



## 119<sup>th</sup> IAMM EQAS Microbiology: Bacteriology/ Serology

CMC MICRO EQAS

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FEBRUARY 2025

119<sup>th</sup> EQAS – SEROLOGY

MEMBER ID:

M | I | I | 7 | 4

Last date for receiving reports: 30<sup>th</sup> APRIL, 2025

**Instructions:**

1. Each individual serum sample to be reconstituted with **0.6mL** of sterile distilled water / deionized water.
2. Please perform required tests and send your results as per the attached tabular format.
3. You are instructed to fill up each column; as this information will be used for assessing your performance.
4. **Do not use tick marks and encircle where ever necessary**
5. Please perform C-reactive protein (CRP) & Syphilis Serology assay on the three serum specimens provided as specified.
6. **C-reactive protein (CRP) levels should be expressed only in mg/L; if other units are used, it will be marked as incorrect.**
7. Separate sheets are provided for entering the results.
8. **Evaluation format for Serology:**

a. **Qualitative (2 marks for each serum)**

- Result have to be given as Positive or Negative only
- Correct interpretation: Full marks (2 marks)
- Wrong Interpretation: Zero mark (0 mark)

b. **Semi quantitative / Quantitative (2 marks for each serum)**

We will assess by robust analysis (as per ISO: 13528:2015) using participants results for different peer groups (Nephelometry, Turbidimetry, etc.,) and marking format as based on Z & Z' score, which is as given below.

**Z & Z' score system for Values**

| Z & Z' Score   | Category          | Marks for values |
|----------------|-------------------|------------------|
| $\leq 2$       | Correct           | 2 marks          |
| $>2$ but $< 3$ | Partially correct | 1 mark           |
| $\geq 3$       | Incorrect         | 0 mark           |

Note: As multiple assays have been requested for SE3, kindly mark 'ND' if the assay is not performed at your centre.

**IMPORTANT!! All sera are potentially infectious. Adequate universal precautions to be used while handling the specimens**

**SE1: C-reactive protein (CRP)**

**SE1: Serum specimen from 52-year female with acute febrile illness with rash.**

| S.no | Subject  | C-reactive protein (CRP)   |                     | Result mg/L*        |
|------|--|--|---------------------|---------------------|
| 1    | <b>Method</b>                                  | Qualitative  | Latex agglutination | Positive / Negative |
|      |  | Semi Quantitative  | Latex agglutination | mg/L                |
|      |  | Quantitative   | Nephelometry        | mg/L                |
|      |  |  | Turbidimetry        | < 5 mg/L            |
|      |  |  | ELISA               | mg/L                |
|      |  |  | CLIA                | mg/L                |
|      |  |  | Others:             | mg/L                |
| 2    | <b>Your Normal Range</b>                       | < 5 mg/L   |                     |                     |
| 3    | <b>Name of the kit used</b>                    | CR P -   |                     |                     |
| 4    | <b>Manufacturer<br/>(Name, City, Country)</b>  | BECKMAN COULTER  |                     |                     |
| 5    | <b>Lot No.</b>                                 | 2579   |                     |                     |
| 6    | <b>Expiry date of kit</b>                      | September 25   |                     |                     |
| 7    | <b>Automation used</b>                         | Yes / No   |                     |                     |
| 8    | <b>If yes, give details of Automation used</b> | Model: BECKMAN COULTER<br>Manufacturer:<br>City: USA<br>Country: |                     |                     |

\*It is understood that the value mentioned is in mg/L only

**IMPORTANT!! All sera are potentially infectious. Adequate universal precautions to be used while handling the specimens**

**SE2: C-reactive protein (CRP)**

**SE2: Serum specimen from 40-year male with acute febrile illness.**

| S.no | Subject  | C-reactive protein (CRP)  |                     | Result mg/L*        |
|------|--|---------------------------|---------------------|---------------------|
| 1    | <b>Method</b>                                  | Qualitative               | Latex agglutination | Positive / Negative |
|      |  | Semi Quantitative         | Latex agglutination | mg/L                |
|      |  | Quantitative              | Nephelometry        | mg/L                |
|      |  |                           | Turbidimetry        | 78.1 mg/L           |
|      |  |                           | ELISA               | mg/L                |
|      |  |                           | CLIA                | mg/L                |
|      |  |                           | Others:             | mg/L                |
| 2    | <b>Your Normal Range</b>                       | < 5 mg/L                  |                     |                     |
| 3    | <b>Name of the kit used</b>                    | CRP                       |                     |                     |
| 4    | <b>Manufacturer<br/>(Name, City, Country)</b>  | Beckman Coulter, USA      |                     |                     |
| 5    | <b>Lot No.</b>                                 | 2579                      |                     |                     |
| 6    | <b>Expiry date of kit</b>                      | September - 25            |                     |                     |
| 7    | <b>Automation used</b>                         | Yes/No                    |                     |                     |
| 8    | <b>If yes, give details of Automation used</b> | Model:<br>Beckman Coulter | City:               | Country: USA        |
|      |  | Manufacturer:             |                     |                     |

\*It is understood that the value mentioned is in mg/L only

**IMPORTANT!! All sera are potentially infectious. Adequate universal precautions to be used while handling the specimens**

**SE3. Syphilis Serology**

**SE3: Serum specimen from 26-year old pregnant woman in 1<sup>st</sup> trimester.**

| S.<br>no | Subject  | Syphilis Serology  |  |  |  |
|----------|--|--|--|--|--|
|          |  | RPR / VDRL   | TPHA   | Syphilis ELISA   | Other Test   |
| 1        | <b>Result</b>                                  | NON REACTIVE   |  |  |  |
| 2        | <b>Units/ Dilution</b>                         | < 1:1 diln.  |  |  |  |
| 3        | <b>Name of the kit used</b>                    | RECKON -<br>RPR.   |  |  |  |
| 4        | <b>Manufacturer Name</b>                       | RECKON   |  |  |  |
|          | <b>City</b>                                    | VADODARA   |  |  |  |
|          | <b>Country</b>                                 | INDIA  |  |  |  |
| 5        | <b>Lot No.</b>                                 | 24H*77V  |  |  |  |
| 6        | <b>Expiry date of kit</b>                      | JULY 2026  |  |  |  |
| 7        | <b>Automation used</b>                         | Yes / <u>No</u>  | Yes / No   | Yes / No   | Yes / No   |
| 8        | <b>If yes, give details of Automation used</b> | Model:<br><br>Manufacturer:<br><br>City:<br><br>Country: | Model:<br><br>Manufacturer:<br><br>City:<br><br>Country: | Model:<br><br>Manufacturer:<br><br>City:<br><br>Country: | Model:<br><br>Manufacturer:<br><br>City:<br><br>Country: |

**Laboratory / Institution Name:**

**Date of Dispatch:**

22/06/2025

**Authorized signatory:**

**Signature:** Sadiya

**Name:** DR. SADIYA SULTANA