abcam

Product datasheet

Recombinant Human IFNGR1 protein ab181887

1 References 1 Image

Description

Product name Recombinant Human IFNGR1 protein

Purity > 95 % SDS-PAGE.

Endotoxin level < 1.000 Eu/µg
Expression system HEK 293 cells

Accession P15260

Protein length Protein fragment

Animal free No.

Nature Recombinant

Species Human

Sequence EMGTADLGPSSVPTPTNVTIESYNMNPIVYWEYQIMPQVP

VFTVEVKNYG

VKNSEWIDACINISHHYCNISDHVGDPSNSLWVRVKARVG

QKESAYAKSE EFAVCRDGKIGPPKLDIRKE EKQIMIDIFHPSVFVNGDEQEVDYDPET

TCYIRVYNVYVRMNGSEIQYKILTQKEDDCDEIQCQLAIPVS SLNSQYCV SAEGVLHVWGVTTEKSKEVCITIFNSSIKG

Predicted molecular weight 52 kDa including tags

Amino acids 18 to 245

Additional sequence information Fused with Fc fragment of Human IgG1 at the C-terminus (AAH05333).

Specifications

Our Abpromise guarantee covers the use of ab181887 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 Lyophilized

Preparation and Storage

Stability and Storage Shipped at 4°C. Store at 4°C prior to reconstitution. Store at -80°C. Avoid freeze / thaw cycle. For

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long term storage it is recommended to add a carrier protein on reconstitution (0.1% HSA or BSA).

pH: 7.4

Constituents: 0.61% Tris, 0.75% Glycine, 5% Trehalose

Reconstitution

Reconstitute with sterile deionized water to a concentration of 400 µg/ml.

General Info

Function

Involvement in disease

Receptor for interferon gamma. Two receptors bind one interferon gamma dimer.

Defects in IFNGR1 are a cause of mendelian susceptibility to mycobacterial disease (MSMD) [MIM:209950]; also known as familial disseminated atypical mycobacterial infection. This rare condition confers predisposition to illness caused by moderately virulent mycobacterial species, such as Bacillus Calmette-Guerin (BCG) vaccine and environmental non-tuberculous mycobacteria, and by the more virulent Mycobacterium tuberculosis. Other microorganisms rarely cause severe clinical disease in individuals with susceptibility to mycobacterial infections, with the exception of Salmonella which infects less than 50% of these individuals. The pathogenic mechanism underlying MSMD is the impairment of interferon-gamma mediated immunity whose severity determines the clinical outcome. Some patients die of overwhelming mycobacterial disease with lepromatous-like lesions in early childhood, whereas others develop, later in life, disseminated but curable infections with tuberculoid granulomas. MSMD is a genetically heterogeneous disease with autosomal recessive, autosomal dominant or X-linked inheritance.

Sequence similarities

Belongs to the type II cytokine receptor family.

Contains 2 fibronectin type-III domains.

Contains 2 lg-like C2-type (immunoglobulin-like) domains.

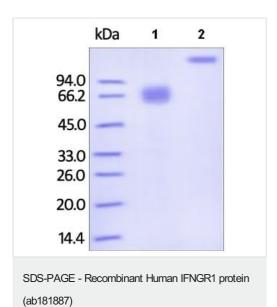
Post-translational modifications

Phosphorylated at Ser/Thr residues.

Cellular localization

Membrane.

Images



SDS-PAGE analysis of ab181887 stained overnight with Coomassie Blue.

Lane 1: DTT-reduced

Lane 2: Non-reduced

As a result of glycosylation, DTT-reduced protein migrates as 75-100 kDa and non-reduced protein migrates as 150-170 kDa.

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