REGISTRATION INSTRUCTIONS & RIQAS POLICIES

CRITERIA FOR PARTICIPATION

This programme is available to any laboratory running the Cardiac assays listed in this document. Quantitative results will be accepted on this programme.

INTRODUCTION

Method questionnaires are available for all routine RIQAS Programmes and are reviewed and updated every month, as indicated by the issue date at the bottom of every page. They are designed to allow you to register for this RIQAS Programme and to inform you of RIQAS protocols and policies. It is important that you read and understand all the information in these introductory pages before completing the enrolment document, which forms the basis of your registration and contract with RIQAS. If you have any questions or concerns about any of the information presented in this document, please contact RIQAS either directly or through your local Randox Laboratories representative.RIQAS Calendar dates and information about the RIQAS portfolio of products can be found on www.randox.com/external-quality-assessment.

REGISTRATION INSTRUCTIONS

NOTE: IF A REGISTERED PARTICIPANT DOES NOT PARTICIPATE FOR A CYCLE, THEY WILL BE EXPECTED TO COMPLETE NEW ENROLMENT DOCUMENTS IN ORDER TO RE-JOIN THE PROGRAMME.

METHOD QUESTIONNAIRE:- To be retained by participant

This method questionnaire should be completed and retained by you for your records. Please ensure that you complete the method questionnaire in full. Your details will help us to classify your results correctly and thus provide you with useful statistical data.

In order to fully complete this questionnaire you will also need a copy of the RIQAS Instruments and Reagent Suppliers which is available to download from the Randox website (www.randox.com/external-quality-assessment.com). Please ensure you have this list available when completing this questionnaire.

Following this introduction section is the method questionnaire which indicates the method codes available for each parameter along with the standard RIQAS unit. On the method questionnaire, for each parameter you wish to run, please tick the method appropriate to you, then state your instrument code, reagent code, and the units that you use in your laboratory if they are different from the RIQAS standard units. If codes are not available for your assay, please state the details of your method clearly in the section at the end of the enrolment document.

NB. For enzymes, it is important for you to record the temperature at which the assay is performed.

Once your method questionnaire has been completed, you must transfer the information onto your enrolment document.

ENROLMENT DOCUMENT:- To be returned to RIQAS

Please be aware that it may take up to 3 weeks to process enrolment documents if you are not entering your own assay details. When registering RIQAS enrolment documents, it is recommended that you state business contact details, rather than personal.

A. LABORATORY REFERENCE NUMBER

On receipt of an enrolment document, each participant is assigned a **laboratory reference number** which consists of a **participant number** which is unique to your laboratory and a **registration letter** which is assigned for each new registration we receive from you. If you are a current or previous participant, please state your **participant number** on the enrolment document. If you do not have a Laboratory Reference Number, this will be generated by RIQAS when you register for the first time. Please quote this number on all correspondence with RIQAS.

B. GROUP REPORTS AND MULTIPLE REGISTRATIONS

Assessment of the same parameters on multiple systems - It is possible to enrol multiple instruments within your laboratory, up to five instruments per programme (volume permitting) can be added at no extra cost for comparative performance assessment. Kindly complete separate enrolment documents for each instrument clearly identifying each instrument in the box provided. A complementary instrument group report is supplied if you have returned results for more than one registration of the same programme. If you intend to enrol laboratories at different sites or if you are part of a group of laboratories, an inter-laboratory group report for each sample can be supplied on receipt of a completed authorisation form from each registered laboratory. Please contact RIQAS for a copy of the official inter-laboratory authorisation form.

C. CYCLE/PRODUCT REQUIREMENTS

Please tick the cycles you wish to subscribe for. If there is more than one kit/product offered for the programme, please also tick the kit you wish to subscribe for.

D. PRIMARY CONTACT DETAILS

It is important to state the full address details of the Quality Assessment Officer or contact person who will receive all correspondence during the cycle. Please also state the company name of the Randox representative who is supplying you with the RIQAS product under 'Randox Office/Distributor'
Please inform RIQAS of any change to contact details as soon as possible.

