

REGISTRATION INSTRUCTIONS & RIQAS POLICIES

CRITERIA FOR PARTICIPATION

This programme is available to any laboratory running the Maternal Screening 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). 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.

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. RIQASNet

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. These are available on RIQASNet. 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 order/enrolment document.

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 Certificate 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



0010

RQ9137 - MATERNAL SCREENING PROGRAMME

METHOD QUESTIONNAIRE

AFP - kU/l

CODE	METHOD
AFAAI	<input type="checkbox"/> Abbott Alinity
AFARC	<input type="checkbox"/> Abbott Architect
AFABX	<input type="checkbox"/> Abbott AxSYM / AxSYM Plus
AFAKB	<input type="checkbox"/> Alkor Bio EIA-AFP
AF-AIC	<input type="checkbox"/> Aptasys Indra CLIA
AFABC	<input type="checkbox"/> Autobio CLIA
AFSAN	<input type="checkbox"/> Beckman Access / LXi725
AFDXI	<input type="checkbox"/> Beckman Dxl 600 / 800
AFBHM	<input type="checkbox"/> Beijing Hotgen MQ60
AFBIV	<input type="checkbox"/> bioMerieux Vidas / Mini Vidas
AFBMR	<input type="checkbox"/> Bio-rad Microplate Reader
AFBMI	<input type="checkbox"/> Boditech Med Inc i-CHROMA
AFBRR	<input type="checkbox"/> Brahms Kryptor
AFCII	<input type="checkbox"/> CIS IRMA
AFCAX	<input type="checkbox"/> Cormay Auryx ECLIA
AFDME	<input type="checkbox"/> DiaMetra ELISA
AFETI	<input type="checkbox"/> Diasorin ETI-MAX 3600
AFLIA	<input type="checkbox"/> Diasorin Liaison
AFLIX	<input type="checkbox"/> Diasorin Liaison XL
AFDIS	<input type="checkbox"/> DIAsource IRMA
AFDRE	<input type="checkbox"/> DRG Elisa
AFCE	<input type="checkbox"/> Fujirebio CanAg EIA
AFGM	<input type="checkbox"/> Gamma Counter
AFGBE	<input type="checkbox"/> General Biologicals ELISA
AFHMC	<input type="checkbox"/> Human HumaCLIA SR
AFHUM	<input type="checkbox"/> Human Humanreader
AFIMI	<input type="checkbox"/> Immunotech IRMA
AFIZI	<input type="checkbox"/> Izotop IRMA
AFMIN	<input type="checkbox"/> Mindray Series
AFOR	<input type="checkbox"/> Orqanon Reader 530
AFVEC	<input type="checkbox"/> Ortho Vitros 3600/5600/ECi/XT 7600
AFDEL	<input type="checkbox"/> Perkin Elmer Delfia Xpress / Autodelfia
AFWW	<input type="checkbox"/> Perkin Elmer Wizard
AFRCE	<input type="checkbox"/> Roche Cobas 4000/e411
AFC6	<input type="checkbox"/> Roche Cobas e601/602
AFE8	<input type="checkbox"/> Roche Cobas e402/e801
AFEYS	<input type="checkbox"/> Roche Elecsys 2010 / 1010
AFRME	<input type="checkbox"/> Roche Modular E170
AFSYI	<input type="checkbox"/> Shenzhen YHLO iFlash Series
AFSAI	<input type="checkbox"/> Siemens Atellica IM
AFCC	<input type="checkbox"/> Siemens/Bayer ACS 180
AFCE	<input type="checkbox"/> Siemens Centaur
AFDD	<input type="checkbox"/> Siemens/Dade Dimension
AFDPI	<input type="checkbox"/> Siemens/DPC Immulite 1000
AFDP2	<input type="checkbox"/> Siemens/DPC Immulite 2000 / 2500
AFDS	<input type="checkbox"/> Siemens Stratus CS
AFSNM	<input type="checkbox"/> SNIBE Maglumi analysers
AFSNM2	<input type="checkbox"/> SNIBE Maglumi analysers II
AFSF	<input type="checkbox"/> Stat Fax Elisa Readers
AFSTR	<input type="checkbox"/> Stratec Immunotech
AFSHI	<input type="checkbox"/> Sysmex HISCL Series
AFTS	<input type="checkbox"/> Tecan Sunrise
AFTOS	<input type="checkbox"/> Tosoh AIA Series
AFTOC	<input type="checkbox"/> Tosoh AIA-CL Series
AFZEN	<input type="checkbox"/> Zentech ELISA

