

IL-15R alpha Protein, Human (HEK293, His-Avi)

Cat. No.:	HY-P78468
Synonyms:	IL-15 R alpha; CD215; IL-15RA; MGC104179
Species:	Human
Source:	HEK293
Accession:	Q13261 (I31-T205)
Gene ID:	3601
Molecular Weight:	50-70 kDa

PROPERTIES

Biological Activity	Immobilized Human IL-15 at 1µg/ml (100µl/Well) on the plate. Dose response curve for Human IL-15RA, His Tag with the EC ₅₀ of 12.1ng/ml determined by ELISA.
Appearance	Lyophilized powder.
Formulation	Lyophilized from a 0.22 µm filtered solution of PBS, pH 7.4. Normally 5% trehalose is added as protectant before lyophilization.
Endotoxin Level	<1 EU/µg, determined by LAL method.
Reconstitution	It is not recommended to reconstitute to a concentration less than 100 µg/mL in ddH ₂ O.
Storage & Stability	Stored at -20°C for 2 years. After reconstitution, it is stable at 4°C for 1 week or -20°C for longer (with carrier protein). It is recommended to freeze aliquots at -20°C or -80°C for extended storage.
Shipping	Room temperature in continental US; may vary elsewhere.

DESCRIPTION

Background

IL-15R alpha is expressed on various cell types, including lymphocytes, myeloid cells, nonlymphoid and nonhematopoietic cells^[4]. IL-15R alpha is down-regulated in Epstein-Barr virus associated gastric cancer (EBVaGC) via promoter hypermethylation^[5].

The sequence of amino acids in IL-15R alpha differs in different species. Human IL-15R alpha shares <55% aa sequence identity with mouse.

IL-15R alpha is required for transporting of IL-15 from the endoplasmic reticulum to the cell surface to bind with β (CD122) and γ (CD132) chains on responding lymphocytes^{[4][6]}. When binding with IL-15, the complex increases the in vivo half-life of IL-15 and enhances binding affinity of IL-15 with IL-15Rβ/γ in NK cells and CD8+ T cells. Thus, the signal transmission improves proliferation and antitumor activities of NK cells and CD8+ T cells^[2]. Moreover, IL-15R alpha on the cancer cell surface induces the malignant phenotype, such as augmented cancer cell growth, migration and invasion, and decreased apoptosis^[5].

IL-15R alpha binds with IL-15 and activates the antitumor functions of NK cells and CD8+ T cells, and is also important in

REFERENCES

- [1]. Yin Guo, et al. Immunobiology of the IL-15/IL-15R α complex as an antitumor and antiviral agent. *Cytokine Growth Factor Rev.* 2017 Dec;38:10-21.
- [2]. Johan Mj Van den Bergh, et al. IL-15 receptor alpha as the magic wand to boost the success of IL-15 antitumor therapies: The upswing of IL-15 transpresentation. *Pharmacol Ther.* 2017 Feb;170:73-79.
- [3]. Spencer W Stonier, et al. Trans-presentation: a novel mechanism regulating IL-15 delivery and responses. *Immunol Lett.* 2010 Jan 4;127(2):85-92.
- [4]. Patrick R Burkett, et al. IL-15R alpha expression on CD8+ T cells is dispensable for T cell memory. *Proc Natl Acad Sci U S A.* 2003 Apr 15;100(8):4724-9.
- [5]. Jing Wei, et al. Tumor cell-expressed IL-15R α drives antagonistic effects on the progression and immune control of gastric cancer and is epigenetically regulated in EBV-positive gastric cancer. *Cell Oncol (Dordr).* 2020 Dec;43(6):1085-1097.
- [6]. Emanuela Romano, et al. Human Langerhans cells use an IL-15R- α /IL-15/pSTAT5-dependent mechanism to break T-cell tolerance against the self-differentiation tumor antigen WT1. *Blood.* 2012 May 31;119(22):5182-90.
- [7]. J G Giri, et al. Identification and cloning of a novel IL-15 binding protein that is structurally related to the alpha chain of the IL-2 receptor.
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Caution: Product has not been fully validated for medical applications. For research use only.

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