HAMA ELISA

Enzyme immunoassay for the quantitative determination of human Anti mouse IgG antibodies (HAMA) in serum and plasma.

REF RE53161

Σ 96

2-8 °C

EU: For research use only.
U.S.: For research use only. Not for use in diagnostic procedures.
1. INTENDED USE

This ELISA (enzyme-linked immunosorbent assay) kit is produced for the quantitative determination of human anti-mouse IgG antibody (HAMA) levels in patient serum or plasma samples. It detects both HAMA-IgG and HAMA-IgM subtypes. The test might be used as an aid for detection of patients with positive HAMA that may affect prescribed diagnosis and treatment involving monoclonal murine IgG.

2. SUMMARY OF PHYSIOLOGY

Clinically, mouse monoclonal antibodies (IgG) and their fragments are used in vivo diagnosis procedure (radionuclides) and treatment for patients with various diseases. In patients, even a single dose injection of murine monoclonal IgG may induce immune response directed against this foreign protein (immunogen). In the circulation, the presence of human antibody against murine IgG would bind to the injected murine IgG and, therefore, diminish the efficacy of either in-vivo diagnosis or treatment. Especially, the HAMA would increase the risk of anaphylactic complications to subsequent administration of the murine IgG based therapy.

The present of HAMA in patient serum or plasma specimens causes both false positive and false negative immunoassay test results depending on assay principles and monoclonal antibodies use in the assay system.

This HAMA ELISA is a ready to use test kit with well-breakable microtiter plate and simple test procedures. It also provides a wide measurement range without high dose “hook” effect.

3. ASSAY PRINCIPLE

This ELISA is designed, developed and produced for the quantitative measurement of HAMA in serum and plasma samples. The assay utilizes the two-site “sandwich” technique with two selected antibodies that bind to HAMA.

Assay standards, controls and patient samples are directly added to wells of a microplate that is coated with murine IgG. After the first incubation period, the HAMA binds to the murine IgG on the wall of microtiter well and unbound proteins in each microtiter well are washed away. Then a horseradish peroxidase (HRP) labeled murine IgG is added to each microtiter well and a “sandwich” of “murine IgG HAMA – murine IgG” is formed. The unbound HRP conjugated murine IgG is removed in the subsequent washing step. For the detection of this immunocomplex, the well is then incubated with a substrate solution in a timed reaction and then measured in a spectrophotometric microplate reader. The enzymatic activity of the immunocomplex bound to HAMA on the wall of the microtiter well is directly proportional to the amount of HAMA in the sample. A standard curve is generated by plotting the absorbance versus the respective HAMA concentration for each standard on point-to-point, cubical scales or 4 parameter curve fit. The concentration of HAMA in test samples is determined directly from this standard curve.

4. REAGENTS: Preparation and Storage

This test kit must be stored at 2 – 8°C upon receipt. For the expiration date of the kit refer to the label on the kit box. All components are stable until this expiration date.

Prior to use allow all reagents to come to room temperature. Reagents from different kit lot numbers should not be combined or interchanged.

<table>
<thead>
<tr>
<th>MTP</th>
<th>Murine IgG Coated Microplate</th>
</tr>
</thead>
<tbody>
<tr>
<td>One well breakable microplate with 12 x eight strips (96 wells total) coated with murine IgG. The plate is framed and sealed in a foil zipper bag with a desiccant. This reagent should be stored at 2 – 8°C and is stable until the expiration date on the kit box.</td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>ENZCONJ</th>
<th>Enzyme Conjugate</th>
</tr>
</thead>
<tbody>
<tr>
<td>One vial containing 0.6 mL HRP labeled murine IgG in a stabilized protein matrix. This reagent must be diluted with Enzyme Conjugate Diluent before use. This reagent should be stored at 2 – 8°C and is stable until the expiration date on the kit box.</td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>ENZCONJDIL</th>
<th>Enzyme Conjugate Diluent</th>
</tr>
</thead>
<tbody>
<tr>
<td>One vial containing 12 mL ready to use buffer. It should be used only for Enzyme Conjugate dilution according to the assay procedures. This reagent should be stored at 2 – 8°C and is stable until the expiration date on the kit box.</td>
<td></td>
</tr>
</tbody>
</table>
**CAL**

**HAMA Standards**
Five vials each containing 0.5 mL of a different level of HAMA in a liquid protein matrix with a non-azide based preservative. Refer to vial for exact concentration for each standard. These reagents should be stored at 2 – 8°C and are stable until the expiration date on the kit box.

