

## Brilliant Violet 421™ anti-human IFN-γ Antibody

<b>Catalog# / Size</b>	502531 / 25 tests 502532 / 100 tests
<b>Clone</b>	4S.B3
<b>Regulatory Status</b>	RUO
<b>Other Names</b>	Interferon-γ, Immune interferon, Type II interferon, T cell interferon, Macrophage-activating factor (MAF), IFN-g, IFN-gamma
<b>Isotype</b>	Mouse IgG1, κ
<b>Description</b>	Interferon-γ is a potent multifunctional cytokine which is secreted primarily by activated NK cells and T cells. Originally characterized based on anti-viral activities, IFN-γ also exerts anti-proliferative, immunoregulatory, and proinflammatory activities. IFN-γ can upregulate MHC class I and II antigen expression by antigen-presenting cells.

### Product Details

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<b>Verified Reactivity</b>	Human
<b>Reported Reactivity</b>	Chimpanzee, Baboon, Cynomolgus, Rhesus
<b>Antibody Type</b>	Monoclonal
<b>Host Species</b>	Mouse
<b>Immunogen</b>	Partially purified, native human IFN-γ
<b>Formulation</b>	Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide and BSA (origin USA).
<b>Preparation</b>	The antibody was purified by affinity chromatography and conjugated with Brilliant Violet 421™ under optimal conditions.
<b>Concentration</b>	Lot-specific (to obtain lot-specific concentration, please enter the lot number in our <a href="#">Concentration and Expiration Lookup</a> or <a href="#">Certificate of Analysis</a> online tools.)
<b>Storage &amp; Handling</b>	The antibody solution should be stored undiluted between 2°C and 8°C, and protected from prolonged exposure to light. <b>Do not freeze.</b>
<b>Application</b>	<a href="#">ICFC - Quality tested</a>
<b>Recommended Usage</b>	<p>Each lot of this antibody is quality control tested by <a href="#">intracellular immunofluorescent staining with flow cytometric analysis</a>. For flow cytometric staining, the suggested use of this reagent is 5 μl per million cells in 100 μl staining volume or 5 μl per 100 μl of whole blood.</p> <p>Brilliant Violet 421™ excites at 405 nm and emits at 421 nm. The standard bandpass filter 450/50 nm is recommended for detection. Brilliant Violet 421™ is a trademark of Sirigen Group Ltd.</p> <p><a href="#">Learn more about Brilliant Violet™.</a></p>

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<b>Excitation Laser</b>	Violet Laser (405 nm)
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<b>Application Notes</b>	<p><b>ELISA or ELISPOT Detection<sup>5</sup>:</b> The biotinylated 4S.B3 antibody is useful as a detection antibody for a sandwich ELISA or ELISPOT assay, when used in conjunction with purified NIB42 antibody (Cat. No. 502402/502404) or purified MD-1 antibody (Cat. No. 507502/507513) as the capture antibody.</p> <p><b>Flow Cytometry<sup>3,4,6-8</sup>:</b> The fluorochrome-labeled 4S.B3 antibody is useful for intracellular immunofluorescent staining and flow cytometric analysis to identify IFN-γ -producing cells within</p>
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mixed cell populations.

**Additional reported applications (for the relevant formats) include:** neutralization<sup>1,2</sup>, Western blotting, immunohistochemical staining of paraformaldehyde-fixed, saponin-treated tissue sections, and immunocytochemistry. The 4S.B3 antibody can neutralize the bioactivity of natural or recombinant IFN- $\gamma$ .

**Note:** For testing human IFN- $\gamma$  in serum or plasma, BioLegend's ELISA Max™ Sets (Cat. No. 430101 to 430106) are specially developed and recommended.

#### Application References

(PubMed link indicates BioLegend citation)

1. Meager A, *et al.* 1984. *J. Interferon Res.* 4:619. (Neut)
2. Meager A, 1987. *Lymphokines and Interferons: A Practical Approach.* IRL Press Ltd, Oxford, p. 105. (Neut)
3. Sester M, *et al.* 2002. *J. Virol.* 76:3748. (ICFC)
4. Infante-Duarte C, *et al.* 2000 *J. Immunol.* 165:6107. (ICFC)
5. Goodier M, *et al.* 2000. *J. Immunol.* 165:139. (ELISA)
6. Chen H, *et al.* 2005. *J. Immunol.* 175:591. (ICFC)
7. Smeltz RB, 2007. *J. Immunol.* 178:4786. (ICFC)
8. Iwamoto S, *et al.* 2007. *J. Immunol.* 179:1449. (ICFC) [PubMed](#)
9. Yoshino N, *et al.* 2000. *Exp. Anim. (Tokyo)* 49:97. (ICFC)

