abcam

Product datasheet

Anti-IFN gamma Receptor beta/AF-1 antibody ab198801

2 References 1 Image

Overview

Product name	Anti-IFN gamma Receptor beta/AF-1 antibody	
Description	Rabbit polyclonal to IFN gamma Receptor beta/AF-1	
Host species	Rabbit	
Tested applications	Suitable for: WB	
Species reactivity	Reacts with: Mouse	
Immunogen	Synthetic peptide corresponding to Human IFN gamma Receptor beta/AF-1. (Near the C terminal; NP_005525). Database link: <u>P38484</u>	
Positive control	Mouse skeletal muscle tissue lysate.	
General notes	The Life Science industry has been in the grips of a reproducibility crisis for a number of years. Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets your needs before purchasing.	
	If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be found below, along with publications, customer reviews and Q&As	

Properties	
Form	Liquid
Storage instructions	Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C long term. Avoid freeze / thaw cycle.
Storage buffer	pH: 7.4 Preservative: 0.05% Sodium azide Constituents: 49% PBS, 50% Glycerol (glycerin, glycerine)
Purity	Immunogen affinity purified
Clonality	Polyclonal
lsotype	lgG

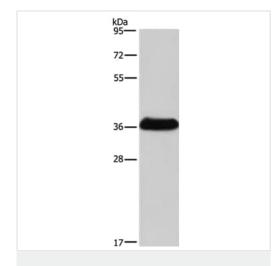
The Abpromise guarantee Our <u>Abpromise guarantee</u> covers the use of ab198801 in the following tested applications.

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

Application	Abreviews	Notes
WB		1/2000 - 1/5000. Predicted molecular weight: 38 kDa.

Target	
Function	Part of the receptor for interferon gamma. Required for signal transduction. This accessory factor is an integral part of the IFN-gamma signal transduction pathway and is likely to interact with GAF, JAK1, and/or JAK2.
Involvement in disease	Defects in IFNGR2 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.
Cellular localization	Membrane.

Images



Western blot - Anti-IFN gamma Receptor beta/AF-1 antibody (ab198801)

Anti-IFN gamma Receptor beta/AF-1 antibody (ab198801) at 1/400 dilution + Mouse skeletal muscle tissue lysate at 40 μg

Predicted band size: 38 kDa

Exposure time: 40 seconds

8% SDS-PAGE.

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