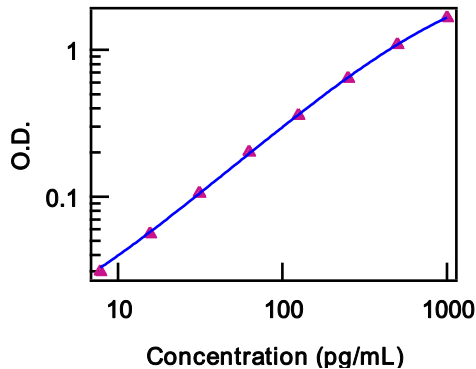


## Mouse TNF alpha ELISA Ready-SET-Go!<sup>®</sup>

**Catalog Number:** 88-7324

**Also known as:** Tumor Necrosis Factor alpha

**RUO: For Research Use Only. Not for use in diagnostic procedures.**



Standard curve of Mouse TNF alpha ELISA Ready-SET-Go!<sup>®</sup>

### Product Information

**Contents:** Mouse TNF alpha ELISA Ready-SET-Go!<sup>®</sup>

**REF** **Catalog Number:** 88-7324

**Sensitivity:** 8 pg/mL

**Standard Curve Range:** 8 - 1000 pg/mL



**Temperature Limitation:** Store at 2-8°C except standard which should be stored at less than or equal to -70°C.



**Batch Code:** Refer to vial



**Use By:** Refer to vial



**Caution, contains Azide**

### Description

This Mouse TNF alpha ELISA Ready-SET-Go! reagent set (with or without high-affinity binding microwell plates) contains the necessary reagents, buffers and diluents for performing quantitative enzyme linked immunosorbent assays (ELISA). This ELISA reagent set is specifically engineered for accurate and precise measurement of mouse TNF alpha protein levels from samples including serum, plasma, tissue homogenate, lavage fluid and supernatants from cell cultures. Recognition of mouse TNF alpha by this assay is not interfered with by 1000x excess soluble TNF alpha receptor type I or II.

### Components

**Capture Antibody.** Pre-titrated, purified antibody

**Detection Antibody.** Pre-titrated, biotin-conjugated antibody

**Standard.** Recombinant cytokine for generating standard curve and calibrating samples

**ELISA/ELISPOT Coating Buffer Powder.** This Ready-Set-Go! ELISA Set may contain ELISA/ELISPOT Coating Buffer Powder (Reconstitute to 1L with dH2O and filter (0.22 µM)) or 10X PBS ELISA Coating Buffer (Dilute 1 part 10X Buffer into 9 parts dH2O).

**Assay Diluent.** 5X concentrated

**Detection enzyme.** Pre-titrated Avidin-HRP

**Substrate Solution.** Tetramethylbenzidine (TMB) Substrate Solution

**Certificate of Analysis.** Lot-specific instructions for dilution of antibodies and standards

**96 Well Plate.** Corning Costar 9018 (included with product Cat. #'s ending in suffixes -22, -44, -76, -86)

### References

Cho HJ, Seon MR, Lee YM, Kim J, Kim JK, Kim SG, Park JH. 3,3'-Diindolylmethane suppresses the inflammatory response to lipopolysaccharide in murine macrophages. *J Nutr.* 2008 Jan;138(1):17-23. (RSG ELISA kit, **TC supernatant**, PubMed)

Nakano T, Araki K, Nakatani H, Kobayashi M, Sugimoto T, Furuya Y, Matsuoka T, Jin T, Hanazaki K. Effects of geldanamycin and thalidomide on the Th1/Th2 cytokine balance in mice subjected to operative trauma. *Surgery.*

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## Mouse TNF alpha ELISA Ready-SET-Go!®

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2007 Apr;141(4):490-500. (RSG ELISA kit, **serum**, PubMed)

Chen L, Lin SX, Agha-Majzoub R, Overbergh L, Mathieu C, Chan LS. CCL27 is a critical factor for the development of atopic dermatitis in the keratin-14 IL-4 transgenic mouse model. *Int Immunol.* 2006 Aug;18(8):1233-42. (RSG ELISA kit, **tissue homogenate**, PubMed)