#### Product Citations

1. Liu Y, *et al.* 2017. *Oncogene.* 10.1038/onc.2017.209. [PubMed](#)
2. Agelidis A, *et al.* 2017. *Cell Rep.* 10.1016/j.celrep.2017.06.041. [PubMed](#)
3. Felices M, *et al.* 2018. *JCI Insight.* 3. [PubMed](#)
4. Chalan P, *et al.* 2016. *J Rheumatol.* 43: 1008 - 1016. [PubMed](#)
5. Siedlik J, *et al.* 2017. *J Immunol Methods.* 10.4049/jimmunol.1700003. [PubMed](#)
6. Langenberg MCC, *et al.* 2020. *Nat Med.* 26:326. [PubMed](#)
7. Wu B, *et al.* 2020. *Cell Metabolism.* 32(6):967-980.e5. [PubMed](#)

#### RRID

AB\_10900083 (BioLegend Cat. No. 502531)  
AB\_2561398 (BioLegend Cat. No. 502532)

## Antigen Details

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<b>Structure</b>	Cytokine; dimer; 20-25 kD (Mammalian)
<b>Bioactivity</b>	Antiviral/antiparasitic activities; inhibits proliferation; enhances MHC class I and II expression on APC
<b>Cell Sources</b>	CD8 <sup>+</sup> and CD4 <sup>+</sup> T cells, NK cells
<b>Cell Targets</b>	T cells, B cells, macrophages, NK cells, endothelial cells, fibroblasts
<b>Receptors</b>	IFN- $\gamma$ R $\alpha$ (CDw119) dimerized with IFN- $\gamma$ R $\beta$ (AF-1)
<b>Cell Type</b>	Tregs
<b>Biology Area</b>	Cell Biology, Immunology, Neuroinflammation, Neuroscience
<b>Molecular Family</b>	Cytokines/Chemokines
<b>Antigen References</b>	<ol style="list-style-type: none"><li>1. Fitzgerald K, <i>et al.</i> Eds. 2001. <i>The Cytokine FactsBook.</i> Academic Press, San Diego.</li><li>2. De Maeyer E, <i>et al.</i> 1992. <i>Curr. Opin. Immunol.</i> 4:321.</li><li>3. Farrar M, <i>et al.</i> 1993. <i>Annu. Rev. Immunol.</i> 11:571.</li><li>4. Gray P, <i>et al.</i> 1987. <i>Lymphokines</i> 13:151.</li></ol>
<b>Regulation</b>	Upregulated by IL-2, FGF-basic, EGF; downregulated by vitamin D3 or DMN; labile at pH2
<b>Gene ID</b>	<a href="#">3458</a>

## Related Protocols

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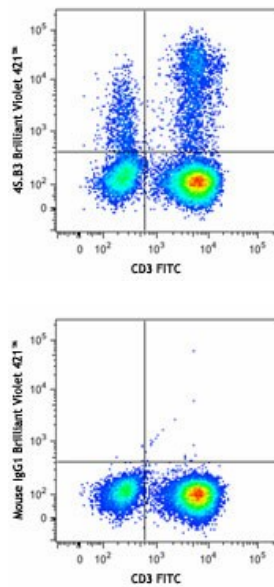
[Intracellular Cytokine Staining Protocol - Video](#)

[Intracellular Flow Cytometry Staining Protocol](#)

## Other Formats

PE anti-human IFN- $\gamma$ , APC anti-human IFN- $\gamma$ , FITC anti-human IFN- $\gamma$ , Biotin anti-human IFN- $\gamma$ , Purified anti-human IFN- $\gamma$ , Alexa Fluor® 488 anti-human IFN- $\gamma$ , Alexa Fluor® 647 anti-human IFN- $\gamma$ , Alexa Fluor® 700 anti-human IFN- $\gamma$ , Pacific Blue™ anti-human IFN- $\gamma$ , PerCP/Cyanine5.5 anti-human IFN- $\gamma$ , APC/Cyanine7 anti-human IFN- $\gamma$ , PE/Cyanine7 anti-human IFN- $\gamma$ , Brilliant Violet 421™ anti-human IFN- $\gamma$ , Brilliant Violet 570™ anti-human IFN- $\gamma$ , Brilliant Violet 605™ anti-human IFN- $\gamma$ , Brilliant Violet 650™ anti-human IFN- $\gamma$ , Brilliant Violet 711™ anti-human IFN- $\gamma$ , Brilliant Violet 785™ anti-human IFN- $\gamma$ , Brilliant Violet 510™ anti-human IFN- $\gamma$ , PE/Dazzle™ 594 anti-human IFN- $\gamma$ , APC/Fire™ 750 anti-human IFN- $\gamma$ , PerCP anti-human IFN- $\gamma$ , Brilliant Violet 750™ anti-human IFN- $\gamma$ , KIRAVIA Blue 520™ anti-human IFN- $\gamma$  Antibody, Spark NIR™ 685 anti-human IFN- $\gamma$  Antibody

## Product Data



Human peripheral blood lymphocytes were stimulated with PMA + Ionomycin for 6 hours (in the presence of monensin), surface stained with CD3 FITC, fixed, permeabilized, and then stained with IFN- $\gamma$  (clone 4S.B3) Brilliant Violet 421™ (top) or mouse IgG1,  $\kappa$  Brilliant Violet 421™ isotype control (bottom).

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