Other methods - Please specify in document

INSTRUMENT CODE

REAGENT CODE

OTHER UNITS SPECIFY

RQ9137 - MATERNAL SCREENING PROGRAMME

METHOD QUESTIONNAIRE

HCG Free Beta - U/I

CODE	METHOD
HCBAAI	<input type="checkbox"/> Abbott Alinity
HCBARC	<input type="checkbox"/> Abbott Architect
HCBABX	<input type="checkbox"/> Abbott Axysm / Axsym Plus
HCBABC	<input type="checkbox"/> Autobio CLIA
HCBSAN	<input type="checkbox"/> Beckman Access
HCBDXI	<input type="checkbox"/> Beckman Dxl 600 / 800
HCBBIV	<input type="checkbox"/> bioMerieux Vidas / Mini Vidas
HCBBRR	<input type="checkbox"/> Brahms Kryptor
HCBCII	<input type="checkbox"/> CIS IRMA
HCBDME	<input type="checkbox"/> Demeditec ELISA
HCBETI	<input type="checkbox"/> Diasorin ETI-MAX 3600
HCBLIA	<input type="checkbox"/> Diasorin Liaison
HCBLIX	<input type="checkbox"/> Diasorin Liaison XL
HCBDRE	<input type="checkbox"/> DRG Elisa
HCBHUM	<input type="checkbox"/> Human Humanreader
HCBIBL	<input type="checkbox"/> IBL, ELISA
HCBIMI	<input type="checkbox"/> Immunotech IRMA
HCBIZI	<input type="checkbox"/> Izotop IRMA
HCBOR	<input type="checkbox"/> Organon Reader 530
HCBVEC	<input type="checkbox"/> Ortho Vitros, 3600 / 5600 / Eci
HCBDEL	<input type="checkbox"/> Perkin Elmer DELFIA Xpress
HCBDEL2	<input type="checkbox"/> Perkin Elmer DELFIA /Auto DELFIA
HCBWW	<input type="checkbox"/> Perkin Elmer Wizard
HCBAQT	<input type="checkbox"/> Radiometer AQT90 Flex
HCBRCE	<input type="checkbox"/> Roche Cobas 4000/e411
HCBC6	<input type="checkbox"/> Roche Cobas e601/602
HCBE8	<input type="checkbox"/> Roche Cobas e402/e801
HCBEYS	<input type="checkbox"/> Roche Elecsys 2010 / 1010
HCBRME	<input type="checkbox"/> Roche Modular E170
HCBSAI	<input type="checkbox"/> Siemens Atellica IM
HCBC	<input type="checkbox"/> Siemens/Bayer ACS 180/180 SE
HCBCEN	<input type="checkbox"/> Siemens Centaur
HCBDD	<input type="checkbox"/> Siemens/Dade Dimension/RxL
HCBDPI	<input type="checkbox"/> Siemens/DPC Immulite 1000
HCBDP2	<input type="checkbox"/> Siemens/DPC Immulite 2000 / 2500
HCBDS	<input type="checkbox"/> Siemens Stratus CS
HCBSNM	<input type="checkbox"/> SNIBE Maglumi analysers
HCBSNM2	<input type="checkbox"/> SNIBE Maglumi analysers II
HCBSF	<input type="checkbox"/> Stat Fax Elisa Readers
HCBSTR	<input type="checkbox"/> Stratec Immunotech
HC BTS	<input type="checkbox"/> Tecan Sunrise
HCC2	<input type="checkbox"/> Wantai Caris 200
HCW2	<input type="checkbox"/> Wantai Wan200+
HCXME	<input type="checkbox"/> Xema Medical EIA
HCTZEN	<input type="checkbox"/> Zentech ELISA

