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Product Information

Product ID M9634 CAS No. 35891-70-4

Chemical Name

Synonym

Formula C₂₁H₃₉NO₆ Formula Wt. 401.54 Melting Point 175-180C Purity ≥99%, TLC

Solubility DMSO (25 mg/ml; heat briefly

in boiling water bath); soluble in dilute base (50 mM NaOH) to ~5 mg/ml and methanol: 2

Store Temp 4°C' Ship Temp Ambient

Description Myriocin is derived from ascomycete *Isaria sinclairii*. Myriocin is the parent compound of fingolimod, an immunosuppressive sphingosine-1 receptor inhibitor. Myriocin exhibits both immunosuppressive and anticancer activities. This compound inhibits serine palmitoyltransferase, preventing sphingolipid formation. In vitro, myriocin decreases expression of cylin B1, cdc2, and

> cdc25C and increase expression of p53 and p21, inducing G2/M phase cell cycle arrest and inhibiting growth of melanoma cells. In animal models, myriocin decreases levels of CD4+ lymphocytes and alters the total lymphocyte populations.

Pricing and Availability

Bulk quanitites available upon request

Product ID	Size	List Price
M9634	1 mg	\$37.30
M9634	5 mg	\$124.00
M9634	10 mg	\$206.60

References Wadsworth JM, Clarke DJ, McMahon SA, et al. The chemical basis of serine palmitoyltransferase inhibition by myriocin. J Am Chem Soc. 2013 Sep 25;135(38):14276-85. PMID: 23957439.

> Lee YS, Choi KM, Choi MH, et al. Serine palmitoyltransferase inhibitor myriocin induces growth inhibition of B16F10 melanoma cells through G(2) /M phase arrest. Cell Prolif. 2011 Aug;44(4):320-9. PMID: 21645154.

Chiba K, Matsuyuki H, Maeda Y, et al. Role of sphingosine 1-phosphate receptor type 1 in lymphocyte egress from secondary lymphoid tissues and thymus. Cell Mol Immunol. 2006 Feb;3(1):11-9. PMID: 16549044.

Johnson VJ. He O. Osuchowski MF, et al. Disruption of sphingolipid homeostasis by myriocin, a mycotoxin, reduces thymic and splenic T-lymphocyte populations. Toxicology. 2004 Sep 1;201(1-3):67-75. PMID: 15297021.

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