

Spark NIR™ 685 anti-mouse IFN-γ Antibody

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| Catalog# / Size | 505861 / 25 µg 505862 / 100 µg |
| Clone | XMG1.2 |
| Regulatory Status | RUO |
| Other Names | Interferon-γ, Immune interferon, Type II interferon, T cell interferon, Macrophage-activating factor (MAF) |
| Isotype | Rat IgG1, κ |
| Description | IFN-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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| Verified Reactivity | Mouse |
| Antibody Type | Monoclonal |
| Host Species | Rat |
| Immunogen | <i>E. coli</i> -expressed, recombinant mouse IFN-γ |
| Formulation | Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide |
| Preparation | The antibody was purified by affinity chromatography and conjugated with Spark NIR™ 685 under optimal conditions. |
| Concentration | 0.5 mg/mL |
| Storage & Handling | The antibody solution should be stored undiluted between 2°C and 8°C, and protected from prolonged exposure to light. Do not freeze . |
| Application | ICFC - Quality tested |
| Recommended Usage | Each lot of this antibody is quality control tested by intracellular immunofluorescent staining with flow cytometric analysis . For flow cytometric staining, the suggested use of this reagent is ≤ 0.25 µg per million cells in 100 µL volume. It is recommended that the reagent be titrated for optimal performance for each application. * Spark NIR™ 685 has a maximum excitation of 665 nm and a maximum emission of 685 nm. |
| Excitation Laser | Red Laser (633 nm) |
| Application Notes | ELISA^{1-4,11,14} or ELISPOT⁵ Detection: The biotinylated XMG1.2 antibody is useful as a detection antibody for a sandwich ELISA or ELISPOT assay, when used in conjunction with purified R4-6A2 antibody (Cat. No. 505702/505706) as the capture antibody and recombinant mouse IFN-γ (Cat. No. 575309) as the standard. ELISA or ELISPOT Capture: The purified XMG1.2 antibody is useful as a capture antibody for a sandwich ELISA or ELISPOT assay, when used in conjunction with biotinylated R4-6A2 antibody (Cat. No. 505704) as the detection antibody and recombinant mouse IFN-γ (Cat. No. 575309) as the standard. The LEAF™ purified antibody is suggested for ELISPOT capture (Cat. No. 505812). Flow Cytometry^{7,8,12,13,16}: The fluorochrome-labeled XMG1.2 antibody is useful for intracellular immunofluorescent staining and flow cytometric analysis to identify IFN-γ-producing cells within mixed cell populations. Neutralization^{1-3,9,10}: The XMG1.2 antibody can neutralize the bioactivity of natural or recombinant IFN-γ. The LEAF™ purified antibody (Endotoxin <0.1 EU/µg, Azide-Free, 0.2 µm filtered) is recommended for neutralization of mouse IFN-γ bioactivity <i>in vivo</i> and <i>in vitro</i> (Cat. No. 505812). For <i>in vivo</i> studies or highly sensitive assays, we recommend Ultra-LEAF™ purified antibody (Cat. No. 505834) with a lower endotoxin limit than standard LEAF™ purified antibodies (Endotoxin <0.01 EU/µg). Additional reported applications (for the relevant formats) include: Western blotting, |

immunohistochemical staining of frozen tissue sections^{6,22,23}, and immunocytochemistry.

Note: For testing mouse IFN- γ in serum, plasma or supernatant, BioLegend's ELISA Max™ Sets (Cat. No. 430801 to 430806) are specially developed and recommended.

Application References

(PubMed link indicates BioLegend citation)

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2. Sander B, *et al.* 1993. *J. Immunol. Meth.* 166:201. (ELISA, Neut)
3. Abrams J, *et al.* 1995. *Curr. Prot. Immunol.* John Wiley and Sons, New York. Unit 6.20. (ELISA, Neut)
4. Yang X, *et al.* 1993. *J. Immunoassay* 14:129. (ELISA)
5. Klinman D, *et al.* 1994. *Curr. Prot. Immunol.* John Wiley and Sons, New York. Unit 6.19. (ELISPOT)
6. Sander B, *et al.* 1991. *Immunol. Rev.* 119:65. (IHC)
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8. Ko SY, *et al.* 2005. *J. Immunol.* 175:3309. (FC) [PubMed](#)
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10. DeKrey GK, *et al.* 1998. *Infect. Immun.* 66:827. (Neut)
11. Dzhagalov I, *et al.* 2007. *J. Immunol.* 178:2113. (ELISA)
12. Lawson BR, *et al.* 2007. *J. Immunol.* 178:5366. (FC)
13. Lee JW, *et al.* 2006. *Nature Immunol.* 8:181. (FC) [PubMed](#)
14. Xu G, *et al.* 2007. *J. Immunol.* 179:5358. (ELISA) [PubMed](#)
15. Montfort M, *et al.* 2004. *J. Immunol.* 173:4084. [PubMed](#)
16. Haring JS, *et al.* 2008. *J. Immunol.* 180:2855. (FC) [PubMed](#)
17. Jordan JM, *et al.* 2008. *Infect Immun.* 76:3717. [PubMed](#)
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19. Charles N, *et al.* 2010. *Nat. Med.* 16:701. (FC) [PubMed](#)
20. Cui Y, *et al.* 2009. *Invest. Ophth. Vis. Sci.* 50:5811. (FC) [PubMed](#)
21. Mykkanen OT, *et al.* 2014. *PLoS One.* 9:114790. [PubMed](#)
22. Yokogawa M, *et al.* 2013. *Mol. Carcinog.* 52:760. (IHC)
23. Mottram PL, *et al.* 1998. *J Immunol.* 161:602. (IHC)

