

Mouse Plasminogen Total Antigen Assay ELISA Kit

Catalog No.	CSI19895A CSI19895B	1 x 96 tests 5 x 96 tests
Intended Use:	This mouse plasminogen total assay is for the quantitative determination of total plasminogen and plasmin in biological fluids.	
Background:	Plasminogen is a single chain glycoprotein zymogen and is the precursor of the fibrinolytic enzyme plasmin. Plasminogen deficiencies are classified as hypoplasminogenemia (Type I) or dysplasminogenemia (Type 2) and are associated with decreased extracellular fibrin clearance leading to mucous membrane lesions and ligneous conjunctivitis.	
Assay Principle:	Mouse plasminogen will bind to the capture antibody coated on the microtiter plate. Plasminogen, plasmin, and plasmin in complex with antiplasmin will react with the antibody on the plate. After appropriate washing steps, polyclonal anti-mouse plasminogen primary antibody binds to the plasminogen. Excess antibody is washed away and bound polyclonal antibody is then reacted with the secondary antibody conjugated to horseradish peroxidase. Following an additional washing step, TMB substrate is used for color development at 450nm. The amount of color development is directly proportional to the concentration of total plasminogen in the sample.	
Reagents Provided:	♦ 96-well (8X12 removable wells) antibody coated microtiter strip plate: containing rat anti-mouse plasminogen antibody dried and blocked on the strip well surface. ♦ 10X Wash Buffer: 1 bottle of 50ml; bring to 1X using DI water ♦ Mouse Plasminogen activity standard: 1 vial lyophilized standard ♦ Anti-plasminogen primary antibody: 1 vial lyophilized polyclonal rabbit anti-mouse antibody ♦ Anti-rabbit horseradish peroxidase secondary antibody: 1 vial concentrated HRP labeled antibody ♦ TMB substrate solution: 1 bottle 10ml solution	
Storage and Stability:	All kit components must be stored at 4°C. Store unopened plate and any unused microtiter strips in the pouch with desiccant. Reconstituted standards and primary may be stored at -70°C for later use. DO NOT freeze/thaw the standards and primary antibody more than once. All other unused kit components must be stored at 4°C. Kit should be used no later than the expiration date.	



Reagents and Equipment Required:

- Pipettes covering 0-10µl and 200-1000µl
- 12-channel pipette covering 30-300µl
- Paper towels or kimwipes
- 50ml tubes
- 1N H₂SO₄
- DI water
- Magnetic stirrer and stir-bars
- Plastic containers
- ELISA plate cover
- TBS buffer
- 3% BSA Blocking buffer
- Microtiter plate spectrophotometer operable at 450nm
- Microtiter plate shaker with uniform horizontally circular movement up to 300rpm.

Warnings:

Avoid skin and eye contact when using TMB One substrate solution since it may be irritating to eyes, skin, and respiratory system. Wear safety goggles and gloves.

Precautions:

- **DO NOT** mix any reagents or components of this kit with any reagents or components of any other kit. This kit is designed to work properly as provided.
- **DO NOT** pipette reagents by mouth.
- Always pour substrate out of the bottle into a clean test tube. **DO NOT** pipette out of the bottle as you could contaminate the substrate.
- Keep plate covered except when adding reagents, washing, or reading.
- All kit components must be kept refrigerated (4°C).
- **DO NOT** smoke, drink, or eat in areas where specimens or reagents are being handled.

Preparation of Reagents:

- **TBS buffer:** 0.1M Tris-HCl 0.15M NaCl, pH 7.4
- **Blocking buffer (BSA):** 3% BSA in TBS buffer

Specimen Collection:

Samples of mouse plasma, serum, urine, cell culture media, or tissue extracts may be applied directly to the plate.

The assay measures total plasminogen in the 0-500 ng/ml range. Samples giving plasminogen levels above 500 ng/ml should be diluted in a similar biological fluid devoid of plasminogen. Blocking buffer may also be used. A dilution of 1:1,000 to 1:10,000 is recommended for measurement of plasminogen in normal murine plasma.

Assay Procedure:

Perform assay at room temperature. Vigorously shake plate (300rpm) at each step of the assay.



Preparation of Standard: Reconstitute standard as directed on vial to give a 500 ng/ml standard solution.

