

Bovine IL-13 ELISA Reagent Set

Cat# NB-06-1116

Technical Notes

This kit is for the quantitative measurement of Bovine IL-13 in cell culture supernatants. If assaying other sample types, an appropriate Sample and Standard Diluent will need to be developed and validated. Any changes to the ELISA protocol may significantly affect the results generated and will require optimization.

Included Components

Description	Quantity	Component Number
Bovine IL-13 Coated Plate	2 each	VS0148B-CP
Bovine IL-13 Standard	2 each	VS0148B-ST
Bovine IL-13 Detection Antibody	2 each	VS0148B-DA
Streptavidin-HRP	1 each	AR0068-001
Plate Sealer	6 each	N/A

Additional Reagents Required

Reagent	Formulation
DPBS	0.008M sodium phosphate, 0.002M potassium phosphate, 0.14M sodium chloride, 0.01M potassium chloride, pH 7.4
Standard and Sample Diluent	Complete cell culture medium used to generate cell culture supernatant samples.
Reagent Diluent	It is critical that this medium contain at least 1% carrier protein. If the medium does not contain carrier protein, use Reagent Diluent to dilute the Standard and samples.
Wash Buffer	4% BSA in DPBS, 0.2 µm filtered
Substrate	0.05% Tween®-20 in DPBS
Stop Solution	3,3',5,5'-tetramethylbenzidine (TMB) Substrate ELISA Accessory 0.18 M Sulfuric Acid ELISA Accessory

Component Preparation

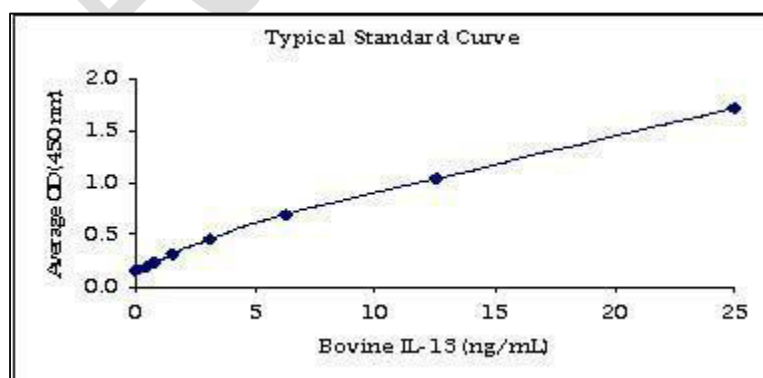
Component	Preparation
Bovine IL-13 Standard	Reconstitute Standard in 1 mL Standard and Sample Diluent. The Standard now has a concentration of 25 ng/mL. Prepare 1:1 serial dilutions of the Standard by mixing 250 µL Standard with 250 µL

Bovine IL-13	Standard and Sample Diluent. Repeat 1:1 serial dilutions until reach a final concentration of 0.39 ng/mL. Use Standard and Sample Diluent as a zero standard.
Detection Antibody Working Solution	Reconstitute Detection Antibody in 500 μ L Reagent Diluent. Dilute the 500 μ L of reconstituted Detection Antibody in 11.5 mL Reagent Diluent.
Streptavidin-HRP Working solution	Dilute 500 μ L of Streptavidin-HRP in 11.5 mL Reagent Diluent.

Elisa procedure

1. Prepare Standard and cell culture supernatant sample dilutions in Standard and Sample Diluent.
2. Add 100 μ L of Standard or sample to appropriate wells.
Note: Run each Standard or sample in duplicate.
3. Cover plate with Plate Sealer and incubate at room temperature (20-25C) for 1 hour.
4. Wash plate FOUR times with Wash Buffer.
Note: Gently squeeze the long sides of plate frame before washing to ensure all strips remain securely in the frame. Empty plate contents. Use a squirt wash bottle to vigorously fill each well completely with 1X Wash Buffer, then empty plate contents. Repeat procedure three additional times for a total of FOUR washes. Blot plate onto paper towels or other absorbent material.
5. Add 100 μ L of Detection Antibody Working Solution to each well.
6. Cover plate with Plate Sealer and incubate at room temperature for 1 hour.
7. Wash plate FOUR times with Wash Buffer as described in step 4.
8. Add 100 μ L of Streptavidin-HRP Working Solution to each well.
9. Cover plate with Plate Sealer and incubate at room temperature for 30 minutes.
10. Wash plate FOUR times with Wash Buffer as described in step 4.
11. Add 100 μ L of TMB Substrate Solution to each well.
12. Develop the plate in the dark at room temperature for 30 minutes.
Note: Do NOT cover plate with Plate Sealer.
13. Stop reaction by adding 100 μ L of Stop Solution to each well.
14. Measure absorbance on a plate reader at 450 nm.

Typical Standard Curve



Data represents a typical standard curve generated using the Neo Biotech Bovine IL-13 ELISA Development Kit. A standard curve should be generated with each assay.

Representative Data

Stimulant Bovine IL-13 (ng/mL) Unstimulated 0.4 Phorbol 12-myristate 13-acetate (PMA; 10 ng/mL) and Ionomycin (500 ng/mL) 1.8	PBMCs harvested by ficoll density gradient from an apparently healthy bovine were suspended in RPMI medium containing 10% serum and stimulated as desired. The cell-free supernatants were harvested following three days stimulation and analyzed in the Neo Biotech Bovine IL-13 ELISA Development Kit.
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