

PRODUCT DATA SHEET

Code No.: BIA-C1015

Pack sizes: 0.5 mg, 2.5 mg

Synonyms :

Cercosporamide

Specifications

CAS # : **131436-22-1**Molecular Formula : **C**₁₆**H**₁₃**NO**₇

Molecular Weight : 331.3

Source : Cercosporidium sp.

Appearance : Light to dark tan solid

Purity : >95% by HPLC

Long Term Storage : -20°C

Solubility : Soluble in ethanol, methanol, DMF or DMSO. Poor water solubility.

Application Notes

Cercosporamide was originally identified as a host-selective phytotoxin and broad spectrum antifungal agent isolated from Cercosporidium henningsii. More recently, cercosporamide was shown to inhibit a cell wall integrity pathway mediated through a serine/threonine protein kinase, Pkc1, that is central to cell wall biosynthesis. It is both a potent (IC50<50nM) and selective inhibitor of Pkc1 kinase.

References

- 1. Discovery of cercosporamide, a known antifungal natural product, as a selective Pkc1 kinase inhibitor through high-throughput screening. Sussman A. et al., Eukaryot Cell. 2004, 3, 932.
- 2. Cloning and characterization of KNR4, a yeast gene involved in (1,3)-beta-glucan synthesis. Hong Z. et al., Mol. Cell Biol. 1994, 14, 1017.
- 3. The structure and biological activity of cercosporamide from Cercosporidium henningsii. Sugawara F. et al., J. Org. Chem. 1991, 56, 909.

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