Llama IFN-γ ELISA development kit

Product Code: 3123-1A-6

CONTENTS, development kit for 6 plates:

Vial 1 (green top)

Monoclonal antibody bIFNγ-I (300 μl)

Concentration: 0.5 mg/ml

Vial 2 (red top)

Biotinylated monoclonal antibody PAN (40 µl)

Concentration: 0.5 mg/ml

Vial 3 (white top)

Streptavidin-Alkaline Phosphatase (80 µl)

Vial 4

Recombinant llama IFN-y standard

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

STORAGE:

Shipped at ambient temperature. On arrival box 1 should be stored refrigerated at 4-8°C and box 2 should be stored frozen at -20°C.

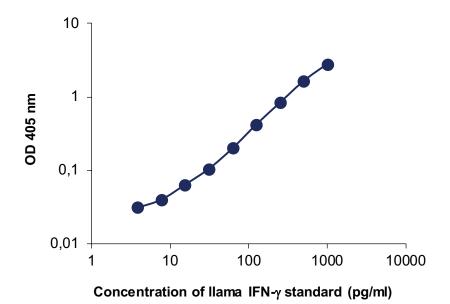
General

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

Reagents: Antibodies are supplied in sterile-filtered (0.2 μ m) PBS with sodium azide (0.02%). Streptavidin-ALP is supplied in 0.1 M Tris buffer with 0.002% Kathon CG.

Standard range: 7-700 pg/ml

Standard calibration: No international standard exists for calibration.



Guidelines for Llama IFN-γ ELISA

- **Day 1** 1. Coat a high protein binding ELISA plate with mAb bIFNγ-I, diluted to 2 μ g/ml in PBS, pH 7.4, by adding 100 μ l/well. Incubate overnight at 4-8°C.
- **Day 2** 2. Wash twice with PBS (200 μl/well).
 - 3. Block plate by adding 200 μ l/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 llama IFN-γ standard by reconstituting the contents of vial 4 in 1 ml PBS with 1% BSA to give a concentration of 0.5 μg/ml. Leave at room temperature for 15 minutes, then vortex the tube and spin down. Use immediately or store in aliquots at -20°C for future use. We recommend the aliquots not to 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 and incubate for 2 hours at room temperature.
 - 7. Wash as in step 4.
 - 8. Add 100 μ l/well of mAb PAN-biotin at 0.2 μ g/ml in incubation buffer. Incubate for 1 hour at room temperature.
 - 9. Wash as in step 4.
 - 10. Add 100 μl/well of Streptavidin-ALP diluted 1:1000 in incubation buffer. Incubate for 1 hour at room temperature.
 - 11. Wash as in step 4.
 - 12. Add 100 μl/well of appropriate substrate solution e.g. p-nitrophenyl-phosphate (pNPP), available from Mabtech product code 3652-P10.
 - 13. Measure the optical density (405 nm for pNPP) in an ELISA reader after suitable developing time.



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





The products are for research use only.

MABTECH shall not be liable for the use or handling of the product or for consequential, special, indirect or incidental damages there from.

Mabtech AB (Head Office)

Sweden Tel: +46 8 716 27 00 mabtech@mabtech.com Mabtech, Inc.

USA Tel: +1 513 871-4500 mabtech.usa@mabtech.com