RIQAS

RANDOX INTERNATIONAL QUALITY ASSESSMENT SCHEME

METHOD QUESTIONNAIRE IMMUNOASSAY SPECIALITY 1 RQ9141

Please be aware that the RIQAS Instrument and reagent supplier codes are now in a separate booklet. Please ensure you have a copy of this in order to complete this document.

REGISTRATION INSTRUCTIONS & RIQAS POLICIES

CRITERIA FOR PARTICIPATION

This programme is available to any laboratory running the 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 at 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 interlaboratory 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. 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 contact your local Randox representative for more information.

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.rigas.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 at 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

Please contact RIQAS at

Tel: +44 (0) 28 9445 4399
E-Mail mail@riqas.com
RIQAS Scheme Co-ordinator: Sally Picton

RANDOX LABORATORIES LTD., 55 Diamond Road, Crumlin, County Antrim, BT29 4QY, United Kingdom



METHOD QUESTIONNAIRE

Anti-TG	(NO	-	
CODE	_	METHOD	
ATGARC	Ш	Abbott Architect/ Alinity	
ATGABX	\vdash	Abbott Axsym	
TGAEA	\vdash	Aida EIA	
TGAKE	\blacksquare	Aesku Diagnostics ELISA	
TGABC	\blacksquare	Autobio CLIA	
rgsan		Beckman Access/LXi725	
TGDXI	\blacksquare	Beckman Dxl 600/800	
TGBDX9	\blacksquare	Beckman Dxl 9000	
TGBHR TGBIV	\blacksquare	Biocode Hycel RIA Biomerieux Vidas	
-	\vdash		
TGBRR TGBRK	\vdash	Brahms RIA Brahms Kryptor	
_	\vdash	71	
TGCDG TGCIR		CDG Q-Strip	
	\vdash	CIS RIA	
TGDME	H	DiaMetra ELISA	
TGLIA		DiaSorin Liaison DiaSorin Liaison XL	
TGLIX TGBYK	Н	DiaSorin RIA	
TGDCH	Н	Diesse Chorus	
TGDRC	H	DIRUI CM Series	
TGEUE	H	Euroimmun ELISA	
TGFJL	H	Fujirebio Lumipulse G Series	
GHUE	H	Human ELISA	
TGHYE	H	Hycor ELISA	
GIEL	H	Inova Microelisa	
GIZR	\vdash	Izotop Anti hTG RIA KIT	
TGSLT	H	Lifotronic eCL	
TGMAI		Maccura I Series	
TGMME	\vdash	Medipan Medizym EIA	
TGMC2		Mindray CL 8000i/6000i/2000i/1200i/1000i	
TGMC3		Mindray CL 900i	
TGORA		Orgentec Alegria	
TGDEL	H	Perkin Elmer DELFIA Xpress/AutoDELFIA	
TGPHU	H	Phadia/ImmunoCAP 100/250	
TGPHE		Phadia ELISA	
TGEVE		Randox Evolution	
TGRCE	H	Roche Cobas 4000 / e411	
TGC6	H	Roche Cobas e601/602	
TGE8	H	Roche Cobas e402/e801	
TGEYS	Ħ	Roche Elecsys	
TGRME	П	Roche Modular E170	
TGSNM		SNIBE Maglumi analysers	
TGSRR	П	SEAC Radim RIA CT	
TGSPA	П	Serodia Particle Agglutination	
TGSAI		Siemens Atellica IM	
TGSA2		Siemens Atellica IM aTgll	
TGCEN		Siemens Centaur	
TGCE2		Siemens Centaur aTgll	
TGDPI		Siemens/DPC Immulite 1000	
TGDP2	П	Siemens/DPC Immulite 2000/2500	
TGTOC		TOSOH AIA CL-Series	
TGTOS		TOSOH AIA Series	
TGZER		ZenTech RIA	
Other me	thod	ds, please specify on enrolment document	
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METHOD QUESTIONNAIRE