E. RIQAS**Net**

RIQASNet is a web-based online method for result entry / method changes and additions of parameters / viewing of released reports. To access RIQASnet go to www.riqas.net. Internet access and login details are required for RIQASNet and Adobe Reader is required for viewing reports. Your initial login information and password will be supplied by RIQAS. Once you have logged in for the first time you will be able to change your RIQASNet password. If you forget your password please follow the 'Forgotten Password' link. Your login information will be based on the 1st email address you supply on your enrolment document. A PDF copy of the report will be sent to this address and can also be sent to 2 other email addresses. These addresses should be stated on your enrolment document.

F. PDF REPORTS

Reports are sent as PDF files. These files can be sent to up to 3 email addresses. Adobe Reader is required to view the reports. The email addresses to which reports are sent can be reviewed and changed on RIQASNet.

G. SUMMARY CSV FILES

Labs can register to receive a csv file which contains a summary of your routine report statistics and performance indicators. This file mirrors the information found on the summary page of your report, except that we have included the calculated SD, SDPA and z-score. Also the PERFORMANCE column will show * in place of the red triangle usually shown on the summary page of your routine report. This can be sent to the 3 email addresses registered to receive the pdf reports. If you wish to receive a summary csv file please indicate this by ticking the box on the enrolment document and include the email addresses to which the reports should be sent. CSV files are also available for Instrument and Inter-Laboratory group reports. Please contact RIQAS for further information.

H. CUSTOMER DECLARATION

The declaration indicates that by submitting your enrolment document to RIQAS, either directly or via your local Randox representative, you have read and understood the RIQAS policies stated in the most recent Method Questionnaire associated with this programme. You understand that the submission of your enrolment document to RIQAS marks the beginning of an on-going agreement, and you will be automatically enrolled in subsequent cycles of this programme until we receive written confirmation of your cancellation. This should be received 12 weeks prior to the month in which the cycle starts. You understand that you must inform RIQAS of any changes to your contact details, assay details or contract status. You authorise Randox Laboratories Ltd. to send communication related to the products and service provided to the e-mail or postal addresses stated on your submitted enrolment document. You understand that you are permitted to request disclosure of, change or erase personal details held by Randox Laboratories Ltd. at any time. Note: Method questionnaires are updated every month and the issue date is stated on every questionnaire and enrolment document.

I. REGISTRATION OF ASSAY DETAILS

Labs can register their assay details using RIQASNet or can complete the 'Registration of Assay Details' section of the enrolment document. Labs should tick the appropriate box under the 'Registration of Assay Details' section of the enrolment document. If a lab wishes RIQAS to register their assay details, they should complete the Registration of Assay Details section using the codes from this method questionnaire and the Instrument/Reagent Supplier Book.

Once a participant has registered they will receive an email containing their RIQASNet login information. Once you have successfully logged in to RIQASNet you will see your various laboratory reference numbers for each registered programme. If you have opted to add parameters/assay details using RIQASNet, please do so as soon as possible (see below).

If no code is available for your assay, please state the details of your method clearly in the section at the end of the enrolment document or follow the instructions on RIQASNet.

For Ortho-Clinical Diagnostics VITROS registrations, please state the 2 digit slide Generation number for each analyte.

If units other than the standard RIQAS units are used, please specify these in the boxes supplied.

ONCE COMPLETED, THE ENROLMENT DOCUMENT SHOULD BE SENT TO RIQAS **FOR REGISTRATION.**

J. UPDATING ASSAY DETAILS

It is possible to change your unit, method, instrument or reagent classification during a cycle.

Method changes via RIQASNet: These can be made in the Assay Details section of the Data Entry menu. A list of your registered laboratory reference numbers will appear on screen. Select the laboratory reference number for which you would like to change the assay details. A current list of assay details will appear, click on the appropriate parameter. To change the details click the arrow box on the appropriate details and select a new one. Save the changes and submit them to RIQAS. Changes will not be instantaneously updated on RIQASNet but will be uploaded onto RIQASNet usually within 3 working days. It is possible to submit results and method changes together as method changes will be made before results are entered in to the RIQAS database.

K. ADDITION OF PARAMETERS / ASSAY DETAILS

Adding Parameters via RIQASNet: Parameters can be added using the Assay Details section of the Data Entry menu. A list of your registered laboratory reference numbers will appear on screen. Select the laboratory reference number for which you would like to add the assay details. At the top of the screen is 'Add Parameter'. Click on this and a list of parameters you are not registered for will appear. Select the parameter you wish to add and click the arrow box on the appropriate details and select your assay details. Save the changes and submit them to RIQAS. As above, additions will be available on RIQASnet usually within 3 working days.

