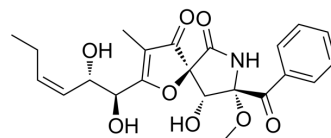


Pseurotin A

Cat. No.:	HY-125916
CAS No.:	58523-30-1
Molecular Formula:	C ₂₂ H ₂₅ NO ₈
Molecular Weight:	431.44
Target:	Endogenous Metabolite
Pathway:	Metabolic Enzyme/Protease
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Pseurotin A, a secondary metabolite of <i>Aspergillus</i> and other fungi, is a competitive inhibitor of chitin synthase and a neuritogenic agent. Pseurotin A inhibits IgE production (IC ₅₀ =3.6 μM). Antitumor activity ^{[1][2][3]} .									
IC₅₀ & Target	Microbial Metabolite									
In Vitro	<p>Pseurotin A inhibits the expression of PCSK9 in HepG2 cells, with an IC₅₀ value of 1.2 μM. Pseurotin A (25-100 μM; 72 hours) significantly suppressed the PCSK9 level in a dose dependent manner in BT-474 and T47D BC cell lines. Pseurotin A (25-100 μM; 72 hours) significantly increases LDLR level in a dose dependent manner^[3].</p> <p>Pseurotin A (40-200 μM; 24-72 hours) shows anti-proliferative activity in the hormone-dependent BC cells^[3]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Viability Assay^[3]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>BT-474 cells, T47D cells</td> </tr> <tr> <td>Concentration:</td> <td>40-200 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>24-72 hours</td> </tr> <tr> <td>Result:</td> <td>Showed weak but time and dose-dependent inhibition of the growth of BT-474 and T47D BC cells, with gradual decrease of the IC₅₀ values over time (260.83-93.64 μM and 267.84-113.08 μM, respectively).</td> </tr> </table>		Cell Line:	BT-474 cells, T47D cells	Concentration:	40-200 μM	Incubation Time:	24-72 hours	Result:	Showed weak but time and dose-dependent inhibition of the growth of BT-474 and T47D BC cells, with gradual decrease of the IC ₅₀ values over time (260.83-93.64 μM and 267.84-113.08 μM, respectively).
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In Vivo	<p>Pseurotin A (10 mg/kg; p.o.; 7X/week, 30 days) shows anti-tumor activity^[3]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <table border="1"> <tr> <td>Animal Model:</td> <td>HFD-fed orthotopic athymic mice (bearing BT-474 tumor cells xenograft mode)^[1]</td> </tr> <tr> <td>Dosage:</td> <td>10 mg/kg</td> </tr> <tr> <td>Administration:</td> <td>P.o.; 7X/week, 30 days</td> </tr> <tr> <td>Result:</td> <td>Significantly suppressed the growth of BT474 tumors.</td> </tr> </table>		Animal Model:	HFD-fed orthotopic athymic mice (bearing BT-474 tumor cells xenograft mode) ^[1]	Dosage:	10 mg/kg	Administration:	P.o.; 7X/week, 30 days	Result:	Significantly suppressed the growth of BT474 tumors.
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REFERENCES

- [1]. Ishikawa M, et al. Pseurotin A and its analogues as inhibitors of immunoglobulin E [correction of immunoglobuline E] production. *Bioorg Med Chem Lett*. 2009;19(5):1457-1460.
- [2]. Maiya S, Grundmann A, Li X, Li SM, Turner G. Identification of a hybrid PKS/NRPS required for pseurotin A biosynthesis in the human pathogen *Aspergillus fumigatus*. *Chembiochem*. 2007;8(14):1736-1743.
- [3]. Abdelwahed KS, et al. Pseurotin A as a novel suppressor of hormone dependent breast cancer progression and recurrence by inhibiting PCSK9 secretion and interaction with LDL receptor. *Pharmacol Res*. 2020;158:104847.
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Caution: Product has not been fully validated for medical applications. For research use only.

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