Instructions for use

Kynurenine ELISA
Kynurenine ELISA

1. Introduction

1.1 Intended use and principle of the test

Enzyme Immunoassay for the quantitative determination of L-Kynurenine in serum and plasma.

After acylation Kynurenine is quantitatively determined by ELISA. The competitive ELISA uses the microtiter plate format. The antigen is bound to the solid phase of the microtiter plate. The acylated standards, controls and samples and the solid phase bound analyte compete for a fixed number of antibody binding sites. When the system is in equilibrium, free antigen and free antigen-antibody complexes are removed by washing. The antibody bound to the solid phase is detected by an anti-rabbit IgG-peroxidase conjugate using TMB as a substrate. The reaction is monitored at 450 nm. Quantification of unknown samples is achieved by comparing their absorbance with a reference curve prepared with known standards.

2. Procedural cautions, guidelines, warnings and limitations

2.1 Procedural cautions, guidelines and warnings

(1) This kit is intended for professional use only. Users should have a thorough understanding of this protocol for the successful use of this kit. Only the test instruction provided with the kit is valid and has to be used to run the assay. Reliable performance will only be attained by strict and careful adherence to the instructions provided.

(2) This assay was validated for a certain type of sample as indicated in Intended Use (please refer to Chapter 1). Any off-label use of this kit is in the responsibility of the user and the manufacturer cannot be held liable.

(3) Reagents of this kit which contain human serum or plasma have been tested and confirmed negative for HIV I/II, HBsAg and HCV by approved procedures. All reagents, however, should be treated as potential biohazards in use and for disposal.

(4) The principles of Good Laboratory Practice (GLP) have to be followed.

(5) In order to reduce exposure to potentially harmful substances, wear lab coats, disposable protective gloves and protective glasses whenever necessary.

(6) All kit reagents and specimens should be brought to room temperature and mixed gently but thoroughly before use. Avoid repeated freezing and thawing of reagents and specimens.

(7) For dilution or reconstitution purposes, use deionized, distilled, or ultra-pure water.

(8) The microplate contains snap-off strips. Unused wells must be stored at 2 °C to 8 °C in the sealed foil pouch with desiccant and used in the frame provided.

(9) Duplicate determination of sample is highly recommended to be able to identify potential pipetting errors.

(10) Once the test has been started, all steps should be completed without interruption. Make sure that the required reagents, materials and devices are prepared ready at the appropriate time.

(11) Incubation times do influence the results. All wells should be handled in the same order and time intervals.

(12) To avoid cross-contamination of reagents, use new disposable pipette tips for dispensing each reagent, sample, standard and control.

(13) A standard curve must be established for each run.

(14) The controls should be included in each run and fall within established confidence limits. The confidence limits are listed in the QC-Report.

(15) Do not mix kit components with different lot numbers within a test and do not use reagents beyond expiry date as shown on the kit labels.

(16) Avoid contact with Stop Solution containing 0.25 M H₂SO₄. It may cause skin irritation and burns. In case of contact with eyes or skin, rinse off immediately with water.

(17) Some reagents contain sodium azide (NaN₃) as preservatives. In case of contact with eyes or skin, rinse off immediately with water. NaN₃ may react with lead and copper plumbing to form explosive metal azides. When disposing reagents, flush with a large volume of water to avoid azide build-up.

(18) TMB substrate has an irritant effect on skin and mucosa. In case of possible contact, wash eyes with an abundant volume of water and skin with soap and abundant water. Wash contaminated objects before reusing them.

(19) For information on hazardous substances included in the kit please refer to Material Safety Data Sheet (MSDS). The Material Safety Data Sheet for this product is made available directly on the website of the manufacturer or upon request.

(20) The expected reference values reported in this test instruction are only indicative. It is recommended that each laboratory establishes its own reference intervals.

(21) The results obtained with this test kit should not be taken as the sole reason for any therapeutic consequence but have to be correlated to other diagnostic tests and clinical observations.

Version: 10.0 Effective: 2014-12-17 2/7
(22) Kit reagents must be regarded as hazardous waste and disposed of according to national regulations.

2.2 Limitations

Any inappropriate handling of samples or modification of this test might influence the results.

2.2.1 Interfering substances

Serum/Plasma

Samples containing precipitates or fibrin strands or which are haemolytic or lipemic might cause inaccurate results.

2.2.2 Drug interferences

There are no known substances (drugs) which ingestion interferes with the measurement of kynurenine level in the sample.

2.2.3 High-Dose-Hook effect

No hook effect was observed in this test.

