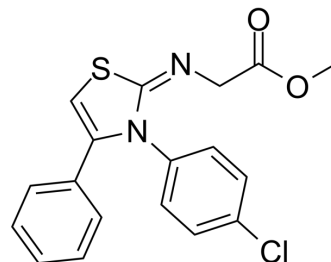


Anticancer agent 122

Cat. No.:	HY-154866
CAS No.:	2924532-50-1
Molecular Formula:	C ₁₈ H ₁₅ ClN ₂ O ₂ S
Molecular Weight:	358.84
Target:	Lactate Dehydrogenase
Pathway:	Metabolic Enzyme/Protease
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Anticancer agent 122, an inhibitor of human lactate dehydrogenase A enzyme (<i>h</i> LDHA), has good anticancer activities and can be used for anticancer research ^[1] .								
In Vitro	<p>Anticancer agent 122 (Compound 8m) (0-100 μM; 72 hours) inhibits liver cancer cell line HepG2 cells and cervical cancer cell line SiHa cells with IC₅₀s of 5.15 μM and 1.65 μM respectively and has not noticeable toxicity in human embryonic kidney cell line HEK293 cells^[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>HEK293, HepG2 and SiHa</td> </tr> <tr> <td>Concentration:</td> <td>0-100 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>72 hours</td> </tr> <tr> <td>Result:</td> <td>Inhibited liver cancer cell line HepG2 cells and cervical cancer cell line SiHa cells with IC₅₀s of 5.15 μM and 1.65 μM respectively. Had not noticeable toxicity in HEK293.</td> </tr> </table>	Cell Line:	HEK293, HepG2 and SiHa	Concentration:	0-100 μM	Incubation Time:	72 hours	Result:	Inhibited liver cancer cell line HepG2 cells and cervical cancer cell line SiHa cells with IC ₅₀ s of 5.15 μM and 1.65 μM respectively. Had not noticeable toxicity in HEK293.
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REFERENCES

[1]. Sharma D, et.al. Design and Synthesis of Thiazole Scaffold-Based Small Molecules as Anticancer Agents Targeting the Human Lactate Dehydrogenase A Enzyme. ACS Omega. 2023 May 10;8(20):17552-17562.

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

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