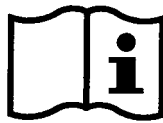


Product information

Information about other products is available at: www.ibl-america.com



User's Manual

Rubella Virus IgM μ -capture ELISA

RUO

REF

IB79897



96 wells

For Research Use Only – Not for Use in Diagnostic Procedures

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1. INTENDED USE

The Rubella Virus IgM μ -capture ELISA is intended for the determination of IgM-class antibodies against Rubella Virus in human serum or plasma (citrate, heparin).

2. PRINCIPLE OF THE ASSAY

The immunoenzymatic determination of specific IgM-class antibodies is based on the ELISA (Enzyme-linked Immunosorbent Assay) μ -capture technique.

Microtiterplates are coated with anti-human IgM-class antibodies to bind the corresponding antibodies of the sample. After washing the wells to remove all unbound sample material, horseradish peroxidase (HRP) labelled antigen is added. This antigen-conjugate binds to the captured specific IgM antibodies. In a second washing step unbound conjugate is removed. The immune complexes are visualized by adding Tetramethylbenzidine (TMB) substrate which gives a blue reaction product. The intensity of this product is proportional to the amount of specific IgM antibodies in the sample. Sulphuric acid is added to stop the reaction. This produces a yellow endpoint colour. Absorbance at 450/620 nm is read using an ELISA Microtiterplate reader.

3. MATERIALS

3.1. Reagents supplied

- SORB** **MT** **Microtiterplate**: 12 break-apart 8-well snap-off strips coated with anti-human IgM-class antibodies; in resealable aluminium foil.
- SAM** **DIL** **Sample Dilution Buffer**: 1 bottle containing 100 mL of phosphate buffer (10 mM) for sample dilution; pH 7.2 \pm 0.2; coloured yellow; ready to use; white cap; \leq 0.0015% (v/v) CMIT/ MIT (3:1).
- STOP** **SOLN** **Stop Solution**: 1 bottle containing 15 mL sulphuric acid, 0.2 mol/l; ready to use; red cap.
- WASH** **SOLN** **20x** **Washing Buffer (20x conc.)**: 1 bottle containing 50 mL of a 20-fold concentrated phosphate buffer (0.2 M), pH 7.2 \pm 0.2, for washing the wells; white cap.
- ENZ** **CONJ** **Conjugate**: 1 bottle containing 12 mL of peroxidase labelled Rubella Virus antigen; coloured yellow; ready to use; black cap.
- SUB** **TMB** **TMB Substrate Solution**: 1 bottle containing 15 mL 3,3',5,5'-tetramethylbenzidine (TMB), $<$ 0.1 %; ready to use; yellow cap.
- CAL** **C** **Positive Control**: 1 vial containing 2 mL control; coloured yellow; ready to use; red cap; \leq 0.02% (v/v) MIT.
- CAL** **B** **Cut-off Control**: 1 vial containing 3 mL control; coloured yellow; ready to use; green cap; \leq 0.02% (v/v) MIT.
- CAL** **A** **Negative Control**: 1 vial containing 2 mL control; coloured yellow; ready to use; blue cap; \leq 0.0015% (v/v) CMIT/ MIT (3:1).

*For hazard and precautionary statements see 12.1

For potential hazardous substances please check the safety data sheet.

3.2. Materials supplied

- 1 Cover foil
- 1 Instruction for use (IFU)
- 1 Plate layout

3.3. Materials and Equipment needed

- ELISA Microtiterplate reader, equipped for the measurement of absorbance at 450/620 nm
- Incubator 37 °C
- Manual or automatic equipment for rinsing Microtiterplate
- Pipettes to deliver volumes between 10 and 1000 μ L
- Vortex tube mixer
- Distilled water
- Disposable tubes

4. STABILITY AND STORAGE

Store the kit at 2...8 °C. The opened reagents are stable up to the expiry date stated on the label when stored at 2...8 °C.

5. REAGENT PREPARATION

It is very important to bring all reagents and samples to room temperature (20...25 °C) and mix them before starting the test run!

5.1. Microtiterplate

The break-apart snap-off strips are coated with anti-human IgM-class antibodies. Immediately after removal of the strips, the remaining strips should be resealed in the aluminium foil along with the desiccant supplied and stored at 2...8 °C.

5.2. Washing Buffer (20x conc.)

Dilute Washing Buffer 1 + 19; e. g. 10 mL Washing Buffer + 190 mL distilled water. The diluted buffer is stable for 5 days at room temperature (20...25 °C). In case crystals appear in the concentrate, warm up the solution to 37 °C e.g. in a water bath. Mix well before dilution.