ORDERING RIQAS PRODUCTS

Please ensure your purchase order for each cycle is placed with your local Randox representative 12 weeks prior to the month in which the cycle starts. This will ensure sufficient time to process and despatch your kit(s) to you. Participants from UK or Ireland may order products directly from RIQAS with an official order number. Orders received within 12 weeks of the start of the cycle will be processed with an additional administration fee. Current prices of RIQAS products are available from your local Randox Laboratories representative.

It may be possible to order RIQAS products during a cycle, subject to availability. Please ask your local Randox representative to check availability before completing the

SHIPPING AND RECEIPT OF RIQAS PRODUCTS

Provided that you have ordered sufficiently in advance, your RIQAS kit(s) will be shipped to you to arrive before the analysis date of the first sample in the kit. If you do not receive your kit(s) before this time, please contact your local Randox representative.

On RIQASNet please access your account and download the relevant Instructions For Use (IFU) document for the programme and cycle purchased. The IFU includes material characteristics, preparation, stability, storage and safety information. On receipt of your RIQAS kit, please check that:

a) it is the product you ordered
b) the correct number of samples are present as indicated on the IFU
c) the samples have the appearance as indicated on the IFU and that none of them are damaged

Please notify your local Randox representative immediately if any of these are incorrect.

Please ensure that the product is immediately stored according to the recommendations on the package labelling.

ASSAY OF SAMPLES & RETURN OF RESULTS

Carefully read the instructions stated on the Instructions for Use (IFU) prior to preparation and assay of RIQAS samples. **These are available on RIQASNet only**. The RIQAS samples should be assayed at the recommended time specified on the IFU. Following appropriate preparation, samples should be treated as routine, unless otherwise stated on the IFU. Please assay the samples on or before the recommended date for analysis and forward your results to RIQAS by no later than **17:00 GMT on the FINAL DATE**, as indicated in the IFU. Results are submitted via RIQASNet, which can be accessed once you have received log in details via email. This will include a link to RIQASNet Instructions for Use.

LATE AND CORRECTED RESULTS

In keeping with the objectives of EQA schemes, participants should be aware that collusion and falsification of results is considered to be unethical and constitutes scientific fraud. RIQAS policies must ensure that a laboratory is unaware of RIQAS means for comparison before submitting their own results. Where a result is not submitted by the final date, a report will be issued, but the missing results will be indicated as “No return” or “N” throughout the RIQAS reports. RIQAS permits the submission of late or corrected results only under the circumstances described below. Requests for the submission of late or corrected results must be submitted in writing and in English on RIQAS Form No. 9277-RQ (either by the participant or their local Randox Representative) and must be approved by RIQAS Management. The form is available on www.riqas.net.

Requests for the submission of late results must be accompanied by evidence that an error has been made, and that the error has not been caused by the participant.

Requests for the correction or removal of erroneous results must be accompanied by evidence that the error was non-analytical, as defined on form 9277-RQ. RIQAS is obliged to inform country-specific regulatory bodies of requests for correction of results (if they request such information for laboratory monitoring purposes).

New reports will be re-issued for late or corrected results only where there has been an error made by Randox Laboratories HQ, Randox representatives or distributors.

LATE RESULTS

In general, late results will not be accepted after the final date.

Late results will only be accepted where there has been an error made by Randox Laboratories HQ, Randox representatives or distributors.

CORRECTED RESULTS

Laboratories may correct results only if it can be determined that the error was non-analytical and where the request for submission is within 4 weeks of the original final date. A laboratory may correct a result under the following circumstances:

- Reconstituting a sample in an incorrect volume before analysis
- Assaying and/or submitting the results for the wrong sample
- Making a transcription error - submission of an analyser print-out indicating that the analysis date was before the final date is required.

DESPATCH OF REPORTS
PDF reports will be emailed within 72 hours of the FINAL DATE and for those registered for RIQASNet the PDF reports will be available on RIQASNet shortly after.

END OF CYCLE REPORTS
At the end of a cycle, a summary report will be issued to all participants. This includes a summary page for each parameter, an Average Absolute SDI report and a Cerificate of Acceptable performance (see below).

