**K-ASSAY Transferrin**

For the Quantitative Determination of Human Transferrin in Serum  

**Cat. No. KAI-023**

**INTENDED USE**

The K-ASSAY Transferrin assay is intended for the quantitative determination of human transferrin in human serum by immunoturbidimetric assay. For IN VITRO DIAGNOSTIC USE.

**SUMMARY**

The K-ASSAY Transferrin assay is intended for the quantitative determination of human transferrin by immunoturbidimetric assay. The antiserum used in the kit was produced against purified human transferrin. The antiserum used in the kit is a goat polyclonal antibody specific for transferrin antibody interacts with the transferrin in the serum forming immune complexes. The immune complexes cause an increase in light scattering, which correlates with the concentration of serum transferrin.

Transferrin has been measured using a variety of methods, including radial immunodiffusion and immunoassay. The K-ASSAY Transferrin assay uses an immunoturbidimetric assay format.

**PRINCIPLE OF TEST**

The K-ASSAY Transferrin assay quantifies the transferrin in the patient's serum based on immunoturbidimetric assay. Calibrators, controls, and patient samples are pipetted into sample cups. Microvolumes of samples and antibody diluent are automatically pipetted into individual cuvettes. Following an initial incubation and measurement of sample blank, antiserum is added to the cuvettes. The samples (antigen) and antiserum are then mixed in the reaction cuvettes. Insoluble antigen-antibody (immune) complexes form. The immune complexes cause an increase in light scattering, which correlates with the concentration of serum transferrin. Following an incubation period lasting approximately 5 min., the absorbance of the solution is measured at 700 nm.

A calibration curve is generated by assaying a series of calibrators with known concentrations of proteins and using the instrument's data reduction capability or manually plotting the change in absorbance versus concentration.

Concentration of the control and patient samples is interpolated from the calibration curve. The antiserum used in the kit is a goat polyclonal antibody specific for human transferrin.

The K-ASSAY Transferrin assay should be run using the K-ASSAY Multi-Analyte Calibrator. Six calibrators are used to prepare a calibration curve for quantifying the levels of transferrin present in the patient's serum sample.

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**KIT COMPOSITION**

**Reagents (Liquid Stable)**

R1: Buffer Reagent 3 x 20 mL
Tris(hydroxymethyl)aminomethane
R2: Antiserum Reagent 2 x 10 mL
Anti-human transferrin goat antiserum (40%)

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**WARNINGS AND PRECAUTIONS**

FOR IN VITRO DIAGNOSTIC USE. Rx only.

Not to be used internally in humans or animals. Normal precautions exercised in handling laboratory reagents should be followed.

Do not mix or use reagents from one test kit with those from a different lot number.

Do not use reagents past their expiration date stated on each reagent container label.

Do not pipette by mouth. Avoid ingestion and contact with skin.

Reagents in this kit contain sodium azide as a preservative. Sodium azide may form explosive compounds in metal drain lines. When disposing of azide salts,” in the Manual Guide-Safety Guide Management No. CDC-22 issued by the Center for Disease Control, Atlanta, Georgia.

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**warnings:** This product can expose you to chemicals including thiourea which is known to the State of California to cause cancer/birth defects or other reproductive harm. For more information go to www.P65Warnings.ca.gov. The R-1 reagent contains 0.009% of thiourea (CAS No. 62-56-6).

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**REAGENT STABILITY**

Opened reagents can be used for one month if stored at 2-8°C. Discard reagents if they become contaminated. Evidence of cloudiness or particulate material in solution is cause to discard.

**INSTRUMENT**

Measurement of absorbance is to be made with an instrument able to accurately read absorbance at 700 nm. Refer to the instrument manual from the manufacturer regarding the following:

- Use or function
- Installation procedures and requirements
- Principles of operation
- Performance characteristics, operating instructions
- Calibration procedures including materials and / or equipment to be used
- Operational precautions, limitations, and hazards
- Service and maintenance information

**SPECIMEN COLLECTION AND PREPARATION**

It is recommended that specimen collection be carried out in accordance with NCCLS document M29-T2. No method can offer complete assurance that human blood samples will not transmit infection. Therefore, all blood samples should be considered potentially infectious.