Other methods - Please speciv in document

INSTRUMENT CODE

REAGENT CODE

OTHER UNITS SPECIFY

RQ9137 - MATERNAL SCREENING PROGRAMME

METHOD QUESTIONNAIRE

HCG Total - U/l	
CODE	METHOD
HCTAAI	<input type="checkbox"/> Abbott Alinity
HCTARC	<input type="checkbox"/> Abbott Architect
HCTABX	<input type="checkbox"/> Abbott Axsym / Axsym Plus
HCAKB	<input type="checkbox"/> Alkor Bio EIA-hCG
HCTABC	<input type="checkbox"/> Autobio CLIA
HCTSAN	<input type="checkbox"/> Beckman Access
HCSA5	<input type="checkbox"/> Beckman Access Total BhCG (5th IS)
HCTAU	<input type="checkbox"/> Beckman Coulter AU 3000i
HCTDXI	<input type="checkbox"/> Beckman Dxl 600 / 800
HCDX5	<input type="checkbox"/> Beckman DXI Total BhCG (5th IS)
HCTBIV	<input type="checkbox"/> bioMerieux Vidas / Mini Vidas
HCTBMI	<input type="checkbox"/> Boditech Med Inc i-CHROMA
HCTBRR	<input type="checkbox"/> Brahms Kryptor
HCTETI	<input type="checkbox"/> Diasorin ETI-MAX 3600
HCTLIA	<input type="checkbox"/> Diasorin Liaison
HCTLIX	<input type="checkbox"/> Diasorin Liaison XL
HCDI5	<input type="checkbox"/> Drawray iStar 500
HCTIMI	<input type="checkbox"/> Immunotech IRMA
HCTIZR	<input type="checkbox"/> Izotop RIA
HCTMIN	<input type="checkbox"/> Mindray Series
HCTVEC	<input type="checkbox"/> Ortho Vitros. 3600 / 5600 / ECI
HCTDEL	<input type="checkbox"/> Perkin Elmer Delfia/Delfia Express / Autodelfia
HCTWW	<input type="checkbox"/> Perkin Elmer Wizard
HCTAQT	<input type="checkbox"/> Radiometer AQT90 Flex
HCTC6	<input type="checkbox"/> Roche Cobas hCG+Beta
HCTE8	<input type="checkbox"/> Roche hCG + Beta e402/e801
HCTEYI	<input type="checkbox"/> Roche hCG STAT(Intact)
HCTSAI	<input type="checkbox"/> Siemens Atellica IM
HCTCC	<input type="checkbox"/> Siemens/Bayer ACS 180 / 180 SE
HCTCEN	<input type="checkbox"/> Siemens Centaur
HCTDD	<input type="checkbox"/> Siemens/Dade Dimension / RxL
HCTDDV	<input type="checkbox"/> Siemens/Dade Dimension Vista
HCTDPI	<input type="checkbox"/> Siemens/DPC Immulite 1000
HCTDP2	<input type="checkbox"/> Siemens/DPC Immulite 2000 / 2500
HCTDST	<input type="checkbox"/> Siemens Stratus CS
HCTSNM	<input type="checkbox"/> SNIBE Maglumi analysers
HCTSNM2	<input type="checkbox"/> SNIBE Maglumi analysers II
HCSF	<input type="checkbox"/> Stat Fax Elisa Readers
HCTSTR	<input type="checkbox"/> Stratec Immunotech
HCTOS	<input type="checkbox"/> Tosoh AIA Series
HCTTOC	<input type="checkbox"/> Tosoh AIA-CL Series
HCTXME	<input type="checkbox"/> Xema Medical EIA
HCTZEN	<input type="checkbox"/> Zentech ELISA