USE OF RIQAS REPORTS
Participants have permission to make copies of their RIQAS reports for internal use and for regulatory purposes only. RIQAS reports must not be duplicated for external use without permission from the RIQAS Scheme Co-ordinator. Under no circumstances should information on RIQAS reports be taken out of context or falsified in any way. Information regarding the format of RIQAS Reports and the monitoring of EQA performance can be found in the RIQAS Brochure on www.randox.com/external-quality-assessment. Information regarding the calculations and scores used to evaluate participants' performance on RIQAS Reports can be found following log in to RIQASNet, in a document entitled "Evaluation of Performance".

CONFIDENTIALITY
Participation in any RIQAS programme is considered to be strictly confidential. Any data transfer or correspondence with participants, either directly or via local Randox representative, will be deemed confidential. Participants should be aware that regulatory authorities have the right to request an assessment of a participant's performance. Where regulatory authorities are to be provided with a participant's results, participants will be notified.

GENERAL DATA PROTECTION REGULATION 2018 & UK DATA PROTECTION ACT 2018
Randox Laboratories Ltd. complies with GDPR and the UK Data Protection Act and holds the minimum information required to maintain the contract with RIQAS customers. Contact details are required in order to effectively provide you with the RIQAS products and services. Participants are not under any obligation to provide personal information to enter into a contract with RIQAS. We recommend that business contact details are provided. All data associated with the provision of RIQAS is collated, stored and processed confidentially and securely, to avoid unlawful processing, accidental loss or damage.

CERTIFICATES OF PARTICIPATION
Complimentary certificates of participation for each RIQAS programme are made available on RIQASNet to participants at the **end of the current cycle**, provided that **at least 50%** of results have been returned. Participants who enrol mid-cycle will be eligible for a Certificate for Participation if they have participated in at least 50% of samples available for the remainder of the cycle since enrolment. The certificate will specify the cycle, programme and the LABORATORY / HOSPITAL NAME which is detailed in the certificate section of RIQASNet. At the end of a cycle, a list of all eligible labs will be exported from RIQASNet and certificates will be created according to these details. Please ensure all certificate details are up to date in your RIQASNet account.

CERTIFICATE OF ACCEPTABLE PERFORMANCE
Participants are also provided with a Certificate of Acceptable Performance within their End-of-Cycle report. Acceptable performance is considered to be a Cycle Average Absolute SDI of less than 2. While all participants receive an end-of-cycle report, participants (including those who enrol mid-cycle) are only eligible for Certificates of Performance if they have returned more than half of the samples in a full cycle.

PERFORMANCE SURVEILLANCE OF UK LABS
RIQAS is obligated to identify and report persistent poor performing UK labs to the National Quality Assessment Advisory Panel. Poor performers are identified as those failing to meet performance criteria agreed with NQAAP. The performance criteria is specified in all performance surveillance correspondence with participants, and is also available on request. Participants are initially informed of poor performance by letter. Failure to improve performance will prompt details to be forwarded to NQAAP. All information sent to participants and NQAAP is strictly confidential. Please contact RIQAS if you require further information on Performance Surveillance.

PARTICIPANT FEEDBACK, COMPLAINTS & APPEALS
In order to ensure that RIQAS provides an appropriate and satisfying service, participants are invited to complete a feedback survey on RIQASNet. You may contact us at any time during the cycle, should you have any requests for additional programmes or parameters or comments regarding existing programmes.

RIQAS makes every effort to ensure that the samples provided are clinically challenging to as many laboratory systems as possible. For details, please contact RIQAS either directly or through your local Randox representative.

Should the need arise, participants may raise requests or enquiries through correspondence with the local Randox Laboratories representative or by contacting RIQAS directly. Participants may appeal against the evaluation of their performance by completing a PARTICIPANT APPEALS FORM, 10770-RQ. Participants may raise a complaint in relation to the product or service provided by completing the PARTICIPANT COMPLAINTS FORM, 10772-RQ. These forms are available on RIQASNet, or on request from RIQAS.

SUB-CONTRACTING
RIQAS sub-contracts aspects of the scheme. RIQAS accepts responsibility for the sub-contractors' work and protocols are in place to ensure that sub-contractors are deemed competent.

OUR COMPETENCE AS A PROFICIENCY TESTING PROVIDER
On request, RIQAS is willing to co-operate with participants seeking evidence of our competence as a proficiency testing provider or information on the design and implementation of RIQAS Programmes.

