

Monkey IFN- γ ELISA development kit

Product Code: 3421M-1H-6

CONTENTS, development kit for 6 plates:

Vial 1 (green top)

Monoclonal antibody MT126L (300 μ l)

Concentration: 0.5 mg/ml

Vial 2 (yellow top)

Biotinylated monoclonal antibody 7-B6-1 (80 μ l)

Concentration: 1 mg/ml

Vial 3 (white top)

Streptavidin-Horseradish Peroxidase (80 μ l)

Vial 4

Recombinant human IFN- γ standard (1 μ g)

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.

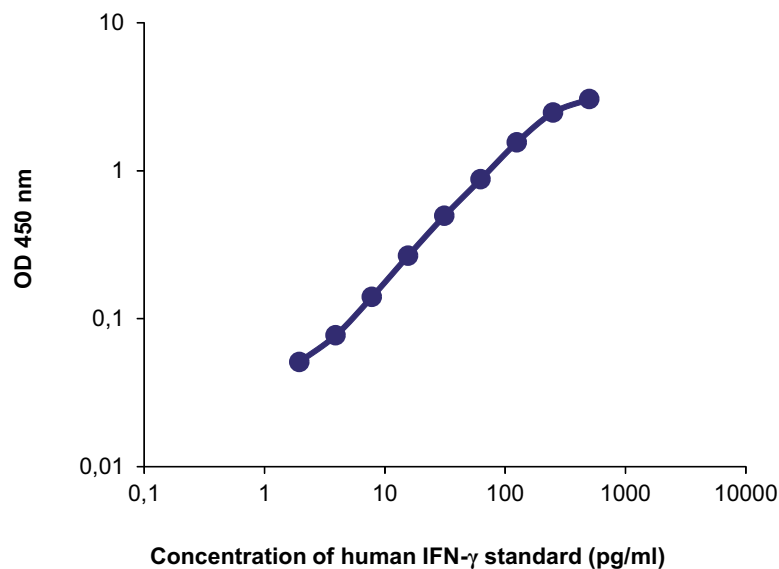
Serum/plasma samples: Please note that determination of analyte in serum/plasma samples by this kit requires the use of ELISA diluent (product code: 3652-D2) for dilution of samples, standard and detection antibody. The diluent prevents false positive read-outs which may be caused by interference of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

Specificity: IFN- γ from human, macaques (cynomolgus, rhesus and pig-tailed), baboon, sooty mangabey, African green monkey, aotus, common marmoset and squirrel monkey.

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: No international standard exists for calibration.



Guidelines for Monkey IFN- γ ELISA

- Day 1**
1. Coat a high protein binding ELISA plate with mAb MT126L, diluted to 2 $\mu\text{g}/\text{ml}$ in PBS, pH 7.4, by adding 100 $\mu\text{l}/\text{well}$. Incubate overnight at 4-8°C.
- Day 2**
2. Wash twice with PBS (200 $\mu\text{l}/\text{well}$).
 3. Block plate by adding 200 $\mu\text{l}/\text{well}$ 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 $\mu\text{g}/\text{ml}$. 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 $\mu\text{l}/\text{well}$ 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 $\mu\text{l}/\text{well}$ of mAb 7-B6-1-biotin at 1 $\mu\text{g}/\text{ml}$ 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 $\mu\text{l}/\text{well}$ 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 $\mu\text{l}/\text{well}$ of appropriate substrate solution e.g. TMB, available from Mabtech product code 3652-F10.
 13. Measure the optical density in an ELISA reader after suitable developing time. If required stop the reaction first.

Developed and manufactured by MABTECH AB, Sweden, whose quality management system complies with the standards ISO 9001:2015 & ISO 13485:2016.



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