## Ultrasensitive C-Reactive Protein (uCRP)

Accreditation Status:	UKAS Schedule of Accreditation	
Date Scheme started:	1999	
Clinical Applicability:	Monitoring of the acute phase response in neonates. Prognostic indicator of cardiovascular disease and risk assessment for coronary artery disease	
Analytes:	Ultrasensitive C-Reactive Protein (SER/028)	
Units for Reporting:	mg/L	
Samples Distributed:	Liquid format. Normal and pathological human serum	
	Additional materials may be produced for specific recovery experiments by the addition of purified CRP to an analyte-free serum matrix	
Number of Distributions per year:	12	
Number of Samples per Distribution:	2	
Frequency of Distributions:	Every month as outlined in the Distribution Schedule	
Schedule of Analysis:	<b>Data entry</b> is via the web for the submission of results. Data analysis is commenced 14 days after sample dispatch. Late returns are accepted and will contribute to the laboratory's cumulative performance statistics	
Data Analysis:	All Laboratory Trimmed Mean (ALTM) with truncation at 2SD, SD, and CV%. Reports also show method specific statistics. Individual laboratory performance is expressed in terms of MRBIS, SDBIS, and MRVIS	
	Chosen Coefficient of Variation for Ultrasensitive C-Reactive Protein is 8%	
Performance Scoring:	MRVIS	
Criteria of Performance:	Laboratory performance is classified in terms of the MRVIS over a running analytical window of 12 Distributions (12 months)	
	Ideal MRVIS Good Adequate Poor	<50 50 - 100 101 - 200 >200 or SDBIS >200
Persistent Poor Performance:	Defined as being in the Poor Performance category for two or more successive Distributions	