

Human IFN-γ CytoSetTM

10 Plate Format

Lot-specific Technical Data Sheet

Lot#: 094604/P **Expiration:** 30.06.2011

Catalog # CHC1233

Coating Antibody: Anti-Human IFN-y (0.250 mg/0.125 mL)

Part Number: Lot Number: 9K13/1

Form: Liquid, 1 vial, contains 0.1% sodium azide Storage: Store at 2-8°C until expiration date.

Recommended Dilution: Dilute to 2 µg/mL with Coating Buffer A (Cat. # CB07100, or see Recommended Buffers). For example, to make

10 mL (enough to coat 1 plate), add 10 μL coating antibody to 9.990 μL Coating Buffer A.

Detection Antibody: Anti- Human IFN-y Biotin (0.025 mg/ 0.125 mL)

Part Number: 58.123.03 Lot Number: 9K13/1

Form: Liquid, 1 vial, contains 0.1% sodium azide Storage: Store at 2-8°C until expiration date.

Recommended Dilution: Dilute to 0.16 µg/mL with Assay Buffer (Cat. # DS98200, or see Recommended Buffers). For example, to make

enough for 1 plate, add 4.4 μL detection antibody to 5.4956 mL Assay Buffer.

Standard: Recombinant Human IFN-y

Part Number: 58.123.10 (additional vials of standard may be purchased using this part number)

Lot Number: 9G11/1

Form: Lyophilized, 5 vials Storage: Store at 2-8°C.

Reconstitute with Assay Buffer (Cat. # DS98200 or see Recommended Buffers) to yield a stock of 10,000 pg/mL. Reconstitution:

After 10 minutes of rehydratation, use the standard stock immediately or aliquot in polypropylene tubes and freeze

at -80°C. Do not store at room temperature or at 4°C and do not subject to more than one freeze-thaw cycle. Standard Curve:

Dilute standard stock to 2,000 pg/mL (120 μL stock plus 480 mL Assay Buffer) with Assay Buffer (Cat. # DS98200 or see Recommended Buffers). Add 300 µL Assay Buffer to 6 tubes and label as as 1000, 500, 250, 125, 62.5 and 31.2 pg/mL. Make serial dilutions starting with 2,000 pg/mL by transferring 300 μL of each

standard to next tube and vortexing each tube. Assay Buffer should be used as the zero standard.

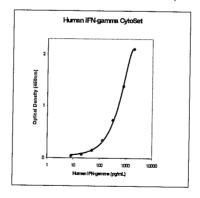
Streptavidin-HRP:

0.250 mL Part Number: 41.000.03 Lot Number: 9F8/1

Form: Liquid, 1vial, contains 0.05% thymol Store at 2-8°C until expiration date. Storage:

Recommended Dilution: Dilute to 1/5000 For example, to make enough for 1 plate, add 2 µL of streptavidin-HRP to 9.998 mL of Assay

Buffer (Cat. # DS98200 or see Recommended Buffers).



Representative standard curve was generated by following the recommended assay procedure, which includes the use of the BioSource CytoSetTM Buffer Set (Cat. # CNB0011)

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Intended Use and Materials Provided

The CytoSet™ for Human IFN-γ contains components required to construct an enzyme-linked immunoassay for the specific and quantitative measurement of IFN-7. Sufficient quantities of all reagents are provided to yield 10 plates of 96 wells if the recommended assay procedure and recommended storage and handling of materials are followed as specified on this insert. The materials provided are FOR RESEARCH USE

Recommended Buffers and Solutions

The BioSource CytoSet™ Buffer Set (Cat. # CNB0011) containing Coating Buffers A and B, Assay Buffer, Substrate Solution (TMB), Stop Solution, and Wash Buffer is recommended.

Coating Buffer A (Cat. # CB07100) from BioSource is recommended. Alternate buffer choice listed below. Coating Buffer A:

8.0 g NaCl, 1.13 g Na₂HPO₄, 0.2 g KH₂PO₄, 0.2 g KCl, 0.1% ProClinTM; q.s. to 1.0 L with distilled H₂O, pH to 7.4.