Yang D, Zhang Y, Nguyen HG, Koupenova M, Chauhan AK, Makitalo M, Jones MR, St Hilaire C, Seldin DC, Toselli P, Lamperti E, Schreiber BM, Gavras H, Wagner DD, Ravid K. The A2B adenosine receptor protects against inflammation and excessive vascular adhesion. *J Clin Invest.* 2006 Jul;116(7):1913-23. (RSG ELISA kit, **plasma**, PubMed)

Mallen-St Clair J, Pham CT, Villalta SA, Caughey GH, Wolters PJ. Mast cell dipeptidyl peptidase I mediates survival from sepsis. *J Clin Invest.* 2004 Feb;113(4):628-34. (RSG ELISA kit, **lavage fluid**, PubMed)

### Related Products

88-7013 Mouse IL-1 beta ELISA Ready-SET-Go!®

88-7121 Mouse IL-12 p70 ELISA Ready-SET-Go!®

88-7234 Mouse IL-23 ELISA Ready-SET-Go!® (Discontinued: Please see 88-7230 (2nd generation assay))

88-8314 Mouse IFNγ 'Femto-HS' High Sensitivity ELISA Ready-Set-Go!

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## Enzyme Linked Immunosorbent Assay (ELISA)

### Research Use Only

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#### Protocol: ELISA Ready-Set-Go!

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The following protocol is a general guideline for the Ready-SET-Go! Sets

##### Materials Provided

- Please refer to the Certificate of Analysis (C of A) for components

##### Other Materials Needed

- Buffers\*
  - Wash Buffer: 1x PBS, 0.05% Tween-20 (or eBioscience ELISA Wash Buffer Powder, Cat. No. 00-0400)
  - Stop Solution: 1M H<sub>3</sub>PO<sub>4</sub> (recommended) or 2N H<sub>2</sub>SO<sub>4</sub>
- Pipettes
- Refrigerator & frost-free -20°C freezer
- 96-well plate (Corning Costar 9018 or NUNC Maxisorp®)  
**NOTE: The use of ELISA plates which are not high affinity protein binding plates will result in suboptimal performance, e.g., low signal or inconsistent data. Do not use tissue culture plates or low protein absorption plates. Use only the Corning Costar 9018 or NUNC Maxisorp (Cat. No. 44-2404) 96-well plates**
- 96-well ELISA plate reader (microplate spectrophotometer)
- ELISA plate washer (highly recommended)

**NOTE: To ensure optimal results from this ELISA Ready-SET-Go! Set, please only use the components included in the set. Exchanging of components is not recommended as a change in signal may occur.**

##### Time Requirements

- 1 overnight incubation
- 4½-hour incubations
- 1 hour washing and analyzing samples

##### Experimental Procedure

1. Coat Corning Costar 9018 (or Nunc Maxisorp®) ELISA plate with 100 µL/well of capture antibody in Coating Buffer (dilute as noted on C of A, which is included with the reagent set). Seal the plate and incubate overnight at 4°C.
2. Aspirate wells and wash 3 times with >250 µL/well Wash Buffer\*. Allowing time for soaking (~1 minute) during each wash step increases the effectiveness of the washes. Blot plate on absorbent paper to remove any residual buffer.
3. Dilute 1 part 5X concentrated Assay Diluent with 4 parts DI water.\* Block wells with 200 µL/well of 1X Assay Diluent. Incubate at room temperature for 1 hour.
4. Optional: Aspirate and wash at least once with Wash Buffer.
5. Using 1X Assay Diluent\*, dilute standards as noted on the C of A to prepare the top concentration of the standard. Add 100 µL/well of top standard concentration to the appropriate

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## Enzyme Linked Immunosorbent Assay (ELISA)

### Research Use Only

wells. Perform 2-fold serial dilutions of the top standards to make the standard curve for a total of 8 points. Add 100 µL/well of your samples to the appropriate wells. Seal the plate and incubate at room temperature for 2 hours (or overnight at 4°C for maximal sensitivity).

6. Aspirate/wash as in step 2. Repeat for a total of 3-5 washes\*\*.
7. Add 100 µL/well of detection antibody diluted in 1X Assay Diluent\* (dilute as noted on C of A). Seal the plate and incubate at room temperature for 1 hour.
8. Aspirate/wash as in step 2. Repeat for a total of 3-5 washes\*\*.
9. Add 100 µL/well of Avidin-HRP\* diluted in 1X Assay Diluent (dilute as noted on C of A). Seal the plate and incubate at room temperature for 30 minutes.
10. Aspirate and wash as in step 2. In this wash step, soak wells in Wash Buffer\* for 1 to 2 minutes prior to aspiration. Repeat for a total of 5-7 washes\*\*.
11. Add 100 µL/well of Substrate Solution to each well. Incubate plate at room temperature for 15 minutes.
12. Add 50 µL of Stop Solution to each well.
13. Read plate at 450 nm. If wavelength subtraction is available, subtract the values of 570 nm from those of 450 nm and analyze data.

