abcam

Product datasheet

Recombinant Human EPO-R protein ab211321

1 Image

Description

Product name Recombinant Human EPO-R protein

Purity > 95 % SDS-PAGE.

Affinity purified

Endotoxin level < 1.000 Eu/µg

Expression system Insect cells

Accession <u>P19235</u>

Protein length Protein fragment

Animal free No

Nature Recombinant

Species Human

Sequence APPPNLPDPKFESKAALLAARGPEELLCFTERLEDLVCF

WEEAASAGVGP

GNYSFSYQLEDEPWKLCRLHQAPTARGAVRFWCSLPTA

DTSSFVPLELRV

TAASGAPRYHRVIHINEVVLLDAPVGLVARLADESGHVVL

RWLPPPETPM

TSHIRYEVDVSAGNGAGSVQRVEILEGRTECVLSNLRGRT

RYTFAVRARM

AEPSFGGFWSAWSEPVSLLTPSDLDPHHHHHH

Predicted molecular weight 26 kDa including tags

Amino acids 25 to 250

Tags His tag C-Terminus

Additional sequence information NP_000112.

Specifications

Our Abpromise guarantee covers the use of ab211321 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Applications SDS-PAGE

Form Liquid

Additional notes Previously labelled as EPO Receptor.

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Preparation and Storage

Stability and Storage

Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C or -

80°C. Avoid freeze / thaw cycle.

pH: 7.40

Constituents: 90% PBS, 10% Glycerol (glycerin, glycerine)

General Info

Function Receptor for erythropoietin. Mediates erythropoietin-induced erythroblast proliferation and

differentiation. Upon EPO stimulation, EPOR dimerizes triggering the JAK2/STAT5 signaling cascade. In some cell types, can also activate STAT1 and STAT3. May also activate the LYN α

tyrosine kinase.

Isoform EPOR-T acts as a dominant-negative receptor of EPOR-mediated signaling.

Tissue specificity

Erythroid cells and erythroid progenitor cells. Isoform EPOR-F is the most abundant form in EPO-dependent erythroleukemia cells and in late-stage erythroid progenitors. Isoform EPOR-S and isoform EPOR-T are the predominant forms in bone marrow. Isoform EPOR-T is the most abundant from in early-stage erythroid progenitor cells.

Involvement in disease

Defects in EPOR are the cause of erythrocytosis familial type 1 (ECYT1) [MIM:133100]. ECYT1 is an autosomal dominant disorder characterized by increased serum red blood cell mass, elevated hemoglobin and hematocrit, hypersensitivity of erythroid progenitors to erythropoietin, erythropoietin low serum levels, and no increase in platelets nor leukocytes. It has a relatively benign course and does not progress to leukemia.

Sequence similarities

Belongs to the type I cytokine receptor family. Type 1 subfamily.

Contains 1 fibronectin type-III domain.

Domain

The WSXWS motif appears to be necessary for proper protein folding and thereby efficient

intracellular transport and cell-surface receptor binding.

The box 1 motif is required for JAK interaction and/or activation.

Contains 1 copy of a cytoplasmic motif that is referred to as the immunoreceptor tyrosine-based

inhibitor motif (ITIM). This motif is involved in modulation of cellular responses. The

phosphorylated ITIM motif can bind the SH2 domain of several SH2-containing phosphatases.

Post-translational modifications

On EPO stimulation, phosphorylated on C-terminal tyrosine residues by JAK2. The phosphotyrosine motifs are also recruitment sites for several SH2-containing proteins and adapter proteins which mediate cell proliferation. Phosphorylation on Tyr-454 is required for PTPN6 interaction, Tyr-426 for PTPN11. Tyr-426 is also required for SOCS3 binding, but Tyr-456 to the professor of t

454/Tyr-456 motif is the preferred binding site.

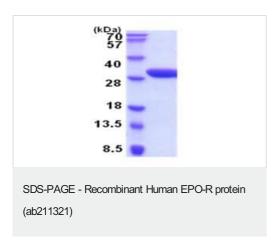
Ubiquitinated by NOSIP; appears to be either multi-monoubiquitinated or polyubiquitinated.

 $\label{thm:continuity} \mbox{ Ubiquitination mediates proliferation and survival of EPO-dependent cells.}$

Cellular localization

Cell membrane and Secreted. Secreted and located to the cell surface.

Images



15% SDS-PAGE analysis of ab211321 (3µg). Molecular weight is 28-40kDa on SDS-PAGE under reducing conditions.

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