Human RBP
ELISA Kit

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Hinweis/Note:
Der Packungsbeileger dient nur als erste Information.
Der relevante Packungsbeileger liegt der Ware bei.

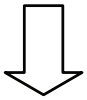
The datasheet is only a first information.
The relevant datasheet is included with the product.

For any questions regarding troubleshooting or performing the assay, please contact our support team at support@assaypro.com.

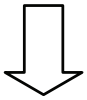
Thank you for choosing Assaypro.

Assay Summary

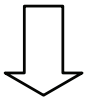
Add 50 µl of Standard/ Sample per well.
Incubate 2 hours.



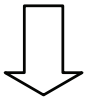
Wash, then add
50 µl of Biotinylated Antibody per well.
Incubate 1 hour.



Wash, then add
50 µl of SP Conjugate per well.
Incubate 30 minutes.



Wash, then add
50 µl of Chromogen Substrate per well.
Incubate 12 minutes.



Add 50 µl of Stop Solution per well.
Read at 450 nm immediately.

Assay Template

	A	B	C	D	E	F	G	H
1								
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AssayMax Retinol-Binding Protein (RBP) ELISA Kit

Catalog No. ER2005-1
Sample Insert/Reference Only

Introduction

Retinol-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, encharged 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) and microvascular complications with type-2 diabetes mellitus (5).

Principle of the Assay

The AssayMax Human RBP ELISA (Enzyme-Linked Immunosorbent Assay) kit is designed for detection of human RBP in urine, milk, saliva, CSF, and cell culture samples. This assay employs a quantitative sandwich enzyme immunoassay technique that measures RBP in less than 4 hours. A polyclonal antibody specific for RBP has been pre-coated onto a 96-well microplate with removable strips. 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

- **Prepare all reagents (working diluent buffer, wash buffer, standard, biotinylated antibody, and SP conjugate) as instructed, prior to running the assay.**
- **Prepare all samples prior to running the assay. The dilution factors for the samples are suggested in this protocol. However, the user should determine the optimal dilution factor.**
- **Spin down the SP conjugate vial and the biotinylated antibody vial before opening and using contents.**
- This kit is for research use only.
- The kit should not be used beyond the expiration date.

- The Stop Solution is an acidic solution.

Reagents

- **Human 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 precut, pressure sensitive sealing tapes, which can be cut to fit the format of the individual assay.
- **Human RBP Standard:** Human RBP in a buffered protein base (2.5 μg, lyophilized).
- **Biotinylated Human RBP Antibody (50x):** A 50-fold biotinylated polyclonal antibody against RBP (140 μl).
- **EIA Diluent Concentrate (10x)**: A 10-fold concentrated buffered protein base (20 ml).
- **Wash Buffer Concentrate (20x)**: A 20-fold concentrated buffered surfactant (30 ml, 2 bottles).
- **Streptavidin-Peroxidase Conjugate (SP Conjugate):** A 100-fold concentrate (80 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

- Upon arrival, immediately store components of the kit at recommended temperatures up to the expiration date.
- Store SP Conjugate and Biotinylated Antibody at -20°C.
- Store Microplate, Diluent Concentrate (10x), Wash Buffer, Stop Solution, and Chromogen Substrate at 2-8°C.
- Unused microplate wells may be returned to the foil pouch with the desiccant packs and resealed. May be stored for up to 30 days in a vacuum desiccator.
- Diluent (1x) may be stored for up to 30 days at 2-8°C.
- Store Standard at 2-8°C before reconstituting with Diluent and at -20°C after reconstituting with Diluent.

Other Supplies Required

- Microplate reader capable of measuring absorbance at 450 nm.
- Pipettes (1-20 μl, 20-200 μl, 200-1000 μl, and multiple channel).
- Deionized or distilled reagent grade water.

Sample Collection and Storage

- **Cell Culture Supernatants:** Centrifuge cell culture media at 3000 x *g* for 10 minutes to remove debris. Collect supernatants and assay. Store samples at -20°C or below. Avoid repeated freeze-thaw cycles.
- **Urine:** Collect urine using sample pot. Centrifuge samples at 800 x *g* for 10 minutes and assay. Store samples at -20°C or below for up to 3 months. Avoid repeated freeze-thaw cycles.
- **Milk:** Collect milk using sample tube. Centrifuge samples at 800 x *g* for 10 minutes and assay. Store samples at -20°C or below for up to 3 months. Avoid repeated freeze-thaw cycles.
- **Saliva:** Collect saliva using sample tube. Centrifuge samples at 800 x *g* for 10 minutes and assay. Store samples at -20°C or below for up to 3 months. Avoid repeated freeze-thaw cycles.
- **CSF:** Collect cerebrospinal fluid (CSF) using sample pot. Centrifuge samples at 3000 x *g* for 10 minutes. Dilute sample 1:5 into EIA Diluent and assay. The undiluted samples can be stored at -80°C 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.
- **EIA Diluent Concentrate (10x)**: If crystals have formed in the concentrate, mix gently until the crystals have completely dissolved. Dilute the EIA Diluent Concentrate 1:10 with reagent grade water. Store for up to 30 days at 2-8°C.
- **Human RBP Standard:** Reconstitute the 2.5 μg of Human RBP Standard with 2.5 ml of EIA Diluent to generate a 1 μg/ml standard solution. Allow the standard to sit for 10 minutes with gentle agitation prior to making dilutions. Prepare duplicate or triplicate standard points by serially diluting the standard solution (1 μg/ml) 1:2 with equal volume of EIA Diluent to produce 0.5, 0.25, 0.125, 0.063, 0.031, 0.016, and 0.008 μg/ml solutions. EIA Diluent serves as the zero standard (0 μg /ml). Any remaining solution should be frozen at -20°C and used within 30 days.