5.3. TMB Substrate Solution

The reagent is ready to use and has to be stored at 2...8 °C, away from the light. The solution should be colourless or could have a slight blue tinge. If the substrate turns into blue, it may have become contaminated and should be thrown away.

6. SAMPLE COLLECTION AND PREPARATION

Use human serum or plasma (citrate, heparin) samples with this assay. If the assay is performed within 5 days after sample collection, the samples should be kept at 2...8 °C; otherwise they should be aliquoted and stored deep-frozen (-70...-20 °C). If samples are stored frozen, mix thawed samples well before testing. Avoid repeated freezing and thawing. Heat inactivation of samples is not recommended.

6.1. Sample Dilution

Before assaying, all samples should be diluted 1+100 with Sample Dilution Buffer. Dispense 10 μ L sample and 1 mL Sample Dilution Buffer into tubes to obtain a 1+100 dilution and thoroughly mix with a Vortex.

7. ASSAY PROCEDURE

Please read the instruction for use carefully **before** performing the assay. Result reliability depends on strict adherence to the instruction for use as described. The following test procedure is only validated for manual procedure. If performing the test on ELISA automatic systems we recommend increasing the washing steps from three up to five and the volume of Washing Buffer from 300 μ L to 350 μ L to avoid washing effects. Pay attention to chapter 12. Prior to commencing the assay, the distribution and identification plan for all samples and standards/controls (duplicates recommended) should be carefully established on the plate layout supplied in the kit. Select the required number of microtiter strips or wells and insert them into the holder.

Perform all assay steps in the order given and without any delays.

A clean, disposable tip should be used for dispensing each standard/control and sample.

Adjust the incubator to 37 \pm 1 °C.

1. Dispense 100 μ L standards/controls and diluted samples into their respective wells. Leave well A1 for the Substrate Blank.
2. Cover wells with the foil supplied in the kit.
3. **Incubate for 1 hour \pm 5 min at 37 \pm 1 °C.**
4. When incubation has been completed, remove the foil, aspirate the content of the wells and wash each well three times with 300 μ L of Washing Buffer. Avoid overflows from the reaction wells. The interval between washing and aspiration should be > 5 sec. At the end carefully remove remaining fluid by tapping strips on tissue paper prior to the next step!
Note: Washing is important! Insufficient washing results in poor precision and false results.
5. Dispense 100 μ L Conjugate into all wells except for the Substrate Blank well A1.
6. **Incubate for 30 min at room temperature (20...25 °C).** Do not expose to direct sunlight.
7. Repeat step 4.

8. Dispense 100 μ L TMB Substrate Solution into all wells.
9. **Incubate for exactly 15 min at room temperature (20...25 °C) in the dark.** A blue colour occurs due to an enzymatic reaction.
10. Dispense 100 μ L Stop Solution into all wells in the same order and at the same rate as for the TMB Substrate Solution, thereby a colour change from blue to yellow occurs.
11. Measure the absorbance at 450/620 nm within 30 min after addition of the Stop Solution.

7.1. Measurement

Adjust the ELISA Microtiterplate reader **to zero** using the **Substrate Blank**.

If - due to technical reasons - the ELISA Microtiterplate reader cannot be adjusted to zero using the Substrate Blank, subtract its absorbance value from all other absorbance values measured in order to obtain reliable results!

Measure the absorbance of all wells at **450 nm** and record the absorbance values for each stand-ard/control and sample in the-plate layout.

Bichromatic measurement using a reference wavelength of 620 nm is recommended.

Where applicable calculate the mean absorbance values of all duplicates.

8. RESULTS

8.1. Run Validation Criteria

In order for an assay run to be considered valid, these Instructions for Use have to be strictly followed and the following criteria must be met:

- **Substrate Blank:** Absorbance value < **0.100**
- **Negative Control:** Absorbance value < **Cut-off**
- **Cut-off Control:** Absorbance value **0.150 – 1.300**
- **Positive Control:** Absorbance value > **Cut-off**

If these criteria are not met, the test is not valid and must be repeated.

8.2. Calculation of Results

The Cut-off is the mean absorbance value of the Cut-off Control determinations.