Other methods - Please speciv in document

INSTRUMENT CODE

REAGENT CODE

OTHER UNITS SPECIFY

RQ9137 - MATERNAL SCREENING PROGRAMME

METHOD QUESTIONNAIRE

INHIBIN A - ng/l

CODE	METHOD
INAAAI	<input type="checkbox"/> Abbott Alinity
INAARC	<input type="checkbox"/> Abbott Architect
INABBX	<input type="checkbox"/> Abbott Axsysm / Axsym Plus
INAAE	<input type="checkbox"/> Beckman Inhibin A Active ELISA
INASAN	<input type="checkbox"/> Beckman Access
INADSX	<input type="checkbox"/> Beckman DSX
INADXI	<input type="checkbox"/> Beckman Dxl 600 / 800
INABDX9	<input type="checkbox"/> Beckman Dxl 9000
INABIV	<input type="checkbox"/> bioMerieux Vidas / Mini Vidas
INABRR	<input type="checkbox"/> Brahms Kryptor
INAETI	<input type="checkbox"/> Diasorin ETI-MAX 3600
INALIA	<input type="checkbox"/> Diasorin Liaison
INALIX	<input type="checkbox"/> Diasorin Liaison XL
INAOE	<input type="checkbox"/> ELISA
INAIMI	<input type="checkbox"/> Immunotech IRMA
INAVEC	<input type="checkbox"/> Ortho Vitros, 3600 / 5600 / ECi
INADEL	<input type="checkbox"/> Perkin Elmer Delfia/Delfia Express / Autodelfia
INAWW	<input type="checkbox"/> Perkin Elmer Wizard
INAEYS	<input type="checkbox"/> Roche Elecsys 2010 / 1010
INARME	<input type="checkbox"/> Roche Modular E170
INACC	<input type="checkbox"/> Siemens/Bayer ACS 180 / 180 SE
INACEN	<input type="checkbox"/> Siemens Centaur
INADD	<input type="checkbox"/> Siemens/Dade Dimension
INADPI	<input type="checkbox"/> Siemens/DPC Immulite 1000
INADP2	<input type="checkbox"/> Siemens/DPC Immulite 2000 / 2500
INADS	<input type="checkbox"/> Siemens Stratus CS
INASF	<input type="checkbox"/> Stat Fax Elisa Readers
INASTR	<input type="checkbox"/> Stratec Immunotech

Other methods - Please specify in document

INSTRUMENT CODE

REAGENT CODE

OTHER UNITS SPECIFY

RQ9137 - MATERNAL SCREENING PROGRAMME

METHOD QUESTIONNAIRE

PAPP-A - U/I

CODE		METHOD
PPAAAI	<input type="checkbox"/>	Abbott Alinity
PPAARC	<input type="checkbox"/>	Abbott Architect
PPAABX	<input type="checkbox"/>	Abbott AxSYM / AxSYM Plus
PPABC	<input type="checkbox"/>	Autobio CLIA
PPSAN	<input type="checkbox"/>	Beckman Access
PPADXI	<input type="checkbox"/>	Beckman Dxl 600 / 800
PPABIV	<input type="checkbox"/>	bioMerieux Vidas / Mini Vidas
PPABRR	<input type="checkbox"/>	Brahms Kryptor
PPADME	<input type="checkbox"/>	Demeditec ELISA
PPAETI	<input type="checkbox"/>	Diasorin ETI-MAX 3600
PPALIA	<input type="checkbox"/>	Diasorin Liaison
PPALIX	<input type="checkbox"/>	Diasorin Liaison XL
PPDRE	<input type="checkbox"/>	DRG Elisa
PPAIBL	<input type="checkbox"/>	IBL, ELISA
PPAIMI	<input type="checkbox"/>	Immunotech IRMA
PPAIZI	<input type="checkbox"/>	Izotop IRMA
PPAVEC	<input type="checkbox"/>	Ortho Vitros, 3600 / 5600 / ECi
PPADEL	<input type="checkbox"/>	Perkin Elmer Delfia/Delfia Express / Autodelfia
PPAWW	<input type="checkbox"/>	Perkin Elmer Wizard
PPARCE	<input type="checkbox"/>	Roche Cobas 4000/e411
PPAC6	<input type="checkbox"/>	Roche Cobas e601/602
PPAE8	<input type="checkbox"/>	Roche Cobas e402/e801
PPAEYS	<input type="checkbox"/>	Roche Elecsys 2010 / 1010
PPARME	<input type="checkbox"/>	Roche Modular E170
PPASAI	<input type="checkbox"/>	Siemens Atellica IM
PPACC	<input type="checkbox"/>	Siemens/Bayer ACS 180 / 180 SE
PPACEN	<input type="checkbox"/>	Siemens Centaur
PPADD	<input type="checkbox"/>	Siemens/Dade Dimension
PPADPI	<input type="checkbox"/>	Siemens/DPC Immulite 1000
PPADP2	<input type="checkbox"/>	Siemens/DPC Immulite 2000 / 2500
PPADS	<input type="checkbox"/>	Siemens Stratus CS
PPASNM	<input type="checkbox"/>	SNIBE Maglumi analysers
PPASNM2	<input type="checkbox"/>	SNIBE Maglumi analysers II
PPASF	<input type="checkbox"/>	Stat Fax Elisa Readers
PPASTR	<input type="checkbox"/>	Stratec Immunotech
PPAZEN	<input type="checkbox"/>	Zentech ELISA

