

**KAMIYA BIOMEDICAL COMPANY**

# Rabbit Transferrin ELISA

**For the quantitative determination of Transferrin  
in rabbit serum or plasma**

**Cat. No. KT-351**

**For Research Use Only.**

## PRODUCT INFORMATION

### Rabbit Transferrin ELISA Cat. No. KT-351

#### INTENDED USE

The Rabbit Transferrin ELISA is a highly sensitive two-site enzyme-linked immunoassay (ELISA) for the quantitative determination of Transferrin in rabbit serum or plasma. For research use only.

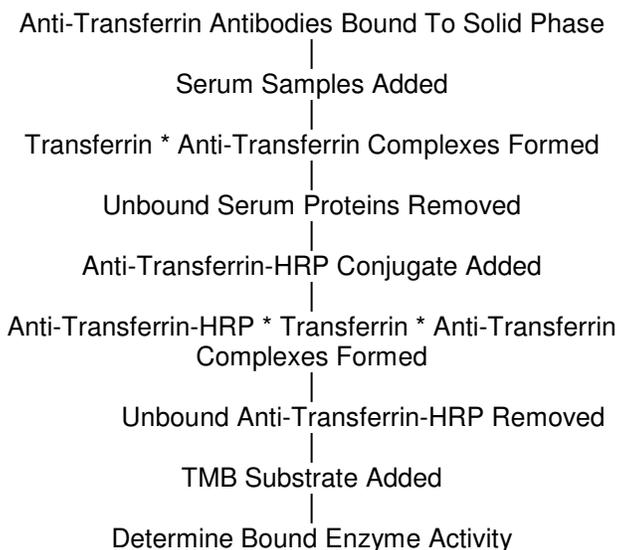
#### INTRODUCTION

Transferrin (TX) is a metal-combining protein that reversibly binds to acid-soluble iron in plasma. It functions to transport iron to the bone marrow and to tissue storage organs such as the liver. Transferrin also participates in the regulation and control of iron absorption and protects against iron intoxication. Like haptoglobin, the carrier of hemoglobin, transferrin is synthesized in the liver, but unlike haptoglobin, transferrin is returned to the circulation after unloading its iron in the reticuloendothelial system. This ELISA can be used to measure transferrin in serum, tissue extracts and other biological fluids.

#### PRINCIPLE

The principle of the double antibody sandwich ELISA is represented in Figure 1. In this assay the transferrin present in serum sample reacts with the anti-transferrin antibodies, which have been adsorbed to the surface of polystyrene microtiter wells. After the removal of unbound serum proteins by washing, anti-transferrin antibodies conjugated with horseradish peroxidase (HRP), are added. This HRP-conjugated antibody forms a complex with the previously bound serum transferrin. Following another washing step, the enzyme bound to the immunosorbent is assayed by the addition of a chromogenic substrate, 3,3',5,5'-tetramethylbenzidine (TMB). The quantity of bound enzyme is proportional to the concentration of transferrin in the sample tested; thus, the absorbance, at 450 nm, is a measure of the concentration of transferrin in the test sample. The quantity of transferrin in the test sample can be interpolated from the calibration curve constructed from the calibrators, and corrected for serum dilution.

Figure 1.



#### COMPONENTS

1. Diluent Concentrate

One bottle containing 50 mL of a 5X concentrated phosphate buffered saline (PBS) solution containing 0.25% Tween, protein stabilizer and 0.25% Proclin 300 as a preservative.

2. Wash Solution Concentrate

One bottle containing 50 mL of a 20X concentrated PBS solution with 1% Tween.

3. Enzyme-Antibody Conjugate Concentrate

One vial containing 200  $\mu$ L of a 100X concentrated affinity-purified anti-rabbit transferrin antibody conjugated with HRP in a stabilizing buffer.

4. TMB Substrate Solution

One vial containing 12 mL of TMB and hydrogen peroxide in citric acid buffer at pH 3.3.

5. Stop Solution

One vial containing 12 mL of 0.3 M sulfuric acid. WARNING: Avoid contact with skin.

6. Microtiter Plate

Twelve removable eight-well strips in well holder frame. Wells are coated with affinity-purified anti-rabbit transferrin.

7. Rabbit Transferrin Calibrator

One vial containing a lyophilized Rabbit Transferrin Calibrator.

