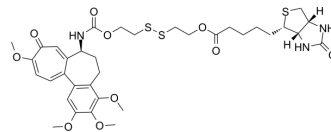


Deac-SS-Biotin

Cat. No.:	HY-144793
Molecular Formula:	C ₃₅ H ₄₅ N ₃ O ₁₀ S ₃
Molecular Weight:	763.94
Target:	Others
Pathway:	Others
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Deac-SS-Biotin is a potent antitumor agent. Deac-SS-Biotin uptakes into the cells through biotin-mediated internalization. Deac-SS-Biotin combined with DTT (Glutathione mimetic) can effectively inhibit microtubule assembly and displays greater antitumor activity ^[1] .																
In Vitro	<p>Deac-SS-Biotin (compound 9) (5 μM; 24 h) has a remarkable reduction-responsive drug release^[1]. Biotin effectively prevents Deac-SS-Biotin from binding to biotin receptor, thereby blocking the cellular uptake of Deac-SS-Biotin^[1].</p> <p>Deac-SS-Biotin (5 μM, 5 μM and 10 μM DTT; 0-80 min) effectively inhibits microtubule assembly and displays greater antitumor activity^[1].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Proliferation Assay^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>SGC-7901, A549, HeLa, L929 cells</td> </tr> <tr> <td>Concentration:</td> <td>0-20 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>72 h</td> </tr> <tr> <td>Result:</td> <td>Showed potent antiproliferative activity with IC₅₀s of 0.124, 0.085, 0.108, 4.22 μM for SGC-7901, A549, HeLa, L929 cells.</td> </tr> </table> <p>Cell Viability Assay^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>A549 cells</td> </tr> <tr> <td>Concentration:</td> <td>0.17 μM</td> </tr> <tr> <td>Incubation Time:</td> <td></td> </tr> <tr> <td>Result:</td> <td>Increased the cell viability from 23.6% to 68.9% with the increase of biotin concentration from 0.075 to 0.60 μM.</td> </tr> </table>	Cell Line:	SGC-7901, A549, HeLa, L929 cells	Concentration:	0-20 μM	Incubation Time:	72 h	Result:	Showed potent antiproliferative activity with IC ₅₀ s of 0.124, 0.085, 0.108, 4.22 μM for SGC-7901, A549, HeLa, L929 cells.	Cell Line:	A549 cells	Concentration:	0.17 μM	Incubation Time:		Result:	Increased the cell viability from 23.6% to 68.9% with the increase of biotin concentration from 0.075 to 0.60 μM.
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REFERENCES

[1]. Wang C, et al. Design, synthesis, and biological evaluation of biotinylated colchicine derivatives as potential antitumor agents. J Enzyme Inhib Med Chem. 2022 Dec;37(1):411-420.

Caution: Product has not been fully validated for medical applications. For research use only.

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA