

GHR Polyclonal Antibody

YT5573 Catalog No:

Human; Mouse; Rat Reactivity:

Applications: WB;ELISA

Target: GHR

Fields: >>Cytokine-cytokine receptor interaction;>>Neuroactive ligand-receptor

interaction;>>PI3K-Akt signaling pathway;>>JAK-STAT signaling

pathway;>>Growth hormone synthesis, secretion and action

Gene Name: **GHR**

Protein Name: Growth hormone receptor

P10912

P16882

Human Gene Id: 2690

Human Swiss Prot

No:

Mouse Gene Id: 14600

Mouse Swiss Prot

No:

Rat Gene Id: 25235

Rat Swiss Prot No: P16310

Immunogen: The antiserum was produced against synthesized peptide derived from the N-

terminal region of human GHR. AA range:21-70

GHR Polyclonal Antibody detects endogenous levels of GHR protein. **Specificity:**

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: 1:10000. Not yet tested in other applications.

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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: 140kD

Cell Pathway: Cytokine-cytokine receptor interaction; Neuroactive ligand-receptor

interaction; Jak_STAT;

Background: This gene encodes a member of the type I cytokine receptor family, which is a

transmembrane receptor for growth hormone. Binding of growth hormone to the receptor leads to receptor dimerization and the activation of an intra- and intercellular signal transduction pathway leading to growth. Mutations in this gene have been associated with Laron syndrome, also known as the growth hormone insensitivity syndrome (GHIS), a disorder characterized by short stature. In humans and rabbits, but not rodents, growth hormone binding protein (GHBP) is

generated by proteolytic cleavage of the extracellular ligand-binding domain from the mature growth hormone receptor protein. Multiple alternatively spliced

transcript variants have been found for this gene.[provided by RefSeg, Jun 2011],

Function : disease:Defects in GHR are a cause of Laron dwarfism [MIM:262500]; also

known as pituitary dwarfism II; Laron-type pituitary dwarfism I (LTD1) or Laron syndrome (LS). It is the most severe form of growth hormone insensitivity (GHI) characterized by growth impairment, dysmorphic facial features and truncal obesity. Levels of GHBP are low or undetectable in patients with Laron syndrome.,disease:Defects in GHR may be a cause of short stature [MIM:604271]. Short stature is defined by a subnormal rate of

growth.,domain:The box 1 motif is required for JAK interaction and/or activation.,domain:The extracellular domain is the ligand-binding domain representing the growth hormone-binding protein (GHBP).,domain:The

ubiquitination-dependent endocytosis motif (UbE) is required for recruitment of the ubiquitin conjugation system on to the receptor and for its

the abiquitin conjugation system on to the receptor and for it

internalization.,domain:The WSXWS motif a

Subcellular Location:

Cell membrane; Single-pass type I membrane protein. On growth hormone binding, GHR is ubiquitinated, internalized, down-regulated and transported into a degradative or non-degradative pathway. .; [Isoform 2]: Cell membrane; Single-pass type I membrane protein. Remains fixed to the cell membrane and is not internalized.; [Growth hormone-binding protein]: Secreted. Complexed to a

substantial fraction of circulating GH. .

Expression: Expressed in various tissues with high expression in liver and skeletal muscle.

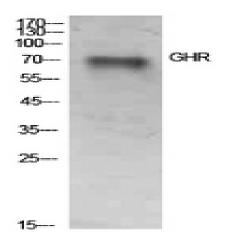
Isoform 4 is predominantly expressed in kidney, bladder, adrenal gland and brain stem. Isoform 1 expression in placenta is predominant in chorion and decidua.



Isoform 4 is highly expressed in placental villi. Isoform 2 is expressed in lung, stomach and muscle. Low levels in liver.

Sort : 6577

Products Images



Western Blot analysis of SKOV3 cells using GHR Polyclonal Antibody. Secondary antibody(catalog#:RS0002) was diluted at 1:20000