

Mouse CXCL10/IP-10 ELISA Kit

Catalog No: CKM049

Quantity: 1 x 96 tests

Introduction

The Mouse IP-10 ELISA (Enzyme-Linked Immunosorbent Assay) kit is an *in vitro* enzyme-linked immunosorbent assay for the quantitative measurement of mouse IP-10 in serum, plasma, cell culture supernatants and urine. This assay employs an antibody specific for mouse IP-10 coated on a 96-well plate. Standards and samples are pipetted into the wells and IP-10 present in a sample is bound to the wells by the immobilized antibody. The wells are washed and biotinylated anti-mouse IP-10 antibody is added. After washing away unbound biotinylated antibody, HRP-conjugated streptavidin is pipetted to the wells. The wells are again washed, a TMB substrate solution is added to the wells and color develops in proportion to the amount of IP-10 bound. The Stop Solution changes the color from blue to yellow, and the intensity of the color is measured at 450 nm.

Reagents and materials supplied with the kit:

Items	Quantity
A. IP-10 Microplate coated with anti-mouse IP-10	96 wells (12 strips x 8 wells)
B. Wash Buffer Concentrate (20x)	25 ml
C. Recombinant mouse IP-10 Standards	2 vials
D. Assay Diluent A for Standard/Sample (serum/plasma) diluent*	30 ml
E. Assay Diluent B (5x) for Standard/Sample (cell culture medium/urine) diluent	15 ml
F. Detection Antibody-Biotinylated anti-mouse IP-10	2 vials (each vial is enough to assay half microplate)
G. Streptavidin-HRP Concentrate (4,000x)	8 µl
H. TMB One-Step Substrate Reagent (TMB in buffered solution)	12 ml
I. Stop Solution (2 M sulfuric acid)	8 ml

* Contains 0.09% sodium azide as preservative. Precaution: Sodium azide is a poisonous and hazardous substance which should be handled by trained staff only.

Storage of Kit Reagents



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Stable for up to 6 months from date of shipment at 2-4°C. Store reconstituted standard (recombinant protein) at -80°C. Opened microplate wells and reagents are stable for 1 month at 2-4°C. Return unused wells to the pouch containing desiccant pack and reseal along the entire edge. If the entire kit is stored at -20°C it is stable for 1 year.

Materials/reagents required but not provided:

- Microplate reader capable of measuring absorbance at 450 nm
- Precision pipettes to deliver 2 µl to 1 ml volumes
- Adjustable 1-25 ml pipettes for reagent preparation
- 100 ml and 1 liter graduated cylinders
- Absorbent paper
- Distilled or deionized water
- Log-log graph paper or computer and software for ELISA data analysis
- Tubes to prepare standard or sample dilutions

Preparation of Kit Reagents

Bring all reagents and samples to room temperature (18 - 25°C) before use.

Sample Dilution

If your samples need to be diluted, use Assay Diluent A (Item D) for dilution of serum/plasma samples, and Assay Diluent B (Item E) for dilution of culture supernatants and urine.

Assay Diluent B

Dilute 5-fold with deionized or distilled water.

IP-10 Standard

- Briefly spin the vial of Item C (Recombinant Mouse IP-10 Standard).
- Add 400 µl Assay Diluent A (for serum/plasma samples) or 1x Assay Diluent B (for cell culture medium and urine) into Item C vial to prepare a 0.1 µg/ml standard.
- Dissolve the powder thoroughly by a gentle mix.
- Add 20 µl IP-10 standard from the vial of Item C, into a tube with 980.0 µl Assay Diluent A or 1x Assay Diluent B to prepare a 2,000 pg/ml stock standard solution.
- Pipette 300 µl Assay Diluent A or 1x Assay Diluent B into each tube. Use the stock standard solution to produce a dilution series (shown below in Figure 1).
- Mix each tube thoroughly before the next transfer.
- Assay Diluent A or 1x Assay Diluent B serves as the zero standard (0 pg/ml).

