

For the quantitative determination concentrations of mouse CCL3/MIP-1 α in cell culture supernatants, serum and plasma.

INTRODUCTION

Macrophage Inflammatory Protein 1 alpha (CCL3/MIP-1 α) is a chemokine belonging to the CC subfamily and is mainly involved in cell adhesion and migration. It plays a crucial role in initiating and maintaining the inflammatory response by recruiting leukocytes to the site of injury or infection. Elevated levels of CCL3/MIP-1 α have been associated with a variety of inflammatory and autoimmune disorders, including rheumatoid arthritis, desiccation syndrome, tuberculosis, and cardiovascular disease. CCL3/MIP-1 α serves as a biomarker for the diagnosis and assessment of the progression of a variety of diseases. Inhibition of CCL3/MIP-1 α is a potential therapeutic strategy.

Tribioscience's Fast Mouse CCL3/MIP-1 α ELISA is designed to quantitatively detect mouse CCL3/MIP-1 α levels in different tissues including skin, muscle, neural, serum, and other biological samples. The main feature is that the kit uses our novel proprietary approaches to combine samples and detections into a one-step instead of the complicated traditional methods. It makes the assay simple, easy, accurate, and fast. The measurement can be finished in 2 hours, with no need for 4-5 hours (Fig. 1). The detection range is from 7 to 500 pg/mL. The levels of mouse CCL3/MIP-1 α samples are parallel to the standard curves obtained using the kit standards linearly. These results indicate that this kit can be used to determine relative mass values for natural mouse CCL3/MIP-1 α protein.

PRINCIPLE OF THE ASSAY

This assay employs the quantitative sandwich enzyme immunoassay technique (See Fig. 1). A monoclonal antibody specific for mouse CCL3/MIP-1 α was pre-coated onto a microplate. Standards and samples are pipetted into the wells, and then, incubated with HRP-conjugated detection antibody specific for mouse CCL3/MIP-1 α . Following a wash to remove any unbound antibody and samples, an **ultra-sensitive TMB substrate solution** is added to the wells for color development. The color intensity is in proportion to the amount of CCL3/MIP-1 α bound in the initial step. The intensity of the color is measured by plate read at 450 nm.

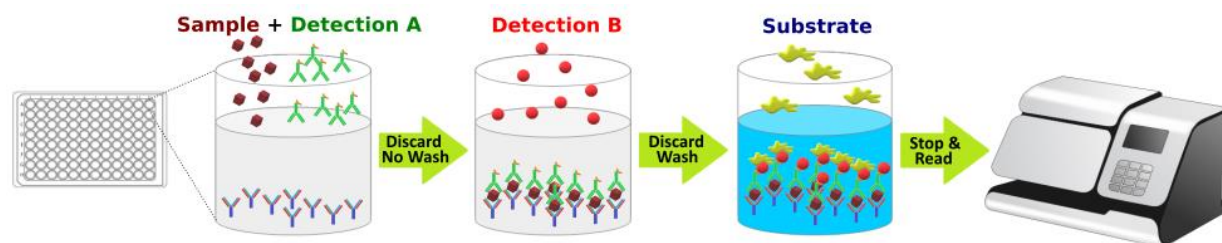


Fig.1: Simple ELISA procedure

KIT CONTENT AND STORAGE CONDITIONS

PART	PART#	DESCRIPTION	STORAGE OF OPENED/ RECONSTITUTED
Mouse CCL3/MIP-1 α Microplate	TBS3082A	96 well polystyrene microplate (12 strips of 8 wells) coated with a polyclonal antibody specific for mouse CCL3/MIP-1 α .	Return unused wells to the foil pouch. Reseal along entire edge of the zip-seal. May be stored for up to 1 month at 2-8 °C.
Mouse CCL3/MIP-1 α Standard	TBS3082B	30 μ L of Recombinant mouse CCL3/MIP-1 α protein (25 ng/mL).	Aliquot and store at -20 °C for up to 1 month in a manual defrost freezer. Avoid repeated freeze-thaw cycles.
Detection A	TBS3082C	2.2 mL of Biotin-mouse CCL3/MIP-1 α antibody.	May be stored for up to 3 months at 2-8 °C.*
Detection B	TBS3082D	300 μ L of Streptavidin-HRP.	
Assay Diluent	TBS3082E	25 mL of a buffered protein base with preservatives.	
Wash Buffer	TBS3000W	12 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.*).

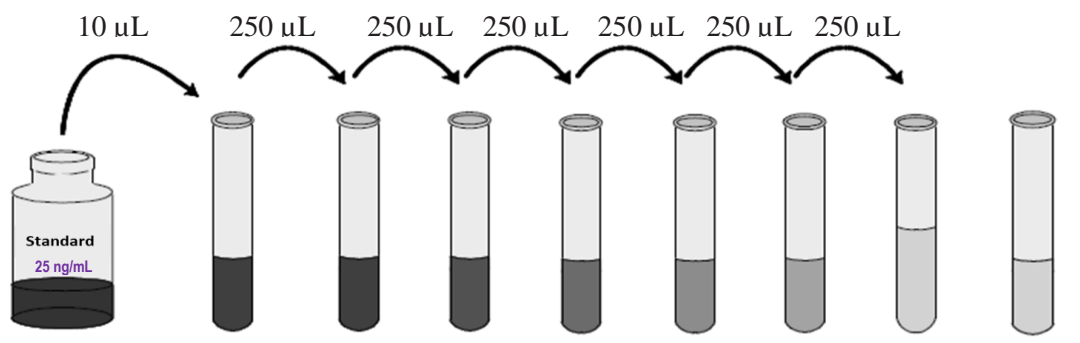
Detection B working solution preparation: Add 240 μ L of **Detection B** streptavidin-HRP to 12 mL Assay Diluent (TBS3050E) to prepare Detection B working solution.

