

Inbakicept

Cat. No.:	HY-P99661
CAS No.:	2135939-52-3
Target:	Interleukin Related
Pathway:	Immunology/Inflammation
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.

BIOLOGICAL ACTIVITY

Description	<p>Inbakicept, also known as N-803 (Nogapendekin alfa inbakicept), contains the IL-15 cytokine antibody Nogapendekin alfa (HY-P99759). Inbakicept is a dimeric human IL-15 receptor alpha (IL-15 Ra) sushi domain/human IgG1 Fc fusion protein and is an IL-15 superagonist complex. Inbakicept amplifies anti-CD20 mAb-mediated NK cell responses and antibody-dependent cellular cytotoxicity (ADCC). Inbakicept also increases degranulation and IFNγ production in cells^[1].</p>									
IC₅₀ & Target	IL-15R α									
In Vitro	<p>Inbakicept (0.01-1 nM; 20 hr) enhances human NK cell cytotoxicity and increases cytotoxic effector molecule expression^[1]. 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>Daudi, Raji, CD19⁺ FL cells</td> </tr> <tr> <td>Concentration:</td> <td>0.01 nM, 0.1 nM, and 1 nM</td> </tr> <tr> <td>Incubation Time:</td> <td>20 hours</td> </tr> <tr> <td>Result:</td> <td>Increased the human NK cell cytotoxicity on Daudi, Raji, CD19⁺ FL cells.</td> </tr> </table>		Cell Line:	Daudi, Raji, CD19 ⁺ FL cells	Concentration:	0.01 nM, 0.1 nM, and 1 nM	Incubation Time:	20 hours	Result:	Increased the human NK cell cytotoxicity on Daudi, Raji, CD19 ⁺ FL cells.
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In Vivo	<p>Inbakicept (0.2 mg/kg; i.v.; twice weekly, for 2 weeks) enhances rituximab-directed protection from a lethal Daudi lymphoma challenge in mouse model^[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>SCID mice with Daudi models^[1]</td> </tr> <tr> <td>Dosage:</td> <td>0.02 mg/kg, 0.05 mg/kg, 0.1 mg/kg, and 0.2 mg/kg</td> </tr> <tr> <td>Administration:</td> <td>Intravenous injection; twice weekly, for 2 weeks</td> </tr> <tr> <td>Result:</td> <td>Enhanced rituximab-directed control of Raji lymphoma challenge.</td> </tr> </table>		Animal Model:	SCID mice with Daudi models ^[1]	Dosage:	0.02 mg/kg, 0.05 mg/kg, 0.1 mg/kg, and 0.2 mg/kg	Administration:	Intravenous injection; twice weekly, for 2 weeks	Result:	Enhanced rituximab-directed control of Raji lymphoma challenge.
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Result:	Enhanced rituximab-directed control of Raji lymphoma challenge.									

REFERENCES

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

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