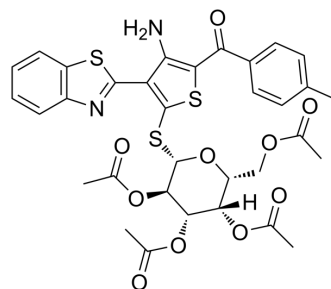


Anticancer agent 90

Cat. No.:	HY-151588
Molecular Formula:	C ₃₃ H ₃₂ N ₂ O ₁₀ S ₃
Molecular Weight:	712.81
Target:	Virus Protease
Pathway:	Anti-infection
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Anticancer agent 90 is a benzothiazole-2-thiophene S-glycoside derivative with antitumor activity. Anticancer agent 90 has high inhibition against the two cell line from ovarian cancer (OVCAR-4), renal cancer (A498) ^[1] .																
In Vitro	<p>Anticancer agent 90 (compound 6d) (0.01 mM; 24 h) has low cytotoxicity, shows high inhibition against two cell lines, SK-MEL-5, OVCAR-4^[1].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Cytotoxicity Assay^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>FRHK-4, Hep2, BGM, Vero, and Huh 7.5 cell lines</td> </tr> <tr> <td>Concentration:</td> <td>0-200 µg/mL</td> </tr> <tr> <td>Incubation Time:</td> <td>24 hours</td> </tr> <tr> <td>Result:</td> <td>Showed nontoxic under the doses of 100 µg/mL against FRHK-4, Hep2, BGM, Vero, and Huh 7.5 cells, respectively.</td> </tr> </table> <p>Cell Viability Assay^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>SK-MEL-5, OVCAR-4, and A498</td> </tr> <tr> <td>Concentration:</td> <td>0.01 mM</td> </tr> <tr> <td>Incubation Time:</td> <td>24 hours</td> </tr> <tr> <td>Result:</td> <td>Inhibited SK-MEL-5 and OVCAR-4 cell viability by 30.64% and 30.71%, respectively.</td> </tr> </table>	Cell Line:	FRHK-4, Hep2, BGM, Vero, and Huh 7.5 cell lines	Concentration:	0-200 µg/mL	Incubation Time:	24 hours	Result:	Showed nontoxic under the doses of 100 µg/mL against FRHK-4, Hep2, BGM, Vero, and Huh 7.5 cells, respectively.	Cell Line:	SK-MEL-5, OVCAR-4, and A498	Concentration:	0.01 mM	Incubation Time:	24 hours	Result:	Inhibited SK-MEL-5 and OVCAR-4 cell viability by 30.64% and 30.71%, respectively.
Cell Line:	FRHK-4, Hep2, BGM, Vero, and Huh 7.5 cell lines																
Concentration:	0-200 µg/mL																
Incubation Time:	24 hours																
Result:	Showed nontoxic under the doses of 100 µg/mL against FRHK-4, Hep2, BGM, Vero, and Huh 7.5 cells, respectively.																
Cell Line:	SK-MEL-5, OVCAR-4, and A498																
Concentration:	0.01 mM																
Incubation Time:	24 hours																
Result:	Inhibited SK-MEL-5 and OVCAR-4 cell viability by 30.64% and 30.71%, respectively.																

REFERENCES

[1]. Azzam et al. Novel Thiophene Thioglycosides Substituted with the Benzothiazole Moiety: Synthesis, Characterization, Antiviral and Anticancer Evaluations, and NS3/4A and USP7 Enzyme Inhibitions. ACS Omega. 2022 Sep 29;7(40):35656-35667.

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