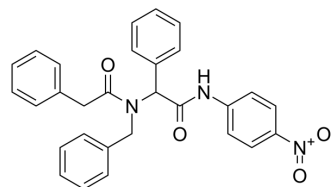


MMP-9-IN-3

Cat. No.:	HY-146216
CAS No.:	2581824-48-6
Molecular Formula:	C ₂₉ H ₂₅ N ₃ O ₄
Molecular Weight:	479.53
Target:	MMP; Akt; Apoptosis
Pathway:	Metabolic Enzyme/Protease; PI3K/Akt/mTOR; Apoptosis
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	MMP-9-IN-3 is a MMP-9 inhibitor (IC ₅₀ : 5.56 nM) that forms hydrogen bond with MMP-9. MMP-9-IN-3 also inhibits AKT activity (IC ₅₀ : 2.11 nM). MMP-9-IN-3 shows cell cytotoxicity and induces cell apoptosis. MMP-9-IN-3 can be used in the research of cancers ^[1] .																	
IC₅₀ & Target	MMP-9 5.56 nM (IC ₅₀)	MMP-1 447 nM (IC ₅₀)	MMP-2 221 nM (IC ₅₀)	MMP13 295 nM (IC ₅₀)														
	AKT 2.11 nM (IC ₅₀)																	
In Vitro	<p>MMP-9-IN-3 (compound 28, 0-1 μM approximately, 72 h) shows cytotoxicity to Wi-38, MCF-7, NFS-60, HepG-2 cells^[1]. MMP-9-IN-3 (72 h) induces apoptosis in MCF-7, NFS-60 and HepG-2 cells at 6.6 nM, 5.8 nM, 5.7 nM respectively^[1]. MMP-9-IN-3 (72 h) induces significant caspase 3/7 activation in MCF-7, NFS-60 and HepG-2 cells at 6.6 nM, 5.8 nM, 5.7 nM respectively^[1]. MMP-9-IN-3 (5.7 nM, 24 h) inhibits cell migration in HepG-2 cells^[1]. 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>Wi-38, MCF-7, NFS-60, HepG-2 cells</td> </tr> <tr> <td>Concentration:</td> <td>0-1 μM approximately</td> </tr> <tr> <td>Incubation Time:</td> <td>72 h</td> </tr> <tr> <td>Result:</td> <td>Showed cytotoxicity with IC₅₀: 314 nM (Wi-38), 6.6 nM (MCF-7), 5.8 nM (NFS-60), 5.7 nM (HepG-2).</td> </tr> </table> <p>Cell Migration Assay^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>HepG-2</td> </tr> <tr> <td>Concentration:</td> <td>5.7 nM</td> </tr> <tr> <td>Incubation Time:</td> <td>24 h</td> </tr> </table>				Cell Line:	Wi-38, MCF-7, NFS-60, HepG-2 cells	Concentration:	0-1 μM approximately	Incubation Time:	72 h	Result:	Showed cytotoxicity with IC ₅₀ : 314 nM (Wi-38), 6.6 nM (MCF-7), 5.8 nM (NFS-60), 5.7 nM (HepG-2).	Cell Line:	HepG-2	Concentration:	5.7 nM	Incubation Time:	24 h
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Cell Line:	HepG-2																	
Concentration:	5.7 nM																	
Incubation Time:	24 h																	

Result:	Inhibited cell migration by 71.25%.
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REFERENCES

[1]. Mohammed Salah Ayoup, et al. Battle tactics against MMP-9; discovery of novel non-hydroxamate MMP-9 inhibitors endowed with PI3K/AKT signaling attenuation and caspase 3/7 activation via Ugi bis-amide synthesis. Eur J Med Chem. 2020 Jan 15;186:111875.

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

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