Mouse CCL3/MIP-1 α Standard Preparation:

Label test tubes as #1 through #8. Pipet 490 μ L of 1x Assay Diluent into tube #1, and 250 μ L into tubes #2 to #8 as **Fig.2 diagram below.**

1. Add 10 μ L of the Mouse CCL3/MIP-1 α Standard stock solution (25 ng/mL) by dilution of 50x to tube #1 and mix.
2. Make 2x serial dilutions using the of 500 pg/mL (tube #1) standard solution from tube #2 through #7 with sequential transfer of 250 μ L to the next concentration. Mix each tube thoroughly before the next transfer. The standard concentration in tube 1 through 7 will be 500, 250, 125, 62.5, 31.25, 15.63, and 7.81 pg/mL. Tube# 8 is blank (0 pg/mL)

Fig.2 Diagram for Mouse CCL3/MIP-1 α standard preparation



	Std1	Std2	Std3	Std4	Std5	Std6	Std7	Std8
Assay Buffer (μL)	490	250	250	250	250	250	250	250
Addition	Stock	Std1	Std2	Std3	Std4	Std5	Std6	
Addition Vol. (μL)	10	250	250	250	250	250	250	0
Final Conc (pg/mL)	500	250	125	62.5	31.25	15.63	7.81	0

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 μ L of standard, sample, or control per well.
2. Add 20 μ L 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 with shaking.**
3. Aspirate each well (no wash). Invert the plate and blot it against clean paper towels.
4. Add 100 μ L of **Detection B working solution** to each well. Incubate at **RT for 1 hour with shaking.**
5. Aspirate each well, and wash for 3 times by filling each well with 300 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 minutes with shaking** (*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 CCL3/MIP-1 α 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 ($R^2=0.9998$) 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 (MOD) of mouse CCL3/MIP-1 α is typically 13 pg/ml.

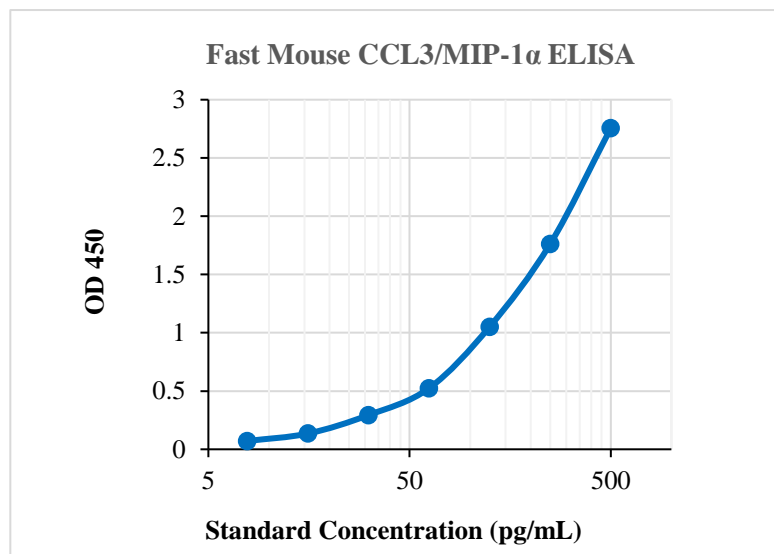
The Intra-assay CV is 3.79% the Inter-assay CV is <10%.

SPECIFICITY

This assay recognizes natural and recombinant mouse CCL3/MIP-1 α .

RELATIVE PRODUCTS

- TBS3030 Fast Mouse IL-1 β ELISA
- TBS3031 Fast Mouse IL-2 ELISA
- TBS3032 Fast Mouse IL-4 ELISA
- TBS3040 Fast Mouse IL-6 ELISA
- TBS3044 Fast Mouse IL-10 ELISA
- TBS3047 Fast Mouse IL-12 p70 ELISA
- TBS3049 Fast Mouse IL-13 ELISA
- TBS3060 Fast Mouse KC ELISA
- TBS3070 Fast Mouse NGF ELISA
- TBS3079 Fast Mouse GM-CSF ELISA
- TBS3080 Fast Mouse G-CSF ELISA
- TBS3084 Fast Mouse IFN- γ ELISA
- TBS3085 Fast Mouse TGF ELISA
- TBS3086 Fast Mouse MCPT-1 ELISA
- TBS3090 Fast Mouse IL-17AF ELISA
- TBS3091 Fast Mouse IL-19 ELISA
- TBS3092 Fast Mouse IL-21 ELISA
- TBS3093 Fast Mouse IL-22 ELISA
- TBS3094 Fast Mouse IL-23 ELISA
- TBS3095 Fast Mouse IL-27 ELISA
- TBS3096 Fast Mouse IL-28B ELISA
- TBS3097 Fast Mouse IL-33 ELISA



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