## **Rheumatoid Antibodies**

Accreditation Status: UKAS Schedule of Accreditation

**Date Scheme started:** General Autoimmune Serology Started 1982, reconfigured 2025

Clinical Applicability: Diagnosis of Rheumatoid Arthritis/Differential diagnosis of inflammatory arthritis

Analytes: Rheumatoid Factor (SER/001)

Citrullinated Proteins (SER/036)

Units for Reporting: IU/ml or U/mL in relation to the appropriate International Reference

Preparations, or titre. Qualitative responses or interpretation of quantitative

results recorded as POSitive or NEGative

Samples Distributed: Liquid format. Normal and pathological human serum

Number of Distributions per year: 6

Number of Samples per Distribution: 2 (1 x RF and 1 x CP)

Frequency of Distributions: Every two months as outlined in the Distribution Schedule

**Schedule of Analysis:** Data entry is via the web for the submission of results. Data analysis

commences 21 days after sample dispatch. Late returns are accepted and will

contribute to the laboratory's cumulative performance statistics

**Data Analysis:** Reports show method or kit related statistics. Qualitative responses are

recorded and assessed in relation to the designated response.

Quantitative results show Method Laboratory Trimmed Mean (MLTM), range of results reported, SD and CV%. MRVIS, MRBIS and SDBIS are also provided for

quantitative results

**Performance Scoring:** MI scoring based on Consensus Designated Response

Criteria of Performance: Laboratory performance is classified in terms of OMIS for

qualitative responses for analytes for which the laboratory is registered over

a running analytical window of 6 Distributions (12 months)

The categories of performance are:

Good Zero
Adequate 1-2
Poor >2

An OMIS of >2 (out of a possible six in the defined time window) for any one

analyte will also be classified as poor performance.

**Persistent Poor Performance:** Defined as being in the Poor Performance category for two or more

successive Distributions