Coating Buffer B: Coating Buffer B (Cat. # CB01100) from BioSource is recommended. Alternate buffer choice listed below.

4.3 g NaHCO₃, 5.3 g Na₂CO₃, 0.1% ProClinTM. q.s. to 1.0 L with distilled H₂O₂, pH to 9.4.

Assay Buffer (Cat. # DS98200) from BioSource is recommended. Alternate buffer choice listed below.

8.0 g NaCl, 1.13 g Na₂HPO₄, 0.2 g KH₂PO₄, 0.2 g KCl, 5.0 g bovine serum albumin (fraction V), 1 mL Tween 20 and 0.5% ProClinTM as a preservative; q.s. to 1.0 L with distilled H_2O , pH to 7.4.

Wash Buffer: Wash Buffer 25x (Cat. # WB01) from BioSource is recommended. Alternate buffer choice listed below.

0.2 g KH₂PO₄ 1.9 g, K₂HPO₄ .3H₂O 0.4 g EDTA, 0.5 mL Tween 20; q.s. to 1.0 L with distilled H₂O, pH to 7.4.

TMB (Cat. # SB01) from BioSource is recommended. Alternate solution choice listed below. Substrate Solution:

Tetramethylbenzidine (TMB) and Hydrogen Peroxide.

6. Stop Solution: Stop Solution (Cat.# SS01100) from BioSource is recommended. Alternate solution choice listed below.

1.8 N H₂SO₄.

Assay Optimization

Assay Buffer:

CytoSctsTM from BioSource are designed to be very flexible for your experiments. Consequently, the assay procedure contains only recommendations. The assay procedure has been optimized for use with tissue culture samples. However, serum and plasma samples may be used but may require that certain assay parameters be modified. Investigators are advised to determine optimal buffer formulations, concentrations and incubation times for individual applications.

Recommended Assay Procedure

- 1. Prepare coating solution by diluting the coating antibody. See "coating antibody" section for the recommended coating antibody
- 2. Coat plates with 100 μ L per well of the coating solution. Cover plates and incubate overnight (12-18 hr.) at 4°C.
- 3. Aspirate wells and wash 1 time with $> 400~\mu L$ of Wash Buffer per well. Following wash, invert and tap on absorbent paper to remove excess liquid.
- 4. Block plate with 300 μ L per well of Assay Buffer for 1 hour at room temperature.
- 5. Aspirate, invert, and tap on absorbent paper to remove excess liquid.
- 6. Prepare standards and sample dilutions in Assay Buffer (or in a diluent that most closely matches the matrix of your sample). For recommended dilutions and storage of the standard, see "standard" section.
- 7. Pipette 100 μL of standards (in duplicate) and samples into designated wells.
- Immediately following step 7, add 50 µL of the working detection antibody into each well. For recommended dilutions, see "detection antibody" section.. Incubate for 2 hours at room temperature with continual shaking (700 rpm).
- 9. Aspirate and wash 5 times using the method in step 3.
- 10. Add 100 μ L of the working streptavidin-HRP solution into each well. For recommended dilutions, see "streptavidin-HRP conjugate" section. Incubate for 30 minutes at room temperature with continual shaking (700 rpm).
- 11. Aspirate and wash 5 times using the method in step 3.
- 12. Add 100 µL of the TMB substrate to each well. Incubate plate for 30 minutes at room temperature with continual shaking (700 rpm).
- 13. Add 100 μL of Stop Solution to each well.
- 14. Measure absorbance at 450 nm (reference absorbance: 650 nm) within 30 minutes of adding Stop Solution. Calculate results using a loglog or 4-parameter curve fit.

Additional Materials Required

- 96 well NUNC MaxiSorp microplates; NUNC Cat. # 434797.
- Pipettes, shaker and timer.
- Microplate reader with a detector that can measure absorbance at 450 nm.
- 1 L graduated cylinder; plate washer or wash bottle.
- Polypropylene tubes for standards and sample dilutions, if needed.

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