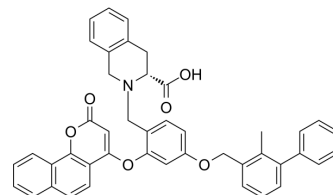


PD-1/PD-L1-IN-27

Cat. No.:	HY-146740
CAS No.:	2891831-47-1
Molecular Formula:	C ₄₄ H ₃₅ NO ₆
Molecular Weight:	673.75
Target:	PD-1/PD-L1
Pathway:	Immunology/Inflammation
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	PD-1/PD-L1-IN-27 is a potent PD-1/PD-L1 inhibitor with an IC ₅₀ value of 134 nM. PD-1/PD-L1-IN-27 shows antitumor effects with low T cell cytotoxicity. PD-1/PD-L1-IN-27 has the ability to activate CD8 ⁺ T cells and reduces T cell exhaustion ^[1] .								
IC₅₀ & Target	IC ₅₀ : 134 nM (PD-1/PD-L1) ^[1]								
In Vitro	<p>PD-1/PD-L1-IN-27 (compound MZ58) (2.5, 5, 10, 50 μM; 48 h) shows antitumor effects with low T cell cytotoxicity^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Proliferation Assay^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>MDA-MB-231 cells</td> </tr> <tr> <td>Concentration:</td> <td>2.5, 5, 10, 50 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>48 h</td> </tr> <tr> <td>Result:</td> <td>Facilitated cell proliferation with low T cell cytotoxicity.</td> </tr> </table>	Cell Line:	MDA-MB-231 cells	Concentration:	2.5, 5, 10, 50 μM	Incubation Time:	48 h	Result:	Facilitated cell proliferation with low T cell cytotoxicity.
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Concentration:	2.5, 5, 10, 50 μM								
Incubation Time:	48 h								
Result:	Facilitated cell proliferation with low T cell cytotoxicity.								
In Vivo	<p>PD-1/PD-L1-IN-27 (10, 25 mg/kg; i.g.) shows antitumor effect^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <table border="1"> <tr> <td>Animal Model:</td> <td>20-22 g, 6-8 weeks, male C57BL/6J mice (MC 38 cell lines)^[1]</td> </tr> <tr> <td>Dosage:</td> <td>10, 25 mg/kg</td> </tr> <tr> <td>Administration:</td> <td>i.g.</td> </tr> <tr> <td>Result:</td> <td>Showed antitumor effect.</td> </tr> </table>	Animal Model:	20-22 g, 6-8 weeks, male C57BL/6J mice (MC 38 cell lines) ^[1]	Dosage:	10, 25 mg/kg	Administration:	i.g.	Result:	Showed antitumor effect.
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REFERENCES

[1]. Zhang M, et al. Molecular hybridization used to design and synthesize neo-tanshinlactone derivatives as PD-1/PD-L1 inhibitors. *Bioorg Med Chem.* 2022; 54:116579.

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

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