

## DATA SHEET

Versions: 01

Revision date: 25/11/2023

### 1. Identification

Product name: ELISA kits for measurement of Mouse TGF- $\beta$ 

Reactions: 96,48 rxns

Cat. No.: PRA-MTGF-96

### 2. Description

Transforming Growth Factor-beta (TGF- $\beta$ ), classified as an anti-inflammatory cytokine, is produced by a diverse array of immune cells, including regulatory T lymphocytes and macrophages. Renowned for its anti-inflammatory properties, TGF- $\beta$  interacts with specific receptors on immune cells. Despite its anti-inflammatory nature, this cytokine assumes a crucial role in the growth and maturation of Th17 lymphocytes, aligning it with an inflammatory function when coupled with IL-2 and IL-6 cytokines. TGF- $\beta$  exhibits potent anti-inflammatory characteristics. Despite its anti-inflammatory nature, TGF- $\beta$  plays a pivotal role in the growth and maturation of Th17 lymphocytes, contributing to its dual role in inflammation. TGF- $\beta$  is instrumental in establishing homeostasis post-microbial infections and preventing the development of autoimmune diseases. As a primarily anti-inflammatory indicator, TGF- $\beta$  finds extensive use in laboratory studies. Its application aids in investigating disease conditions and evaluating the inflammatory or anti-inflammatory effects of pharmaceutical agents. The current kit, meticulously designed and produced, utilizes both mouse and human anti-TGF- $\beta$  monoclonal antibodies. It's important to note that this kit is specifically intended for measuring TGF- $\beta$  levels in mice and humans and may not be suitable for similar animal cases. This comprehensive overview underscores the complex and dual nature of TGF- $\beta$ , highlighting its significance in both anti-inflammatory responses and inflammatory processes, making it a key focus in experimental investigations into various disease states.



## DATA SHEET

Versions: 01

Revision date: 25/11/2023

### 3. Kit Contents

Component	Cat. no	Quantity
Human anti-MTGF- $\beta$ pre-coated plate	PRA-MTGFP	96 vials
Standards	PRA-MTGFSN1-4	200 $\mu$ l
HRP-Avidin buffer	PRA-HA	5 ml
HRP	PRA-HAA	540 $\mu$ l
Substrate	PRA-SU	5 ml
Stopping	PRA-ST	7 ml
10X washing buffer	PRA-WB	40 ml
Detection Ab	PRA-MTGFD	5 ml

### 4. Storage specifications

All components of the ELISA kits can be stored at 4°C temperature.

### 5. Applications

Detection of both inflammatory and anti-inflammatory factors through ELISA.

### 6. Assay Procedure

How to Prepare Solutions:

1. Washing Buffer:

- To prepare the washing solution, dilute it with distilled water at a 1:10 ratio.

2. HRP-Avidin:

- Spin the HRP vial using a microfuge device, then add all its contents to the HRP-Avidin buffer vial.

- For quantities less than 48 assays, mix 416  $\mu$ L of HRP-Avidin, 41  $\mu$ L of HRP per 8-well row.



## DATA SHEET

Versions: 01

Revision date: 25/11/2023

### Working with the Kit:

#### 1. Plate Preparation:

- Remove the plate from its package and let it reach room temperature in a dry environment.

- Add 50  $\mu$ L of standards #4 to #1 in the first to fourth wells.

#### 2. Sample Incubation:

- Add 50  $\mu$ l of the desired sample to the remaining wells.

- Incubate for 60 minutes on a 200 RPM shaker at room temperature.

#### 3. Plate Washing:

- After proper incubation, wash the plates three times using the washing solution (after adding the washing solution, incubate the plates for approximately 1 minute at room temperature and then drain).

#### 4. Conjugated Antibody Addition:

- Add 50  $\mu$ l of conjugated antibody (Detection ab) to all wells.

- Incubate for 60 minutes on a 200 RPM shaker at room temperature.

#### 5. Plate Washing (Again):

- After proper incubation, wash the plates three times using washing solution.

#### 6. HRP-Avidin Addition:

- Add 50  $\mu$ l of HRP-Avidin solution to all wells.

- Incubate for 30 minutes on a shaker (at least at RPM 200).

#### 7. Plate Washing (Again):

- After proper incubation, wash the plates five times using washing solution.

#### 8. Substrate Addition:

- Add 50  $\mu$ l of substrate to all wells and incubate for 15 minutes. Note that 15 minutes is enough for incubation, but if the amount of color produced is low, the time can be increased to 20 minutes.

#### 9. Stopping Reaction:

- Add 25  $\mu$ l of the stopping solution to all the wells.

#### 10. Measurement:

- Measure the absorbance of the samples in an ELISA reader at a wavelength of 450 nm.



## DATA SHEET

Versions: 01

Revision date: 25/11/2023

### 7. Safety

- The solutions used in the kit have oxidizing and acidic properties.
- Avoid direct contact with skin and eyes.
- In case of contact with the mentioned tissues, wash with plenty of water and go to the nearest medical center.

### 8. Quality Certifications

### 9. Further information

- This product is developed, designed, and sold exclusively only for research purposes use.
- The product was not tested for use in diagnostics or for drug development.
- It is not suitable for administration to humans or animals.

### 10. Other Kits

Other ELISA kits:

Human:

IL-1 $\beta$ , IL-2, IL-4, IL-6, IL-8, IL-10, IL-12, IL-13, IL-18, IL-23, IL-29, IL-17A, TGF- $\beta$ , VEGF, TNF- $\alpha$ , IFN- $\gamma$ , CCL2 (MCP-1), CCL3 (MIP-1-alpha), CXCL10 (IP-10), CXCL12 (SDF-1), CCL21

Mouse:

IL-1 $\beta$ , IL-2, IL-4, IL-6, IL-10, IL-13, IL-33, IL-18, TNF- $\alpha$ , TGF- $\beta$ , CCL3, IFN- $\gamma$ , Total IgG, Total IgE

Rat:

TNF- $\alpha$ , IL-1 $\beta$ , IL-6, IL-10, IL-17A



## DATA SHEET

Versions: 01

Revision date: 25/11/2023

### NOTE

All products have been manufactured by Karmania Pars Gene Company in the Iran.

