



# Bovine IFN- $\gamma$ ELISpot Development Module

Catalog Number: SEL2300

## REAGENTS PROVIDED

**Bovine IFN- $\gamma$  Capture Antibody Concentrate** (Part # 843915 - 1 vial of lyophilized anti-bovine IFN- $\gamma$ ).

**Bovine IFN- $\gamma$  Detection Antibody Concentrate** (Part # 843916) - 1 vial of lyophilized biotinylated antibody specific for bovine IFN- $\gamma$ .

\*Each vial contains sufficient antibodies to run ELISpot assays on approximately five 96-well microplates, when using the protocol provided.

## REAGENT PREPARATION & STORAGE

**Capture Antibody Concentrate** - Reconstitute with 1 mL of PBS. After reconstitution, store at 2-8 °C for up to 60 days or aliquot and store at -20 °C to -70 °C for up to 6 months.

**Detection Antibody Concentrate** - Reconstitute with 1 mL of Reagent Diluent. After reconstitution, store at 2-8 °C for up to 60 days or aliquot and store at -20 °C to -70 °C for up to 6 months.

**Note:** For optimal performance, prepare the working dilutions of the Capture and Detection Antibodies immediately before use.

## OTHER SUPPLIES REQUIRED

- ELISpot Blue Color Module or equivalent (R&D Systems, Catalog # SEL002)
- PBS (137 mM NaCl, 2.7 mM KCl, 8.1 mM Na<sub>2</sub>HPO<sub>4</sub>, 1.5 mM KH<sub>2</sub>PO<sub>4</sub>, pH 7.2-7.4, 0.2  $\mu$ m filtered).
- Wash Buffer (i.e. 0.05% Tween<sup>®</sup> 20 in PBS).
- Blocking Buffer (i.e. 1% BSA, 5% Sucrose in PBS).
- Reagent Diluent (i.e. 1% BSA in PBS, pH 7.2-7.4, 0.2  $\mu$ m filtered).
- 37 °C CO<sub>2</sub> incubator.
- Deionized H<sub>2</sub>O.
- Positive Control - Use recombinant bovine IFN- $\gamma$  (R&D Systems, Catalog # 2300-BG) or cells known to secrete bovine IFN- $\gamma$ .
- 96-well plates (Nitrocellulose-bottom plates, PVDF-bottom Immunospot<sup>®</sup> plates, or flat-bottom polystyrene Immulon<sup>®</sup> ELISA plates).
- Squirrt bottle, manifold dispenser, or automated microplate washer.
- Dissection microscope or an automated ELISpot Reader.

## PROTOCOL

When a 96-well PVDF microplate is used, a 1:60 dilution of the Capture and Detection antibodies is recommended. **Each investigator should determine the optimal working dilution of the antibodies depending on the type of microplate, Wash Buffer and Blocking Buffer used.**

1. Calculate the total volume of Capture Antibody needed and dilute to the working concentration using PBS.
2. Immediately add 100  $\mu$ L of the diluted Capture Antibody per well. Cover the plate with the lid and incubate overnight at 2-8 °C.
3. Aspirate capture antibody from each well and wash 3 times with Wash Buffer or PBS (350  $\mu$ L/well) using either a squirt bottle, manifold dispenser, or autowasher. After the final wash, remove any remaining liquid by inverting the plate and blotting it against a clean paper towel.  
**Note:** Do not touch the membranes during washing to avoid damage.
4. Block membranes by adding 200  $\mu$ L of Blocking Buffer to each well. Incubate for 2 hours at room temperature.
5. Aspirate Blocking Buffer as described in step 3. Rinse with the same media in which the cells will be cultured.  
**Note:** Do not discard the culture media until cells are ready to be plated.
6. Aspirate culture media from the plate and immediately fill appropriate wells with 100  $\mu$ L of culture media containing bovine IFN- $\gamma$  secreting cells. Incubate at 37 °C in a 5% CO<sub>2</sub> incubator. Incubation time must be determined empirically.  
**Note:** We recommend running a positive control (recombinant protein), negative control (same number of unstimulated cells as stimulated cells), and background control (sterile culture media) with each assay.
7. Wash plate 4 times with Wash Buffer. Remove any remaining Wash Buffer by inverting the plate and blotting it against a clean paper towel.
8. Calculate the total volume of Detection Antibody needed and dilute to the working concentration using Reagent Diluent.
9. Add 100  $\mu$ L of the diluted Detection Antibody per well. Cover the plate with the lid and incubate overnight at 2-8 °C.
10. Aspirate Detection Antibody and wash as described in step 3. Microplates are ready for color development.

## COLOR DEVELOPMENT

Color development may be done using the ELISpot Blue Color Module that may be purchased separately. Alternatively, another chromogen of choice may be used.

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