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Keliximab

Cat. No.:	HY-P99680
CAS No.:	174722-30-6
Target:	Interleukin Related
Pathway:	Immunology/Inflammation
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.

BIOLOGICAL ACTIVITY

Description	Keliximab (SB-210396) is a chimeric human/macaque IgG1 anti-CD4 monoclonal antibody with a K_i value of 1.0 nM for soluble CD4. Keliximab blocks T cell proliferation and inhibits IL-2 production. Keliximab can be used for cancer research ^[1] [2].									
In Vitro	<p>Keliximab shows a K_d value of 1.0 nM for binding with soluble CD4 (sCD4)^[2].</p> <p>Keliximab (0.01-10 $\mu\text{g}/\text{mL}$; 30 min) dose-dependently binds to the CD4⁺ thymoma cell line SupT1 with an ED₅₀ of 0.2 $\mu\text{g}/\text{mL}$ [1].</p> <p>Keliximab (0-1000 ng/mL) blocks T cell proliferation and IL-2 production with IC₅₀s of 10-30 ng/mL in primary human plasmablastic lymphomas (PBLs)^[1].</p> <p>Keliximab (1-1000 ng/mL; 5 min) dose-dependently mediates cell-cell adhesion with an ED₅₀ value of 20 ng/mL^[1].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p>									
In Vivo	<p>Keliximab (5-125 mg/kg; i.v., once) decreases CD4⁺ T cells^[1].</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>Transgenic mice bearing human CD4 molecules on their T cells^[1]</td> </tr> <tr> <td>Dosage:</td> <td>5-125 mg/kg</td> </tr> <tr> <td>Administration:</td> <td>Intravenous injection, 5-125 mg/kg, once</td> </tr> <tr> <td>Result:</td> <td>Dose-dependently down-modulated CD4 and rapidly declined the number of circulating CD4⁺ T cells compared with clenoliximab.</td> </tr> </table>		Animal Model:	Transgenic mice bearing human CD4 molecules on their T cells ^[1]	Dosage:	5-125 mg/kg	Administration:	Intravenous injection, 5-125 mg/kg, once	Result:	Dose-dependently down-modulated CD4 and rapidly declined the number of circulating CD4 ⁺ T cells compared with clenoliximab.
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REFERENCES

- [1]. Sharma A, et al. Comparative pharmacodynamics of keliximab and clenoliximab in transgenic mice bearing human CD4. *J Pharmacol Exp Ther.* 2000 Apr;293(1):33-41.
- [2]. Anderson D, et al. A primatized MAb to human CD4 causes receptor modulation, without marked reduction in CD4⁺ T cells in chimpanzees: in vitro and in vivo characterization of a MAb (IDEC-CE9.1) to human CD4. *Clin Immunol Immunopathol.* 1997 Jul;84(1):73-84.

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

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