

For the quantitative determination of mouse CD8 concentrations in cell culture supernates, serum, and plasma.

INTRODUCTION

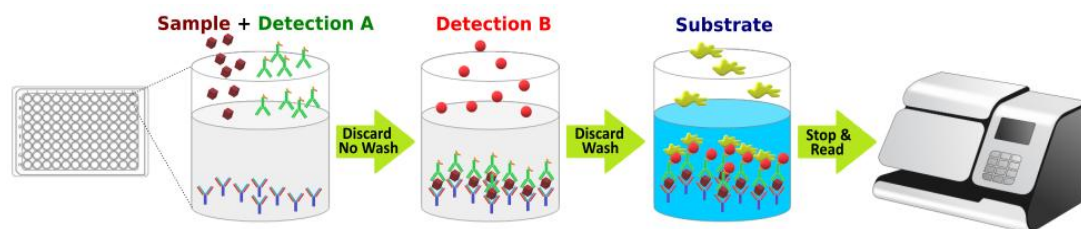
CD8 is a protein that expressed on cytotoxic T-lymphocytes (T-cells), thymocytes, peripheral blood lymphocytes, and a population of natural killer (NK) cells. CD8⁺ T cells recognize antigens presented by MHC class I molecules on the surface of target cells, leading to the activation and execution of their cytotoxic functions. CD8⁺ T cells are involved in various disease processes, including chronic infections multiple sclerosis, rheumatoid arthritis, ulcerative colitis, type 1 diabetes mellitus, cancer, and autoimmune diseases. Their activation and function are critical for the development of effective vaccines and immunotherapies.

Tribio Fast Mouse CD8 ELISA is designed to quantitatively detect Mouse CD8 levels in a variety of tissues, including skin, muscle, neural tissue, serum, and other biological samples. **The key feature of this kit is the use of novel proprietary technology that integrates sample processing and detection into a simplified one-step procedure, instead of the complex multi-step workflows used in traditional ELISA methods. This significantly simplifies the assay, making it easy to perform, highly accurate, and time efficient. The entire measurement can be completed within 3 hours, rather than the conventional 4–5 hours (Fig. 1).** The detection range spans from 3 to 200 ng/mL. Mouse CD8 sample levels show linear parallelism with the standard curve generated using kit standards, indicating that this assay can be used for the relative quantification of native Mouse CD8 protein in biological samples.

PRINCIPLE OF THE ASSAY

This assay employs our novel proprietary sandwich enzyme immunoassay techniques (Fig. 1). A monoclonal antibody specific to mouse CD8 is pre-coated onto a microplate. Standards or samples and a biotin-conjugated detection antibody are pipetted into the wells and concurrently incubated to form a sandwich complex in one step. Simply aspirate each well without washing and directly add Streptavidin-HRP into the complex. Following a wash, an **ultra-sensitive TMB substrate solution** is added to the wells for color development. The color intensity is proportional to the amount of CD8 bound in the initial step. The intensity of the color is measured by plate reading at 450 nm.

Fig. 1: Assay Principle:



KIT CONTENT AND STORAGE CONDITIONS

PART	PART#	DESCRIPTION	STORAGE OF OPENED/ RECONSTITUTED
Mouse CD8 Microplate	TBS3004A	96 well polystyrene microplate (12 strips of 8 wells) coated with a polyclonal antibody specific for mouse CD8.	Return unused wells to the foil pouch. Reseal along the entire edge of the zip-seal. May be stored for up to 1 month at 2-8 °C.
Mouse CD8 Standard	TBS3004B	30 µL of Recombinant mouse CD8 protein (10 µg/mL).	Aliquot and store at -20 °C for up to 1 month in a manual defrost the freezer. Avoid repeated freeze-thaw cycles.
Detection A	TBS3004C	220 µL of Biotin-Mouse CD8 antibody (10x).	May be stored for up to 3 months at 2-8 °C.*
Detection B	TBS3004D	300 µL of Streptavidin-HRP (50x).	
Assay Diluent	TBS3004E	15 mL of a buffered protein base with preservatives.	
Wash Buffer	TBS3000W	15 mL of concentrated solution (10x).	
TMB Substrate	TBS3000T	12 mL of ultra-sensitive TMB substrate.	
Stop Solution	TBS3000S	6 mL of 2 N sulfuric acid.	

