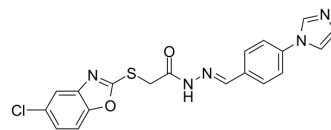


AKT-IN-18

Cat. No.:	HY-149842
Molecular Formula:	C ₁₉ H ₁₄ ClN ₅ O ₂ S
Molecular Weight:	411.86
Target:	Akt; Apoptosis
Pathway:	PI3K/Akt/mTOR; Apoptosis
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	AKT-IN-18, an inhibitor of Akt, inhibits Akt with an IC ₅₀ of 69.45 μM in A549 cells. AKT-IN-18 induces apoptosis and can be used in non-small cell lung cancer study ^[1] .								
IC₅₀ & Target	Akt 69.45 μM (IC ₅₀)								
In Vitro	<p>AKT-IN-18 (Compound 3i) (0-500 μM; 24 hours) exerts cytotoxic activity against A549 cells, and induces apoptosis^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Apoptosis Analysis^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>A549, L929 cells</td> </tr> <tr> <td>Concentration:</td> <td>0-500 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>24 hours</td> </tr> <tr> <td>Result:</td> <td>Exerted cytotoxic activity against A549 cells with IC₅₀ of 83.59 μM without influencing normal (L929) cells at effective doses. Induced early apoptosis caused at 2.7% and late apoptotic cells at 6.04%.</td> </tr> </table>	Cell Line:	A549, L929 cells	Concentration:	0-500 μM	Incubation Time:	24 hours	Result:	Exerted cytotoxic activity against A549 cells with IC ₅₀ of 83.59 μM without influencing normal (L929) cells at effective doses. Induced early apoptosis caused at 2.7% and late apoptotic cells at 6.04%.
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REFERENCES

[1]. Erdönmez B, et.al. Design, Synthesis, and Evaluation of a New Series of Hydrazones as Small-Molecule Akt Inhibitors for NSCLC Therapy. ACS Omega. 2023 May 24;8(22):20056-20065.

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

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