Serum is required for this assay. Soon after the blood is drawn, it should be allowed to clot, centrifuged, and the serum separated from the clot to a plastic tube (not glass) within 2 hours. Serum should be stored refrigerated (2-8°C) and can be used within one week or should be stored frozen for up to 2 months.

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**USE PLASTIC TUBES FOR STORING THE SAMPLE, DO NOT USE GLASS.**

**AUTOMATED ANALYZER APPLICATION**

Suitable for two-reagent automated analyzers that use a multi-point calibration method.

**PROCEDURE**

**Materials Supplied**

Reagent 1 (R-1) Buffer Reagent 3 x 20 mL
Reagent 2 (R-2) Antiserum Reagent 2 x 10 mL

**Materials Required But Not Supplied**

Calibrators: K-ASSAY Multi-Analyte Calibrator, Cat. No. KAI-016C. (6 calibrators containing human serum with known levels of transferrin).

Two Reagent Clinical Chemistry Analyzer:

- Capable of accurately dispensing the required volumes
- Capable of maintaining 37°C

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**ASSAY PROCEDURE**

Note: Allow all reagents and specimens to warm to room temperature. Mix all reagents gently before using.

An example of automated application (Hitachi 717):

<table>
<thead>
<tr>
<th>Sample</th>
<th>3 μL</th>
</tr>
</thead>
<tbody>
<tr>
<td>R1 (Buffer Reagent)</td>
<td>300 μL</td>
</tr>
<tr>
<td>R2 (Antiserum Reagent)</td>
<td>100 μL</td>
</tr>
</tbody>
</table>

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**Automated Method (Example)**

Chemistry Parameters for Automatic Analyzers

**INSTRUMENT**

Roche / Hitachi 717

**TEMPERATURE**

37°C

**TEST**

(TF)

**ASSAY CODE**

(2 POINT) : (24) - (50)

**SAMPLE VOLUME**

(3) ( )

**R-1 VOLUME**

(300) ( ) (NO)

**R-2 VOLUME**

(100) ( ) (NO)

**WAVELENGTH**

(700) ( )

**CALIB. METHOD**

(NONLINEAR) (1) (6)

**STD.(1) Conc.-POS.**

(1) ( - )

**STD.(2) Conc.-POS.**

(2) ( - )

**STD.(3) Conc.-POS.**

(3) ( - )

**STD.(4) Conc.-POS.**

(4) ( - )

**STD.(5) Conc.-POS.**

(5) ( - )

**STD.(6) Conc.-POS.**

(6) ( - )

**SD LIMIT**

(999)

**DUPLICATE LIMIT**

(10000)

**SENSITIVITY LIMIT**

(0)

**ABS. LIMIT (SLOPE)**

(32000) ( INCREASE )

**PROZONE LIMIT**

(320000) ( LOWER )

**EXPECTED VALUE**

(-99999) (99999)

**PANIC VALUE**

(-99999) (99999)

**INSTRUMENT FACTOR**

(1.00)

*1-6: Input concentration of calibrators.

Parameters for other automated analyzers are available.

**CALIBRATION**

It is recommended that transferrin levels be determined using a multi-point calibration curve prepared using the K-ASSAY Multi-Analyte Calibrator. It is recommended that the user determine calibration curve frequency as this depends on the instrument and type/number of other assays being performed. Initially, calibration should be performed each day.

**QUALITY CONTROL**

A quality control program is recommended for all clinical testing laboratories. It is recommended that control sera, both normal and abnormal, be run with each batch of samples to monitor the procedure.
LIMITATIONS OF PROCEDURE

The measurable range for transferrin is between 45 to 455 mg/dL. Grossly lipemic samples and samples with very high triglyceride concentrations should be diluted 1 part sample with 1 part isotonic saline or filtered to decrease nonspecific light scattering. Multiply results by 2 to compensate for the dilution.

