

Anti-Growth hormone receptor/GHR Picoband™ Antibody

Catalog Number: A00698-1

About Ghr

Growth hormone receptor is a protein that in humans is encoded by the GHR gene. It is mapped to 15 A1; 15 1.84 cM. 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.

Overview

Product Name	Anti-Growth hormone receptor/GHR Picoband™ Antibody
Reactive Species	Mouse, Rat
Description	Boster Bio Anti-Growth hormone receptor/GHR Picoband™ Antibody catalog # A00698-1. Tested in ELISA, Flow Cytometry, WB applications. This antibody reacts with Mouse, Rat.
Application	ELISA, Flow Cytometry, WB
Clonality	Polyclonal
Formulation	Each vial contains 4mg Trehalose, 0.9mg NaCl, 0.2mg Na ₂ HPO ₄ , 0.05mg NaN ₃ .
Storage Instructions	Store at -20°C for one year from date of receipt. After reconstitution, at 4°C for one month. It can also be aliquotted and stored frozen at -20°C for six months. Avoid repeated freeze-thaw cycles.
Host	Rabbit
Uniprot ID	P16882

Technical Details

Immunogen	E.coli-derived mouse Growth hormone receptor/GHR recombinant protein (Position: T25-Q650).
Recommended Detection Systems	Boster recommends Enhanced Chemiluminescent Kit with anti-Rabbit IgG (EK1002) for Western blot.
Cross Reactivity	No cross reactivity with other proteins.
Isotype	Rabbit IgG
Form	Lyophilized
Concentration	Adding 0.2 ml of distilled water will yield a concentration of 500 µg/ml.

Purification	Immunogen affinity purified.
Suggested Dilutions	<p>Dilute the sample so that the expected range of concentrations fall within the detection range of this kit.</p> <p>If the expected range of concentration is unknown, a pilot test should be conducted to decide the optimal dilution ratio for your samples.</p> <p>Some PubMed article(s) citing the expression level of this target are as follows: Boster Bio's internal QC testing used Western blot, 0.25-0.5μg/ml, Mouse, Rat Flow Cytometry, 1-3μg/1x10⁶ cells, Mouse, Rat Direct ELISA, 0.1-0.5μg/ml, Mouse</p> <p>For protocols, please visit https://www.bosterbio.com/protocol-and-troubleshooting/</p>

Anti-Growth hormone receptor/GHR Picoband™ Antibody (A00698-1) Images

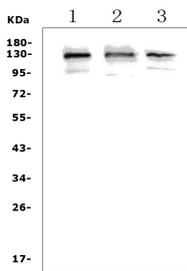


Figure 1. Western blot analysis of Ghr using anti-Ghr antibody (A00698-1).

Electrophoresis was performed on a 5-20% SDS-PAGE gel at 70V (Stacking gel) / 90V (Resolving gel) for 2-3 hours. The sample well of each lane was loaded with 50ug of sample under reducing conditions.

Lane 1: rat C6 whole cell lysates,
 Lane 2: mouse RAW246.7 whole cell lysates,
 Lane 3: mouse Neuro-2a whole cell lysates.

After Electrophoresis, proteins were transferred to a Nitrocellulose membrane at 150mA for 50-90 minutes. Blocked the membrane with 5% Non-fat Milk/ TBS for 1.5 hour at RT. The membrane was incubated with rabbit anti-Ghr antigen affinity purified polyclonal antibody (Catalog # A00698-1) at 0.5 ug/mL overnight at 4°C, then washed with TBS-0.1%Tween 3 times with 5 minutes each and probed with a goat anti-rabbit IgG-HRP secondary antibody at a dilution of 1:5000 for 1.5 hour at RT. The signal is developed using an Enhanced Chemiluminescent detection (ECL) kit (Catalog # EK1002) with Tanon 5200 system. A specific band was detected for Ghr at approximately 130KD. The expected band size for Ghr is at 72KD.

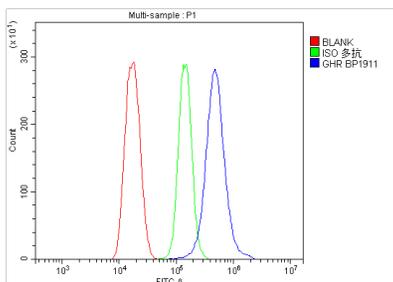


Figure 2. Flow Cytometry analysis of HEPA1-6 cells using anti-Ghr antibody (A00698-1).

Overlay histogram showing HEPA1-6 cells stained with A00698-1 (Blue line). The cells were blocked with 10% normal goat serum. And then incubated with rabbit anti-Ghr Antibody (A00698-1, 1ug/1x10⁶ cells) for 30 min at 20°C. DyLight®488 conjugated goat anti-rabbit IgG (BA1127, 5-10ug/1x10⁶ cells) was used as secondary antibody for 30 minutes at 20°C. Isotype control antibody (Green line) was rabbit IgG (1ug/1x10⁶) used under the same conditions. Unlabelled sample (Red line) was also used as a control.

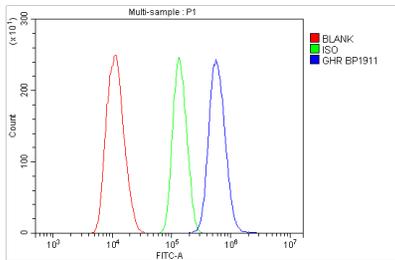


Figure 3. Flow Cytometry analysis of NRK cells using anti-Ghr antibody (A00698-1).

Overlay histogram showing NRK cells stained with A00698-1 (Blue line). The cells were blocked with 10% normal goat serum. And then incubated with rabbit anti-Ghr Antibody (A00698-1, 1ug/1x10⁶ cells) for 30 min at 20°C. DyLight® 488 conjugated goat anti-rabbit IgG (BA1127, 5-10ug/1x10⁶ cells) was used as secondary antibody for 30 minutes at 20°C. Isotype control antibody (Green line) was rabbit IgG (1ug/1x10⁶) used under the same conditions. Unlabelled sample (Red line) was also used as a control.

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