RRID

AB_2894641 (BioLegend Cat. No. 505861)

AB_2894641 (BioLegend Cat. No. 505862)

Antigen Details

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| Structure | Cytokine; dimer; 40-80 kD (Mammalian) |
| Bioactivity | Antiviral/antiparasitic activities; inhibits proliferation; enhances MHC class I and II expression on APCs |
| Cell Sources | CD8 ⁺ and CD4 ⁺ T cells, NK cells |
| Cell Targets | T cells, B cells, macrophages, NK cells, endothelial cells, fibroblasts |
| Receptors | IFN- γ R α (CDw119) dimerized with IFN- γ R β (AF-1) |
| Cell Type | Tregs |
| Biology Area | Cell Biology, Immunology, Neuroinflammation, Neuroscience |
| Molecular Family | Cytokines/Chemokines |
| Antigen References | <ol style="list-style-type: none">1. Fitzgerald K, <i>et al.</i> Eds. 2001. The Cytokine FactsBook. Academic Press, San Diego.2. De Maeyer E, <i>et al.</i> 1992. <i>Curr. Opin. Immunol.</i> 4:321.3. Farrar M, <i>et al.</i> 1993. <i>Annu. Rev. Immunol.</i> 11:571.4. Gray P, <i>et al.</i> 1987. <i>Lymphokines</i> 13:151. |
| Regulation | Upregulated by IL-2, FGF-basic, EGF; downregulated by 1- α -25-Dihydroxy vitamin D3, dexamethasone |
| Gene ID | 15978 |

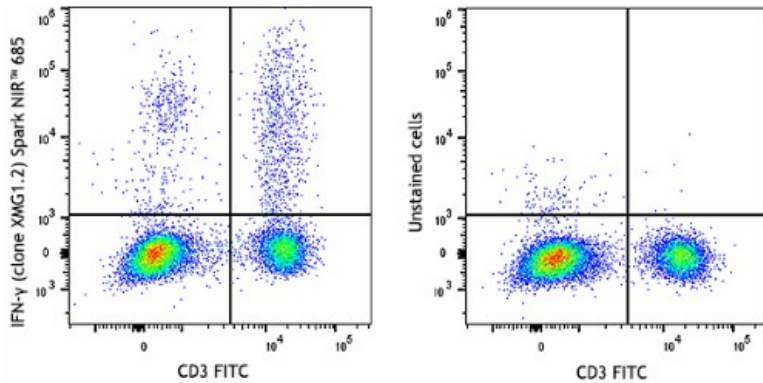
Related Protocols

[Intracellular Flow Cytometry Staining Protocol](#)

Other Formats

APC anti-mouse IFN- γ , Biotin anti-mouse IFN- γ , FITC anti-mouse IFN- γ , PE anti-mouse IFN- γ , Purified anti-mouse IFN- γ , Alexa Fluor® 488 anti-mouse IFN- γ , Alexa Fluor® 647 anti-mouse IFN- γ , Pacific Blue™ anti-mouse IFN- γ , PerCP/Cyanine5.5 anti-mouse IFN- γ , PE/Cyanine7 anti-mouse IFN- γ , Brilliant Violet 421™ anti-mouse IFN- γ , Brilliant Violet 650™ anti-mouse IFN- γ , Ultra-LEAF™ Purified anti-mouse IFN- γ , Brilliant Violet 711™ anti-mouse IFN- γ , Brilliant Violet 785™ anti-mouse IFN- γ , Brilliant Violet 605™ anti-mouse IFN- γ , Brilliant Violet 510™ anti-mouse IFN- γ , Purified anti-mouse IFN- γ (Maxpar® Ready), PE/Dazzle™ 594 anti-mouse IFN- γ , Alexa Fluor® 700 anti-mouse IFN- γ , APC/Cyanine7 anti-mouse IFN- γ , GolnVivo™ Purified anti-mouse IFN- γ , APC/Fire™ 750 anti-mouse IFN- γ , Spark NIR™ 685 anti-mouse IFN- γ

Product Data



C57BL/6 mouse splenocytes stimulated with PMA + Ionomycin in the presence of monensin (6 hrs) were stained with CD3 (clone 145-2C11) FITC, then fixed, permeabilized, and intracellularly stained with IFN- γ (clone XMG1.2) Spark NIR™ 685 (left) or CD3 FITC (right).

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