3. Storage and stability

Store the unopened reagents at 2 - 8 °C until expiration date. Do not use components beyond the expiry date indicated on the kit labels. Once opened the reagents are stable for 1 month when stored at 2 – 8 °C. Once the resealable pouch has been opened, care should be taken to close it tightly with desiccant again.

4. Materials

4.1 Content of the kit

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
<th>Content/Volume</th>
</tr>
</thead>
<tbody>
<tr>
<td>BA D-0033</td>
<td>Macrotiter Plate - Ready to use</td>
<td>2 x 48 well plate, empty</td>
</tr>
<tr>
<td>BA D-0090</td>
<td>Adhesive Foil - Ready to use</td>
<td>1 x 4 foils</td>
</tr>
<tr>
<td>BA E-0030</td>
<td>Wash Buffer Concentrate - Concentrated 50x</td>
<td>Buffer with a non-ionic detergent and physiological pH</td>
</tr>
<tr>
<td>BA E-0040</td>
<td>Enzyme Conjugate - Ready to use</td>
<td>Goat anti-rabbit immunoglobulins conjugated with peroxidase</td>
</tr>
<tr>
<td>BA E-0055</td>
<td>Substrate - Ready to use</td>
<td>Chromogenic substrate containing tetramethylbenzidine, substrate buffer and hydrogen peroxide</td>
</tr>
<tr>
<td>BA E-0080</td>
<td>Stop Solution - Ready to use</td>
<td>0.25 M sulfuric acid</td>
</tr>
<tr>
<td>BA E-2231</td>
<td>Kynurenine Microtiter Strips - Ready to use</td>
<td>1 x 96 well (12x8) antigen precoated microwell plate in a resealable pouch with desiccant</td>
</tr>
<tr>
<td>BA E-2210</td>
<td>Kynurenine Antiserum - Ready to use</td>
<td>Rabbit anti-kynurenine antibody, blue coloured</td>
</tr>
</tbody>
</table>

Version: 10.0   Effective: 2014-12-17
### Standards and Controls - Ready to use

<table>
<thead>
<tr>
<th>Cat. no.</th>
<th>Component</th>
<th>Colour/Cap</th>
<th>Concentration ng/ml</th>
<th>Concentration nmol/l</th>
<th>Volume/Vial</th>
</tr>
</thead>
<tbody>
<tr>
<td>BA E-2201</td>
<td>STANDARD A</td>
<td>white</td>
<td>0</td>
<td>0</td>
<td>4 ml</td>
</tr>
<tr>
<td>BA E-2202</td>
<td>STANDARD B</td>
<td>light yellow</td>
<td>100</td>
<td>480</td>
<td>4 ml</td>
</tr>
<tr>
<td>BA E-2203</td>
<td>STANDARD C</td>
<td>orange</td>
<td>300</td>
<td>1440</td>
<td>4 ml</td>
</tr>
<tr>
<td>BA E-2204</td>
<td>STANDARD D</td>
<td>dark blue</td>
<td>1000</td>
<td>4800</td>
<td>4 ml</td>
</tr>
<tr>
<td>BA E-2205</td>
<td>STANDARD E</td>
<td>light grey</td>
<td>3000</td>
<td>14400</td>
<td>4 ml</td>
</tr>
<tr>
<td>BA E-2206</td>
<td>STANDARD F</td>
<td>black</td>
<td>10000</td>
<td>48000</td>
<td>4 ml</td>
</tr>
<tr>
<td>BA E-2251</td>
<td>CONTROL 1</td>
<td>light green</td>
<td>Refer to QC-Report for expected value and acceptable range!</td>
<td></td>
<td>4 ml</td>
</tr>
<tr>
<td>BA E-2252</td>
<td>CONTROL 2</td>
<td>dark red</td>
<td></td>
<td></td>
<td>4 ml</td>
</tr>
</tbody>
</table>

Conversion: Kynurenine (ng/ml) x 4.80 = Kynurenine (nmol/l)

Content: TRIS buffer with non-mercury stabilizer, spiked with defined quantity of kynurenine

**BA E-2211**

**Acylation Buffer** - Ready to use

Content: 2-(N-Morpholino)ethanesulfonic acid (MES) buffer

Volume: 1 x 55 ml/vial, brown cap

**BA E-2212**

**Acylation Reagent** - Ready to use

Content: acylation reagent in dimethylsulfoxide (DMSO)

Volume: 1 x 6 ml/vial, green cap

### 4.2 Additional materials and equipment required but not provided in the kit

- Calibrated precision pipettes to dispense volumes between 20 – 500 µl
- Microtiter plate washing device (manual, semi-automated or automated)
- ELISA reader capable of reading absorbance at 450 nm and if possible 620 – 650 nm
- Temperature controlled incubator (37 °C) or similar heating device
- Microtiter plate shaker (shaking amplitude 3 mm; approx. 600 rpm)
- Absorbent material (paper towel)
- Water (deionized, distilled, or ultra-pure)
- Vortex mixer

### 5. Sample collection and storage

**Plasma**

Whole blood should be collected into centrifuge tubes containing EDTA as anti-coagulant (Monovette™ or Vacuette™) and centrifuged according to manufacturer’s instruction immediately after collection. Haemolytic and lipemic samples should not be used for the assay.