Example: Absorbance value Cut-off Control 0.44 + absorbance value Cut-off control 0.42 = 0.86 / 2 = 0.43

$$\text{Cut-off} = 0.43$$

8.2.1. Results in Units [U]

$\frac{\text{Sample (mean) absorbance value} \times 10}{\text{Cut-off}} = [\text{Units} = \text{U}]$

Example: $\frac{1.591 \times 10}{0.43} = 37 \text{ U}$

8.3. Interpretation of Results

| | | |
|-----------|----------|--|
| Cut-off | 10 U | - |
| Positive | > 11 U | Antibodies against the pathogen are present. There has been a contact with the antigen (pathogen resp. vaccine). |
| Equivocal | 9 – 11 U | Antibodies against the pathogen could not be detected clearly. It is recommended to repeat the test with a fresh sample in 2 to 4 weeks. If the result is equivocal again the sample is judged as negative . |
| Negative | < 9 U | The sample contains no antibodies against the pathogen. A previous contact with the antigen (pathogen resp. vaccine) is unlikely. |

9. SPECIFIC PERFORMANCE CHARACTERISTICS

The results refer to the groups of samples investigated; these are not guaranteed specifications. For further information about the specific performance characteristics please contact IBL-America.

9.1. Precision

| Intraassay | n | Mean (E) | CV (%) |
|------------|----|----------|--------|
| #1 | 24 | 0.821 | 5.40 |
| #2 | 24 | 1.276 | 3.56 |

| #3 | 24 | 2.174 | 2.05 |
|-------------------|----------|-----------------|---------------|
| Interassay | n | Mean (U) | CV (%) |
| #1 | 12 | 15.93 | 4.49 |
| #2 | 12 | 25.94 | 5.84 |
| #3 | 12 | 5.17 | 7.32 |

9.2. Specificity

The specificity is defined as the probability of the assay of scoring negative in the absence of the specific analyte. It is 99.4% (95% confidence interval: 96.71% - 99.98%)

9.3. Sensitivity

The sensitivity is defined as the probability of the assay of scoring positive in the presence of the specific analyte. It is 100% (95% confidence interval: 88.43% - 100%).

9.4. Interferences

Interferences with hemolytic, lipemic or icteric samples are not observed up to a concentration of 10 mg/mL hemoglobin, 5 mg/mL triglycerides and 0.5 mg/mL bilirubin.

9.5. Cross Reactivity

Investigation of a sample panel with antibody activities to potentially cross-reacting parameters did not reveal evidence of false-positive results due to cross-reactions.

10. LIMITATIONS OF THE PROCEDURE

Bacterial contamination or repeated freeze-thaw cycles of the sample may affect the absorbance values.

11. PRECAUTIONS AND WARNINGS

- The test procedure, the information, the precautions and warnings in the instructions for use have to be strictly followed. The use of the testkits with analyzers and similar equipment has to be validated. Any change in design, composition and test procedure as well as for any use in combination with other products not approved by the manufacturer is not authorized; the user himself is responsible for such changes. The manufacturer is not liable for false results and incidents for these reasons. The manufacturer is not liable for any results by visual analysis of the samples.
- Only for research use.
- All materials of human or animal origin should be regarded and handled as potentially infectious.
- All components of human origin used for the production of these reagents have been tested for anti-HIV antibodies, anti-HCV antibodies and HBsAg and have been found to be non-reactive.
- Do not interchange reagents or Microtiterplates of different production lots.
- No reagents of other manufacturers should be used along with reagents of this test kit.
- Do not use reagents after expiry date stated on the label.
- Use only clean pipette tips, dispensers, and lab ware.
- Do not interchange screw caps of reagent vials to avoid cross-contamination.
- Close reagent vials tightly immediately after use to avoid evaporation and microbial contamination.
- After first opening and subsequent storage check conjugate and standard/control vials for microbial contamination prior to further use.
- To avoid cross-contamination and falsely elevated results pipette samples and dispense reagents without splashing accurately into the wells.
- The ELISA is only designed for qualified personnel following the standards of good laboratory practice (GLP).
- For further internal quality control each laboratory should additionally use known samples.

11.1. Safety note for reagents containing hazardous substances

Reagents may contain CMIT/MIT (3:1) or MIT (see 4.1 oder refers to 4.1)

Therefore, the following hazard and precautionary statements apply.



| | | |
|----------------|-----------|---|
| Warning | H317 | May cause an allergic skin reaction. |
| | P261 | Avoid breathing spray |
| | P280 | Wear protective gloves/ protective clothing. |
| | P302+P352 | IF ON SKIN: Wash with plenty of soap and water. |
| | P333+P313 | If skin irritation or rash occurs: Get medical advice/ attention. |

P362+P364 Take off contaminated and Wash it before reuse.

