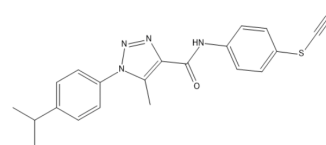


Anticancer agent 83

Cat. No.:	HY-151426
CAS No.:	904815-29-8
Molecular Formula:	C ₂₀ H ₁₉ N ₅ OS
Molecular Weight:	377.46
Target:	Apoptosis
Pathway:	Apoptosis
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Anticancer agent 83 is a potent anticancer agent, inhibits LOX IMVI cells growth with a GI ₅₀ value of 0.15 mM. Anticancer agent 83 reduces mitochondrial membrane potential and induces DNA damage to induces leukemia cells apoptosis ^[1] .																
In Vitro	<p>Anticancer agent 83 (compound 4a) (0.01-100 μM; 24 h) showing strong activity towards human colon carcinoma HCT116 p53^{-/-} cells with deletion of P53 gene (GI₅₀=8.4 μM), human epidermoid cervix carcinoma KB3-1 (GI₅₀=7.4 μM), human ovarian carcinoma Skov 3 cells (GI₅₀=10 μM), and human chronic myelogenous leukemia K562 cells (GI₅₀=5.4 μM)^[1]. Anticancer agent 83 (0.5 μM; 24 h) interrupts DNA stability and induction of apoptosis in Jurkat cells, and decreases mitochondrial membrane potential^[1].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Viability Assay^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>Human colon carcinoma HCT116 p53^{-/-} cells with deletion of P53 gene, human epidermoid cervix carcinoma KB3-1, human ovarian carcinoma Skov 3 cells, and human chronic myelogenous leukemia K562 cells</td> </tr> <tr> <td>Concentration:</td> <td>0.01, 0.1, 1, 10, 100 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>24 hours</td> </tr> <tr> <td>Result:</td> <td>Inhibited cell viability in different cells with GI₅₀s of 8.4 M (HCT116 p53^{-/-}), 7.4 M (KB3-1), 10 M (Skov 3), and 5.4 M (K562), respectively.</td> </tr> </table> <p>Immunofluorescence^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>Jurkat cells</td> </tr> <tr> <td>Concentration:</td> <td>0.5 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>24 hours</td> </tr> <tr> <td>Result:</td> <td>Induced morphological changes (including apoptotic bodies, membrane blebbing, chromatin condensation), and DNA fragmentation in Jurkat T-cells.</td> </tr> </table>	Cell Line:	Human colon carcinoma HCT116 p53 ^{-/-} cells with deletion of P53 gene, human epidermoid cervix carcinoma KB3-1, human ovarian carcinoma Skov 3 cells, and human chronic myelogenous leukemia K562 cells	Concentration:	0.01, 0.1, 1, 10, 100 μM	Incubation Time:	24 hours	Result:	Inhibited cell viability in different cells with GI ₅₀ s of 8.4 M (HCT116 p53 ^{-/-}), 7.4 M (KB3-1), 10 M (Skov 3), and 5.4 M (K562), respectively.	Cell Line:	Jurkat cells	Concentration:	0.5 μM	Incubation Time:	24 hours	Result:	Induced morphological changes (including apoptotic bodies, membrane blebbing, chromatin condensation), and DNA fragmentation in Jurkat T-cells.
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Incubation Time:	24 hours																
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REFERENCES

[1]. Pokhodylo N, et al. Novel N-(4-thiocyanatophenyl)-1H-1,2,3-triazole-4-carboxamides exhibit selective cytotoxic activity at nanomolar doses towards human leukemic T-cells. Eur J Med Chem. 2022 Nov 5;241:114633.

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

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