DEVIATION FROM EXISTING POLICIES/SERVICE
If there is any deviation from the existing policies or service, participants will be notified either directly or via their local Randox representative.

COMMUNICATION
As part of the service provided by Randox Laboratories Ltd., participants may be contacted by e-mail regarding updates and new products, in line with Randox Laboratories Ltd. privacy policy, as stated in www.randox.com.

This programme is accredited by UKAS TO ISO/IEC 17043:2010 via Fixed Scope

Please contact RIQAS at
Tel: +44 (0) 28 9445 4399
E-Mail mail@riqas.com
RIQAS Scheme Co-ordinator: Sarah Fleck
RANDOX LABORATORIES LTD., 55 Diamond Road, Crumlin, County Antrim, BT29 4QY, United Kingdom

UKAS PROFICIENCY TESTING
0010

CK, Total U/l

CODE		METHOD
CKTD	☐	Dithioerythritol (DTE)
CKDIF	☐	Dithioerythritol (DTE) IFCC correlated
CKTM	☐	Monothioglycerol
CKTIF	☐	CK-NAC (IFCC)
CKTSE	☐	CK-NAC serum start
CKTSU	☐	CK-NAC substrate start
CKTDC	☐	Ortho Vitros MicroSlide Systems
CKTDT	☐	Vitros DT60/DT60 II/DTSC II
		Vitros Slide Generation Number ☐☐
		Other methods, please specify on enrolment document

INSTRUMENT CODE ☐

REAGENT CODE ☐

RESULTS REPORTED AT 25°C ☐ 30°C ☐ 37°C ☐

OTHER UNITS, SPECIFY ☐

CK-MB, Activity U/l

CODE		METHOD
CKMAB	☐	Abbott Immunoinhibition
CKMIF	☐	Immunoinhibition (IFCC)
CKMSE	☐	Immunoinhibition, serum start
CKMSU	☐	Immunoinhibition, substrate start
CKMDC	☐	Ortho Vitros MicroSlide Systems
CKMDT	☐	Vitros DT60/DT60 II/DTSC II
		Vitros Slide Generation Number ☐☐
		Other methods, please specify on enrolment document

INSTRUMENT CODE ☐

REAGENT CODE ☐

RESULTS REPORTED AT 25°C ☐ 30°C ☐ 37°C ☐

OTHER UNITS, SPECIFY ☐

RQ9127 - CARDIAC PROGRAMME
METHOD QUESTIONNAIRE

CK-MB, Mass ng/ml

CODE		METHOD
CKAAI	☐	Abbott Alinity
CKARC	☐	Abbott Architect
CKABX	☐	Abbott Axsym
CKABB	☐	Abbott IMx
CKAIS	☐	Abbott i-STAT
CKALO	☐	Autobio CLIA
CKSAN	☐	Beckman Access
CKDXI	☐	Beckman DxI 600 / 800
CKBHM	☐	Beijing Hotgen MQ60
CKVID	☐	bioMerieux, VIDAS
CKBIC	☐	Boditech ichroma
CKCAX	☐	Cormay Auryx ECLIA
CKCXS	☐	Cormay Auryx CK-MB STAT
CKLIA	☐	Diasorin Liaison
CKLIX	☐	Diasorin Liaison XL
CKFIA	☐	Finecare FIA
CKFJL	☐	Fujirebio Lumipulse G Series
CKGEI	☐	Getein 1100 IF Quan Analyser
CKGE1	☐	Getein 1600 Analyzer
CKGEF	☐	Getein FIA8000 Analyser
CKHIT	☐	Hitachi
CKMIP	☐	Micropoint CK-MB
CKAQT	☐	Radiometer AQT90 Flex
CKEV	☐	Randox Evidence / Investigator
CKMBS	☐	Mindray BS Series
CKMC2	☐	Mindray CL Series
CKMP	☐	Mitsubishi Pathfast
CKVEC	☐	Ortho Vitros 3600/5600/ECi/XT 7600
CKTRI	☐	Quidel Triage Meter Plus
CKEVE	☐	Randox Evolution
CKRX	☐	Randox RX Series
CKRAM	☐	Response Biomedical Ramp
CKRCR	☐	Roche Cardiac Reader
CKEY8	☐	Roche Cobas e402/e801
CKRH	☐	Roche Cobas h232
CKEYS	☐	Roche Elecsys / E170 / c6000 / e411
CKSL	☐	Samsung Labgeo IB10
CKSDC	☐	SD Biosensor CKMB FIA
CKSAI	☐	Siemens Atellica IM
CKCC	☐	Siemens ACS180
CKBAY	☐	Siemens Immuno 1
CKCEN	☐	Siemens Centaur
CKDD	☐	Siemens Dimension
CKDO	☐	Siemens Opus
CKDPI	☐	Siemens Immulite 1000
CKDP2	☐	Siemens Immulite 2000
CKDP5	☐	Siemens Immulite 2500
CKSYU	☐	Shenzhen YHLO UNICELL CK-MB
CKSIS	☐	Siemens Stratus CS
CKSNM	☐	SNIBE Maglumi Analysers
CKSNM2	☐	SNIBE Maglumi Analysers II
CKTSC	☐	Tisenc Accre CLIA
CKTOC	☐	TOSOH AIA CL-Series
CKTOS	☐	Tosoh AIA Series
CKXBA	☐	Xiamen Biotime FIA Analysers