Storage: up to 48 hours at 2 - 8 °C, for longer period (up to 6 month) at -20 °C.

Repeated freezing and thawing should be avoided.

**Serum**

Collect blood by venipuncture (Monovette™ or Vacuette™ for serum), allow to clot, and separate serum by centrifugation according to manufacturer’s instruction. Do not centrifuge before complete clotting has occurred. Patients receiving anticoagulant therapy may require increased clotting time.

Haemolytic and lipemic samples should not be used for the assay.

Storage: up to 48 hours at 2 - 8 °C, for longer period (up to 6 month) at -20 °C.

Repeated freezing and thawing should be avoided.

### 6. Test procedure

Allow all reagents and samples to reach room temperature and mix thoroughly by gentle inversion before use. Duplicate determinations are recommended.

The binding of the antisera and of the enzyme conjugate and the activity of the enzyme are temperature dependent, and the absorption values may vary if a thermostat is not used. The higher the temperature, the higher the absorption values will be. Varying incubation times will have similar influences on the absorbance. The optimal temperature during the Enzyme Immunoassay is between 20 – 25 °C.
In case of overflow, read the absorbance of the solution in the wells within 10 minutes, using a microplate reader set to 405 nm.

6.1 Preparation of reagents

Wash Buffer
Dilute the 20 ml Wash Buffer Concentrate with water (deionized, distilled, or ultra-pure) to a final volume of 1000 ml.
Storage: 1 month 2 – 8 °C.

Acylation Reagent
The Acylation Reagent has a freezing point of 18.5 °C. To ensure that the Acylation Reagent forms a homogeneous, crystal-free solution when being used, it must have reached room temperature.

6.2 Acylation

1. Pipette 20 µl of the standards, controls and samples into the appropriate wells of the Macrotiter Plate.
2. Add 500 µl of the Acylation Buffer to all wells.
3. Add 50 µl of the Acylation Reagent to all wells and mix shortly.
4. Cover the plate with Adhesive Foil and incubate 90 min at 37 °C.
5. Use 20 µl for the ELISA!

6.3 Kynurenine ELISA

1. Pipette 20 µl of the prepared standards, controls and samples into the appropriate wells of the Kynurenine Microtiter Strips.
2. Pipette 50 µl of the Kynurenine Antiserum into all wells and mix shortly.
3. Cover plate with Adhesive Foil and incubate for 15 - 20 h (overnight) at 2 – 8 °C.
4. Remove the foil. Discard or aspirate the contents of the wells. Wash the plate 4 x by adding 300 µl of Wash Buffer, discarding the content and blotting dry each time by tapping the inverted plate on absorbent material.
5. Pipette 100 µl of the Enzyme Conjugate into all wells.
6. Incubate for 30 min at RT (20 – 25 °C) on a shaker (approx. 600 rpm).
7. Discard or aspirate the contents of the wells. Wash the plate 4 x by adding 300 µl of Wash Buffer, discarding the content and blotting dry each time by tapping the inverted plate on absorbent material.
8. Pipette 100 µl of the Substrate into all wells and incubate for 20 - 30 min at RT (20 – 25 °C) on a shaker (approx. 600 rpm). Avoid exposure to direct sunlight!
9. Add 100 µl of the Stop Solution to each well and shake the microtiter plate to ensure a homogeneous distribution of the solution.
10. Read the absorbance of the solution in the wells within 10 minutes, using a microplate reader set to 450 nm (if available a reference wavelength between 620 nm and 650 nm is recommended).

7. Calculation of results

<table>
<thead>
<tr>
<th>Measuring range</th>
<th>Kynurenine</th>
</tr>
</thead>
<tbody>
<tr>
<td>63.3 – 10 000 ng/ml</td>
<td></td>
</tr>
</tbody>
</table>

The standard curve is obtained by plotting the absorbance readings (calculate the mean absorbance) of the standards (linear, y-axis) against the corresponding standard concentrations (logarithmic, x-axis). Use non-linear regression for curve fitting (e.g. spline, 4- parameter, akima).