Other methods - Please specify in document

INSTRUMENT CODE	<input type="text"/>
REAGENT CODE	<input type="text"/>
OTHER UNITS SPECIFY	<input type="text"/>

RQ9137 - MATERNAL SCREENING PROGRAMME

METHOD QUESTIONNAIRE

UNCONJUGATED ESTRIOL - nmol/l

CODE	METHOD
UE3ARC	<input type="checkbox"/> Abbott Architect I2000 / 8200
UE3ABX	<input type="checkbox"/> Abbott AxSYM / AxSYM Plus
UE3ABC	<input type="checkbox"/> Autobio CLIA
UE3SAN	<input type="checkbox"/> Beckman Access (33570)
UE3SA2	<input type="checkbox"/> Beckman Access (C22255)
UE3DXI	<input type="checkbox"/> Beckman Dxl 600 / 800 (33570)
UE3BIV	<input type="checkbox"/> bioMerieux Vidas / Mini Vidas
UE3BRR	<input type="checkbox"/> Brahms Kryptor
UE3CAL	<input type="checkbox"/> Calbiotech ELISA
UE3DME	<input type="checkbox"/> Demeditec ELISA
UE3DM	<input type="checkbox"/> DiaMetra ELISA
UE3ETI	<input type="checkbox"/> Diasorin ETI-MAX 3600
UE3LIA	<input type="checkbox"/> Diasorin Liaison
UE3LIX	<input type="checkbox"/> Diasorin Liaison XL
UE3DRE	<input type="checkbox"/> DRG Elisa
UE3DSL	<input type="checkbox"/> DSL gammacounter
UE3HUP	<input type="checkbox"/> Human Plate
UE3IBL	<input type="checkbox"/> IBL, ELISA
UE3ICE	<input type="checkbox"/> Immunospec Corporation ELISA
UE3IMI	<input type="checkbox"/> Immunotech IRMA
UE3IZR	<input type="checkbox"/> Izotop RIA
UE3LDE	<input type="checkbox"/> Labor Diagnostike Nord ELISA
UE3OR	<input type="checkbox"/> Organon Reader 530
UE3VEC	<input type="checkbox"/> Ortho Vitros 3600 / 5600 / ECI
UE3DEL	<input type="checkbox"/> Perkin elmer auto DELFIA
UE3DEJ	<input type="checkbox"/> Perkin elmer DELFIA Epress
UE3WW	<input type="checkbox"/> Perkin Elmer Wizard
UE3EYS	<input type="checkbox"/> Roche Elecsys 2010 / 1010
UE3RME	<input type="checkbox"/> Roche Modular E170
UE3CC	<input type="checkbox"/> Siemens/Bayer ACS 180 / 180 SE
UE3CEN	<input type="checkbox"/> Siemens Centaur
UE3DD	<input type="checkbox"/> Siemens/Dade Dimension
UE3DPI	<input type="checkbox"/> Siemens/DPC Immulite 1000
UE3DP2	<input type="checkbox"/> Siemens/DPC Immulite 2000 / 2500
UE3DS	<input type="checkbox"/> Siemens Stratus CS
UE3SNM	<input type="checkbox"/> SNIBE Maglumi analysers
UE3SNM2	<input type="checkbox"/> SNIBE Maglumi analysers II
UE3SF	<input type="checkbox"/> Stat Fax Elisa Readers
UE3STR	<input type="checkbox"/> Stratec Immunotech
UE3TS	<input type="checkbox"/> Tecan Sunrise
UE3ZEN	<input type="checkbox"/> Zentech ELISA

Other methods - Please specify in document

INSTRUMENT CODE

REAGENT CODE

OTHER UNITS SPECIFY