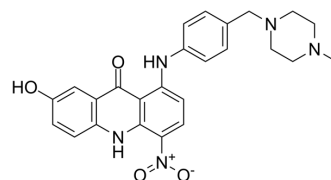


Topoisomerase II inhibitor 4

Cat. No.:	HY-143280
CAS No.:	2560590-49-8
Molecular Formula:	C ₂₅ H ₂₅ N ₅ O ₄
Molecular Weight:	459.5
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 II inhibitor 4 (compound E17) is a potent Topoisomerase II inhibitor. Topoisomerase II inhibitor 4 triggers G2/M cell cycle arrest and shows anti-tumor activity with strong cytotoxic and anti-proliferative effect ^[1] .																
IC₅₀ & Target	Topoisomerase II																
In Vitro	<p>Topoisomerase II inhibitor 4 (compound E17) inhibits cancer cells with IC₅₀ values of 4.55 μM (MDA-MB-231), 6.61 μM (A549), 2.18 μM (KG1)^[1].</p> <p>Topoisomerase II inhibitor 4 (1 μM; 0-22 h) induces G2/M cell cycle arrest^[2].</p> <p>Topoisomerase II inhibitor 4 (5 μM; 4 h) inhibits topo II-mediated chromosomes condensation in CRC cells, while also (100 μM; 0.5, 1, 2 h) induces topo II-DNA complex accumulation without degradation of topo II^[2].</p> <p>Topoisomerase II inhibitor 4 (1 μM; 4 h) exerts anti-tumor activity without increasing γH2AX levels in HCT116, while no DNA damage and apoptosis as well^[2].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Migration Assay^[2]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>HCT116, and WI-38 cells</td> </tr> <tr> <td>Concentration:</td> <td>0.4, 0.6, 0.8, 1.0, 1.2 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>0, 12, 24, 48 hours</td> </tr> <tr> <td>Result:</td> <td>Resulted migration inhibition in HCT116 and SW480 cells.</td> </tr> </table> <p>Cell Viability Assay^[2]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>HCT116, and WI-38 cells</td> </tr> <tr> <td>Concentration:</td> <td>0.39, 0.78, 1.56, 3.12, 6.25, 12.5, 25 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>72 hours</td> </tr> <tr> <td>Result:</td> <td>Exerted anti-proliferative effect in a dose-dependent manner.</td> </tr> </table>	Cell Line:	HCT116, and WI-38 cells	Concentration:	0.4, 0.6, 0.8, 1.0, 1.2 μM	Incubation Time:	0, 12, 24, 48 hours	Result:	Resulted migration inhibition in HCT116 and SW480 cells.	Cell Line:	HCT116, and WI-38 cells	Concentration:	0.39, 0.78, 1.56, 3.12, 6.25, 12.5, 25 μM	Incubation Time:	72 hours	Result:	Exerted anti-proliferative effect in a dose-dependent manner.
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REFERENCES

- [1]. Li ZY, et al. Structural optimizations and bioevaluation of N-substituted acridone derivatives as strong topoisomerase II inhibitors. *Bioorg Chem.* 2022 Feb. 119:105543.
- [2]. Chen JN, et al. E17 exerts anti-tumor activity through inhibiting topo II-mediated chromosomes condensation in CRC cells. *Biochem Biophys Res Commun.* 2019 May 28. 513(2):313-318.
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Caution: Product has not been fully validated for medical applications. For research use only.

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