

DESCRIPTION

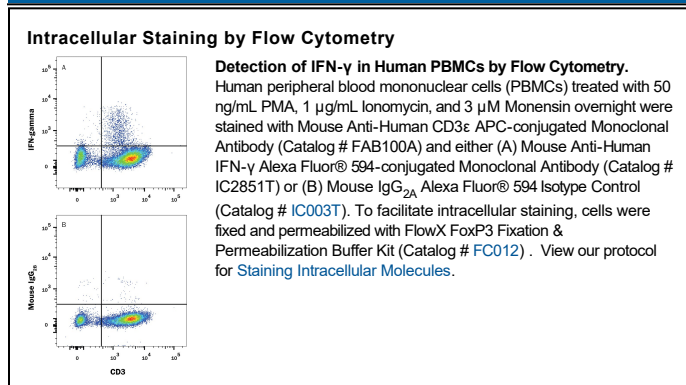
Species Reactivity	Human
Specificity	Detects human IFN- γ in direct ELISAs. In direct ELISAs, no cross-reactivity with recombinant mouse IFN- γ , recombinant rat IFN- γ , or recombinant porcine IFN- γ is observed.
Source	Monoclonal Mouse IgG _{2A} Clone # 25718
Purification	Protein A or G purified from hybridoma culture supernatant
Immunogen	<i>E. coli</i> -derived recombinant human IFN- γ Gln24-Gln166 Accession # AAP20098.1
Conjugate	Alexa Fluor 594 Excitation Wavelength: 590 nm Emission Wavelength: 617 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



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- γ), also known as type II or immune interferon, exerts a wide range of immunoregulatory activities and is considered to be the prototype proinflammatory cytokine (1, 2). Mature human IFN- γ exists as a non-covalently linked homodimer of 20-25 kDa variably glycosylated subunits (3). It shares 90% amino acid (aa) sequence identity with rhesus IFN- γ , 59-64% with bovine, canine, equine, feline, and porcine IFN- γ , and 37-43% with cotton rat, mouse, and rat IFN- γ . IFN- γ dimers bind to IFN- γ RI (alpha subunits) which then interact with IFN- γ RII (beta subunits) to form the functional receptor complex of two α and two β subunits. Inclusion of IFN- γ RII increases the binding affinity for ligand and the efficiency of signal transduction (4, 5). IFN- γ is produced by a variety of immune cells including monocytes, NK cells, $\gamma\delta$ T cells, CD8⁺ T cells, multiple T cell subsets (including Th1, CCR7⁻ T_{EM} and CD103⁺CD69⁺ T_{RM} cells plus proinflammatory FoxP3 regulatory T cells (6-12). It plays a key role in host defense by promoting the development and activation of Th1 cells, chemoattraction and activation of monocytes and macrophages, upregulation of antigen presentation molecules, and immunoglobulin class switching in B cells. It also exhibits antiviral, antiproliferative, and apoptotic effects (6, 13, 14). In addition, IFN- γ functions as an anti-inflammatory mediator by promoting the development of regulatory T cells and inhibiting Th17 cell differentiation (15, 16). The pleiotropic effects of IFN- γ contribute to the development of multiple aspects of atherosclerosis (7). Finally, IFN- γ regulates blood cell production, particularly during immune challenge. In particular, IFN- γ appears to promote monocyte production while depressing neutrophil, B cell and eosinophil production (17).

References:

1. Billiau, A. and P. Matthys (2009) Cytokine Growth Factor Rev. **20**:97.
2. Pestka, S. *et al.* (2004) Immunol. Rev. **202**:8.
3. Gray, P.W. and D.V. Goeddel (1982) Nature **298**:859.
4. Marsters, S.A. *et al.* (1995) Proc. Natl. Acad. Sci. USA **92**:5401.
5. Krause, C.D. *et al.* (2000) J. Biol. Chem. **275**:22995.
6. Schroder, K. *et al.* (2004) J. Leukoc. Biol. **75**:163.
7. Yamaguchi, R. *et al.* (2015) Blood Cells Mol. Dis. **55**:127.
8. Paolini, R. *et al.* (2015) Cytokine Growth Factor Rev. **26**:113.
9. Hou, L. *et al.* (2015) Int. Immunopharmacol. **28**:887.
10. de Graujo-Souza, P.S. *et al.* (2015) J. Immunol Res. **2015**:849573.
11. Geginat, J. *et al.* (2014) Front. Immunol. **5**:630.
12. Padiyan, P. and J. Zhu (2015) Cytokine **76**:13.
13. McLaren, J.E. and D.P. Ramji (2009) Cytokine Growth Factor Rev. **20**:125.
14. Zhu, L. *et al.* (2014) Int. Rev. Immunol. **34**:82.
15. Muhl, H. and J. Pfeilschifter (2003) Int. Immunopharmacol. **3**:1247.
16. Kelchtermans, H. *et al.* (2008) Trends Immunol. **29**:479.
17. de Bruin, A.M. *et al.* (2014) Blood **124**:2479.

PRODUCT SPECIFIC NOTICES

This product is provided under an agreement between Life Technologies Corporation and R&D Systems, Inc, and the manufacture, use, sale or import of this product is subject to one or more US patents and corresponding non-US equivalents, owned by Life Technologies Corporation and its affiliates. The purchase of this product conveys to the buyer the non-transferable right to use the purchased amount of the product and components of the product only in research conducted by the buyer (whether the buyer is an academic or for-profit entity). The sale of this product is expressly conditioned on the buyer not using the product or its components (1) in manufacturing; (2) to provide a service, information, or data to an unaffiliated third party for payment; (3) for therapeutic, diagnostic or prophylactic purposes; (4) to resell, sell, or otherwise transfer this product or its components to any third party, or for any other commercial purpose. Life Technologies Corporation will not assert a claim against the buyer of the infringement of the above patents based on the manufacture, use or sale of a commercial product developed in research by the buyer in which this product or its components was employed, provided that neither this product nor any of its components was used in the manufacture of such product. For information on purchasing a license to this product for purposes other than research, contact Life Technologies Corporation, Cell Analysis Business Unit, Business Development, 29851 Willow Creek Road, Eugene, OR 97402, Tel: (541) 465-8300. Fax: (541) 335-0354.