

Product datasheet

Recombinant human IGF2 protein ab155617

2 Images

Description

Product name	Recombinant human IGF2 protein
Biological activity	Measured in a serum-free cell proliferation assay using MCF7 Human breast cancer cells. The ED ₅₀ for this effect is typically 2-10 ng/ml.
Purity	> 95 % SDS-PAGE. Lyophilized from 0.22µm filtered solution
Endotoxin level	< 1.000 Eu/µg
Expression system	HEK 293 cells
Accession	<u>P01344-1</u>
Protein length	Protein fragment
Animal free	No
Nature	Recombinant
Species	Human
Sequence	AYRPSETLCGGELVDTLQFVCGDRGFYFSRPASRVSRRS RGVEECCFRS CDLALLETYCATPAKSE
Predicted molecular weight	34 kDa
Molecular weight information	The protein migrates as 35-38 kDa under reducing (R) condition (SDS-PAGE) due to glycosylation.
Amino acids	25 to 91
Tags	Fc tag N-Terminus
Additional sequence information	This protein carries a human IgG1 Fc tag at the N-terminus.

Specifications

Our **Abpromise guarantee** covers the use of **ab155617** in the following tested applications.

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

Applications	Functional Studies SDS-PAGE
Form	Lyophilized

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: 7.4

Constituents: 5% Trehalose, 0.75% Glycine, 0.605% Tris

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

Reconstitution

Reconstitute with sterile deionized water to a concentration of 1 mg/ml.

General Info

Function

The insulin-like growth factors possess growth-promoting activity. In vitro, they are potent mitogens for cultured cells. IGF-II is influenced by placental lactogen and may play a role in fetal development.

Preptin undergoes glucose-mediated co-secretion with insulin, and acts as physiological amplifier of glucose-mediated insulin secretion. Exhibits osteogenic properties by increasing osteoblast mitogenic activity through phosphoactivation of MAPK1 and MAPK3.

Involvement in disease

Epigenetic changes of DNA hypomethylation in IGF2 are a cause of Silver-Russell syndrome (SIRS) [MIM:180860]. SIRS is a clinically heterogeneous condition characterized by severe intrauterine growth retardation, poor postnatal growth, craniofacial features such as a triangular shaped face and a broad forehead, body asymmetry, and a variety of minor malformations.

Sequence similarities

Belongs to the insulin family.

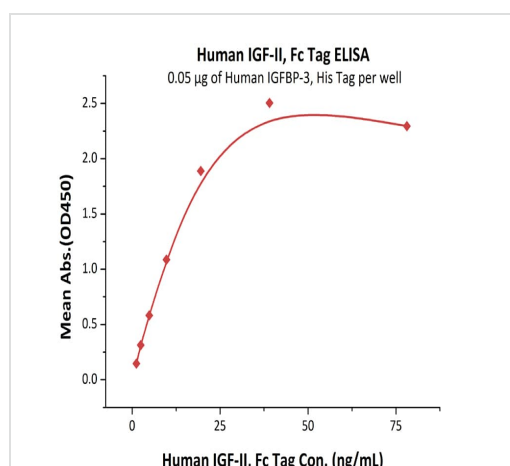
Post-translational modifications

O-glycosylated with a core 1 or possibly core 8 glycan.

Cellular localization

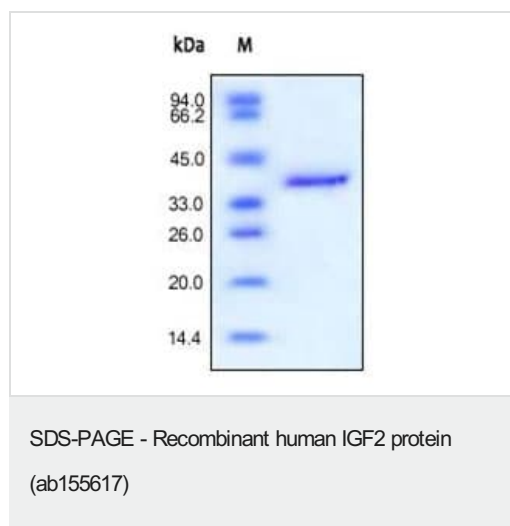
Secreted.

Images



Immobilized Human IGFBP-3, His Tag at 0.5 µg/mL (100 µL/well) can bind Human IGF-II, Fc Tag with a linear range of 1-20 ng/mL.

Functional Studies - Recombinant human IGF2 protein (ab155617)



SDS-PAGE analysis of reduced ab155617 stained overnight with Coomassie Blue.

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

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