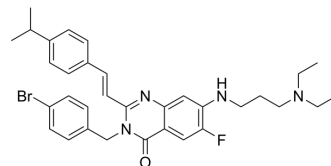


BLM-IN-2

Cat. No.:	HY-151939
Molecular Formula:	C ₃₃ H ₃₈ BrFN ₄ O
Molecular Weight:	605.58
Target:	DNA/RNA Synthesis; Apoptosis
Pathway:	Cell Cycle/DNA Damage; Apoptosis
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	BLM-IN-2 is a Bloom's Syndrome Protein (BLM) inhibitor with an IC ₅₀ value of 0.8 μM. BLM-IN-2 effectively suppresses the proliferation, invasion, cell cycle arrest and apoptosis of CRC cells. BLM-IN-2 can be used for the research of colorectal cancer (CRC) ^[1] .																
IC₅₀ & Target	IC ₅₀ : 0.8 μM (BLM) ^[1]																
In Vitro	<p>BLM-IN-2 (0-20 μM) has good inhibitory effect on BLM unwinding and binding DNA with IC₅₀ values of 0.8 μM and 2.3 μM, respectively^[1].</p> <p>BLM-IN-2 exhibits the potent BLM-dependent cytotoxicity against the CRC cells but weak against normal cells^[1].</p> <p>BLM-IN-2 (3 μM; 48 h) disrupts the HRR level while inhibiting BLM located on the DSB site and trigger DNA damage in the CRC cells^[1].</p> <p>BLM-IN-2 (0-5 μM; 48 h) effectively suppresses the proliferation and invasion of CRC cells, along with cell cycle arrest and apoptosis^[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>HCT116 cells; HCT116, SW480 and RKO cells</td> </tr> <tr> <td>Concentration:</td> <td>0-5 μM; 0.5, 1, 2 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>48 h; 10 days</td> </tr> <tr> <td>Result:</td> <td>Induced proliferation arrest. Completely inhibited the growth of cancer cells at the concentration around 2 μM, had a good anti-CRC activity.</td> </tr> </table> <p>Apoptosis Analysis^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>HCT116, SW480 and RKO cells</td> </tr> <tr> <td>Concentration:</td> <td>1 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>48 h</td> </tr> <tr> <td>Result:</td> <td>Induced apoptosis and necrosis in HCT116, SW480 and RKO.</td> </tr> </table>	Cell Line:	HCT116 cells; HCT116, SW480 and RKO cells	Concentration:	0-5 μM; 0.5, 1, 2 μM	Incubation Time:	48 h; 10 days	Result:	Induced proliferation arrest. Completely inhibited the growth of cancer cells at the concentration around 2 μM, had a good anti-CRC activity.	Cell Line:	HCT116, SW480 and RKO cells	Concentration:	1 μM	Incubation Time:	48 h	Result:	Induced apoptosis and necrosis in HCT116, SW480 and RKO.
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Cell Cycle Analysis^[1]

Cell Line:	HCT116, SW480 and RKO cells
Concentration:	4 μ M
Incubation Time:	48 h
Result:	Changed the cell proportion of the S or G2/M phase in CRC cells, arrested the cell cycle at the S phase in HCT116 and SW480 and arrested the cell cycle of HCT116, SW480 and RKO at the G2/M phase.

Cell Invasion Assay^[1]

Cell Line:	HCT116 cells
Concentration:	0.25, 0.5, 1, 2, 4 μ M
Incubation Time:	48 h
Result:	Obviously decreased the invasion in HCT116 cells with an IC ₅₀ value of 1.0 μ M and had inhibitory on CRC invasion.

REFERENCES

[1]. Jia-Li Tu, et al. Design, synthesis and evaluation of N3-substituted quinazolinone derivatives as potential Bloom's Syndrome protein (BLM) helicase inhibitor for sensitization treatment of colorectal cancer. Eur J Med Chem. 2022 Nov 21;246:114944.

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

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