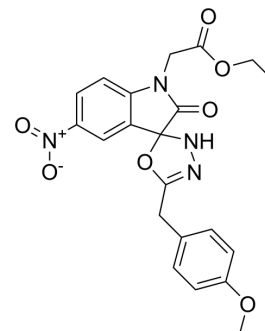


## Aldose reductase-IN-7

<b>Cat. No.:</b>	HY-161472
<b>Molecular Formula:</b>	C <sub>21</sub> H <sub>20</sub> N <sub>4</sub> O <sub>7</sub>
<b>Molecular Weight:</b>	440.41
<b>Target:</b>	Aldose Reductase
<b>Pathway:</b>	Metabolic Enzyme/Protease
<b>Storage:</b>	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	Aldose reductase-IN-7 (Compound 6k) targets Aldose reductase. Aldose reductase-IN-7 exhibits potent enzyme inhibitory activity ( $K_i = 0.186 \pm 0.020 \mu\text{M}$ ), showing superiority to Epalrestat (HY-66009), which is currently in clinical use. Aldose reductase-IN-7 is less cytotoxic and possesses potent anticancer activity <sup>[1]</sup> .								
<b>In Vitro</b>	<p>Aldose reductase-IN-7 (24 h) shows low cytotoxicity in L929 cells (<math>IC_{50} = 569.58 \pm 0.80 \mu\text{M}</math>). Significant anticancer activity can be observed in MCF-7 cells (<math>IC_{50} = 110.87 \pm 0.42 \mu\text{M}</math>)<sup>[1]</sup>.</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Cytotoxicity Assay<sup>[1]</sup></p> <table border="1"> <tr> <td>Cell Line:</td> <td>L929 cells and MCF-7 cells</td> </tr> <tr> <td>Concentration:</td> <td></td> </tr> <tr> <td>Incubation Time:</td> <td>24h</td> </tr> <tr> <td>Result:</td> <td>Ideally, killed cancer cells without harming healthy cells (<math>S_1</math> value of 5.13). Specifically inhibited the proliferation of cancer cells compared to non-cancerous cells.</td> </tr> </table>	Cell Line:	L929 cells and MCF-7 cells	Concentration:		Incubation Time:	24h	Result:	Ideally, killed cancer cells without harming healthy cells ( $S_1$ value of 5.13). Specifically inhibited the proliferation of cancer cells compared to non-cancerous cells.
Cell Line:	L929 cells and MCF-7 cells								
Concentration:									
Incubation Time:	24h								
Result:	Ideally, killed cancer cells without harming healthy cells ( $S_1$ value of 5.13). Specifically inhibited the proliferation of cancer cells compared to non-cancerous cells.								

### REFERENCES

[1]. Güleç Ö, et al. Novel spiroindoline derivatives targeting aldose reductase against diabetic complications: Bioactivity, cytotoxicity, and molecular modeling studies. Bioorg Chem. 2024 Apr;145:107221.

**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