Insulin

For the Quantitative Determination of Human Insulin in Serum or Plasma

Cat. No. KAI-071

INTENDED USE
For the quantitative determination of human insulin in serum and plasma by immunoturbidimetric assay. FOR IN VITRO DIAGNOSTIC USE.

SUMMARY
Insulin is a peptide hormone with approximate molecular weight of 5,800 daltons. Secreted from β cells of the islet of Langerhans in the pancreas, insulin acts to reduce the blood sugar level. Since the blood insulin level reflects the function of β cells, insulin has been widely used as an important diagnostic tool for diabetes mellitus. The K-ASSAY® Insulin test is a highly specific assay for insulin in serum or plasma.

PRINCIPLE OF TEST
Latex particles coated with antibody specific to human insulin form immune complexes in the presence of insulin from the sample. The immune complexes cause an increase in light scattering that is proportional to the concentration of insulin in the serum or plasma sample. The light scattering is measured by reading turbidity at 600 nm primary, 800 nm secondary. The sample insulin concentration is determined versus insulin calibrators of known concentrations.

KIT COMPOSITION
Reagents (Liquid Stable)
R1: Buffer Reagent, pH 8.2
Tris(hydroxymethyl)aminomethane (100 mM)
R2: Latex Suspension
Anti-human insulin mouse monoclonal antibody (~1 mg/mL)

WARNINGS AND PRECAUTIONS
For In Vitro Diagnostic Use Only. It 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.

SPECIMEN COLLECTION AND PREPARATION
Serum
Blood should be collected from a patient and the serum separated as soon as possible. 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. It is recommended that specimen collection be carried out in accordance with NCCCLS document M29-A2.
Avoid repeated freeze/thaw cycles.

Plasma
Whole blood is collected in sodium citrate, sodium EDTA and sodium fluoride anticoagulant. After collection, immediately centrifuge. In the U.S., follow NCCCLS guideline H-3-A2. Avoid repeated freeze/thaw cycles.
Serum or plasma may be stored refrigerated (2-8°C) for up to a week. For long-term storage, keep at -20°C or below.

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 2 x 13.5 mL
Reagent 2 (R-2) Latex Suspension 2 x 5 mL

Materials Required But Not Supplied
Calibrators: K-ASSAY® Insulin Calibrator, Cat. No. KAI-072C
Purified water
Clinical chemistry analyzer capable of accurately reading at 600 nm (main) and 800 nm (sub), accurately dispensing the required volumes, and maintaining 37°C.

Assay Procedure
Note: Allow all reagents and specimens to warm to room temperature. Mix all reagents gently before using.
An example of automated application:
Sample 12 µL
- 1.5 x R1 (Buffer Reagent) 135 µL 37 °C, 5 min.
- 2 x R2 (Latex Suspension) 50 µL 37 °C, 5 min.
2 Point End at 600nm main, 800nm sub (if available)

Automated Method (Example)

<table>
<thead>
<tr>
<th>INSTRUMENT</th>
<th>Roche / Hitachi 917</th>
</tr>
</thead>
<tbody>
<tr>
<td>TEMPERATURE</td>
<td>37°C</td>
</tr>
<tr>
<td>TEST</td>
<td>( INS )</td>
</tr>
<tr>
<td>ASSAY CODE</td>
<td>(2 POINT END</td>
</tr>
<tr>
<td>WAVELENGTH</td>
<td>(19)</td>
</tr>
<tr>
<td>SAMPLE VOLUME, µL</td>
<td>(12.0)</td>
</tr>
<tr>
<td>REAGENT VOL (R1)</td>
<td>(135)</td>
</tr>
<tr>
<td>REAGENT VOL (R2)</td>
<td>(0)</td>
</tr>
<tr>
<td>REAGENT VOL (R3)</td>
<td>(50)</td>
</tr>
<tr>
<td>REAGENT VOL (R4)</td>
<td>(0)</td>
</tr>
<tr>
<td>ABS. LIMIT (SLOPE)</td>
<td>(3200)</td>
</tr>
<tr>
<td>PROZONE LIMIT</td>
<td>(-32000)</td>
</tr>
<tr>
<td>CALIB. TYPE</td>
<td>(SPLINE)</td>
</tr>
<tr>
<td>POINT</td>
<td>(6)</td>
</tr>
<tr>
<td>SPAN POINT</td>
<td>(6)</td>
</tr>
<tr>
<td>SD LIMIT</td>
<td>(999)</td>
</tr>
<tr>
<td>DUPLICATE LIMIT</td>
<td>10000</td>
</tr>
<tr>
<td>SENSITIVITY LIMIT</td>
<td>(0)</td>
</tr>
<tr>
<td>S1ABS RANGE</td>
<td>(3200)</td>
</tr>
<tr>
<td>INSTRUMENT FACTOR</td>
<td>a=1.0</td>
</tr>
<tr>
<td>b=0.0</td>
<td></td>
</tr>
<tr>
<td>UNIT</td>
<td>(µIU/mL)</td>
</tr>
<tr>
<td>STD.(1) Conc.-POS.</td>
<td>0.0</td>
</tr>
<tr>
<td>STD.(2) Conc.-POS.</td>
<td>2.0</td>
</tr>
<tr>
<td>STD.(3) Conc.-POS.</td>
<td>3.0</td>
</tr>
<tr>
<td>STD.(4) Conc.-POS.</td>
<td>4.0</td>
</tr>
<tr>
<td>STD.(5) Conc.-POS.</td>
<td>5.0</td>
</tr>
<tr>
<td>STD.(6) Conc.-POS.</td>
<td>6.0</td>
</tr>
</tbody>
</table>

Use DI water for standard 1. Use Insulin calibrators 1-5 for standards 2-6.