## MATERIALS REQUIRED BUT NOT PROVIDED

- Test tubes
- Precision pipette (2  $\mu$ L to 200  $\mu$ L)
- Microplate washer/aspirator
- Distilled or de-ionized H<sub>2</sub>O
- Microplate reader
- Assorted glassware for the preparation of reagents and buffer solutions
- Timer
- Vortex mixer

## PRECAUTIONS

1. Read the instructions carefully before beginning the assay.
2. This kit is for research use only.
3. Great care has been taken to ensure the quality and reliability of this product. However, it is possible that in certain cases, unusual results may be obtained due to high levels of interfering factors.
4. Preservatives
  - Diluent contains 0.25% Proclin 300 as a preservative.
5. No additives or preservatives are necessary to maintain the integrity of the specimen. Avoid azide contamination.
6. Azide and thimerosal at concentrations higher than 0.1% inhibit the enzyme reaction.
7. Other precautions:
  - Do not interchange kit components from different lots.
  - Do not use kit components beyond the expiration date.
  - Protect reagents from direct sunlight.
  - Do not pipette by mouth.
  - Do not eat, drink, smoke or apply cosmetics where reagents are used.
  - Avoid all contact with the reagents by using gloves.
  - Stop solution contains diluted sulfuric acid. Irritation to eyes and skin is possible. Flush with water after contact.

## REAGENT PREPARATION

1. Diluent Concentrate

The Diluent solution supplied is a 5X concentrate and must be diluted 1:5 with distilled or de-ionized water.

2. Wash Solution Concentrate

The Wash Solution supplied is a 20X concentrate and must be diluted 1:20 with distilled or de-ionized water. Crystal formation in the concentrate is not uncommon when storage temperatures are low. Warming of the concentrate to 30-35°C before dilution can dissolve crystals.

3. Enzyme-Antibody Conjugate Concentrate

The Enzyme-Antibody Conjugate supplied is a 100X concentrate and must be diluted 1:100. The required amount of working conjugate solution for each microtiter plate is prepared by adding 100  $\mu$ L Enzyme-Antibody Conjugate to 10 mL of 1X Diluent. Mix uniformly, but gently. Avoid foaming.

4. TMB Substrate Solution  
Ready to use as supplied.
5. Stop Solution  
Ready to use as supplied.
6. Microtiter Plate  
Ready to use as supplied.

7. Rabbit Transferrin Calibrator

The Rabbit Transferrin Calibrator stock solution is 2.0 mL at 145 ng/mL. Rabbit Transferrin Calibrators need to be prepared immediately prior to use (see chart below). Mix well between each step. Avoid foaming. For samples containing lower levels of Transferrin, it is possible to extend the utility of the lower detection limit of this assay by making further serial 2-fold dilution of calibrator 5.

Calibrator	Concentration (ng/mL)	Calibrator Volume added to 1X Diluent	Volume of 1X Diluent
0	1450	10 $\mu$ L Rabbit Transferrin Calibrator	990 $\mu$ L
1	100	40 $\mu$ L Calibrator 0	540 $\mu$ L
2	50	0.3 mL Calibrator 1	0.3 mL
3	25	0.3 mL Calibrator 2	0.3 mL
4	12.5	0.3 mL Calibrator 3	0.3 mL
5	6.25	0.3 mL Calibrator 4	0.3 mL
6	3.125	0.3 mL Calibrator 5	0.3 mL

## STORAGE AND STABILITY

1. Complete Kit

The expiration date for the kit is stated on the outer label. The recommended storage temperature is 4°C. **Note: See long-term storage recommendations below for the Rabbit Transferrin Calibrator.**

2. Diluent Concentrate

The 5X Diluent Concentrate is stable until the expiration date. The 1X working solution is stable for at least one week from the date of preparation. Both solutions should be stored at 4°C.

3. Wash Solution Concentrate

The 20X Wash Solution Concentrate is stable until the expiration date. The 1X working solution is stable for at least one week from the date of preparation. Both solutions can be stored at room temperature (RT, 16-25°C) or at 4°C.

4. Enzyme-Antibody Conjugate Concentrate

Undiluted anti-Transferrin-HRP conjugate should be stored at 4°C and diluted immediately prior to use. The working conjugate solution is stable for one day at 4°C.

5. TMB Substrate Solution

The TMB Substrate Solution should be stored at 4°C and is stable until the expiration date.

6. Stop Solution

The Stop Solution should be stored at 4°C and is stable until the expiration date.

7. Microtiter Plate

Anti-rabbit transferrin coated wells are stable until the expiration date and should be stored at 4°C in the sealed foil pouch with a desiccant pack.

8. Rabbit Transferrin Calibrator

The lyophilized Rabbit Transferrin Calibrator should be stored at 4°C or frozen until the expiration date. The reconstituted calibrator should be aliquoted and stored frozen. Avoid multiple freeze/thaw cycles. The Rabbit Transferrin Calibrator should be stable at 4°C for up to 7 days. The working calibrator solutions should be prepared immediately prior to use and are stable for one day.

