

IFN-γ (PN0458) Nb-FC recombinant antibody

Catalog No: YA0605

Reactivity: Human

Applications: ELISA

Target: IFNy

Fields: >>Proteasome;>>Cytokine-cytokine receptor interaction;>>HIF-1 signaling

pathway;>>Necroptosis;>>TGF-beta signaling pathway;>>Osteoclast

differentiation;>>Antigen processing and presentation;>>JAK-STAT signaling

pathway;>>Natural killer cell mediated cytotoxicity;>>IL-17 signaling

pathway;>>Th1 and Th2 cell differentiation;>>Th17 cell differentiation;>>T cell

receptor signaling pathway;>>Type I diabetes

mellitus;>>Leishmaniasis;>>Chagas disease;>>African trypanosomiasis;>>Malari a;>>Toxoplasmosis;>>Amoebiasis;>>Tuberculosis;>>Hepatitis C;>>Influenza A;>>Herpes simplex virus 1 infection;>>Pathways in cancer;>>PD-L1 expression

and PD-1 checkpoint pathway in cancer;>>Inflammatory bowel

disease;>>Systemic lupus erythematosus;>>Rheumatoid arthritis;>>Allograft rejection;>>Graft-versus-host disease;>>Fluid shear stress and atherosclerosis

Gene Name: IFNG

Protein Name: Interferon gamma

Human Gene Id: 3458

Human Swiss Prot

No:

Immunogen: Purified recombinant Human IFNy

P01579

Specificity: This recombinant monoclonal antibody can detects endogenous levels of IFNy

protein.

Formulation: Phosphate-buffered solution

Source: Camel, chimeric fusion of Nanobody (VHH) and mouse IgG1 Fc domain,

recombinantly produced from 293F cell



Dilution: ELISA 1:5000-100000

Purification: Recombinant Expression and Affinity purified

Concentration: Please check the information on the tube

Storage Stability: -15°C to -25°C/1 year(Avoid freeze / thaw cycles)

Cell Pathway: Proteasome; Cytokine-cytokine receptor interaction; Regulation of

autophagy;TGF-beta;Jak_STAT;Natural killer cell mediated

cytotoxicity; T_Cell_Receptor; Type I diabetes mellitus; Systemic lupus erythemato

Background: This gene encodes a soluble cytokine that is a member of the type II interferon

class. The encoded protein is secreted by cells of both the innate and adaptive immune systems. The active protein is a homodimer that binds to the interferon gamma receptor which triggers a cellular response to viral and microbial

infections. Mutations in this gene are associated with an increased susceptibility to viral, bacterial and parasitic infections and to several autoimmune diseases.

[provided by RefSeq, Dec 2015],

Function: disease:In Caucasians, genetic variation in IFNG is associated with the risk of

aplastic anemia (AA) [MIM:609135]. AA is a rare disease in which the reduction of the circulating blood cells results from damage to the stem cell pool in bone marrow. In most patients, the stem cell lesion is caused by an autoimmune attack. T-lymphocytes, activated by an endogenous or exogenous, and most often

unknown antigenic stimulus, secrete cytokines, including IFN-gamma, which would in turn be able to suppress hematopoiesis..function:Produced by

lymphocytes activated by specific antigens or mitogens. IFN-gamma, in addition to having antiviral activity, has important immunoregulatory functions. It is a potent activator of macrophages, it has antiproliferative effects on transformed

cells and it can potentiate the antiviral and antitumor effects of the type $\ensuremath{\mathsf{I}}$

interferons.,online information:Interferon ga

Subcellular Location:

Secreted.

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