2.6: Initial concentration of calibrators (using one decimal place [KX]).

# = User Defined

Parameters for other automated analyzers are available.

CALIBRATION
It is recommended that a multi-point calibration curve be made using the K-ASSAY® Insulin Calibrator (KAI-072C). 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
Normal and abnormal controls of known concentration should be included in each assay performed. These controls should fall within the ranges established by each lab for the particular lot of controls. The validity of the assay is in question if the value for the controls generated by the assay’s calibration curve does not fall within the stated range. Recalibrate if the value determined for the controls falls outside the established recovery range.
LIMITATIONS OF PROCEDURE

The measurable range for insulin is 1.0 to 100 μU/mL. If the insulin value of a sample is greater than highest calibrator value, dilute 1 part sample with 3 parts isotonic saline and re-assay. Multiply results by 4 to compensate for the dilution.

PERFORMANCE

Specificity

When a sample with a known value is assayed, the result is within ± 15% of the assigned value.

Precision

When a sample is assayed 5 times (within-run), the absorbance C.V. is ≤ 10%. (Within Run)
The following results were obtained on a Hitachi 917 analyzer with human serum:

<table>
<thead>
<tr>
<th>Sample</th>
<th>Sample II</th>
<th>Sample III</th>
</tr>
</thead>
<tbody>
<tr>
<td>N</td>
<td>10</td>
<td>10</td>
</tr>
<tr>
<td>Mean (μU/mL)</td>
<td>18.81</td>
<td>27.82</td>
</tr>
<tr>
<td>Std. Dev.</td>
<td>0.224</td>
<td>0.192</td>
</tr>
<tr>
<td>C.V. %</td>
<td>1.19</td>
<td>0.69</td>
</tr>
</tbody>
</table>

(Between Runs)
The following results were obtained on a Hitachi 917 analyzer with human serum:

<table>
<thead>
<tr>
<th>Sample IV</th>
<th>Sample V</th>
<th>Sample VI</th>
</tr>
</thead>
<tbody>
<tr>
<td>N</td>
<td>10</td>
<td>10</td>
</tr>
<tr>
<td>Mean (μU/mL)</td>
<td>12.21</td>
<td>25.90</td>
</tr>
<tr>
<td>Std. Dev.</td>
<td>0.530</td>
<td>0.777</td>
</tr>
<tr>
<td>C.V. %</td>
<td>4.34</td>
<td>3.00</td>
</tr>
</tbody>
</table>

Accuracy / Correlation

A comparison of the K-ASSAY® Insulin and another company’s Insulin EIA was performed with the following results:

Serum Sample

y = 0.8061x + 2.8674
r = 0.997
n = 32
x = another company’s Insulin assay
y = K-ASSAY® Insulin Assay

Plasma Sample

y = 0.8161x + 4.5552
r = 0.986
n = 47
x = another company’s Insulin assay
y = K-ASSAY® Insulin Assay

Assay Range

1.0 to 100 μU/mL (or value of highest calibration point)

Lower Limit of Detection

The analytical sensitivity is 1 μU/mL. This means that when saline and serum containing 1 μU/mL of insulin are tested 10 times, ± 2.6 SD of the respective results do not overlap each other.

INTERFERENCE

No cross-reactivity with pro-insulin was observed. Hemoglobin, bile or rheumatoid factor did not interfere with the assay.

Bilirubin, Conjugated  No interference up to 19.9 mg/dL
Bilirubin, Unconjugated No interference up to 19.3 mg/dL
Hemoglobin         No interference up to 450 mg/dL
Lipemia            No interference up to a formazin turbidity of 1,550
Rheumatoid Factor  No interference up to 4500 IU/L

PROZONE

No hook effect seen up to at least 1,000 μU/mL.

EXPECTED VALUES

The expected range for fasting insulin concentration has been reported to be up to 20 to 35 μU/mL (RIA).¹

REFERENCES


LABELING SYMBOLS

- Catalog Number
- Expiration or “Use By” Date
- Lot Number
- Consult Package Insert for Instructions for Use
- For In Vitro Diagnostic Use
- CE Mark Registered
- For Prescription Use Only
- Temperature Limitation. Store between 2 and 8 degrees C
- Manufacturer
- Authorized Representative in the European Community

EU AUTHORIZED REPRESENTATIVE

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