

## **IL-7R Polyclonal Antibody**

Catalog No: YT2341

**Reactivity:** Human; Mouse; Monkey

**Applications:** WB;ELISA

Target: IL-7R

Fields: >>Cytokine-cytokine receptor interaction;>>FoxO signaling pathway;>>PI3K-Akt

signaling pathway;>>JAK-STAT signaling pathway;>>Hematopoietic cell

lineage;>>Pathways in cancer;>>Primary immunodeficiency

Gene Name: IL7R

**Protein Name:** Interleukin-7 receptor subunit alpha

P16871

P16872

Human Gene Id: 3575

**Human Swiss Prot** 

No:

Mouse Gene Id: 16197

**Mouse Swiss Prot** 

No:

**Immunogen:** The antiserum was produced against synthesized peptide derived from human

IL-7R/CD127. AA range:410-459

**Specificity:** IL-7R Polyclonal Antibody detects endogenous levels of IL-7R protein.

**Formulation :** Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.

Source: Polyclonal, Rabbit, IgG

**Dilution:** WB 1:500 - 1:2000. ELISA: 1:40000. Not yet tested in other applications.

**Purification:** The antibody was affinity-purified from rabbit antiserum by affinity-

chromatography using epitope-specific immunogen.



Concentration: 1 mg/ml

**Storage Stability:** -15°C to -25°C/1 year(Do not lower than -25°C)

Observed Band: 52 60kD

**Cell Pathway:** Cytokine-cytokine receptor interaction; Jak\_STAT; Hematopoietic cell

lineage; Primary immunodeficiency;

**Background:** The protein encoded by this gene is a receptor for interleukin 7 (IL7). The

function of this receptor requires the interleukin 2 receptor, gamma chain (IL2RG), which is a common gamma chain shared by the receptors of various cytokines, including interleukins 2, 4, 7, 9, and 15. This protein has been shown to play a critical role in V(D)J recombination during lymphocyte development. Defects in this gene may be associated with severe combined immunodeficiency (SCID). Alternatively spliced transcript variants have been found. [provided by

RefSeg, Dec 2015],

**Function:** disease:A genetic variation in transmembrane domain of IL7R is associated with

susceptibility to multiple sclerosis (MS) [MIM:126200]. Overtransmission of the major 'C' allele coding for Thr-244 are detected in offspring affected with multiple sclerosis. In vitro analysis of transcripts from minigenes containing either 'C' allele (Thr-244) or 'T' allele (Ile-244) shows that the 'C' allele results in an approximately two-fold increase in the skipping of exon 6, leading to increased production of a soluble form of IL7R. Thus, the multiple sclerosis associated 'C' risk allele of IL7R would probably decrease membrane-bound expression of IL7R. As this risk allele is common in the general population, some additional triggers are probably required for the development and progression of MS., disease:Defects in IL7R are

a cause of autosomal recessive severe combined immunodeficiency T-cell-

negativ

Subcellular [Isoform 1]: Cell me Location : Cell membrane; Sing

[Isoform 1]: Cell membrane; Single-pass type I membrane protein.; [Isoform 3]: Cell membrane; Single-pass type I membrane protein.; [Isoform 4]: Secreted.

**Expression :** B-cell, Epithelium, Spleen, Testis,

Tag: orthogonal

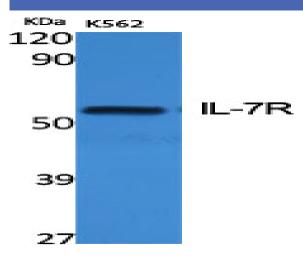
**Sort :** 8528

**No4:** 1

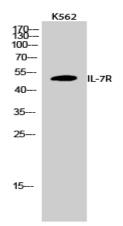
Host: Rabbit

Modifications: Unmodified

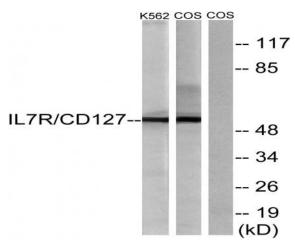
## **Products Images**



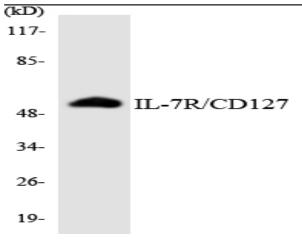
Western Blot analysis of various cells using IL-7R Polyclonal Antibody



Western Blot analysis of K562 cells using IL-7R Polyclonal Antibody



Western blot analysis of lysates from K562 and COS cells, treated with insulin 0.01U/ml 15', using IL-7R/CD127 Antibody. The lane on the right is blocked with the synthesized peptide.



Western blot analysis of the lysates from Jurkat cells using IL-7R/CD127 antibody.