**CONTROL**

**HAMA Controls**
Two vials each containing 0.5 mL of a different levels of HAMA in a liquid protein matrix with a non-azide based preservative. Refer to vials for exact concentration range for each control. Both controls should be stored at 2 – 8°C and are stable until the expiration date on the kit box.

**ASSAYBUF**

**Assay Buffer**
One bottle containing 30 mL of ready to use phosphate buffered saline based assay buffer with bovine serum albumin added. This reagent should be stored at 2 – 8°C and is stable until the expiration date on the kit box.

**WASHBUF CONC**

**Wash Buffer Concentrate**
One bottle contains 30 mL of 30 fold concentrate. Before use the contents must be diluted with 870 mL of distilled water and mix well. Upon dilution this yields a working wash solution containing a surfactant in phosphate buffered saline with a non-azide and non-mercury based preservative. The diluted wash buffer should be stored at room temperature and is stable until the expiration date on the kit box.

**TMB SUBS**

**TMB Substrate**
One bottle contains 12 mL of tetramethylbenzidine (TMB) with stabilized hydrogen peroxide. This reagent should be stored at 2 – 8°C and is stable until the expiration date on the kit box.

**TMB STOP**

**TMB Stop Solution**
One bottle contains 12 mL of 0.5 M sulfuric acid. This reagent should be stored at 2 – 8°C or room temperature and is stable until the expiration date on the kit box.

5. **SAFETY PRECAUTIONS**
The reagents must be used in a professional laboratory environment and are for research use only. Source material (e.g. highly purified bovine serum albumin) of bovine serum was derived in the contiguous 48 United States. It was obtained only from donor health animals maintained under veterinary supervision and found free of contagious diseases. Wear gloves while performing this assay and handle these reagents as if they are potential infectious. Avoid contact with reagents containing TMB, hydrogen peroxide, or sulfuric acid. TMB may cause irritation to skin and mucous membranes and cause an allergic skin reaction. TMB is a suspected carcinogen. Sulfuric acid may cause severe irritation on contact with skin. Do not get in eyes, on skin, or on clothing. Do not ingest or inhale fumes. On contact, flush with copious amounts of water for at least 15 minutes. Use Good Laboratory Practices.

6. **MATERIALS REQUIRED BUT NOT PROVIDED**
1. Precision single channel pipettes capable of delivering 25 µL, 50 µL, 100 µL, and 1000 µL etc.
2. Repeating dispenser suitable for delivering 100 µL.
3. Disposable pipette tips suitable for above volume dispensing.
4. Disposable 12 x 75 mm or 13 x 100 glass tubes.
5. Disposable plastic 100 mL and 1000 mL bottle with caps.
6. Aluminum foil.
7. Deionized or distilled water.
8. Plastic microtiter well cover or polyethylene film.
9. ELISA multichannel wash bottle or automatic (semi-automatic) washing system.
10. Spectrophotometric microplate reader capable of reading absorbance at 450 nm.

7. **SPECIMEN COLLECTION**
Only 50 µL of human serum or plasma is required for HAMA measurement in duplicate. No special preparation of individual is necessary prior to specimen collection. In the case of serum, whole blood should be collected and must be allowed to clot for a minimum of 30 minutes at room temperature before the serum is separated by centrifugation (850 – 1500xg for 10 minutes). The serum should be separated from the clot within three hours of blood collection and transferred to a clean test tube. Serum or plasma samples should be stored at 2 - 8°C if the assay is to be performed within 72 hours. Otherwise, patient samples should be stored at -20°C or below until measurement. Avoid repeated (more than three times) freezing and thawing of specimen.
8. ASSAY PROCEDURE

Reagent Preparation
(1) Prior to use allow all reagents to come to room temperature. Regents from different kit lot numbers should not be combined or interchanged.
(2) ELISA Wash Concentrate must be diluted to working solution prior use. Please see REAGENTS section for details.

Assay Procedure
(1) Place a sufficient number of murine IgG coated microwell strips/wells in a holder to run HAMA standards, controls and unknown samples in duplicate.
(2) Test Configuration

