



AssayMax Retinol-Binding Protein (RBP) ELISA Kit

Catalog Number ER2005-1

Introduction

Retinoid-binding protein (RBP) is a transport protein that acts by solubilizing and protecting its labile ligands in aqueous spaces. It also has diverse and specific functions in regulating the disposition, metabolism and activities of retinoids (1). Retinol-binding protein is the specific plasma carrier of retinol, and in charge of the vitamin transport from the liver to target cells (2). Lower serum RBP level associates with diarrhea (3). High level of RBP in urine could be a good indicator of renal damage (4), microvascular complications with type-2 diabetes mellitus (5).

Principal of the Assay

The AssayMax Human RBP ELISA kit is designed for detection of human RBP in urine and cell culture supernatants. This assay employs a quantitative sandwich enzyme immunoassay technique that measures RBP in 3.5 hours. A polyclonal antibody specific for RBP has been pre-coated onto a microplate. RBP in standards and samples is sandwiched by the immobilized antibody and biotinylated polyclonal antibody specific for RBP, 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

- **RBP Microplate:** A 96-well polystyrene microplate (12 strips of 8 wells) coated with a polyclonal antibody against human RBP.
- **Sealing Tapes:** Each kit contains 3 pre-cut, pressure-sensitive sealing tapes, which can be cut to fit the format of the individual assay.
- **RBP Standard:** Recombinant human RBP in a buffered protein base (6 µg, lyophilized).
- **Biotinylated RBP Antibody (100x):** A 100-fold biotinylated polyclonal antibody against RBP (80 µl).
- **EIA Diluent Concentrate (10x):** A 10-fold concentrated buffered protein base (30 ml).
- **Wash Buffer Concentrate (20x):** A 20-fold concentrated buffered surfactant (30 ml).
- **Streptavidin-Peroxidase Conjugate (SP Conjugate):** A 100-fold concentrate (90 µl).
- **Chromogen Substrate:** A ready-to-use stabilized peroxidase chromogen substrate tetramethylbenzidine (8 ml).

- **Stop Solution:** A 0.5 N hydrochloric acid to stop the chromogen substrate reaction (12 ml).

Storage Condition

- Store unopened kit at 2 - 8⁰C up to expiration date.
- Opened EIA Diluent may be stored for up to 1 month at 2 - 8⁰C. Store reconstituted reagents 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, 200-1000µl and multiple channel).
- Deionized or distilled reagent grade water.

Sample Collection and Storage

- **Plasma:** Collect plasma using one-tenth volume of 0.1 M sodium citrate as an anticoagulant. Centrifuge samples at 2000 x g for 10 minutes and assay. Dilute samples 1:500 into EIA Diluent. The undiluted samples can be stored at -20⁰C or below for up to 3 months. Avoid repeated freeze-thaw cycles.
- **Serum:** Samples should be collected into a serum separator tube. After clot formation, centrifuge samples at 2000 x g for 10 minutes. Remove serum and assay. Dilute samples 1:500 into EIA Diluent. The undiluted samples can be stored at -20⁰C or below for up to 3 months. Avoid repeated freeze-thaw cycles.
- **Cell Culture Supernatants:** Centrifuge cell culture media at 2000 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 600 x g for 10 minutes and assay. Dilute samples 1:4 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.
- **EIA Diluent Concentrate (10x):** Dilute the EIA Diluent Concentrate 1:10 with reagent grade water. Store for up to 1 month at 2 - 8⁰C.
- **RBP Standard:** Reconstitute the 6 µg of human RBP Standard with 3 ml of EIA Diluent to generate a stock solution of 2 µg/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 (2 µg/ml) with equal volume EIA Diluent to produce 1, 0.5, 0.25, 0.125, 0.063, and 0.031 µg/ml. EIA Diluent serves as the zero standard (0 µg /ml). Any remaining solution should be frozen at -20⁰C.

Standard Point	Dilution	[RBP] ($\mu\text{g/ml}$)
P1	1 part Standard (2 $\mu\text{g/ml}$)	2.000
P2	1 part P1 + 1 part EIA Diluent	1.000
P3	1 part P1 + 1 part EIA Diluent	0.500
P4	1 part P1 + 1 part EIA Diluent	0.250
P5	1 part P1 + 1 part EIA Diluent	0.125
P6	1 part P1 + 1 part EIA Diluent	0.063
P7	1 part P1 + 1 part EIA Diluent	0.031
P8	EIA Diluent	0.000

- **Biotinylated RBP Antibody (100x):** Spin down the antibody briefly and dilute the desired amount of the antibody 1:100 with EIA Diluent. Any remaining solution should be frozen at -20°C .
- **Wash Buffer Concentrate (20x):** Dilute the Wash Buffer Concentrate 1:20 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. Any remaining solution should be frozen at -20°C .

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 ($20-30^{\circ}\text{C}$).
- 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. Cover wells with a sealing tape and incubate for two 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 RBP Antibody to each well and incubate for one hour.
- Wash five times with 200 μl of Wash Buffer.
- Add 50 μl of Streptavidin-Peroxidase Conjugate per 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 9 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 some unstable black particles may be generated at high concentration points after stopping the reaction for about 10 minutes, which will reduce the readings.

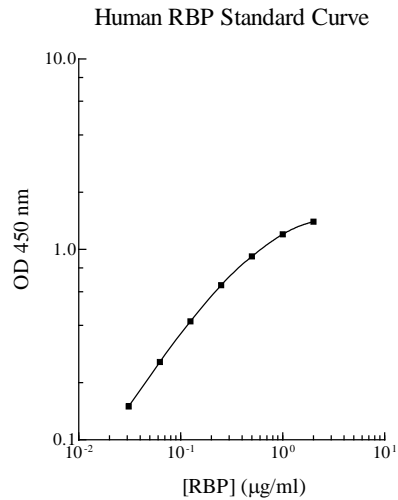
Data Analysis

- Calculate the mean value of the triplicate readings for each standard and sample.
- To generate a Standard Curve, plot the graph using the standard concentrations on the x-axis 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 four-parameter logistic curve-fit.

- Determine the unknown sample concentration from the Standard Curve and multiply the value by the dilution factor.

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 RBP is typically 8 ng/ml.
- Intra-assay and inter-assay coefficients of variation were 4.2 % and 7.1% respectively.

Linearity

Sample Dilution	Average Percentage of Expected Value	
	Plasma	Serum
1:250	91%	92%
1:500	98%	101%
1:1000	105%	108%

Sample Dilution	Average Percentage of Expected Value
	Urine
No Dilution	96%
1:2	102%
1:4	110%

Sample Dilution	Average Percentage of Expected Value
	Cell Culture Media
1:2	97%
1:4	105%
1:8	107%

Recovery

Standard Added Value	0.05 – 0.5 ug/ml
Recovery %	82-115 %
Average Recovery %	98.5 %

Cross-Reactivity

Species	% Cross Reactivity
Beagle	None
Bovine	None
Monkey	< 10
Mouse	< 1
Rat	None
Swine	< 0.5

- 10% FBS in culture media will not affect the assay.

References

- (1) Noy N. (2000) *Biochem. J.* 348, 481-495
- (2) Bellovino D *et. al* (2003) *Mol Aspects Med.* 24(6):411-20
- (3) Mitra AK *et. al* (2002) *J Health Popul Nutr.* 20(1): 12-7
- (4) Corso A *et. al.* (1999) *Ann Hematol.* 78(8): 371-5
- (5) Hong CY *et. al* (2000) *J Diabetes Complications* 14(5):259-65

Related Products

- ER3005-1 AssayMax Human RBP4 ELISA Kit

Version 2.5