Further information can be found in the safety data sheet.

11.2. Disposal Considerations

Residues of chemicals and preparations are generally considered as hazardous waste. The disposal of this kind of waste is regulated through national and regional laws and regulations. Contact your local authorities or waste management companies which will give advice on how to dispose hazardous waste.

ABBREVIATIONS

| | |
|------|--|
| CMIT | 5-chloro-2-methyl-4-isothiazolin-3-one |
| MIT | 2-methyl-2H-isothiazol-3-one |

SUMMARY OF TEST PROCEDURE

SCHEME OF THE ASSAY Rubella Virus IgM μ -capture ELISA












Test Preparation

| |
|--|
| <p>Prepare reagents and samples as described. Establish the distribution and identification plan for all samples and standards/controls on the plate layout supplied in the kit. Select the required number of microtiter strips or wells and insert them into the holder.</p> |
|--|

Assay Procedure

| | Substrate Blank (A1) | Negative Control | Cut-off Control | Positive Control | Sample (diluted 1+100) |
|--|----------------------|------------------|-----------------|------------------|------------------------|
| Negative Control | - | 100 μ L | - | - | - |
| Cut-off Control | - | - | 100 μ L | - | - |
| Positive Control | - | - | - | 100 μ L | - |
| Sample (diluted 1+100) | - | - | - | - | 100 μ L |
| <p>Cover wells with foil supplied in the kit Incubate for 1 h at 37 \pm 1 $^{\circ}$C Wash each well three times with 300 μL of Washing Buffer</p> | | | | | |
| Conjugate | - | 100 μ L | 100 μ L | 100 μ L | 100 μ L |
| <p>Incubate for 30 min at room temperature (20...25 $^{\circ}$C) Do not expose to direct sunlight Wash each well three times with 300 μL of Washing Buffer</p> | | | | | |
| TMB Substrate Solution | 100 μ L | 100 μ L | 100 μ L | 100 μ L | 100 μ L |
| <p>Incubate for exactly 15 min at room temperature (20...25 $^{\circ}$C) in the dark</p> | | | | | |
| Stop Solution | 100 μ L | 100 μ L | 100 μ L | 100 μ L | 100 μ L |
| <p>Photometric measurement at 450 nm (reference wavelength: 620 nm)</p> | | | | | |

SYMBOLS USED WITH IBL-AMERICA ASSAYS

| Symbol | English | Deutsch | Français | Espanol | Italiano |
|---|------------------------------------|--|--|---|-------------------------------------|
|  | European Conformity | CE-Konformitätskennzeichnung | Conforme aux normes européennes | Conformidad europea | Conformità europea |
|  | Consult instructions for use | Gebrauchsanweisung beachten | Consulter les instructions d'utilisation | Consulte las Instrucciones | Consultare le istruzioni per l'uso |
|  | In vitro diagnostic device | In-vitro-Diagnostikum | Usage Diagnostic in vitro | Diagnóstico in vitro | Per uso Diagnostica in vitro |
|  | For research use only | Nur für Forschungszwecke | Seulement dans le cadre de recherches | Sólo para uso en investigación | Solo a scopo di ricerca |
|  | Catalogue number | Katalog-Nr. | Référence | Número de catálogo | No. di Cat. |
|  | Lot. No. / Batch code | Chargen-Nr. | No. de lot | Número de lote | Lotto no |
|  | Contains sufficient for <n> tests/ | Ausreichend für "n" Ansätze | Contenu suffisant pour "n" tests | Contenido suficiente para <n> ensayos | Contenuto sufficiente per "n" saggi |
|  | Note warnings and precautions | Warnhinweise und Vorsichtsmaßnahmen beachten | Avertissements et mesures de précaution font attention | Tiene en cuenta advertencias y precauciones | Annoti avvisi e le precauzioni |
|  | Storage Temperature | Lagerungstemperatur | Temperature de conservation | Temperatura de conservacion | Temperatura di conservazione |
|  | Expiration Date | Mindesthaltbarkeitsdatum | Date limite d'utilisation | Fecha de caducidad | Data di scadenza |
|  | Legal Manufacturer | Hersteller | Fabricant | Fabricante | Fabbricante |
| <i>Distributed by</i> | Distributor | Vertreiber | Distributeur | Distribuidor | Distributore |