

## Recombinant Human IL-21 (carrier-free)

<b>Catalog# / Size</b>	571202 / 10 µg 571204 / 25 µg 571206 / 100 µg 571208 / 500 µg
<b>Regulatory Status</b>	RUO
<b>Other Names</b>	Interleukin 21
<b>Description</b>	IL-21 was identified by a functional screening from conditioned medium of activated T cells using Baf3-IL-21R transfectants. In normal B cells, IL-21 can mediate cell proliferation, growth arrest, and terminal differentiation, or apoptosis, depending on their activation status. IL-21 enhances B cell proliferation following incubation with an activating CD40 antibody. This result suggests that IL-21 enhances B cell function following T:B cell interactions. Nevertheless, IL-21 inhibits the proliferation of murine and human B cells stimulated with anti-IgM antibodies. In addition, IL-21 enhances proliferation, cytotoxic activity, and IFN $\gamma$ production by CD8-effector T cells, and induces terminal differentiation of activated natural killer (NK) cells. Also, IL-21 can drive Th17 responses in conjunction with TGF- $\beta$ . Nevertheless, IL-21 is not essential for the differentiation of Th17 cells <i>in vitro</i> or <i>in vivo</i> , as it was showed using IL-21 and IL-21R-deficient mice. IL-6 induces the production of IL-21 from CD4 T cells upon TCR stimulation. In addition, IL-6 and IL-21 are key players in the Tfh differentiation, characterized by increased protein expression of both Bcl-6 and CXCR5. IL-21R complex is formed with the IL-21R-alpha chain and the common subunits ( $\gamma$ c) shared with other interleukins, such as IL-2, IL-4, IL-7, IL-9, and IL-15.

### Product Details

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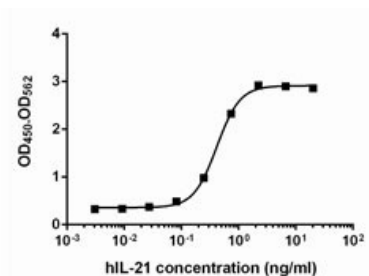
<b>Source</b>	Human IL-21, amino acids Gln30-Ser162 (Accession# NM_021803) was expressed in <i>E. coli</i> .
<b>Molecular Mass</b>	The 133 amino acid recombinant protein has a predicted molecular mass of approximately 15.5 kD. The DTT-reduced and non-reduced protein migrate at approximately 16 kD by SDS-PAGE. The N-terminal amino acid is Glutamine.
<b>Purity</b>	> 95%, as determined by Coomassie stained SDS-PAGE.
<b>Formulation</b>	0.22 µm filtered protein solution is in PBS and 1 mM EDTA.
<b>Endotoxin Level</b>	Less than 0.01 ng per µg cytokine as determined by the LAL method.
<b>Concentration</b>	10 and 25 µg sizes are bottled at 200 µg/mL. 100 µg size and larger sizes are lot-specific and bottled at the concentration indicated on the vial. To obtain lot-specific concentration, please enter the lot number in our <a href="#">Concentration and Expiration Lookup</a> or <a href="#">Certificate of Analysis</a> online tools.
<b>Storage &amp; Handling</b>	Unopened vial can be stored between 2°C and 8°C for up to 2 weeks, at -20°C for up to six months, or at -70°C or colder until the expiration date. For maximum results, quick spin vial prior to opening. The protein can be aliquoted and stored at -20°C or colder. Stock solutions can also be prepared at 50 - 100 µg/mL in appropriate sterile buffer, carrier protein such as 0.2 - 1% BSA or HSA can be added when preparing the stock solution. Aliquots can be stored between 2°C and 8°C for up to one week and stored at -20°C or colder for up to 3 months. <b>Avoid repeated freeze/thaw cycles.</b>
<b>Activity</b>	The ED <sub>50</sub> is 0.6-3.0 ng/ml, corresponding to a specific activity of 8.3 x 10 <sup>5</sup> - 3.3 x 10 <sup>6</sup> units/mg as determined by a dose-dependent induction of IFN $\gamma$ secretion by NK-92 cells.
<b>Application</b>	<a href="#">Bioassay</a>
<b>Application Notes</b>	BioLegend carrier-free recombinant proteins provided in liquid format are shipped on blue-ice. Our comparison testing data indicates that when handled and stored as recommended, the liquid format has equal or better stability and shelf-life compared to commercially available lyophilized proteins after reconstitution. Our liquid proteins are verified in-house to maintain activity after shipping on blue ice and are backed by our <a href="#">100% satisfaction guarantee</a> . If you have any concerns, contact us at <a href="mailto:tech@biolegend.com">tech@biolegend.com</a> .
<b>Product Citations</b>	1. Dai Q, <i>et al.</i> 2020. Front Immunol. 11:539654. <a href="#">PubMed</a>

2. Khanam A, *et al.* 2021. *Front Immunol.* 11:599648. [PubMed](#)
3. Zorro MM, *et al.* 2020. *J Autoimmun.* 108:102422. [PubMed](#)
4. Guo Q, *et al.* 2022. *Blood Adv.* 6:5668. [PubMed](#)
5. Jenks SA *et al.* 2018. *Immunity.* 49(4):725-739. [PubMed](#)
6. Vidard L, *et al.* 2019. *J Immunol.* 203:676. [PubMed](#)

## Antigen Details

<b>Structure</b>	Interleukin
<b>Distribution</b>	CD4 activated T cells, Th17 cells, NK T cells
<b>Function</b>	IL-21 induces T and B cell proliferation, B cell Ig class switching to IgG production, decreases dendritic cell function, enhances differentiation of effector and central memory T cells, and regulates T cell and hematopoietic progenitor cell homeostasis.
<b>Interaction</b>	T cells, B cells, NK cells, and dendritic cells as well as some nonimmune cells, such as fibroblasts and epithelial cells.
<b>Ligand/Receptor</b>	IL-21R and the common $\gamma$ c subunit (CD132) are components of the IL-21R complex.
<b>Cell Type</b>	B cells, Dendritic cells, Hematopoietic stem and progenitors, Tregs
<b>Biology Area</b>	Cell Biology, Immunology, Innate Immunity, Stem Cells
<b>Molecular Family</b>	Cytokines/Chemokines
<b>Antigen References</b>	<ol style="list-style-type: none"> <li>1. Parrish-Novak J, <i>et al.</i> 2000. <i>Nature</i> 408:57.</li> <li>2. De Toter D, <i>et al.</i> 2006. <i>Blood</i> 107:3708.</li> <li>3. Korn T, <i>et al.</i> 2007. <i>Nature</i> 448:484.</li> <li>4. Coquet JM, <i>et al.</i> 2008. <i>J. Immunol.</i> 180:7097.</li> <li>5. Dienz O, <i>et al.</i> 2009. <i>J. Exp. Med.</i> 206:69.</li> <li>6. Kaplan MH, <i>et al.</i> 2011. <i>Blood</i> 117:6198.</li> <li>7. Eto D, <i>et al.</i> 2011. <i>PLoS One</i> 6:e17739.</li> </ol>
<b>Gene ID</b>	<a href="#">59067</a>

## Product Data



Human IL-21 induced-production of IFN $\gamma$  by NK-92 cells.

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