

ELISA Flex: Mouse IFN- α (ALP)

3326-1A-6 | 3326-1A-20

ELISA Flex kit for quantitative determination of native and recombinant mouse IFN- α in solution, e.g. cell supernatant and serum/plasma.

The kit includes		3326-1A-6 for 6 plates	3326-1A-20 for 20 plates
Capture mAbs:	MT24A/44A/104 (0.5 mg/ml)	350 μ l	1200 μ l
Detection mAbs:	MT9L/14A/113, biotin (0.5 mg/ml)	150 μ l	500 μ l
	Streptavidin-ALP	80 μ l	250 μ l
	Recombinant mouse IFN- α ELISA standard	1 vial	1 vial
	Standard reconstitution buffer A5	1 ml	1 ml

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

Shipping and storage

Shipped at ambient temperature. All reagents should be stored at 4-8 °C upon receipt, except the standard which should be stored at -20 °C. Antibodies are supplied in sterile-filtered PBS with sodium azide (0.02%). Streptavidin-ALP is supplied in 0.1 M Tris buffer with 0.002% Kathon CG. The expiry date indicates how long unopened products, stored according to instructions, are recommended for use.

General and Preparations

Specificity

The kit contains matched monoclonal antibodies (mAbs) specific for native and recombinant mouse IFN- α . Subtypes 1, 2, 4, 5, 6T, 7/10 (129/Sv), 8/6, 9, 11, 12, 13, 14, 15 (A), and B are detected.

Standard range

8-1000 pg/ml

Calibration

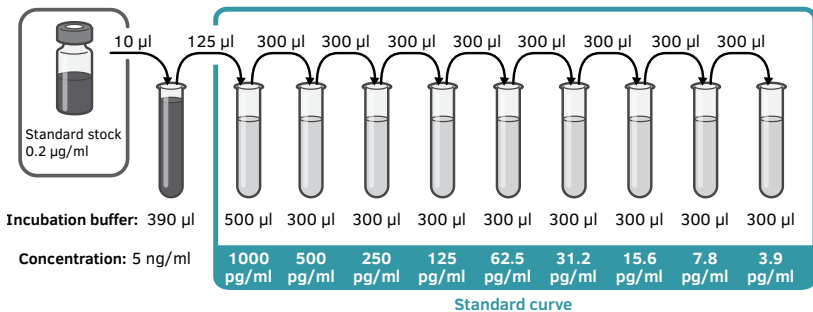
No international standard exists for calibration.

Reconstitution of ELISA standard

Reconstitute the ELISA standard to a stock solution of 0.2 $\mu\text{g/ml}$ by adding 1 ml of the standard reconstitution buffer. Allow the standard to dissolve for 5 minutes and mix thoroughly. The standard should be kept in aliquots at -20°C . Avoid repeated freeze-thaw cycles.

Preparation of standard curve

Prepare within 30 minutes of use. Volumes are sufficient for duplicates.



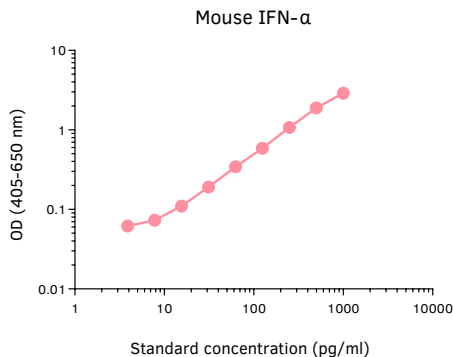
Protocol

Day 1

1. Add 100 μ l/well of capture mAbs MT24A/44A/104 diluted to 3 μ g/ml in PBS, pH 7.4. Use high protein binding ELISA plates. Incubate overnight at 4-8 $^{\circ}$ C.

Day 2

2. Empty the plate and add 200 μ l/well of PBS with 0.05% Tween 20 and 0.1% BSA (incubation buffer) to block the plate. Incubate for 1 hour at room temperature.
3. Wash the plate 5 times with PBS containing 0.05% Tween 20 (300 μ l/well).
4. Add 100 μ l/well of samples or standards diluted in incubation buffer. Include assay background control, i.e. wells without standard. Incubate for 2 hours at room temperature.
5. Wash as above.
6. Add 100 μ l/well of detection mAbs MT9L/14A/113-biotin diluted to 1 μ g/ml in incubation buffer. Incubate for 1 hour at room temperature.
7. Wash as above.
8. Add 100 μ l/well of Streptavidin-ALP diluted 1:1000 in incubation buffer. Incubate for 1 hour at room temperature.
9. Wash as above.
10. Add 100 μ l/well of pNPP substrate (product code: 3652-P10) and incubate at room temperature protected from direct light for approximately 60 minutes.
11. Measure the optical density in an ELISA reader at 405 nm. Preferably use a reader capable of subtracting a reference wavelength of between 570 and 650 nm. Representative standard curve shown below.



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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