If the transferrin concentration of a patient sample is greater than the highest calibrator, dilute 1 part sample with 4 parts isotonic saline and reassay. Multiply results by 5 to compensate for the dilution.

PERFORMANCE

Precision

The precision for the K-ASSAY® Transferrin assay was determined using packaged reagents, pooled human serum, and a Hitachi 717 analyzer.

Precision Assay: Within Run

<table>
<thead>
<tr>
<th>Sample I</th>
<th>Sample II</th>
<th>Sample III</th>
</tr>
</thead>
<tbody>
<tr>
<td>N = 20</td>
<td>N = 20</td>
<td>N = 20</td>
</tr>
<tr>
<td>Mean = 70.7</td>
<td>Mean = 300.75</td>
<td>Mean = 526.4</td>
</tr>
<tr>
<td>SD = 0.92</td>
<td>SD = 5.02</td>
<td>SD = 5.77</td>
</tr>
<tr>
<td>CV = 1.3%</td>
<td>CV = 1.67%</td>
<td>CV = 1.097%</td>
</tr>
</tbody>
</table>

Precision Assay: Between Runs

<table>
<thead>
<tr>
<th>Sample I</th>
<th>Sample II</th>
</tr>
</thead>
<tbody>
<tr>
<td>N = 8</td>
<td>N = 8</td>
</tr>
<tr>
<td>Mean = 175.38</td>
<td>Mean = 340.9</td>
</tr>
<tr>
<td>SD = 1.6</td>
<td>SD = 1.46</td>
</tr>
<tr>
<td>CV = 0.91%</td>
<td>CV = 0.4%</td>
</tr>
</tbody>
</table>

Accuracy / Correlation

A comparison of the K-ASSAY® Transferrin assay and an Incstar Transferrin Test Kit was performed using a Hitachi 717. The test results provided the following data:

\[ y = 1.064x - 50.124 \]
\[ r = 0.982 \]
\[ n = 60 \]
\[ x = \text{Incstar Transferrin Test Kit} \]
\[ y = \text{K-ASSAY® Transferrin assay} \]

<table>
<thead>
<tr>
<th>x min</th>
<th>y min</th>
<th>x max</th>
<th>y max</th>
<th>mean</th>
<th>x mean</th>
<th>y mean</th>
</tr>
</thead>
<tbody>
<tr>
<td>148</td>
<td>110</td>
<td>440</td>
<td>426</td>
<td>246</td>
<td>212</td>
<td></td>
</tr>
</tbody>
</table>

Assay Range

45 - 455 mg/dL

INTERFERENCE

<table>
<thead>
<tr>
<th>Ascorbic Acid</th>
<th>No interference up to 50 mg/dL</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bilirubin C</td>
<td>No interference up to 20 mg/dL</td>
</tr>
<tr>
<td>Bilirubin F</td>
<td>No interference up to 20 mg/dL</td>
</tr>
<tr>
<td>Hemoglobin</td>
<td>No interference up to 500 mg/dL</td>
</tr>
<tr>
<td>Intralipid</td>
<td>No interference up to 500 mg/dL</td>
</tr>
</tbody>
</table>

EXPECTED VALUE

The expected value as reported is between 260 to 400 mg/dL. Each laboratory should establish its own expected values using this kit.

REFERENCES


LABELING SYMBOLS

<table>
<thead>
<tr>
<th>LOT</th>
<th>Reagent</th>
</tr>
</thead>
<tbody>
<tr>
<td>RGT</td>
<td>Expiration or “Use By” Date</td>
</tr>
<tr>
<td>REF</td>
<td>Catalog Number</td>
</tr>
<tr>
<td>IVD</td>
<td>For In Vitro Diagnostics Use</td>
</tr>
<tr>
<td>2-8 °C</td>
<td>Temperature Limitation, Store between 2 and 8 degrees C</td>
</tr>
<tr>
<td>Manufacturer</td>
<td></td>
</tr>
<tr>
<td>Authorized Representative for In Vitro Diagnostics Use</td>
<td></td>
</tr>
</tbody>
</table>

EU AUTHORIZED REPRESENTATIVE

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