Monkey IFN-γ ELISA development kit

Product Code: 3420M-1H-6

CONTENTS, development kit for 6 plates:

Vial 1 (green top) Monoclonal antibody GZ-4 (150 µl) Concentration: 1 mg/ml

Vial 2 (yellow top) Biotinylated monoclonal antibody 7-B6-1 (80 µl) Concentration: 1 mg/ml

Vial 3 (white top) Streptavidin-Horseradish Peroxidase (80 μl)

Vial 4 Recombinant human IFN-γ standard (1μg)

To ensure total recovery of stated quantity, vials have been overfilled.

STORAGE:

Shipped at ambient temperature. On arrival vials should be stored refrigerated at 4-8°C.

General

Intended use: For quantitative determination of native and recombinant monkey IFN- γ in solution, e.g. cell culture supernatant.

Serum/plasma samples: Please note that determination of analyte in serum/plasma samples by this kit requires the use of ELISA diluent (product code: 3652-D2) for dilution of samples, standard and detection antibody. The diluent prevents false positive read-outs which may be caused by interference of heterophilic antibodies commonly found in human serum/plasma and possibly also in monkey serum/plasma. Please contact Mabtech for further information.

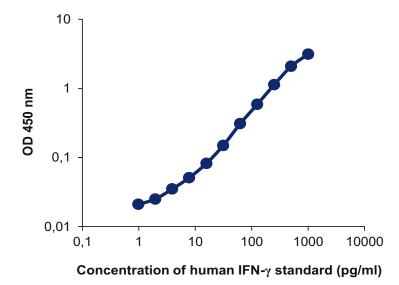
Specificity: Native and recombinant human IFN- γ and native IFN- γ from rhesus and cynomolgus macaques. Inquire for reactivities with other monkey species. For detection of chimpanzee IFN- γ the ELISA kit for human IFN- γ can be used.

Reagents: Antibodies are supplied in sterile-filtered (0.2 μ m) PBS with sodium azide (0.02%). Streptavidin-HRP is supplied in PBS with 1% BSA and 0.002% Kathon CG.

Standard range: 4-400 pg/ml

Standard calibration: 1 ng of supplied standard equals 176 U of Gxg01-902-535 NIAID*-standard according to repeated calibrations. Calibration is batch-specific.

*National Institute of Allergy and Infectious Diseases, USA.



Guidelines for Monkey IFN-y ELISA

- **Day 1** 1. Coat a high protein binding ELISA plate with mAb GZ-4, diluted to 2 μg/ml in PBS, pH 7.4, by adding 100 μl/well. Incubate overnight at 4-8°C.
- Day 2 2. Wash twice with PBS ($200 \mu l/well$).
 - 3. Block plate by adding 200 μl/well of PBS with 0.05% Tween 20 containing 0.1% BSA (incubation buffer). Incubate for 1 hour at room temperature.
 - 4. Wash five times with PBS containing 0.05% Tween.
 - 5. Prepare hIFN- γ standard by reconstituting contents of vial 4 in 1 ml PBS with 1% BSA to a concentration of 1 µg/ml. Leave at room temperature for 15 minutes and then vortex the tube. The stock solution should be used immediately or stored in aliquots at -20°C for future use. We recommend the aliquots not be refrozen after initial use. For the test, prepare dilutions of the stock using the standard range as a guideline.
 - 6. Add 100 μl/well of samples or standards diluted in incubation buffer or ELISA diluent for serum/plasma samples and incubate for 2 hours at room temperature.
 - 7. Wash as in step 4.
 - 8. Add 100 μl/well of mAb 7-B6-1-biotin at 1 μg/ml in incubation buffer or ELISA diluent for serum/plasma samples. Incubate for 1 hour at room temperature.
 - 9. Wash as in step 4.
 - 10. Add 100 μl/well of Streptavidin-HRP diluted 1:1000 in incubation buffer. Incubate for 1 hour at room temperature. **Please note that sodium azide used in buffers will inhibit HRP activity.**
 - 11. Wash as in step 4.
 - 12. Add 100 µl/well of appropriate substrate solution.
 - 13. Measure the optical density in an ELISA reader after suitable developing time. If required stop the reaction first.

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