Dilution table for preparation of mouse plasminogen standards:

Plasminogen concentration (ng/mL)	Dilutions
500	100µl from standard vial
250	500µL (BSA) + 500µL (500ng/mL)
100	600µL (BSA) + 400µL (250ng/mL)
50	500µL (BSA) + 500µL (100ng/mL)
20	600µL (BSA) + 400µL (50ng/mL)
10	500µL (BSA) + 500µL (20ng/mL)
5	500µL (BSA) + 500µL (10ng/mL)
2.5	500µL (BSA) + 500µL (5ng/mL)
0	500µL (BSA) Zero point to determine background

NOTE: DILUTIONS FOR THE STANDARD CURVE AND ZERO STANDARD MUST BE MADE AND APPLIED TO THE PLATE IMMEDIATELY.

Standard and Unknown Addition:

Add 100µl of plasminogen standards (in duplicate) and unknowns to wells. Carefully record position of standards and unknowns. Shake plate at 300rpm for 30 minutes. Wash wells three times with 300µl wash buffer. Remove excess wash by gently tapping plate on paper towel or kimwipe. NOTE: If the unknown is thought to have high plasminogen levels, dilutions may be made in plasma devoid of plasminogen, or in blocking buffer. A 1:10,000 dilution, generated by two serial dilutions of 1:100 each, is recommended for the measurement of plasminogen in normal murine plasma.

Primary Antibody Addition:

Reconstitute primary antibody as directed on vial and agitate gently to completely dissolve contents. Add 100µl to all wells. Shake plate at 300rpm for 30 minutes. Wash wells three times with 300µl wash buffer. Remove excess wash by gently tapping plate on paper towel or kimwipe.

Secondary Antibody Addition:

Dilute 2.5µl conjugated secondary antibody in 10ml of 3% BSA blocking buffer and add 100µl to all wells. Shake plate at 300rpm for 30 minutes. Wash wells three times with 300µl wash buffer. Remove excess wash by gently tapping plate on paper towel or kimwipe.



Substrate Incubation:

Add 100µl TMB substrate to all wells and shake plate for 2-10 minutes. Substrate will change from colorless to different strengths of blue. Quench reaction by adding 50µl of 1N H₂SO₄ stop solution to all wells when samples are visually in the same range as the standards. Add stop solution to wells in the same order as substrate upon which color will change from blue to yellow. Mix thoroughly and read final absorbance values at 450nm. For best results read plate immediately

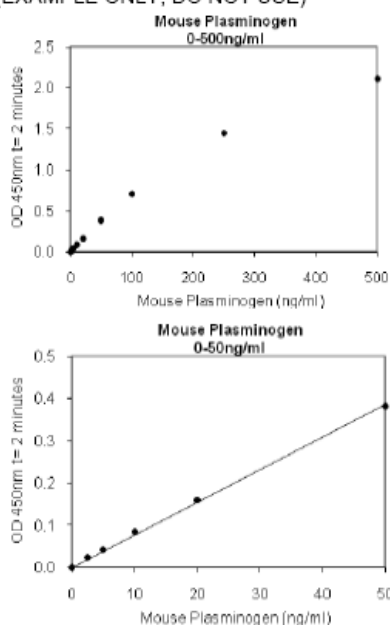
Measurement:

Set the absorbance at 450nm in a microtiter plate spectrophotometer. Measure the absorbance in all wells at 450nm. Subtract zero point from all standards and unknowns to determine corrected absorbance (A_{450}).

Assay Calibration:

Plot A_{450} against the amount of plasminogen in the standards. Fit a straight line through the points using a linear fit procedure. The plasminogen in the unknowns can be determined from this curve.

A typical standard curve.
(EXAMPLE ONLY, DO NOT USE)



Expected Values:

The concentration of human plasminogen in pooled donor plasma from normal individuals was found to be 195 ± 10 $\mu\text{g/ml}$. Normal values of plasminogen in mouse plasma have not been conclusively determined but are believed to be similar to human plasma. Plasminogen antigen was found to be 84 ± 8 $\mu\text{g/ml}$ in a small sample (n=4) of normal mice. Hypercoagulable states may be induced by genetically decreased plasminogen or abnormal plasminogen.

Disclaimer:

This information is believed to be correct but does not claim to be all-inclusive and shall be used only as a guide. The supplier of this kit shall not be held liable for any damage resulting from handling or from contact with the above product.

96 Well Plate Layout

Standards: 18 wells

Samples: 78 wells

	1	2	3	4	5	6	7	8	9	10	11	12
A	0	2.5ng/ml	5ng/ml	10ng/ml	20ng/ml	50ng/ml	100ng/ml	250ng/ml	500ng/ml			
B	0	2.5ng/ml	5ng/ml	10ng/ml	20ng/ml	50ng/ml	100ng/ml	250ng/ml	500ng/ml			
C												
D												
E												
F												
G												
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