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Product Information

Anti-Interferon-y

produced in goat, affinity isolated antiserum

Catalog Number 19141

Synonym: Anti-IFN-γ

Product Description

Anti-Interferon- γ was produced in goat using recombinant, mouse IFN- γ , expressed in *E. coli*, as the immunogen. Affinity isolated antibody is obtained from goat anti-interferon- γ antiserum by immunospecific purification which removes essentially all goat serum proteins, including immunoglobulins, which do not specifically bind to mouse interferon- γ .

Interferon-γ exerts a variety of biological effects including antiviral activity, inhibition of cell or tumor growth^{2,3} and promotion of differentiation of B cells into immunoglobulin-producing cells.^{4,5} In addition to antiviral activity, human IFN-γ is a potent modulator of immune response and modifies cellular processes. 6 IFN-γ is classified as immune interferon. 6 IFN-γ functions as an activating factor to prime macrophages (MAF) for non-specific tumoricidal activity and activates monocytes to exert enhanced cytotoxicity against tumor cells.8 IFN-γ acts a signal for major histocompatibility antigen expression. FN-γ boosts cytotoxicity of natural killer cells and stimulates T cell cytotoxicity. The species specificity of IFN-y resides in the interaction of IFN-γ with its receptor. 10 Human IFN-γ does not bind specifically to mouse, hamster or bovine cells. 10

Reagent

Supplied as a lyophilized powder from 0.2 μ m-filtered phosphate buffered saline (pH 7.4) with 5% trehalose.

Precautions and Disclaimer

This product is for R&D use only, not for drug, household, or other uses. Please consult the Material Safety Data Sheet for information regarding hazards and safe handling practices.

Storage

Prior to reconstitution, store at $-20\,^{\circ}$ C. Reconstituted product may be stored 2-8 $^{\circ}$ C for a maximum of one month. For prolonged storage, freeze in working aliquots at $-20\,^{\circ}$ C. Avoid repeated freezing and thawing.

Reconstitution and Use

To one vial of lyophilized powder, add 1 ml of 0.2 μ m-filtered PBS to produce a 100 μ g/ml stock solution of Anti-IFN- γ . If aseptic technique is used, no further filtration should be needed for use in cell culture environments.

Product Profile

Neutralization: Anti-IFN- γ is tested for its ability to neutralize the biological activity of rmIFN-y on L929 cells. The ND₅₀ of the antibody is defined as the concentration of antibody resulting in a one-half maximal inhibition of bioactivity of rmIFN-y that is present at a concentration just high enough to elicit a maximum response. In this bioassay, rmIFN-γ was mixed with various dilutions of the antibody and the antigen-antibody mixture was added to confluent cultures of L929 cells in a 96-well plate. The assay mixture was incubated at 37 °C for 20-24 hours in a humidified CO₂ incubator. After incubation, the medium was aspirated from all wells and encephalomyocarditis virus (EMCV) was added to each test well. The 96-well plate was incubated for an additional 20-24 hours. The cells were fixed and examined for cytopathic effect by measurement of optical densities in a microplate reader at 540 nm.

Indirect Immunoblotting: 0.1-0.2 μ g/ml antibody detects 2 ng/lane of recombinant, mouse IFN- γ under reducing and non-reducing conditions.

The antibody may also be used in ELISA. By ELISA and immunoblotting, the antibody shows <1% cross-reactivity with recombinant human IFN-γ. In addition, by direct ELISA, the antibody does not cross-react with other cytokines tested.*

References

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ADM,PHC 06/11-1

* rhANG, rhAnnexin V, rhAR, rhB7-1, rhB7-2, rmB7-2, rhBTC, rh β -NGF, rhBDNF, rmC10, rhCD4, rhCD8, rhCD28, rhCNTF, rrCNTF, rhEGF, rhENA-78, rhEPO, rhFGFa, rhFGFb, rhFGF-4, rhFGF-5, rhFGF-6, rhFGF-7, rhFGF-9, rhG-CSF, rhG-CSF, rhG-CSF, rhGDNF, rhGM-CSF, rhGM-CSF, rhGRO(, rhGRO α , rhGRO β , rhHB-EGF, rhHRG- α , rhHGF, rhI-309, rhIGF-I, rhIGF-IR, rhIGF-II, rhIL-1 α , rhIL-1 RI, rhIL-1 RII, rmIL-1 α , rhIL-1 β , rmIL-1 β , rrIL-1 β , rrIL-1 β , rhIL-1 ra, rmIL-2, rhIL-2 sR α , rhIL-2 sR β , rhIL-2 sR(, rmIL-2, rhIL-3, rhIL-3 sR α , rmIL-3, rhIL-4, rhIL-4 sR, rmIL-4, rhIL-5, rhIL-5 sR α , rmIL-5 sR β , rmIL-5, rhIL-6, rhIL-6 sR, rmIL-6, rhIL-7, rhIL-7, rhIL-7, rhIL-8, rhIL-9, rmIL-9, rhIL-10 sR, rmIL-10, rmIL-10 sR, rhIL-11, rhIL-12, rmIL-12, rmIL-13, rmIL-13, rhIL-15, rhIP-10, rhJAK-1, rmJAK-2, rmJE, rmKC, rhLIF, rhLIF R, rmLIF, rhM-CSF, rmM-CSF, rhMCP-1, rhMCP-1 R, rhMCP-2, rhMCP-3, rhMidkine, rhMIF, rhMIP-1 α , rmMIP-1 α , rhMIP-1 α , rmMIP-1 α , rmMIP-2, rhNT-3, rhNT-4, rhOSM, rhPD-ECGF, hPDGF, pPDGF, rhPDGF-AA, rhPDGF-AB, rhPDGF-BB, rhPDGF R α , rhPIGF, rhPTN, rhRANTES, rhSCF, rmSCF, rhsgp130, rhSLPI, rhSTAT-1, rmSTAT-3, rmSTAT-4, hTfR, rhTGF- α , rhTGF- β 3, raTGF- β 5, rhLAP (TGF- β 1), rhLatent TGF- β 1, rhTGF- β 5 sRIII, rhTNF- α , rmTNF- α , rrTNF- α , rhTNF- β , rhsTNF RII, rhSTNF RII, rhTPO, rmTPO, rhVEGF, rmVEGF.