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Product Information

Product ID C3208 CAS No. 41621-49-2

Chemical Name

Synonym

Formula C₁₂H₁₇NO₂ • C₂H₇NO

Formula Wt. 268.36 Melting Point 143°C Purity ≥98%

Solubility DMSO (<1 mg/mL), Water (<1

mg/mL), Ethanol (50 mg/mL), Ethanol/Water 1:1 (100 mg/mL), Ethanol/Water 2:1

Store Temp Ambient Ship Temp Ambient

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Pricing and Availability

Bulk quanitites available upon request

Product ID	Size	List Price
C3208	1 g	\$60.20
C3208	5 g	\$191.10

Description Ciclopirox is a hydroxypyridone compound that exhibits antifungal, anti-inflammatory, anti-angiogenic, and anticancer chemotherapeutic activities. This compound acts as a metal ion chelator, preventing peroxide degradation. Ciclopirox modulates generation of ROS in a PKA/Ras1/Ras2-dependent manner, inducing DNA damage and cell death in Candida and Saccharomyces. Ciclopirox also inhibits mTOR, enhancing anticancer activity of other compounds. In breast cancer, colon adenocarcinoma, and rhabdomyosarcoma cells, this compound downregulates expression of cyclins A, B1, D1, and E, suppresses CDK2 and CDK4, and upregulates expression of p21, inducing G0/G1 phase cell cycle arrest and caspase-mediated apoptosis. In animal models, ciclopirox inhibits tumor growth of breast cancer xenografts. Additionally, ciclopirox inhibits expression of VEGFR3, preventing activation of ERK1/2 and tube formation in vitro. TEST!!!!!!

References Sen S, Hassane DC, Corbett C, et al. Novel mTOR inhibitory activity of ciclopirox enhances parthenolide antileukemia activity. Exp Hematol. 2013 Sep;41(9):799-807.e4. PMID: 23660068.

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> Luo Y, Zhou H, Liu L, et al. The fungicide ciclopirox inhibits lymphatic endothelial cell tube formation by suppressing VEGFR-3mediated ERK signaling pathway. Oncogene. 2011 May 5;30(18):2098-107. PMID: 21217783.

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Caution: This product is intended for laboratory and research use only. It is not for human or drug use.