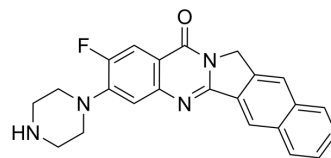


Topoisomerase I inhibitor 4

Cat. No.:	HY-143301
CAS No.:	2485135-31-5
Molecular Formula:	C ₂₃ H ₁₉ FN ₄ O
Molecular Weight:	386.42
Target:	Topoisomerase
Pathway:	Cell Cycle/DNA Damage
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Topoisomerase I inhibitor 4 (compound 7a) is a topoisomerase I inhibitor. Topoisomerase I inhibitor 4 inhibits HepG2, A549, MCF-7 and HeLa cancer cells proliferation with IC ₅₀ s of 1.20, 2.09, 1.56 and 1.92 μM, respectively. Topoisomerase I inhibitor 4 can be used for the research of cancer ^[1] .								
IC₅₀ & Target	IC ₅₀ : 1.20 μM (HepG2), 2.09 μM (A549), 1.56 μM (MCF-7), 1.92 μM (HeLa) ^[1]								
In Vitro	<p>Topoisomerase I inhibitor 4 (1-80 μM) shows Top1 inhibitory activity^[1].</p> <p>Topoisomerase I inhibitor 4 (0-40 μM; 48 h) inhibits proliferation of cancer cells^[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>HepG2, A549, MCF-7 and HeLa cell lines</td> </tr> <tr> <td>Concentration:</td> <td>0-40 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>48 hours</td> </tr> <tr> <td>Result:</td> <td>Showed antiproliferative activities to HepG2, A549, MCF-7 and HeLa cells with IC₅₀s of 1.20, 2.09, 1.56 and 1.92 μM, respectively.</td> </tr> </table>	Cell Line:	HepG2, A549, MCF-7 and HeLa cell lines	Concentration:	0-40 μM	Incubation Time:	48 hours	Result:	Showed antiproliferative activities to HepG2, A549, MCF-7 and HeLa cells with IC ₅₀ s of 1.20, 2.09, 1.56 and 1.92 μM, respectively.
Cell Line:	HepG2, A549, MCF-7 and HeLa cell lines								
Concentration:	0-40 μM								
Incubation Time:	48 hours								
Result:	Showed antiproliferative activities to HepG2, A549, MCF-7 and HeLa cells with IC ₅₀ s of 1.20, 2.09, 1.56 and 1.92 μM, respectively.								

REFERENCES

[1]. Xiang, Y, et al. Design, synthesis, and anticancer activities of 8,9-substituted Luotonin A analogs as novel topoisomerase I inhibitors. Med Chem Res 30, 1512–1522 (2021).

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