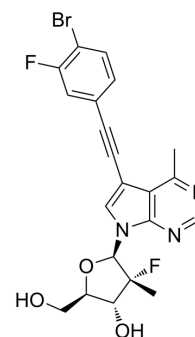


ZIKV-IN-1

Cat. No.:	HY-146957
CAS No.:	2762166-06-1
Molecular Formula:	C ₂₁ H ₁₈ BrF ₂ N ₃ O ₃
Molecular Weight:	478.29
Target:	Virus Protease; Flavivirus
Pathway:	Anti-infection
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	ZIKV-IN-1 is a potent zika virus inhibitor with an EC ₅₀ of 2.8 μM and EC ₉₀ of 6.8 μM. ZIKV-IN-1 shows anti-ZIKV activity with low cytotoxicity. ZIKV-IN-1 shows a strong affinity to ZIKV RdRp domain ^[1] . ZIKV-IN-1 is a click chemistry reagent, it contains an Alkyne group and can undergo copper-catalyzed azide-alkyne cycloaddition (CuAAC) with molecules containing Azide groups.																		
IC₅₀ & Target	EC ₅₀ : 2.8 μM (ZIKV) ^[1]																		
In Vitro	<p>ZIKV-IN-1 (compound 38) (0-80 μM) shows anti-ZIKV activity in a dose-dependent manner in A549 cells^[1]. ZIKV-IN-1 (5, 10, 25 μM; 48 h) decreases the expression of ZIKV E and cleaved caspase 3 protein^[1]. ZIKV-IN-1 (1.25, 2.5, 5, 10, 20 μM; 0-250 s) shows strong affinity to ZIKV RdRp domain (K_d=1.87 μM) in A549 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>A549 cells</td> </tr> <tr> <td>Concentration:</td> <td>0-50 μM</td> </tr> <tr> <td>Incubation Time:</td> <td></td> </tr> <tr> <td>Result:</td> <td>Showed anti-ZIKV activity with an EC₅₀ of 2.8 μM, EC₉₀ of 6.8 μM, and CC₅₀ of >50 μM.</td> </tr> </table> <p>Western Blot Analysis^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>A549, Vero cells</td> </tr> <tr> <td>Concentration:</td> <td>5, 10, 25 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>48 h</td> </tr> <tr> <td>Result:</td> <td>Decreased the expression of ZIKV E and cleaved caspase 3 protein.</td> </tr> </table> <p>Cell Cytotoxicity Assay^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>Vero, SNB19, Huh7, A549 cells</td> </tr> </table>	Cell Line:	A549 cells	Concentration:	0-50 μM	Incubation Time:		Result:	Showed anti-ZIKV activity with an EC ₅₀ of 2.8 μM, EC ₉₀ of 6.8 μM, and CC ₅₀ of >50 μM.	Cell Line:	A549, Vero cells	Concentration:	5, 10, 25 μM	Incubation Time:	48 h	Result:	Decreased the expression of ZIKV E and cleaved caspase 3 protein.	Cell Line:	Vero, SNB19, Huh7, A549 cells
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Result:	Decreased the expression of ZIKV E and cleaved caspase 3 protein.																		
Cell Line:	Vero, SNB19, Huh7, A549 cells																		

Concentration:	0-80 μ M
Incubation Time:	
Result:	Showed low cytotoxicity with CC ₅₀ s of 46.8, 49.4, 43.8, 54.1 μ M for ZG01 strain of Vero, SNB19, Huh7, A549 cells, and CC ₅₀ s of 470.5, 485.2, 512.8, 160.8 μ M for MR766 strain of Vero, SNB19, Huh7, A549 cells, respectively.

REFERENCES

[1]. Yao G, et al. Design, synthesis, and biological evaluation of novel 2'-methyl-2'-fluoro-6-methyl-7-alkynyl-7-deazapurine nucleoside analogs as anti-Zika virus agents. Eur J Med Chem. 2022 Apr 15;234:114275.

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

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