Store the unopened kit at 2-8 °C. Do not use past kit expiration date.
The kit contains sufficient materials to run an ELISA on one 96 well plate.

PRECAUTIONS

Wear protective gloves, clothing, eye, and face protection. Wash hands thoroughly after handling.

REAGENT PREPARATION

Bring all reagents to room temperature before use.

Wash Buffer: Add 12 mL of Wash Buffer Concentrate (10x) to 108 mL of deionized distilled water to prepare 120 mL of Wash Buffer (*If crystals have formed in the concentrate, warm to room temperature and mix gently until the crystals have completely dissolved*).

Detection A working solution preparation: Add 210 µL of **Detection A Stock** (Biotin-mouse CD8 antibody) to 1890 µL Assay Diluent to prepare Detection A working solution. Add 20 µL to each well.

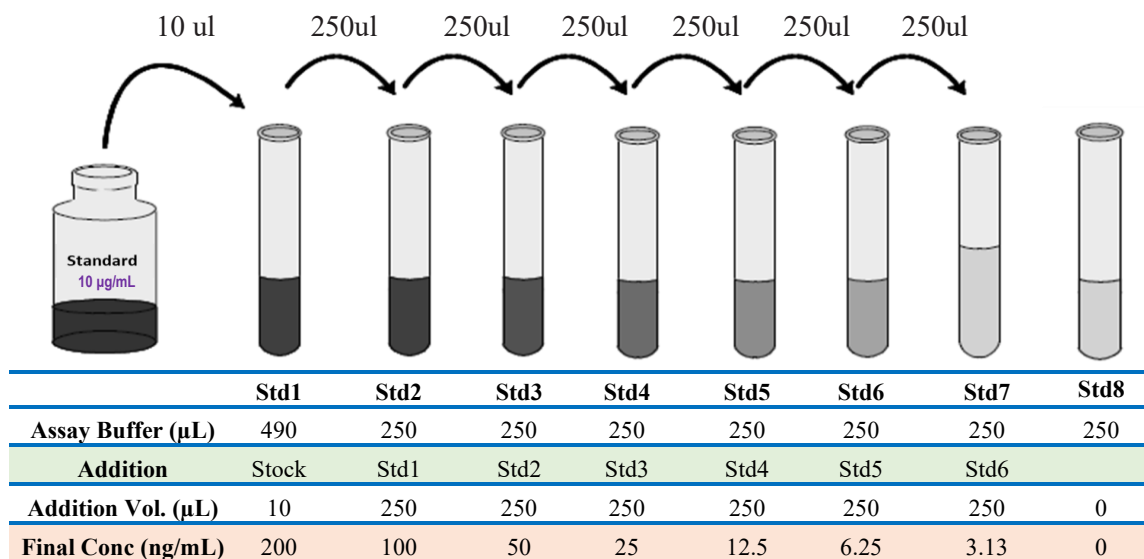
Detection B working solution preparation: Add 240 µL of **Detection B** streptavidin-HRP to 12 mL Assay Diluent to prepare Detection B working solution. Add 100 µL to each well.

Mouse CD8 Standard Preparation:

Label test tubes as #1 through #8. Pipet 490 µL of 1x Assay Diluent into tube #1, and 250 µL into tubes #2 to #7 as diagram below (Fig.2).

1. Add 10 µL of the Mouse CD8 Standard stock solution (10 µg/mL) by dilution of 50X to tube #1 (200 ng/mL), and mix.
2. Make 2x serial dilutions of the standard using the 200 ng/mL standard solution (tube #1) from tube #2 through #7 with sequential transfer of 250 µL to the next concentration. Mix each tube thoroughly before the next transfer. The standard concentration in tubes 1 through 7 will be 200, 100, 50, 25, 12.5, 6.25, and 3.13 ng/mL. Tube# 8 is Standard 0.

Fig.2 Diagram for Mouse CD8 standard preparation



ASSAY PROCEDURE

Bring all reagents and samples to room temperature before use. It is recommended that all standards, controls, and samples be assayed in duplicate.