<table>
<thead>
<tr>
<th>ROW</th>
<th>STRIP 1</th>
<th>STRIP 2</th>
<th>STRIP 3</th>
</tr>
</thead>
<tbody>
<tr>
<td>A</td>
<td>STD 1</td>
<td>STD 5</td>
<td>SAMPLE 2</td>
</tr>
<tr>
<td>B</td>
<td>STD 1</td>
<td>STD 5</td>
<td>SAMPLE 2</td>
</tr>
<tr>
<td>C</td>
<td>STD 2</td>
<td>C 1</td>
<td>SAMPLE 3</td>
</tr>
<tr>
<td>D</td>
<td>STD 2</td>
<td>C 1</td>
<td>SAMPLE 3</td>
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<tr>
<td>E</td>
<td>STD 3</td>
<td>C 2</td>
<td>SAMPLE 4</td>
</tr>
<tr>
<td>F</td>
<td>STD 3</td>
<td>C 2</td>
<td>SAMPLE 4</td>
</tr>
<tr>
<td>G</td>
<td>STD 4</td>
<td></td>
<td>SAMPLE 1</td>
</tr>
<tr>
<td>H</td>
<td>STD 4</td>
<td></td>
<td>SAMPLE 1</td>
</tr>
</tbody>
</table>

(3) Add 25 µL of standards, controls and patient samples into the designated microwell.
(4) Add 100 µL of assay buffer to each well.
(5) Cover the plate with one plate sealer and incubate plate at room temperature for 1 hour.
(6) Prepare Enzyme Conjugate working solution by 1:21 fold dilution of the Enzyme Conjugate with the Enzyme Conjugate Diluent. For each strip, it is required to mix 1 mL of the Enzyme Conjugate diluent with 50 µL of the Enzyme Conjugate in a clean test tube.
(7) Remove plate sealer. Aspirate the contents of each well. Wash each well 5 times by dispensing 350 µL of working wash solution into each well and then completely aspirating the contents. Alternatively, an automated microplate washer can be used.
(8) Add 100 µL of above diluted Enzyme Conjugate working solution to each of the wells.
(9) Cover the plate with the plate sealer and incubate plate at room temperature for 30 min.
(10) Remove plate sealer. Aspirate the contents of each well. Wash each well 5 times by dispensing 350 µL of working wash solution into each well and then completely aspirating the contents. Alternatively, an automated microplate washer can be used.
(11) Add 100 µL of TMB Substrate into each of the wells.
(12) Cover the plate with one plate sealer and also with aluminum foil to avoid exposure to light.
(13) Incubate plate at room temperature for 20 min.
(14) Remove the aluminum foil and plate sealer. Add 100 µL of TMB Stop Solution into each of the wells. Mix gently.
(15) Read the absorbance at 450 nm within 10 minutes in a microplate reader.

NOTE: to reduce the background, one can set the instrument to dual wavelength measurement at 450 nm with background wavelength correction set at 595 nm or 620 nm or 630 nm.
9. PROCEDURAL NOTES
1. It is recommended that all standards, controls and unknown samples be assayed in duplicate. The average absorbance reading of each duplicate should be used for data reduction and the calculation of results.
2. For patient samples with higher than a level 5 standard, it is recommended to measure diluted the specimen with assay buffer at 1:10, 1:100, etc. for a more accurate report.
3. Keep light sensitive reagents in the original amber bottles.
4. Store any unused murine IgG coated strips in the foil Ziploc bag with desiccant to protect from moisture.
5. Careful technique and use of properly calibrated pipetting devices are necessary to ensure reproducibility of the test.
6. Incubation times or temperatures other than those stated in this insert may affect the results.
7. Avoid air bubbles in the microwell as this could result in lower binding efficiency and higher CV% of duplicate reading.
8. All reagents should be mixed gently and thoroughly prior use. Avoid foaming.

10. INTERPRETATION OF RESULTS
1. Calculate the average absorbance for each pair of duplicate test results.
2. Subtract the average absorbance of the STD 1 (0 ng/mL) from the average absorbance of all other readings to obtain corrected absorbance.
3. The standard curve is generated by the corrected absorbance of all standard levels on the ordinate against the standard concentration on the abscissa using point-to-point or log-log paper. Appropriate computer assisted data reduction programs may also be used for the calculation of results. We recommend using Quadratic curve fit.

The HAMA concentrations for the controls and patient samples are read directly from the standard curve using their respective corrected absorbance.

11. EXAMPLE DATA AND STANDARD CURVE
A typical absorbance data and the resulting standard curve from this HAMA ELISA are represented. This curve should not be used in lieu of standard curve run with each assay.

