UK NEQAS

Immunology, Immunochemistry & Allergy

Allergen Specific IgE

Accreditation Status: UKAS Schedule of Accreditation

Date Scheme started: 1988

Clinical Applicability: Diagnosis and management of allergic disease

Analytes: The programme includes the assessment of common or

clinically important, individual IgE specificities, for example:

D1 Dermatophagoides pteronyssinus

E1 Cat epithelium **E**5 Dog dander F1 Egg white F2 Cow's milk F13 Peanut F17 Hazel nut G6 Timothy grass 11 Bee venom 13 Wasp venom

K82 Latex

M3 Aspergillus fumigatus M6 Alternaria alternata

T3 Birch W6 Mugwort

Other allergen specificities may be included, subject to the availability of

clinically validated donor serum units

Units for Reporting: Grade and kU/L (arbitrary)

Samples Distributed: Liquid format. Normal and pathological human

Number of Distributions per year: 6

Number of Samples per Distribution: 2

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

specific IgE tests will be analysed on each specimen

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

is commenced 21 days after sample dispatch. Late returns are accepted and will contribute to the laboratory's cumulative performance statistics

Data Analysis: Analysis by grade shows the overall response and the method

specific responses. Analysis of the quantitative responses in Units shows the All Laboratory Trimmed Mean (ALTM) with truncation at 2SD, SD and CV%. Reports show method specific statistics. Individual laboratory performance is expressed in terms of MRBIS, SDBIS and MRVIS, the DV for calculation of VI being taken from the MLTM

Chosen Coefficient of Variation for Allergen Specific IgE is 12%

Performance Scoring: Cumulative performance scores are based on the quantitative response

with MRVIS scoring over a running window of twelve samples or twelve

nonths

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Criteria of Performance:

Performance assessment is allergen specific. Quantitative performance is assessed for each allergen, and is over a running period of 6 distributions containing that allergen (12 months)

 Ideal
 MRVIS
 <50</th>

 Good
 50 - 100

 Adequate
 101 - 200

Poor >200 or SDBIS >200

The overall quantitative performance is expressed as the OMRVIS, the mean of all the individual allergen specific MRVIS

The semiquantitative Grades are assessed by MI scoring in relation to the Consensus Designated Response (CONDR). (For this purpose, Grades 2-6 are considered as CLEAR POSITIVE)

Good OMIS Zero
Adequate 1 - 3
Poor >4

Overall MIS (OMIS) greater than 4 will also be considered as poor performance

Persistent Poor Performance:

Defined as being in the Poor Performance category for two or more successive distributions