

# Acetylcholine Receptor Antibody

<b>Accreditation Status:</b>	<b>UKAS Schedule of Accreditation</b>								
<b>Date Scheme started:</b>	1991								
<b>Clinical Applicability:</b>	Diagnosis and monitoring of Myasthenia Gravis								
<b>Analytes:</b>	ACR ( <b>SER/008</b> )								
<b>Units for Reporting:</b>	nmol/L								
<b>Samples Distributed:</b>	Liquid format. Normal and pathological human serum Additional materials may be produced for specific recovery experiments by the addition of a reference serum to an analyte-free serum matrix								
<b>Number of Distributions per year:</b>	4								
<b>Number of Samples per Distribution:</b>	3								
<b>Frequency of Distributions:</b>	Every three months as outlined in the <b>Distribution Schedule</b>								
<b>Schedule of Analysis:</b>	<b>Data entry</b> 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								
<b>Data Analysis:</b>	Qualitative responses are recorded for each analyte and assessed in relation to the designated response. Reports show method or kit related statistics in terms of Method Laboratory Trimmed Mean (MLTM) and range of results reported								
<b>Performance Scoring:</b>	MI scoring								
<b>Criteria of Performance:</b>	Laboratory performance is assessed over a running analytical window of 4 Distributions (12 months)  The categories of performance are: <table><thead><tr><th></th><th><u>Total MIS</u></th></tr></thead><tbody><tr><td>Good</td><td>Zero</td></tr><tr><td>Adequate</td><td>1</td></tr><tr><td>Poor</td><td>&gt;1</td></tr></tbody></table>		<u>Total MIS</u>	Good	Zero	Adequate	1	Poor	>1
	<u>Total MIS</u>								
Good	Zero								
Adequate	1								
Poor	>1								
<b>Persistent Poor Performance:</b>	Defined as being in the Poor Performance category for two or more successive Distributions								