The concentrations of the samples and controls can be read directly from the standard curve.

Conversion
Kynurenine (ng/ml) x 4.80 = Kynurenine (nmol/l)
**Expected reference values**

It is strongly recommended that each laboratory should determine its own reference value.

In a study conducted with apparently normal healthy adults, using the Kynurenine ELISA the following value is observed:

<table>
<thead>
<tr>
<th>Plasma / Serum</th>
</tr>
</thead>
<tbody>
<tr>
<td>237.4 – 754.2 ng/ml</td>
</tr>
</tbody>
</table>

### 7.1 Quality control

It is recommended to use control samples according to national regulations. Use controls at both normal and pathological levels. The kit or other commercial controls should fall within established confidence limits. The confidence limits of the kit controls are indicated on the QC-Report.

### 7.2 Typical standard curve

*Example, do not use for calculation!

![Typical Standard Curve](image)

### 8. Assay characteristics

#### Analytical Sensitivity

<table>
<thead>
<tr>
<th>Substance</th>
<th>Cross Reactivity (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>L-Kynurenine</td>
<td>100</td>
</tr>
<tr>
<td>5-Hydroxy-DL-Tryptophan</td>
<td>0.05</td>
</tr>
<tr>
<td>Tyrosin, Phenylalanin, Serotonin, L-Asparagin, Kynurenic Acid</td>
<td></td>
</tr>
</tbody>
</table>

#### Analytical Specificity

<table>
<thead>
<tr>
<th>Substance</th>
<th>Cross Reactivity (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Tryptophan</td>
<td>0.25</td>
</tr>
<tr>
<td>3-Hydroxy-DL-Kynurenin</td>
<td>0.36</td>
</tr>
</tbody>
</table>

#### Precision Serum

<table>
<thead>
<tr>
<th></th>
<th>Intra-Assay</th>
<th>Inter-Assay</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sample</td>
<td>(ng/ml)</td>
<td>SD</td>
</tr>
<tr>
<td>1 (n = 20)</td>
<td>382.3</td>
<td>49.4</td>
</tr>
<tr>
<td>2 (n = 20)</td>
<td>963.0</td>
<td>99.3</td>
</tr>
<tr>
<td>3 (n = 20)</td>
<td>2242.0</td>
<td>244.8</td>
</tr>
<tr>
<td>Sample</td>
<td>Mean (ng/ml)</td>
<td>SD (ng/ml)</td>
</tr>
<tr>
<td>--------</td>
<td>-------------</td>
<td>------------</td>
</tr>
<tr>
<td>1 (n = 20)</td>
<td>386.5</td>
<td>57.1</td>
</tr>
<tr>
<td>2 (n = 20)</td>
<td>986.9</td>
<td>90.4</td>
</tr>
<tr>
<td>3 (n = 20)</td>
<td>2383.8</td>
<td>278.4</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Linearity</th>
<th>Range Linearity %</th>
<th>Mean Linearity %</th>
<th>Serial dilution up to</th>
</tr>
</thead>
<tbody>
<tr>
<td>Serum</td>
<td>90 - 104</td>
<td>95</td>
<td>1:128</td>
</tr>
<tr>
<td>Plasma</td>
<td>89 - 102</td>
<td>94</td>
<td>1:128</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Recovery</th>
<th>Range Recovery (%)</th>
<th>Mean Recovery (%)</th>
<th>% Recovery after spiking</th>
</tr>
</thead>
<tbody>
<tr>
<td>Serum</td>
<td>Sample 1: 90 - 109</td>
<td>101</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Sample 2: 90 - 96</td>
<td>93</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Sample 3: 95 - 118</td>
<td>109</td>
<td></td>
</tr>
<tr>
<td>Plasma</td>
<td>Sample 1: 82 - 106</td>
<td>96</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Sample 2: 90 - 104</td>
<td>99</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Sample 3: 97 - 110</td>
<td>103</td>
<td></td>
</tr>
</tbody>
</table>

| Method Comparison: ELISA vs. LC-MS/MS | Plasma | LC-MS/MS = 0.9x+71.5 | R²=0.9355; N = 30 |

9. References/Literature


For updated literature or any other information please contact your local supplier.

Symbols:

- **Storage temperature**
- **Manufacturer**
- **Contains sufficient for <n> tests**
- **Expiry date**
- **Batch code**
- **For in-vitro diagnostic use only!**
- **Consult instructions for use**
- **Content**
- **CE labelled**
- **Caution**
- **Catalogue number**
- **RUO**
- **For research use only!**