



AssayMax Mouse Albumin ELISA Kit

Catalog Number EMA3201-1

Introduction

Albumin is serum hepatic protein, the most abundant protein in serum and contributes to the maintenance of oncotic pressure as well as to transport of hydrophobic molecules (1). Serum albumin level has been linked in clinical practice to several diseases. Low albumin levels can suggest liver (2), kidneys disease (3), inflammation (4), shock (5), and malnutrition (6). On the other hand, high albumin levels usually reflect dehydration (7).

Principal of the Assay

The AssayMax Mouse Albumin ELISA (Enzyme-Linked Immunosorbent Assay) kit employs a quantitative sandwich enzyme immunoassay technique that measures mouse plasma albumin in less than 2.5 hours. A polyclonal antibody specific for mouse albumin has been pre-coated onto a 96-well microplate with removable strips. Albumin in standards and samples is sandwiched by the immobilized polyclonal antibody and biotinylated polyclonal antibody specific for mouse Albumin, which is recognized by a streptavidin-peroxidase conjugate. All unbound material is then washed away and a peroxidase enzyme substrate is added. The color development is stopped and the intensity of the color is measured.

Caution and Warning

- This kit is for research use only.
- The kit should not be used beyond the expiration date.
- The Stop Solution is an acid solution.

Reagents

- **Mouse Albumin Microplate:** A 96-well polystyrene microplate (12 strips of 8 wells) coated with a polyclonal antibody against mouse albumin.
- **Sealing Tapes:** Each kit contains 3 pre-cut, pressure-sensitive sealing tapes that can be cut to fit the format of the individual assay.
- **Mouse Albumin Standard:** Mouse Albumin in a buffered protein base (400 ng, lyophilized).

- **Biotinylated Mouse Albumin Antibody (100x):** A 100-fold concentrated biotinylated polyclonal antibody against mouse Albumin (80 µl).
- **Streptavidin-Peroxidase Conjugate (SP Conjugate):** A 100-fold concentrate (120 µl).
- **EIA Diluent Concentrate (10x):** A 10-fold concentrated buffered protein base (30 ml).
- **Wash Buffer Concentrate (10x):** A 10-fold concentrated buffered surfactant (2 x 30 ml).
- **Chromogen Substrate:** A ready-to-use stabilized peroxidase chromogen substrate tetramethylbenzidine (8 ml).
- **Stop Solution:** A 0.5 N hydroxychloric acid (12 ml) to stop the chromogen substrate reaction.

Storage Condition

- Store unopened kit at 2-8⁰C up to expiration date.
- Opened reagents may be stored for up to 1 month at 2-8⁰C. Store reconstituted standard and Biotinylated Albumin at -20⁰C or below.
- Opened unused strip wells may return to the foil pouch with the desiccant pack, reseal along zip-seal. May be stored for up to 1 month in a vacuum desiccator.

Other Supplies Required

- Microplate reader capable of measuring absorbance at 450 nm.
- Pipettes (1-20 µl, 20-200 µl, and multiple channel).
- Deionized or distilled reagent grade water.

Sample Collection, Preparation and Storage

- **Cell Culture Supernatants:** Centrifuge cell culture media at 3,000 x g for 10 minutes to remove debris. Collect supernatants and assay. Store samples at -20⁰C or below. Avoid repeated freeze-thaw cycles.
- **Urine:** Collect urine using sample pot. Centrifuge samples at 800 x g for 10 minutes and assay. Dilute samples 1:100 into EIA Diluent. Store samples at -20⁰C or below for up to 3 months. Avoid repeated freeze-thaw cycles.

Reagent Preparation

- Freshly dilute all reagents and bring all reagents to room temperature before use. If crystals have formed in the concentrate, mix gently until the crystals have completely dissolved.
- **Standard Curve:** Reconstitute the 400 ng of albumin standard with 2 ml of EIA Diluent to generate a solution of 200 ng/ml. Allow the standard to sit for 10 minutes with gentle agitation prior to making dilutions. Prepare triplicate standard points by serially diluting the standard solution (200 ng/ml) 1:4 with EIA Diluent to produce 50, 12.5, 3.125, 1.56, 0.781, and 0.195 ng/ml solutions. EIA Diluent serves as the zero standard (0 µg/ml). Any remaining solution should be frozen at < -20⁰C.

