated good activity against MSSA (methicillin-susceptible <i>Staphylococcus aureus</i>), MRSA (methicillin-resistant <i>S. aureus</i>) and VISA (vancomycin-intermediate <i>S. aureus</i>), including strains resistant to mupirocin or fusidic acid. Indolmycin MICs for 20 strains ranged from 8 to 32 mg/L, whereas a single strain exhibited high-level resistance (MIC 128 mg/L) ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. |

REFERENCES

- [1]. R G Werner, et al. Indolmycin inhibits prokaryotic tryptophanyl-tRNA ligase. *Eur J Biochem.* 1976 Sep;68(1):1-3.
- [2]. Julian G Hurdle, et al. Anti-staphylococcal activity of indolmycin, a potential topical agent for control of staphylococcal infections. *J Antimicrob Chemother.* 2004 Aug;54(2):549-52.
- [3]. M R Harnden, et al. Thiazolinone analogues of indolmycin with antiviral and antibacterial activity. *J Med Chem.* 1978 Jan;21(1):82-7.

Caution: Product has not been fully validated for medical applications. For research use only.

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