Other methods, please specify on enrolment document

INSTRUMENT CODE ☐

REAGENT CODE ☐

OTHER UNITS, SPECIFY ☐

RQ9127 - CARDIAC PROGRAMME
METHOD QUESTIONNAIRE

HOMOCYSTEINE µmol/l

CODE		METHOD
HOAAI	☐	Abbott Alinity
HOARC	☐	Abbott Architect
HOABX	☐	Abbott Axsym
HOABB	☐	Abbott IMx
HOALO	☐	Autobio CLIA
HOAXS	☐	Axis Shield
HOAU4	☐	Beckman AU Instruments
HOBEK	☐	Beckman Synchron LX20
HOBIO	☐	Biosino Bio-Technology
HODIA	☐	Diazyme Homocysteine
HOFH	☐	Fleg Healthcare Reagents
HOGES	☐	Gesan Reagents
HOHIT	☐	Hitachi
HOHP	☐	HPLC
HOACLE	☐	IL ACL Elite/Elite Pro/8/9/10000
HOLM	☐	LC/MS
HOEIA	☐	Manual - E.I.A.
HOVEC	☐	Ortho Vitros 3600/5600/Eci/XT 7600
HOREA	☐	Randox Enzymatic Assay
HORAM	☐	Response Biomedical Ramp
HOC6	☐	Roche Cobas c501/502/503/303/311/701/702
HOINT	☐	Roche Integra
HOSBS	☐	Snibe Bioassays
HOSNM	☐	Snibe Maglumi
HOSAI	☐	Siemens Atellica IM
HOCEN	☐	Siemens Centaur
HONEP	☐	Siemens Nephelometer
HODP2	☐	Siemens Immulite 2000 / 2500
HOTOS	☐	Tosoh AIA Series
HOW3V	☐	Weifang 3V Reagents