Standard Point	Dilution	[mAlbumin] (ng/ml)
P1	Standard (200 ng/ml)	200.000
P2	1 part P1 + 3 parts EIA Diluent	50.000
P3	1 part P2 + 3 parts EIA Diluent	12.500
P4	1 part P3 + 3 parts EIA Diluent	3.125
P5	1 part P4 + 3 parts EIA Diluent	0.781
P6	1 part P5 + 3 parts EIA Diluent	0.195
P7	EIA Diluent	0.000

- **Biotinylated Mouse Albumin Antibody (100x):** Spin down the antibody briefly and dilute the desired amount of the antibody 1:100 with EIA Diluent.
- **EIA Diluent Concentrate (10x):** Dilute the EIA Diluent 1:10 with reagent grade water.
- **Wash Buffer Concentrate (10x):** Dilute the Wash Buffer Concentrate 1:10 with reagent grade water.
- **SP Conjugate (100x):** Spin down the SP Conjugate briefly and dilute the desired amount of the conjugate 1:100 with EIA Diluent.

Assay Procedure

- Prepare all reagents, working standards and samples as instructed. Bring all reagents to room temperature before use. The assay is performed at room temperature (20-30°C).
- Remove excess microplate strips from the plate frame and return them immediately to the foil pouch with desiccant inside. Reseal the pouch securely to minimize exposure to water vapor and store in a vacuum desiccator.
- Add 50 µl of standard or sample per well, and cover wells and incubate for 1 hours. Start the timer after the last sample addition.
- Wash five times with 200 µl of Wash Buffer. Invert the plate and decant the contents, and hit it 4-5 times on absorbent paper towel to completely remove liquid at each step.
- Add 50 µl of Biotinylated Mouse Albumin Antibody to each well and incubate for 30 minutes.
- Wash five times with 200 µl of Wash Buffer.
- Add 50 µl of Streptavidin-Peroxidase Conjugate to each well and incubate for 30 minutes. Turn on the microplate reader and set up the program in advance.
- Wash five times with 200 µl of Wash Buffer.
- Add 50 µl of Chromogen Substrate per well and incubate for about 7 minutes or till the optimal blue color density develops. Gently tap plate to ensure thorough mixing and break the bubbles in the well with pipette tip.
- Add 50 µl of Stop Solution to each well. The color will change from blue to yellow.
- Read the absorbance on a microplate reader at a wavelength of 450 nm **immediately**. Please note that after the reaction is stopped for about 10 minutes, some black particles may be generated at high concentration point, which will reduce the readings.

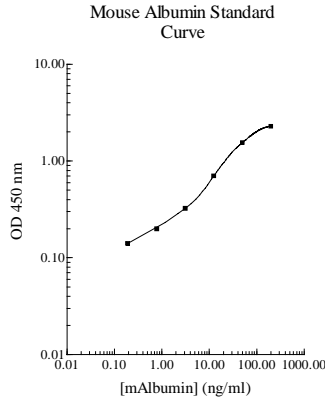
Data Analysis

- Calculate the mean value of the triplicate readings for each standard and sample.

- To generate a ξ and the corresponding mean 450 nm absorbance on the y-axis. The best-fit line can be determined by regression analysis using log-log or 4-parameter curve fit.
- Determine the unknown sample concentration from the Standard Curve.

Standard Curve

- The curve is provided for illustration only. A standard curve should be generated each time the assay is performed.



Performance Characteristics

- The minimum detectable dose of Albumin is typically 150 pg/ml.
- Intra-assay and inter-assay coefficients of variation were 4.5% and 7.3% respectively.
- No significant cross-reactivity or interference was observed.

References

- (1) Gekle M. (2004) *Annu Rev Physiol.*
- (2) Schindler C *et al.* (1999) *J Hepatol.* 31(6):1132
- (3) Hemmeler MH *et al.* (1997) *Nephrol Dial Transplant.* 12 Suppl 2:57-62
- (4) Sesmilo G *et al.* (2004) *Ann Intern Med.* 133(2):111-22
- (5) Wettstein R *et al.* (2004) *Shock.* 22(4):351-357
- (6) Saito T *et al.* (1991) *Jpn J Surg.* 21(4):402-11
- (7) Strand TA (2004) *Am J Clin Nutr.* 79(3):451-6

Revision 2.1

Related Products

- EA2201-1 AssayMax Human Albumin ELISA Kit (for plasma/serum samples)
- EA3201-1 AssayMax Human Albumin ELISA Kit (for cell culture supernatant)
- EMA2201-1 AssayMax Mouse Albumin ELISA Kit (for plasma/serum samples)
- ERA2201-1 AssayMax Rat Albumin ELISA Kit (for plasma/serum samples)
- ERA3201-1 AssayMax Rat Albumin ELISA Kit (for cell culture supernatant)