<table>
<thead>
<tr>
<th>Well I.D.</th>
<th>OD 450 nm Absorbance Results ng/mL</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Readings</td>
</tr>
<tr>
<td>0 ng/mL</td>
<td>0.051</td>
</tr>
<tr>
<td>40 ng/mL</td>
<td>0.182</td>
</tr>
<tr>
<td>135 ng/mL</td>
<td>0.561</td>
</tr>
<tr>
<td>450 ng/mL</td>
<td>1.737</td>
</tr>
<tr>
<td>1500 ng/mL</td>
<td>3.230</td>
</tr>
<tr>
<td>Control 1</td>
<td>0.284</td>
</tr>
<tr>
<td></td>
<td>0.309</td>
</tr>
<tr>
<td>Control 2</td>
<td>1.166</td>
</tr>
<tr>
<td></td>
<td>1.109</td>
</tr>
</tbody>
</table>

HAMA ELISA Standard Curve

HAMA Standards (ng/mL)

HAMA ELISA (RE53161 )

Version 8  2014-09
12. **EXPECTED VALUES**
One hundred seventy normal adult sera were measured with this HAMA ELISA. One hundred sixty sera showed the OD reading very close to the zero calibrator. The 99% confidence normal cut-off is **25 ng/mL**. It is highly recommend that each laboratory establish its own normal cut off level.
One positive sample with HAMA level of 64 ng/ml was further tested with dilution of this sample in 1:2, 1:4 and 1:8. A linear HAMA dilution result was observed and indicated HAMA specific activity of this sample.

13. **LIMITATION OF THE PROCEDURE**
1. Since there is no Gold Standard concentration or international standard available for HAMA measurement, the values of assay standards were established and validated. Results obtained with different assay methods or kits cannot be used interchangeably.
2. For unknown sample values read directly from the assay that is greater than 1500 ng/mL, it is recommended to measure a further diluted sample for a more accurate measurement.
3. Bacterial or fungal contamination of serum specimens or reagents, or cross contamination between reagents may cause erroneous results.
4. Water deionized with polyester resins may inactivate the horseradish peroxidase enzyme.

14. **QUALITY CONTROL**
To assure the validity of the results each assay should include adequate controls with known HAMA levels. We recommend that all assays include the laboratory’s own HAMA controls in addition to those provided with this kit.

15. **PERFORMANCE CHARACTERISTICS**

**Sensitivity**
The sensitivity of this HAMA ELISA as determined by the 95% confidence limit on 20 duplicate determination of zero standard is about 2 ng/mL.

**High Dose “hook” effect**
This assay has showed that it did not have any high dose “hook” effect up to 1,000,000 ng/mL.

**Precision**
The intra-assay precision was validated by measuring one control sample in a single assay with eight-replicate determinations.

<table>
<thead>
<tr>
<th>Mean HAMA Value (ng/mL)</th>
<th>CV (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>51.66</td>
<td>5.1</td>
</tr>
</tbody>
</table>

The inter-assay precision is validated by measuring one control sample in duplicate in 6 individual assays.

<table>
<thead>
<tr>
<th>Mean HAMA Value (ng/mL)</th>
<th>CV (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>52.12</td>
<td>5.8</td>
</tr>
</tbody>
</table>

**Linearity**
Two serum samples were diluted with assay buffer and assayed. The results in the value of ng/mL are as follows:

<table>
<thead>
<tr>
<th>#</th>
<th>DILUTION</th>
<th>OBSERVED VALUE</th>
<th>RECOVERY %</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Neat</td>
<td>88.51</td>
<td>-</td>
</tr>
<tr>
<td></td>
<td>1:2</td>
<td>44.98</td>
<td>101</td>
</tr>
<tr>
<td></td>
<td>1:4</td>
<td>22.85</td>
<td>103</td>
</tr>
<tr>
<td></td>
<td>1:8</td>
<td>14.14</td>
<td>113</td>
</tr>
<tr>
<td>2</td>
<td>Neat</td>
<td>298.12</td>
<td>-</td>
</tr>
<tr>
<td></td>
<td>1:2</td>
<td>141.93</td>
<td>95</td>
</tr>
<tr>
<td></td>
<td>1:4</td>
<td>66.78</td>
<td>90</td>
</tr>
<tr>
<td></td>
<td>1:8</td>
<td>37.15</td>
<td>100</td>
</tr>
</tbody>
</table>
16. REFERENCES


17. Short Assay Protocol:
- Add 25 µl of standards, control and patient sample
- Add 100 µl of assay buffer
- Incubate 1 hour at RT
- Wash strips with diluted wash buffer
- Add 100 µl Enzyme Conjugate
- Incubate 30 min at RT
- Wash strips with diluted wash buffer
- Add 100 µl TMB substrate
- Incubate 20 min at RT
- Add 100 µl TMB stop solution
- Read strips at OD 450 nm
### Symbols