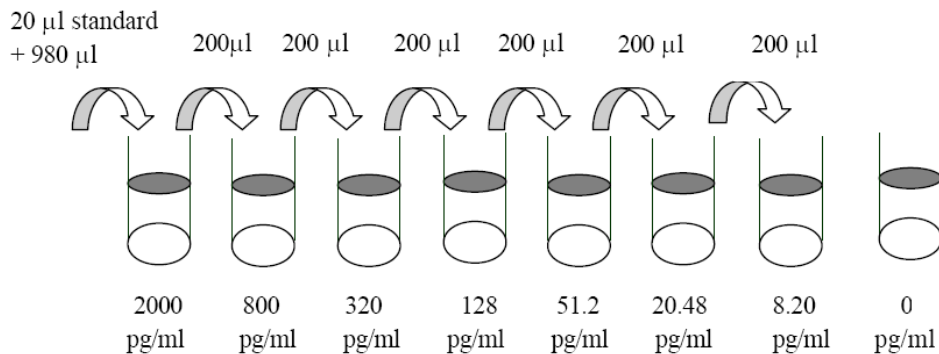
Figure 1



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Wash Buffer Concentrate

- If the Wash Concentrate (20x) (Item B) contains visible crystals, warm to room temperature and mix gently until dissolved.
- Dilute 20 ml of Wash Buffer Concentrate into deionized or distilled water to yield 400 ml of 1x Wash Buffer.

Detection Antibody

- Briefly spin Detection Antibody vial (Item F) before use.
- Add 100 µl of 1x Assay Diluent B into the vial to prepare a detection antibody concentrate.
- Pipette up and down to mix gently (the concentrate can be stored at 2-4°C for 5 days).
- The detection antibody concentrate should be diluted 80-fold with 1x Assay Diluent B and used in step 4 of the **ELISA Method**.

Streptavidin-HRP Concentrate

- Briefly spin Streptavidin-HRP Concentrate vial (Item G) and pipette up and down to mix gently before use.
- Streptavidin-HRP concentrate should be diluted 4,000-fold with 1x Assay Diluent B.
- *For example: Briefly spin the vial (Item G) and pipette up and down to mix gently. Add 3 µl of Streptavidin-HRP concentrate into a tube with 12 ml 1x Assay Diluent B to prepare a 4,000-fold diluted Streptavidin-HRP solution (don't store the diluted solution for next day use).*

ELISA Method

Be sure to read 'Preparation of Kit Reagents' before carrying out the assay.

1. Bring all reagents and samples to room temperature (18 - 25°C) before use. It is recommended that all standards and samples be run at least in duplicate.
2. Add 100 µl of each standard (see **Preparation of Kit Reagents: IP-10 Standard**) and sample into appropriate wells. Cover well and incubate for 2.5 hours at room temperature or overnight at 2-4°C with gentle shaking.



3. Discard the solution and wash 4 times with 1x Wash Solution. Wash by filling each well with Wash Buffer (300 μ l) using a multi-channel pipette or autowasher. 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.
4. Add 100 μ l of 1x prepared biotinylated antibody (see **Preparation of Kit Reagents: Detection Antibody**) to each well. Incubate for 1 hour at room temperature with gentle shaking.
5. Discard the solution and wash as in step 3 above.
6. Add 100 μ l of prepared Streptavidin solution (see **Preparation of Kit Reagents: Streptavidin-HRP Concentrate**) to each well. Incubate for 45 minutes at room temperature with gentle shaking.
7. Discard the solution and wash as in step 3 above.
8. Add 100 μ l of TMB One-Step Substrate Reagent (Item H) to each well. Incubate for 30 minutes at room temperature in the dark with gentle shaking.
9. Add 50 μ l of Stop Solution (Item I) to each well. Read at 450 nm immediately.