Other methods, please specify on enrolment document

INSTRUMENT CODE ☐

REAGENT CODE ☐

OTHER UNITS, SPECIFY ☐

RQ9127 - CARDIAC PROGRAMME
METHOD QUESTIONNAIRE

MYOGLOBIN ng/ml

CODE		METHOD
MYAAQ	☐	Abbott Alinity c
MYAAI	☐	Abbott Alinity i (STAT)
MYARQ	☐	Abbott Architect c (QUANTIA)
MYARC	☐	Abbott Architect i (STAT kit)
MYABX	☐	Abbott Axsym
MYALO	☐	Autobio CLIA
MYSAN	☐	Beckman Access
MYOLY	☐	Beckman AU Instruments
MYDXI	☐	Beckman DxI 600 / 800
MYBHM	☐	Beijing Hotgen MQ60
MYBIM	☐	Boditech ichroma Myoglobin
MYVID	☐	bioMerieux, VIDAS
MYBIZ	☐	Biozek DCR 1000
MYCAX	☐	Cormay Auryx ECLIA
MYCXS	☐	Cormay Auryx Myoglobin STAT
MYLIA	☐	Diasorin Liaison
MYLIX	☐	Diasorin Liaison XL
MYFIA	☐	Finecare FIA
MYFJL	☐	Fujirebio Lumipulse G Series
MYGEI	☐	Getein 1100 IF Quan Analyser
MYGE1	☐	Getein 1600 Analyzer
MYGEF	☐	Getein FIA8000 Analyser
MYSLT	☐	Lifotronic eCL
MYMIP	☐	Micropoint Myoglobin
MYMIN	☐	Mindray BS Series
MYMC2	☐	Mindray CL Series
MYMP	☐	Mitsubishi Pathfast
MYVEC	☐	Ortho Vitros 3600/5600/ECi/XT 7600
MYTRI	☐	Quidel Triage Meter Plus
MYAQT	☐	Radiometer AQT90 Flex
MYRDX	☐	Randox Daytona / Imola
MYEV	☐	Randox Evidence / Investigator
MYEVE	☐	Randox Evolution
MYRAM	☐	Response Biomedical RAMP
MYRCR	☐	Roche Cardiac Reader
MYEY8	☐	Roche Cobas e402/e801
MYRH	☐	Roche Cobas h232
MYEYS	☐	Roche Elecsys / E170 / e601 / e602 / e411
MYHIT	☐	Roche Hitachi/Cobas c303/311/501/502/503
MYINT	☐	Roche Integra
MYSL	☐	Samsung Labgeo IB10
MYSA	☐	Siemens Advia 1200/1650/1800/2400
MYSAI	☐	Siemens Atellica IM
MYCEN	☐	Siemens Centaur
MYDD	☐	Siemens Dimension
MYBN	☐	Siemens Nephelometer
MYDO	☐	Siemens Opus
MYDPI	☐	Siemens Immulite 1000
MYDP2	☐	Siemens Immulite 2000
MYDP5	☐	Siemens Immulite 2500
MYST	☐	Siemens Stratus CS
MYSMI	☐	Snibe Maglumi analysers
MYSMI2	☐	Snibe Maglumi analysers II
MYTOC	☐	TOSOH AIA CL-Series
MYTOS	☐	Tosoh AIA Series
MYXBA	☐	Xiamen Biotime FIA Analysers

Other methods, please specify on enrolment document

INSTRUMENT CODE ☐

REAGENT CODE ☐

OTHER UNITS, SPECIFY ☐

RQ9127 - CARDIAC PROGRAMME
METHOD QUESTIONNAIRE

TROPONIN I ng/ml

CODE		METHOD
TIAAI	☐	Abbott Alinity i (STAThs)
TIARC	☐	Abbott Architect STAT
TIARS	☐	Abbott Architect STAT hs
TIABX	☐	Abbott Axsym
TIABA	☐	Abbott Axsym ADV
TIAIS	☐	Abbott i-STAT
TIABC	☐	Autobio CLIA
TISA2	☐	Beckman Access - A78803
TISA3	☐	Beckman Access - AccuTnI+3
TISAH	☐	Beckman Access 2 / DxC600i Hs
TISAN	☐	Beckman Access Ref 33340
TIDX2	☐	Beckman DxI - A78803
TIDX3	☐	Beckman DxI - AccuTnI+3
TIDX9	☐	Beckman DxI 9000
TIDXH	☐	Beckman DxI Hs
TIDXI	☐	Beckman DxI Ref 33340
TIBHM	☐	Beijing Hotgen MQ60
TIVIH	☐	BioMerieux VIDAS hs Troponin I
TIVIU	☐	BioMerieux VIDAS Ultra
TIBAF	☐	Boditech AFIAS Tn-I Plus
TIBIC	☐	Boditech ichroma Tn-I
TICAX	☐	Cormay Auryx ECLIA
TILIA	☐	Diasorin Liaison
TILIX	☐	Diasorin Liaison XL
TIEC3	☐	Edan C3 TnI Rapid Test
TIEUR	☐	Eurolyser
TIFIA	☐	Finecare FIA
TIFJL	☐	Fujirebio Lumipulse G Series
TIGEI	☐	Getein 1100 IF Quan Analyser
TIGE1	☐	Getein 1600 Analyzer
TIGEF	☐	Getein FIA8000 Analyser
TIGE2	☐	Getein hs-cTnI Fast Test Kit
TIHCT	☐	Hipro cTnI Test
TISLT	☐	Lifotronic eCL
TILFT	☐	Lifotronic Troponin I
TIMTC	☐	Medcaptain Troponin I CLIA
TIMIP	☐	Micropoint Troponin I
TIMC2	☐	Mindray CL 8000i/6000i/2000i/1200i/1000i
TIMC3	☐	Mindray CL 900i
TIMC4	☐	Mindray CL 8/6/2/12/1000i Hs
TIMC5	☐	Mindray CL 900i Hs
TIMP	☐	Mitsubishi Pathfast
TIMPS	☐	Mitsubishi Pathfast Hs
TIVEC	☐	Ortho Vitros
TIVE2	☐	Ortho Vitros hs TnI
TITRI	☐	Quidel Triage Meter Plus
TIAQT	☐	Radiometer AQT90 Flex
TIEV	☐	Randox Evidence / Investigator

