

ODN 2088

Cat. No.:	HY-150738	
CAS No.:	1146541-45-8	
Molecular Weight:	4878	
Target:	Toll-like Receptor (TLR)	DNA, d(P-thio)(T-C-C-T-G-G-C-G-G-G-A-A-G-T)
Pathway:	Immunology/Inflammation	
Storage:	-20°C, sealed storage, away from moisture	
	* In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture)	

BIOLOGICAL ACTIVITY

Description	ODN 2088 is a potent TLR3, TLR7 and TLR9 inhibitor. ODN 2088 shows no cytotoxic. ODN 2088 inhibits the release of IFN- α and IL-6 ^{[1][2][3]} .								
In Vitro	<p>ODN 2088 (0.01, 0.1, 1, 10 μM; 48 h) shows no cytotoxic for for human PBMCs^[1].</p> <p>ODN 2088 (0.01, 0.1, 1, 10 μM; 24 h) inhibits the release of IFN-α in CpG-ODN 2216 (3 μM) and TLR7-ligand RNA-ORN 22075 (5 μM) stimulated human PBMCs^[1].</p> <p>ODN 2088 (0.01, 0.1, 1, 10 μM; 48 h) hardly inhibits the IL-6 release stimulated with CpG-ODN 2006 (100 nM) but inhibits the IL-6 release stimulated with imiquimod (5 μg/ml) in human PBMCs^[1].</p> <p>ODN 2088 (0.1, 1, 10 μM; 24 h) hardly inhibits IL-6 release by B-cells stimulated with CpG-DNA 2006 (100 nM) but inhibits the IL-6 release by imiquimod (5 μg/ml) stimulated human B-cells^[1].</p> <p>ODN 2088 (1, 10 μM; 48 h) increases the expression of CD86 and HLA-DR in the absence of CpG-ODN 2006 or imiquimod in CD20⁺ B-cells^[1].</p> <p>ODN 2088 presumably impairs TLR9-induced signaling induces by CpG-ODNs by competitive binding to the C-terminal fragment of TLR9^[1].</p> <p>ODN 2088 (0.001, 0.01, 0.1, 1, 10 μM; 24 h) inhibits the TNF-α secretion in a dose-dependent manner in CpG-ODN 1826 (100 nM) stimulated BMDMs and shows weekly inhibits when stimulated by the TLR7-agonist imiquimod^[3].</p> <p>ODN 2088 (10 μM) stimulates B cells to proliferate^[3].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Cytotoxicity Assay^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>Human PBMCs</td> </tr> <tr> <td>Concentration:</td> <td>0.01, 0.1, 1, 10 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>48 h</td> </tr> <tr> <td>Result:</td> <td>Showed no cytotoxic for for human PBMCs.</td> </tr> </table>	Cell Line:	Human PBMCs	Concentration:	0.01, 0.1, 1, 10 μ M	Incubation Time:	48 h	Result:	Showed no cytotoxic for for human PBMCs.
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REFERENCES

[1]. Römmeler F, et al. Guanine-modified inhibitory oligonucleotides efficiently impair TLR7- and TLR9-mediated immune responses of human immune cells. PLoS One. 2015 Feb 19;10(2):e0116703.

[2]. Duramad O, et al. Inhibitors of TLR-9 act on multiple cell subsets in mouse and man in vitro and prevent death in vivo from systemic inflammation. J Immunol. 2005 May 1;174(9):5193-200.

[3]. Römmler F, et al. Guanine modification of inhibitory oligonucleotides potentiates their suppressive function. J Immunol. 2013 Sep 15;191(6):3240-53.

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

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