abcam

Product datasheet

Recombinant human Growth Hormone protein (Active) ab51232

1 References 2 Images

Description

Product name Recombinant human Growth Hormone protein (Active)

Biological activity Measured in a cell proliferation assay using Nb2-11 Rat lymphoma cells. The ED50 range ≤

0.2ng/ml.

Purity > 95 % SDS-PAGE.

Recombinant human Growth Hormone was purified by FPLC gel-filtration chromatography, after

refolding of the isolated inclusion bodies in a renaturation buffer.

Endotoxin level < 1.000 Eu/µg

Expression system Escherichia coli

Accession <u>P01241</u>

Protein length Full length protein

Animal free No

Nature Recombinant

Species Human

Sequence FPTIPLSRLFDNAMLRAHRLHQLAFDTYQEFEEAYIPKEQK

YSFLQNPQT

SLCFSESIPTPSNREETQQKSNLELLRISLLLIQSWLEPVQF

LRSVFANS

 ${\tt LVYGASDSNVYDLLKDLEEGIQTLMGRLEDGSPRTGQIFK}$

QTYSKFDTNS

 ${\tt HNDDALLKNYGLLYCFRKDMDKVETFLRIVQCRSVEGSC}$

GF

Predicted molecular weight 22 kDa

Amino acids 27 to 217

Additional sequence information Full length mature chain without signal peptide. NP_000506.

Specifications

Our **Abpromise guarantee** covers the use of **ab51232** in the following tested applications.

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

Applications Western blot

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SDS-PAGE

Functional Studies

Form Liquid

Preparation and Storage

Stability and Storage Shipped at 4°C. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles.

pH: 7.40

Constituents: 0.0134% PBS, 1.6016% Mannitol

This product is an active protein and may elicit a biological response in vivo, handle with caution.

General Info

Function Plays an important role in growth control. Its major role in stimulating body growth is to stimulate

the liver and other tissues to secrete IGF-1. It stimulates both the differentiation and proliferation of myoblasts. It also stimulates amino acid uptake and protein synthesis in muscle and other tissues.

Involvement in disease Defects in GH1 are a cause of growth hormone deficiency isolated type 1A (IGHD1A)

[MIM:262400]; also known as pituitary dwarfism I. IGHD1A is an autosomal recessive deficiency of GH which causes short stature. IGHD1A patients have an absence of GH with severe dwarfism

and often develop anti-GH antibodies when given exogenous GH.

Defects in GH1 are a cause of growth hormone deficiency isolated type 1B (IGHD1B)

[MIM:612781]; also known as dwarfism of Sindh. IGHD1B is an autosomal recessive deficiency of GH which causes short stature. IGHD1B patients have low but detectable levels of GH. Dwarfism

is less severe than in IGHD1A and patients usually respond well to exogenous GH.

Defects in GH1 are the cause of Kowarski syndrome (KWKS) [MIM:262650]; also known as

pituitary dwarfism VI.

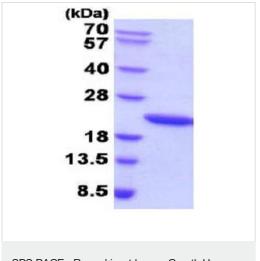
Defects in GH1 are a cause of growth hormone deficiency isolated type 2 (IGHD2) [MIM:173100]. IGHD2 is an autosomal dominant deficiency of GH which causes short stature. Clinical severity is variable. Patients have a positive response and immunologic tolerance to growth hormone

therapy.

Sequence similaritiesBelongs to the somatotropin/prolactin family.

Cellular localization Secreted.

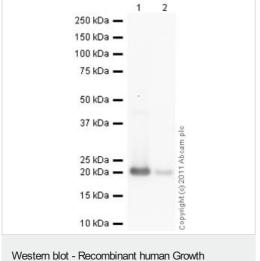
Images



15% SDS-PAGE - Recombinant human Growth Hormone protein (Active) (ab51232, 3 μg).

Reducing conditions. Visualized by coomassie blue stain.





Western blot - Recombinant human Growth Hormone protein (Active) (ab51232) **All lanes :** Anti-Growth Hormone antibody [GH-1] (<u>ab9821</u>) at 1/500 dilution

Lane 1 : Recombinant human Growth Hormone protein (Active) (ab51232) at 0.1 μg

Lane 2 : Recombinant human Growth Hormone protein (Active) (ab51232) at 0.01 μg

Secondary

All lanes : Goat Anti-Mouse IgG H&L (HRP) preadsorbed (ab97040) at 1/5000 dilution

Developed using the ECL technique.

Performed under reducing conditions.

Exposure time: 2 minutes

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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