1. Add 80 µL of standard, sample, or control per well.
2. Add 20 µL of **Detection A** to the above standard and sample of each well, thoroughly mix. Cover with the adhesive sealer. Incubate at **RT for 2 hours**.
3. Aspirate each well (no wash). Invert the plate and blot it against clean paper towels.
4. Add 100 µL of **Detection B** to each well. Incubate at **RT for 1 hour**.

5. Aspirate each well, and wash 3 times by filling each well with 200 μ L Wash Buffer (*Complete removal of liquid at each step is essential to good performance*). After the last wash, remove any remaining Wash Buffer by aspirating or decanting. Invert the plate and blot it against clean paper towels.
6. Add 100 μ L of **TMB Substrate** to each well. Incubate **at RT for 10-20 min** (*Protect from light*). The color becomes blue.
7. Add 50 μ L of **Stop Solution** to each well. The color in the well should change from blue to yellow (gently tap the plate to ensure thorough mixing).
8. Determine the optical density of each well within 20 minutes, using a microplate reader at 450 nm. If wavelength correction is available, set to 540 nm or 570 nm. If wavelength correction is not available, subtract readings at 540 nm or 570 nm from the readings at 450 nm. This subtraction will correct for optical imperfections in the plate. Readings made directly at 450 nm without correction may be higher and less accurate.

CALCULATION OF RESULTS

Average the duplicate readings for each standard, control, and sample subtract the average zero standard optical density (O.D.).

Create a standard curve using computer software capable of generating a four-parameter logistic (4-PL) curve-fit. As an alternative, construct a standard curve by plotting the mean absorbance for each standard on the Y-axis against the concentration on the X-axis and draw a best fit curve through the points on the graph. The data may be linearized by plotting the log of the mouse CD8 concentrations versus the log of the O.D. and the best fit line can be determined by regression analysis. This procedure will produce an adequate but less precise fit of the data.

TYPICAL DATA

This standard curve ($R^2=0.9995$) is provided for demonstration only. A standard curve should be generated for each set of samples assayed. Fig. 3 is an example of typical Data.

SENSITIVITY

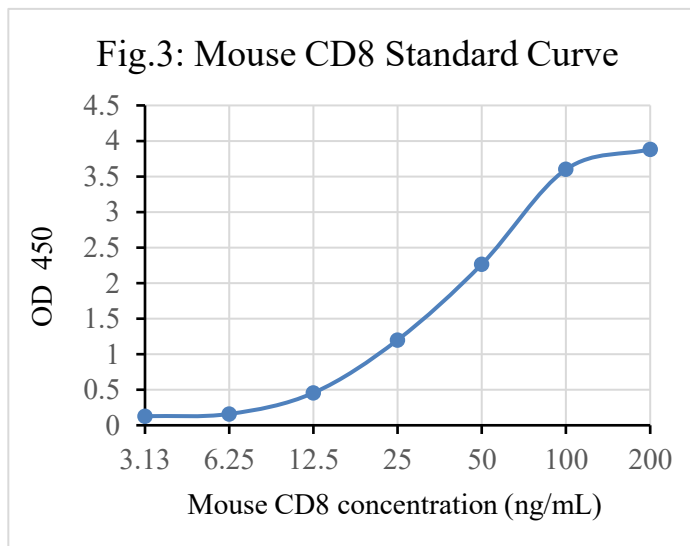
The minimum detectable dose (MDD) of mouse CD8 is typically 10 ng/mL.
The Intra-assay CV is 3.79% the Inter-assay CV is <10%.

SPECIFICITY

This assay recognizes natural and recombinant mouse CD8.

RELATIVE PRODUCTS

- Human IL-1 β ELISA (TBS3219)
- Human IL-2 ELISA (TBS3220)
- Human IL-4 ELISA (TBS3221)
- Human IL-6 ELISA (TBS3223)
- Human IL-7 ELISA (TBS3224)
- Human IL-8 ELISA (TBS3225)
- Human IL-10 ELISA (TBS3226)
- Human IL-13 ELISA (TBS3227)
- Human IL-17 ELISA (TBS3228)
- Human IL-22 ELISA (TBS3229)
- Human IFN-gamma ELISA (TBS3230)
- Human TGF- β 1 ELISA (TBS3232)
- Human GM-CSF ELISA (TBS3233)
- Human MIP-1 α ELISA (TBS3234)



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