- **Cat.-No.**: / **Kat.-Nr.**: / **No.- Cat.**: / **N.º Cat.**: / **Αριθµός-Κατ.**:
- **Lot-No.**: / **Chargen-Bez.**: / **No. Lot.**: / **Lot-No.**: / **Lote N.º**: / **Λοτο ηλικία**:
- **Use by**: / **Verwendbar bis**: / **Utiliser à**: / **Usado por**: / **Usar até**: / **Da utilizarse entro**: / **Χρησιµοποιείται από**:
- **No. of Tests**: / **Kitgröße**: / **Nb. de Tests**: / **No. de Determ.**: / **Quantità dei testi**: / **Αριθµός εξέτάσεων**:
- **CONC**: Concentrate / Konzentrat / Concentré / Concentrar / Concentrado / Concentrato / Συµπύκνωµα
- **LYO**: Lyophilized / Lyophilisat / Lyophilisé / Liofilizado / Liofilizado / Liofilizzato / Λυοφιλιασµένο
- **IVD**: In Vitro Diagnostic Medical Device. / In-vitro-Diagnostikum. / Appareil Médical pour Diagnostics In Vitro. / Dispositivo Médico para Diagnóstico In Vitro. / Equipamento Médico de Diagnóstico In Vitro. / Dispositivo Medico Diagnostic In vitro. / Ιατρική συσκευή για In-Vitro Διάγνωση.
- **Evaluation kit.**: / **Nur für Leistungsbewertungszwecke.**: / **Kit pour évaluation.**: / **Juego de Reactivos para Evaluación.**: / **Kit de avaliação.**: / **Kit di valutazione.** / **Κιτ Αξιολόγησης.**
- **Read instructions before use.**: / **Arbeitsanleitung lesen.**: / **Lire la fiche technique avant emploi.**: / **Lea las instrucciones antes de usar.**: / **Ler as instruções antes de usar.**: / **Leggere le istruzioni prima dell’uso.**: / **Διαβάστε τις οδηγίες πριν την χρήση.**
- **Keep away from heat or direct sun light.**: / **Vor Hitze und direkter Sonneneinstrahlung schützen.**: / **Garder à l’abri de la chaleur et de toute exposition lumineuse.**: / **Manténgase alejado del calor o la luz solar directa.**: / **Non esporre ai raggi solari.**: / **Να φυλάσσεται µακριά από θερµότητα και άµεση επαφή µε το φως του ηλίου.**
- **Store at**: / **Lagern bei**: / **Stocker à**: / **Almacene a**: / **Armazenar a**: / **Conservare a**: / **Αποθήκευση στους**:
- **Manufacturer**: / **Hersteller**: / **Fabricant**: / **Productor**: / **Fabricante**: / **Fabbricante**: / **Παραγωγός**:
- **Caution!**: / **Vorsicht!**: / **Attention!**: / **¡Precaución!**: / **Cuidado!**: / **Attenzione!**: / **Προσοχή!**

Symbols of the kit components see MATERIALS SUPPLIED.

Die Symbole der Komponenten sind im Kapitel KOMPONENTEN DES KITS beschrieben.

Voir MATERIEL FOURNI pour les symbôles des composants du kit.

Símbolos de los componentes del juego de reactivos, vea MATERIALES SUMINISTRADOS.

Para símbolos dos componentes do kit ver MATERIAIS FORNECIDOS.

Per i simboli dei componenti del kit si veda COMPONENTI DEL KIT.

Για τα σύµβολα των συστατικών του κιτ συµβουλευτείτε το ΠΑΡΕΧΟΜΕΝΑ ΥΛΙΚΑ.

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**IBL AFFILIATES WORLDWIDE**

<table>
<thead>
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<th>IBL International GmbH</th>
<th>IBL International Corp.</th>
</tr>
</thead>
<tbody>
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</tr>
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<td>Tel.: + 49 (0) 40 532891 -0 Fax: -11</td>
<td>Tel.: +1 (416) 645 -1703 Fax: -1704</td>
</tr>
<tr>
<td>E-MAIL: <a href="mailto:IBL@IBL-International.com">IBL@IBL-International.com</a></td>
<td>E-MAIL: <a href="mailto:Sales@IBL-International.com">Sales@IBL-International.com</a></td>
</tr>
</tbody>
</table>

**LIABILITY**: Complaints will be accepted in each mode –written or vocal. Preferred is that the complaint is accompanied with the test performance and results. Any modification of the test procedure or exchange or mixing of components of different lots could negatively affect the results. These cases invalidate any claim for replacement. Regardless, in the event of any claim, the manufacturer’s liability is not to exceed the value of the test kit. Any damage caused to the kit during transportation is not subject to the liability of the manufacturer.

Symbols Version 3.5 / 2012-01-20