## INDICATIONS OF INSTABILITY

If the test is performing correctly, the results observed with the calibrator solutions should be within 20% of the expected values.

## SPECIMEN COLLECTION AND HANDLING

Blood should be collected by venipuncture and the serum separated from the cells, after clot formation, by centrifugation. Specimens may be shipped at room temperature (RT) and then stored refrigerated at 4°C if testing is to take place within one week after collection. If testing is to take place later than one week, specimens should be stored at -20°C. Avoid repeated freezing/thawing.

For any sample that might contain pathogens, care must be taken to prevent contact with open wounds. No additives or preservatives are necessary to maintain the integrity of the specimen. Avoid azide contamination.

## ASSAY PROTOCOL

### Dilution of Serum or Plasma Samples

Due to the high-sensitive nature of the assay, each serum or plasma sample should be diluted before use for a normal assay. A dilution of serum/plasma at 1:50,000 is appropriate for most samples. For absolute quantification of samples that yield results outside the range of the calibration curve a lesser or greater dilution might be required.

To prepare a 1:50,000 dilution of sample, transfer 5 µL of sample to 495 µL of 1X Diluent. This gives you a 1:100 dilution. Next, dilute the 1:100 sample by transferring 2 µL, to 998 µL of 1X Diluent. You now have a 1:50,000 dilution of your sample. Mix thoroughly at each stage.

### Procedure

Bring all reagents to RT before use.

1. Add 100 µL of 1X Diluent to each of the wells in A1 & A2. These will serve for an evaluation of the background associated with the assay.
2. Pipette 100 µL of
  - Calibrator 1 (100 ng/mL) into wells B1 & B2
  - Calibrator 2 (50 ng/mL) into wells C1 & C2
  - Calibrator 3 (25 ng/mL) into wells D1 & D2
  - Calibrator 4 (12.5 ng/mL) into wells E1 & E2
  - Calibrator 5 (6.25 ng/mL) into wells F1 & F2
  - Calibrator 6 (3.13 ng/mL) into wells G1 & G2
3. Pipette 100 µL of diluted serum/plasma sample (test sample 1) into wells H1 & H2. The next sample goes in wells A3 & A4, the next in B3 & B4 and so on.
4. Incubate the Microtiter Plate at 22°C (RT) for sixty (60 ± 2) minutes. Keep plate level during incubation.
5. Following incubation, aspirate the contents of the wells.
6. Completely fill each well with appropriately diluted Wash Solution and aspirate. Repeat three times, for a total of four washes. If washing manually: completely fill wells with diluted Wash Solution, invert the plate and pour/shake out the contents in a waste container. Follow this by sharply striking the wells on absorbent paper to remove residual Wash Solution. Repeat three times for a total of four washes.
7. Pipette 100 µL of appropriately diluted Enzyme-Antibody Conjugate to each well. Incubate at 22°C (RT) for ten (10 ± 2) minutes.
8. Wash and blot the wells as described in Steps 5 and 6.
9. Pipette 100 µL of TMB Substrate Solution into each well.
10. Incubate at RT for precisely ten (10) minutes.
11. After ten (10) minutes, add 100 µL of Stop Solution to each well.

12. Determine the absorbance at 450 nm of the contents of each well. Zero the plate reader to air.

The absorbance of the final reaction mixture can be measured up to two hours after the addition of the Stop Solution. However, good laboratory practice dictates that the measurement be made as soon as possible.

## RESULTS

1. Subtract the average background value from the test values for each sample.
2. Using the results observed for the calibrators construct a calibration curve. The appropriate curve fit is that of a four-parameter logistics curve. A second order polynomial (quadratic) or other curve fits may also be used.
3. Interpolate test sample values from the calibration curve. Correct for serum dilution factor to arrive at transferrin concentration in original sample.

## QUALITY CONTROL

In accord with good laboratory practice, the assays for specific transferrin require meticulous quality control. Each laboratory should use routine quality control procedures to establish inter- and intra-assay precision and performance characteristics.

## LIMITATION OF THE PROCEDURE

1. Reliable and reproducible results will be obtained when the assay procedure is carried out with a complete understanding of the information contained in the package insert instructions and with adherence to good laboratory practice.
2. Factors that might affect the performance of the assay include proper instrument function, cleanliness of glassware, quality of distilled or de-ionized water, washing thoroughly and accuracy of reagent and sample pipetting.

## **FOR RESEARCH USE ONLY**

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