

IL-12B Polyclonal Antibody

YT2312 Catalog No:

Reactivity: Human; Mouse

WB;ELISA **Applications:**

Target: IL-12B

Fields: >>Cytokine-cytokine receptor interaction;>>Toll-like receptor signaling

> pathway;>>RIG-I-like receptor signaling pathway;>>C-type lectin receptor signaling pathway;>>JAK-STAT signaling pathway;>>Th1 and Th2 cell

differentiation;>>Alcoholic liver disease;>>Type I diabetes

mellitus;>>Pertussis;>>Legionellosis;>>Leishmaniasis;>>Chagas

disease;>>African trypanosomiasis;>>Toxoplasmosis;>>Amoebiasis;>>Tubercul osis;>>Measles;>>Influenza A;>>Herpes simplex virus 1 infection;>>Coronavirus

disease - COVID-19;>>Pathways in cancer;>>Proteoglycans in

cancer;>>Inflammatory bowel disease;>>Allograft rejection;>>Lipid and

atherosclerosis

IL12B Gene Name:

Protein Name: Interleukin-12 subunit beta

Human Gene Id: 3593

Human Swiss Prot

P29460

No:

Mouse Swiss Prot P43432

No:

The antiserum was produced against synthesized peptide derived from human Immunogen:

IL-12B. AA range:228-277

IL-12B Polyclonal Antibody detects endogenous levels of IL-12B protein. **Specificity:**

Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide. Formulation:

Polyclonal, Rabbit, IgG Source:

WB 1:500 - 1:2000. ELISA: 1:5000. Not yet tested in other applications. **Dilution:**



Purification: The antibody was affinity-purified from rabbit antiserum by affinity-

chromatography using epitope-specific immunogen.

Concentration: 1 mg/ml

Storage Stability: -15°C to -25°C/1 year(Do not lower than -25°C)

Observed Band: 40kD

Cell Pathway: Cytokine-cytokine receptor interaction; Toll_Like; RIG-I-like

receptor; Jak_STAT; Type I diabetes mellitus; Allograft rejection;

Background: This gene encodes a subunit of interleukin 12, a cytokine that acts on T and

natural killer cells, and has a broad array of biological activities. Interleukin 12 is a disulfide-linked heterodimer composed of the 40 kD cytokine receptor like subunit encoded by this gene, and a 35 kD subunit encoded by IL12A. This cytokine is expressed by activated macrophages that serve as an essential inducer of Th1 cells development. This cytokine has been found to be important for sustaining a sufficient number of memory/effector Th1 cells to mediate long-term protection to an intracellular pathogen. Overexpression of this gene was observed in the central nervous system of patients with multiple sclerosis (MS), suggesting a role of this cytokine in the pathogenesis of the disease. The promoter polymorphism of this gene has been reported to be associated with the severity of atopic and non-

atopic asthma in children. [provi

Function: disease:Defects in IL12B 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

Subcellular Location : Secreted.

Expression:

Brain,

Products Images

