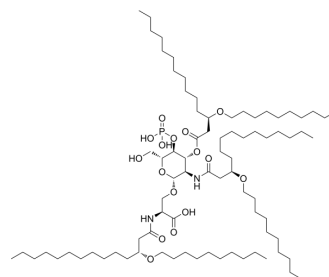


GSK1795091

Cat. No.:	HY-111792
CAS No.:	1233589-81-5
Molecular Formula:	C ₈₁ H ₁₅₇ N ₂ O ₁₆ P
Molecular Weight:	1446.09
Target:	Toll-like Receptor (TLR)
Pathway:	Immunology/Inflammation
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	GSK1795091 (CRX-601), an immunologic stimulator, is a synthetic TLR4 agonist. Antitumor activity. GSK1795091 can be used as a vaccine adjuvant to enhance both mucosal and systemic immunity to influenza virus vaccines. Not only, GSK1795091 inhibits tumor growth and increases the survival in mice model, but results in long term survival in influenza challenge model in mice ^{[1][2][3]} .								
In Vivo	<p>GSK1795091 (CRX-601) (0.1 µg /mouse, intranasal administration) as an adjuvant combined with detergent split-influenza antigen (H3N2) induces strong both mucosal and systemic immune responses in mice^[2].</p> <p>GSK1795091 (25 µg/mouse; iv; once weekly for 3 doses) inhibits tumor growth and results in long term survival in tumor model in mice^[3].</p> <p>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>Female BALB/c mice primed with H3N2^[1]</td> </tr> <tr> <td>Dosage:</td> <td>0.1 µg /mouse</td> </tr> <tr> <td>Administration:</td> <td>Intranasal administration</td> </tr> <tr> <td>Result:</td> <td> <p>As an adjuvant combined with detergent split-influenza antigen (H3N2) generated strong local and systemic immunity against co-administered influenza antigens while exhibiting high efficacy against two heterotypic influenza challenges.</p> <p>Mice receiving adjuvanted vaccines had significantly higher IgA titers than non-adjuvanted (vehicle) controls in an adjuvant dose-dependent manner.</p> <p>Adjuvanted vaccines promoted antigen-specific IgG and IgA antibody responses and the generation of polyfunctional antigen-specific Th17 cells.</p> </td> </tr> </table>	Animal Model:	Female BALB/c mice primed with H3N2 ^[1]	Dosage:	0.1 µg /mouse	Administration:	Intranasal administration	Result:	<p>As an adjuvant combined with detergent split-influenza antigen (H3N2) generated strong local and systemic immunity against co-administered influenza antigens while exhibiting high efficacy against two heterotypic influenza challenges.</p> <p>Mice receiving adjuvanted vaccines had significantly higher IgA titers than non-adjuvanted (vehicle) controls in an adjuvant dose-dependent manner.</p> <p>Adjuvanted vaccines promoted antigen-specific IgG and IgA antibody responses and the generation of polyfunctional antigen-specific Th17 cells.</p>
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REFERENCES

- [1]. Maroof A, et al. Intranasal vaccination promotes detrimental Th17-mediated immunity against influenza infection. PLoS Pathog. 2014 Jan;10(1):e1003875.
- [2]. Cebada J, et al. OX40 agonists for cancer treatment: a patent review. Expert Opin Ther Pat. 2021 Jan;31(1):81-90.
- [3]. Hug BA, et al. Safety, Pharmacokinetics, and Pharmacodynamics of the TLR4 Agonist GSK1795091 in Healthy Individuals: Results from a Randomized, Double-blind,

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

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