CODE		METHOD
TIRAM	☐	Response Biomedical Ramp
TIEYS	☐	Roche Elecsys / E170 / c6000 / e411
TIE8	☐	Roche Elecsys Troponin I
TISL	☐	Samsung Labgeo IB10
TISDS	☐	SD Biosensor TnI FIA
TISBT	☐	Shenzhen Superbio Troponin I
TISYU	☐	Shenzhen YHLO UNICELL cTnI
TISAI	☐	Siemens Atellica IM
TISAV	☐	Siemens Atellica VTLi HS
TICC	☐	Siemens ACS 180
TIBAY	☐	Siemens Immuno1
TICEN	☐	Siemens Centaur
TICEH	☐	Siemens Centaur TNIH
TIDD	☐	Siemens Dimension
TIDDE	☐	Siemens Dimension Exl LOCI
TIDDEH	☐	Siemens Dimension Exl LOCI Hs
TIDDV	☐	Siemens Dimension Vista LOCI
TIDDVH	☐	Siemens Dimension Vista LOCI Hs
TIDO	☐	Siemens Opus
TIDPI	☐	Siemens Immulite 1000
TIDP2	☐	Siemens Immulite 2000
TIDP5	☐	Siemens Immulite 2500
TIST	☐	Siemens Stratus CS
TISNM	☐	SNIBE Maglumi Analysers
TISNM2	☐	SNIBE Maglumi Analysers II
TISNMH	☐	SNIBE Maglumi hs-cTnI
TITSC	☐	Tisenc Accre CLIA
TITOC	☐	Tosoh AIA CL Series
TITOS	☐	Tosoh AIA Series
TIVC1	☐	Vedalab Check-1
TIWSL	☐	Wondfo SmarLumi Series
TIXBB	☐	Xiamen Biotime cTnI Rapid Test

Other methods, please specify on enrolment document

INSTRUMENT CODE ☐

REAGENT CODE ☐

OTHER UNITS, SPECIFY ☐

RQ9127 - CARDIAC PROGRAMME
METHOD QUESTIONNAIRE

TROPONIN T ng/ml

CODE		METHOD
TTALO	☐	Autobio CLIA
TTACM	☐	Aikang CLIA-mate TnT Kit
TTCXS	☐	Cormay Auryx hs Troponin T STAT
TTFIN	☐	Finecare FIA
TTSLT	☐	Lifotronic eCL
TTNPB	☐	Nanjing Poclight c5000
TTAQT	☐	Radiometer AQT90
TTRCR	☐	Roche Cardiac Reader
TTRH	☐	Roche Cobas h232
TTEY8	☐	Roche Cobas e402/e801 TnT kits
TTRH8	☐	Roche Cobas e402/e801 TnT hs kits
TTRHS8	☐	Roche Cobas e402/e801 TnT hs STAT
TTRH86	☐	Roche cobas e402/e801 TnT hs STAT Gen 6
TTEYS	☐	Roche Cobas Troponin T kits
TTRHS	☐	Roche Cobas Troponin T hs kits
TTRHSS	☐	Roche Cobas Troponin T hs STAT
TTSH	☐	Sysmex HISCL
TTWSL	☐	Wondfo SmarLumi Series
TTTSC	☐	Wondfo Tisenc Accre CLIA

When your RIQAS sample gives a result of less than 0.1 ng/ml or less than 0.03 ng/ml ("negative", "<0.1" or "< 0.03"), please submit these values (<0.03 or <0.1)

Other methods, please specify on enrolment document

INSTRUMENT CODE ☐

REAGENT CODE ☐

OTHER UNITS, SPECIFY ☐