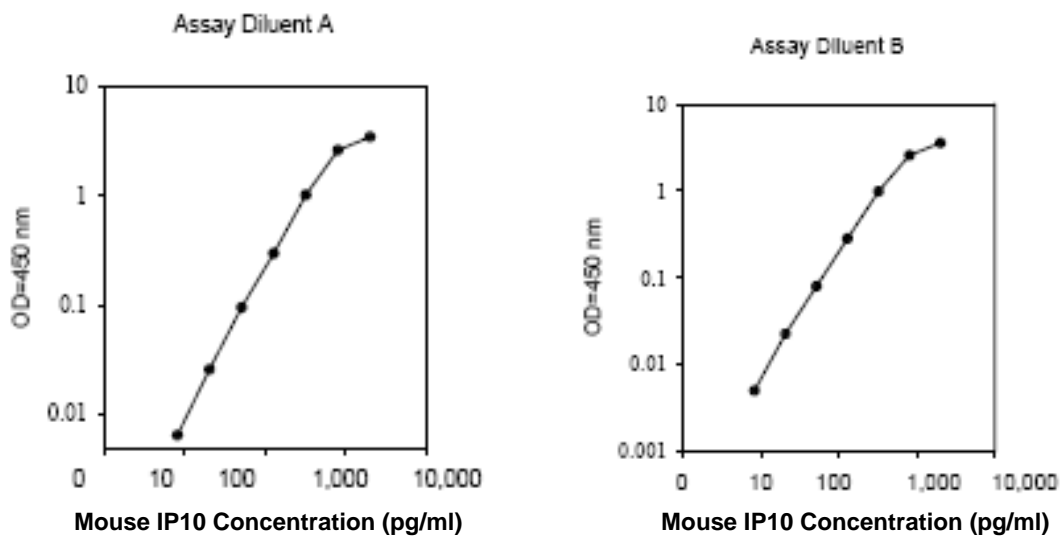
Calculation of Results

Calculate the mean absorbance for each set of duplicate standards, controls and samples, and subtract the average zero standard optical density. Plot the standard curve on log-log graph paper or using Sigma plot software, with standard concentration on the x-axis and absorbance on the y-axis. Draw the best-fit straight line through the standard points.

Figure 2

Typical Data

These standard curves are for demonstration only. A standard curve must be run with each assay.



Performance Characteristics

Sensitivity:

The minimum detectable dose of IP-10 is typically less than 15 pg/ml.

Recovery:

Recovery was determined by spiking various levels of mouse IP-10 into mouse serum, plasma and cell culture media. Mean recoveries are as follows:

Sample Type	Average % Recovery	Range (%)
Serum	97.68	86-106
Plasma	96.42	85-104
Cell culture media	98.86	88-107

Linearity:

Sample Type		Serum	Plasma	Cell Culture Media
1:2	Average % of Expected	96	93	97
	Range (%)	88-104	86-105	85-106
1:4	Average % of Expected	96	95	96
	Range (%)	89-105	87-106	90-106
1:8	Average % of Expected	97	95	98
	Range (%)	87-105	88-107	89-106

Reproducibility:

Intra-assay: CV<10%

Inter-assay: CV<12%

Specificity:

Cross Reactivity: This ELISA kit shows no cross-reactivity with any of the cytokines tested (e.g., Mouse CD30, L CD30, T CD40, CTACK, CXCL16, Eotaxin, Eotaxin-2, Fas Ligand, Fractalkine, GM-CFS, IGFBP3, IGFBP5, IGFBP6, IL1a, IL1 β , IL2, IL3, IL3 Rb, IL4, IL5, IL6, IL9, IL10, IL12 p40/p70, IL12 p70, IL13, IL17, KC, Leptin R, LEPTIN(OB), LIX, L-Selectin, Lymphotoxin, MCP1, MCP5, M-CSF, MIG, MIP1a, MIP2, MIP3 β , MIP3a, PF4, P-Selectin, RANTES, SCF, SDF1a, TARC, TCA3, TECK, TIMP1, TNF-a, TNF RI, TNF RII, TPO, VCAM1, VEGF).



Troubleshooting Guide:

Problem	Cause	Solution
1. Poor standard curve	1. Inaccurate pipetting 2. Improper standard dilution	1. Check pipettes 2. Ensure a brief spin of Item C and dissolve the powder thoroughly by a gentle mix.
2. Low signal	1. Too brief incubation times 2. Inadequate reagent volumes or improper dilution	1. Ensure sufficient incubation time; ELISA Method Step 2 may change to overnight. 2. Check pipettes and ensure correct preparation.
3. Large CV	1. Inaccurate pipetting	1. Check pipettes.
4. High background	1. Plate is insufficiently washed 2. Contaminated wash buffer	1. Review the manual for proper wash. If using a plate washer, check that all ports are unobstructed. 2. Make fresh wash buffer.
5. Low sensitivity	1. Improper storage of the ELISA Kit 2. Stop solution	1. Store your standard at < -20°C after reconstitution, others at 2-4°C. Keep substrate solution protected from light. 2. Stop solution should be added to each well before measure.

NOT FOR HUMAN USE. FOR RESEARCH USE ONLY. NOT FOR DIAGNOSTIC OR THERAPEUTIC USE.



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