

DESCRIPTION

Species Reactivity	Human
Specificity	Detects human IFN- γ in direct ELISAs. In direct ELISAs, no cross-reactivity with recombinant mouse IFN- γ , recombinant rat IFN- γ , or recombinant porcine IFN- γ is observed.
Source	Monoclonal Mouse IgG _{2B} Clone # 25723
Purification	Protein A or G purified from hybridoma culture supernatant
Immunogen	<i>E. coli</i> -derived recombinant human IFN- γ Gln24-Gln166 Accession # AAP20098.1
Conjugate	Alexa Fluor 488 Excitation Wavelength: 488 nm Emission Wavelength: 515-545 nm
Formulation	Supplied in a saline solution containing BSA and Sodium Azide. See Certificate of Analysis for details. *Contains <0.1% Sodium Azide, which is not hazardous at this concentration according to GHS classifications. Refer to the Safety Data Sheet (SDS) for additional information and handling instructions.

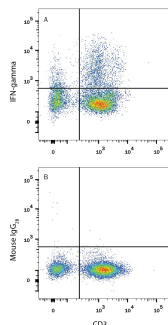
APPLICATIONS

Please Note: Optimal dilutions should be determined by each laboratory for each application. [General Protocols](#) are available in the Technical Information section on our website.

	Recommended Concentration	Sample
Intracellular Staining by Flow Cytometry	5 μ L/10 ⁶ cells	See Below

DATA

Intracellular Staining by Flow Cytometry



Detection of IFN- γ in Human PBMCs by Flow Cytometry.

Human peripheral blood mononuclear cells (PBMCs) treated with 50 ng/mL PMA, 1 μ g/mL Ionomycin, and 3 μ M Monensin overnight were stained with Mouse Anti-Human CD3 ϵ APC-conjugated Monoclonal Antibody (Catalog # [FAB100A](#)) and either (A) Mouse Anti-Human IFN- γ Alexa Fluor® 488-conjugated Monoclonal Antibody (Catalog # IC285G) or (B) Mouse IgG_{2B} Alexa Fluor 488 Isotype Control (Catalog # [IC0041G](#)). To facilitate intracellular staining, cells were fixed and permeabilized with FlowX FoxP3 Fixation & Permeabilization Buffer Kit (Catalog # [FC012](#)). View our protocol for [Staining Intracellular Molecules](#).

PREPARATION AND STORAGE

Shipping	The product is shipped with polar packs. Upon receipt, store it immediately at the temperature recommended below.
Stability & Storage	Protect from light. Do not freeze. <ul style="list-style-type: none"> 12 months from date of receipt, 2 to 8 °C as supplied.

BACKGROUND

Interferon-gamma (IFN-gamma, IFNG), also known as type II or Immune Interferon, exerts a wide range of immunoregulatory activities and is considered to be the prototype proinflammatory cytokine. Mature human IFN-gamma exists as a non-covalently linked homodimer of 20-25 kDa molecular weight variably glycosylated subunits. Glycosylation of IFN-gamma at sites Asn25 and Asn97 is critical for protease resistance. It shares 90% amino acid (aa) sequence identity with rhesus IFN-gamma, 59-64% with bovine, canine, equine, feline, and porcine IFN-gamma, and 37-43% with cotton rat, mouse, and rat IFN-gamma. IFN-gamma dimers bind to IFN-gamma RI (alpha subunits) which then interact with IFN-gamma RII (beta subunits) to form the functional receptor complex of two alpha and two beta subunits. Inclusion of IFN-gamma RII increases the binding affinity for ligand and the efficiency of signal transduction. IFN-gamma is produced by a variety of immune cells under inflammatory conditions, notably by T cells and NK cells. It plays a key function in host defense by promoting the development and activation of Th1 cells, chemoattraction and activation of monocytes and macrophages, up-regulation of antigen presentation molecules, and immunoglobulin class switching in B cells. It also exhibits antiviral, antiproliferative, and apoptotic effects. In addition, IFN-gamma functions as an anti-inflammatory mediator by promoting the development of regulatory T cells and inhibiting Th17 cell differentiation. The pleiotropic effects of IFN-gamma contribute to the development of multiple aspects of atherosclerosis.

Human IFN- γ Alexa Fluor® 488-conjugated Antibody

Monoclonal Mouse IgG_{2B} Clone # 25723

Catalog Number: IC285G

100 Tests, 25 Tests

PRODUCT SPECIFIC NOTICES

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