#### **NOTES:**

**\* Be certain that no sodium azide is present in the solutions used in this assay, as this inhibits HRP enzyme activity.**

**\*\*The number of washes in the protocol was adapted to an automatic plate washer. This can be decreased when using other methods but should be tested empirically. Allowing time for soaking (~ 1 minute) during each wash step increases the effectiveness of the washes.**

## Enzyme Linked Immunosorbent Assay (ELISA)

### Research Use Only

#### Quick Guide: Standard Calibration

The following table indicates the protein standard contained in the Ready-SET-Go! is calibrated against NIBSC standards.

Table of Standard Calibration				
Cytokine	ng of eB standard	ng of NIBSC standard	U of NIBSC standard	NIBSC Lot #
hIL-2	1	1.1	14.6	86/564
hIL-4	1	2.2	22	88/656
hIL-5	1	2.2	22	90/586
hIL-6	1	1.7	170	89/548
hIL-8	1	1.8	180	89/520
hIL-10	1	0.8	4	93/722
hIL-12	1	0.8	8	95/544
hIL-17A	1	0.9	9000	01/420
hIFN-g	1	1.1	22	87/586
hTNF-a	1	0.9	36	87/650
mIL-2	1	3.1	310	93/566
mIL-4	1	3	30	91/656
mIL-6	1	8.5	850	93/730
mIFN-g*	1		4.5	Gg02-901-533
mTNF-a	1	1.7	340	88/532

\* Mouse IFN-g is calibrated using NIH standard (Lot Gg02-901-533) and is measured in Units (U)

#### ELISA Troubleshooting Guide

Problem	Possibility	Solution
A. High background	1. Improper and inefficient washing	1. Improve efficiency of washing. Fill plates completely, soak for 1 minute per wash, as directed
	2. Cross contamination from other specimens or positive control	2. Repeat ELISA being careful when washing and pipetting
	3. Contaminated substrate	3. Substrate should be colorless. Replace
	4. Incorrect dilutions, e.g., conjugate concentration was too high	4. Repeat using correct dilutions
B. No signal	1. Improper, low protein binding capacity plates were used	1. Repeat ELISA using recommended high binding capacity plates
	2. Wrong substrate was used	2. Repeat ELISA using the correct substrate
	3. Enzyme inhibitor present in buffers; e.g., sodium azide in the washing buffer and Assay Diluent inhibits peroxidase activity	3. Repeat ELISA making no enzyme inhibitor is present in any buffers.

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## Enzyme Linked Immunosorbent Assay (ELISA)

### Research Use Only

	4. Coated capture antibody in Assay Diluent rather than Coating Buffer	4. Repeat ELISA using Coating Buffer contained in the set as the diluent for the capture antibody.
C. Very weak signal	1. Improper and inefficient washing	1. Make sure washing procedure is done correctly, with a soak time.
	2. Incorrect dilutions of standard	2. Follow recommendations of standard preparation exactly as written on the C of A
	3. Insufficient incubation time	3. Repeat ELISA following the protocol carefully for each step
	4. Incorrect storage of reagents	4. Store reagents at the correct temperature as indicated on the Technical Data Sheet. Freezing certain components will severely impact results. Do not re-use the standards.
	5. Wrong filter in ELISA reader was used	5. Use the correct wavelength setting
	6. Wrong plate used	6. Use the recommended Corning Costar 9018 or NUNC Maxisorp flat bottom 96 well plates
D. Variation amongst replicates	1. Improper and inefficient washing	1. Make sure washing procedure is done correctly; see C of A. Edge effects can be avoided by moving samples and standards in from the edge of the plate.
	2. Poor mixing of samples	2. Mix samples and reagents gently and equilibrate to proper temperature
	3. Plates not clean	3. Plates should be wiped on bottom before measuring absorbance
	4. Reagents have expired	4. Order a new Ready-Set-Go ELISA.