Standard Point	Dilution	[RBP] (μg/ml)
P1	1 part Standard (1 μg/ml) + 1 part EIA Diluent	0.500
P2	1 part P1 + 1 part EIA Diluent	0.250
P3	1 part P2 + 1 part EIA Diluent	0.125
P4	1 part P3 + 1 part EIA Diluent	0.063
P5	1 part P4 + 1 part EIA Diluent	0.031
P6	1 part P5 + 1 part EIA Diluent	0.016
P7	1 part P6 + 1 part EIA Diluent	0.008
P8	EIA Diluent	0.000

- **Biotinylated Human RBP Antibody (50x):** Spin down the antibody briefly and dilute the desired amount of the antibody 1:50 with EIA Diluent. Any remaining solution should be frozen at -20°C.
- **Wash Buffer Concentrate (20x):** If crystals have formed in the concentrate, mix gently until the crystals have completely dissolved. 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, standard solutions, and samples as instructed. Bring all reagents to room temperature before use. The assay is performed at room temperature (20-25°C).
- Remove excess microplate strips from the plate frame and return them immediately to the foil pouch with desiccants inside. Reseal the pouch securely to minimize exposure to water vapor and store in a vacuum desiccator.
- Add 50 μl of Human RBP Standard or sample per well. Cover wells with a sealing tape and incubate for 2 hours. Start the timer after the last addition.
- Wash five times with 200 μl of Wash Buffer manually. Invert the plate each time and decant the contents; hit 4-5 times on absorbent material to completely remove the liquid. If using a machine, wash six times with 300 μl of Wash Buffer and then invert the plate, decanting the contents; hit 4-5 times on absorbent material to completely remove the liquid.
- Add 50 μl of Biotinylated Human RBP Antibody to each well and incubate for 1 hour.
- Wash the microplate as described above.
- 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.

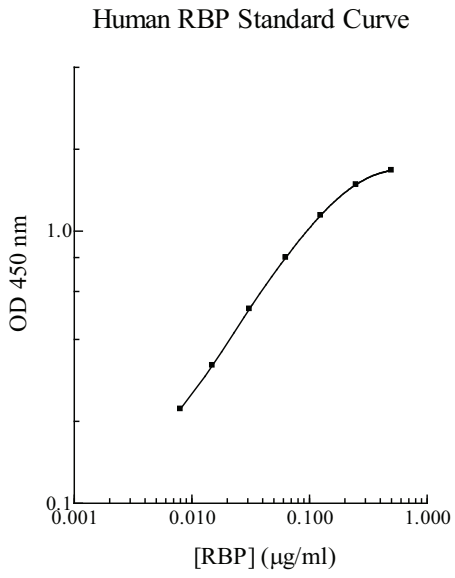
- Wash the microplate as described above.
- Add 50 µl of Chromogen Substrate per well and incubate for about 12 minutes or till the optimal blue color density develops. Gently tap plate to ensure thorough mixing and break the bubbles in the well with pipette tip.
- Add 50 µl of Stop Solution to each well. The color will change from blue to yellow.
- Read the absorbance on a microplate reader at a wavelength of 450 nm **immediately**. If wavelength correction is available, subtract readings at 570 nm from those at 450 nm to correct optical imperfections. Otherwise, read the plate at 450 nm only. 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 duplicate or 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 ~ 0.008 µg/ml.
- Intra-assay and inter-assay coefficients of variation were 4.6% and 7.2% respectively.

Linearity

Average Percentage of Expected Value			
Sample Dilution	**Urine**	**Milk**	**Saliva**
No Dilution	97%	95%	99%
1:2	101%	97%	97%
1:4	110%	108%	107%

Recovery

Standard Added Value	0.01 – 0.1 µg/ml
Recovery %	84 – 114%
Average Recovery %	98%

Cross-Reactivity

Species	% Cross Reactivity
Canine	None
Bovine	None
Monkey	<10%
Mouse	None
Rat	None
Swine	None

- 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

Version 3.2

Related Products

- ER1005-1 AssayMax Human RBP ELISA Kit (Plasma and Serum samples)
- ER3005-1 AssayMax Human RBP4 ELISA Kit (Plasma, Serum, Urine, Milk, Saliva, and Cell Culture samples)
- ECR3005-1 AssayMax Canine RBP4 ELISA Kit (Plasma, Serum, Urine, and Cell Culture samples)
- EMR3005-1 AssayMax Mouse RBP4 ELISA Kit (Plasma, Serum, Urine, and Cell Culture samples)