## PRODUCT INFORMATION



LL-Z 1640-4 Item No. 19159

CAS Registry No.: 66018-41-5

Formal Name: 3,4,7,8,9,10-hexahydro-7,8,9,16-

tetrahydroxy-14-methoxy-3-methyl-1H-2-

benzoxacyclotetradecin-1-one

Synonyms: Antibiotic LL-Z 1640-4, (5Z)-Zeaenol

MF:  $C_{19}H_{24}O_7$ FW: 364.4 ≥99% **Purity:** Supplied as: A solid Storage: -20°C Stability: ≥4 years

Item Origin: Fungus/Curvularia sp.

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

## **Laboratory Procedures**

LL-Z 1640-4 is supplied as a solid. A stock solution may be made by dissolving the LL-Z 1640-4 in the solvent of choice, which should be purged with an inert gas. LL-Z 1640-4 is soluble in organic solvents such as ethanol, methanol, DMSO, and dimethyl formamide.

## Description

LL-Z 1640-4 is a cis-enol resorcylic acid lactone that has been shown to exhibit both antiviral and antiprotozoan activity. While LL-Z 1640-4 is inactive against JNK/p38 kinase signaling, its companion antibiotic, (5Z)-7-oxo zeaenol (Item No. 17459), has been identified as an irreversible inhibitor of the MAP3K TAK1, blocking T cell activation.<sup>2</sup> Thus, LL-Z 1640-4 is useful as a negative control to help dissect the selectivity of this MAP3K inhibitor.

## References

- 1. Ellestad, G.A., Lovell, F.M., Perkinson, N.A., et al. New zearalenone related macrolides and isocoumarins from an unidentified fungus. J. Org. Chem. 43(12), 2339-2343 (1978).
- 2. Sellès, P. and Lett, R. Convergent stereospecific synthesis of LL-Z1640-2 (or C292), hypothemycin and related macrolides. Part 2. Tetrahedron Lett. 43(26), 4627-4631 (2002).

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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