METHOD	QU	ESTIONNAIRE	
Anti-TP0) (k	U/I)	
CODE	- (METHOD	
ATPARC	П	Abbott Architect/ Alinity	
ATPABX	H	Abbott Axsym	
ATPAEA		Aida EIA	
ATPAKE	\vdash	Aesku Diagnostics ELISA	
ATPARE		Autobio CLIA	
ATPSAN	Н	Beckman Access/LXi725	
ATPDXI		Beckman Dxl 600/800	
ATPBDX9	\vdash	Beckman Dxl 9000	
ATPBUX9		Biomerieux Vidas	
ATPBRR ATPBRK	\vdash	Brahms RIA	
ATPCDG		Brahms Kryptor CDG Q-Strip	
		CIS RIA	
ATPCIR		DiaSorin Liaison	
ATPLIA	Ш		
ATPLIX		DiaSorin Liaison XL	
ATPBYK		DiaSorin RIA	
ATPDCH	Ш	Diesse Chorus	
ATPDRC	Ш	DIRUI CM Series	
ATPEPE		Epitope Diagnostics ELISA	
ATPEUE		Euroimmun ELISA	
ATPFJL	\vdash	Fujirebio Lumipulse G Series Hycor ELISA	
ATPHYE		Inova Microelisa	
ATPIEL			
ATPIZER		Izotop Anti hTPO RIA KIT Lifotronic eCL	
ATPSLT		Maccura I Series	
ATPMAI ATPMME		Medipan Medizym EIA	
ATPMC2		Mindray 8000i/6000i/2000i/1200i/1000i	
	\vdash		
ATPMC3		Mindray CL900i	
ATPORA ATPDEL		Orgentec Alegria Perkin Elmer DELFIA Xpress/AutoDELFIA	
ATPPHU		Phadia/ImmunoCAP 100/250	1
ATPPHE		Phadia ELISA	
ATPEVE		Randox Evolution	
ATPRCE		Roche Cobas 4000 / e411	
ATPC6	Н	Roche Cobas e601 / 602	
ATPC8	\vdash	Roche Cobas e402/e801	
ATPRME		Roche Modular E170	
ATPEYS		Roche Elecsys	
ATPSRR		SEAC Radim RIA CT	
ATPSAI		Siemens Atellica IM	
ATPCEN		Siemens Centaur	
ATPDPI		Siemens/DPC Immulite 1000	
ATPDP2		Siemens/DPC Immulite 2000/2500	
ATPTOC		Tosoh AIA-CL Series	
ATPTOS		Tosoh AIA Series	
ATPZER	Ш	ZenTech RIA	
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Other me	ethod	s, please specify on enrolment document	
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REAGENT	COD	E	
OTHER UN	ITC	SPECIEV	
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METHOD QUESTIONNAIRE

METHOD			
C-Peptic	n ex	METHOD	
CPTARC	П	Abbott Architect/ Alinity	
CPTADR		Adaltis RIA	
CPABC		Autobio CLIA	
CPSAN		Beckman Access C-Peptide	
CPCDG CPTCIR	Н	CDG Q-Strip	
CPTCIR		CIS RIA CIS BIO IRMA	
CPTLIA	H	DiaSorin Liaison	
CPTLIX	Н	DiaSorin Liaison XL	
CPTDIR		DIAsource RIA	
CPDRC		DIRUI CM Series	
CPTDRG		DRG ELISA	
CPTDSL CPTFJL		DSL RIA Fujirebio Lumipulse G Series	
CPTILM	\vdash	ILMA	
CPTIRM	Н	Immunotech IRMA	
CPTMC2		Mindray CL 8000i/6000i/2000i/1200i/1000	i
CPTMC3		Mindray CL900i	
CPTMOE		Monobind Inc. ELISA	
CPTVEC CPTRAA	Н	Ortho Vitros 3600/5600/ECi/XT 7600 Radim Alisei	
CPTRAD	Н	RADIM RIA	
CPTEV		Randox Evidence / Investigator	
CPTRCE	Н	Roche Cobas 4000 / e411	
CPTC6		Roche Cobas e601/602	
CPTE8		Roche Cobas e402/e801	
CPTRME	Н	Roche Modular E170	
CPTEYS CPSAI	Н	Roche Elecsys	
CPSA12		Siemens Atellica IM Siemens Atel IM (Rgt lot 207 &cal lot 09&	un)
CPTCEN	\vdash	Siemens Centaur	υ ρ)
CPCEN2		Siemens Cen (Rgt lot 206 & cal lot 08&up)
CPTDPI		Siemens/DPC Immulite 1000	•
CPTDP2		Siemens/DPC Immulite 2000/2500	
CPTTOS	Ш	Tosoh AIA Series	
CPTTOC CPVBE		Tosoh AIA-CL Series Vector Best ELISA	
CPC2	Н	Wantai Caris 200	
CPW2		Wantai Wan200+	
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RQ9141 Method Questionnaire 7/11 Revised Mar 2024

METHOD QUESTIONNAIRE

Insulin u	ıU/r	nl	
CODE	N	IETHOD	
INAAI	Ш	Abbott Alinity I	
INARC	Ш	Abbott Architect	
INABX INABB	\vdash	Abbott AxSym Abbott Imx	
INADE	\vdash	Alfa Prime ELISA	
INABC	H	Autobio CLIA	
INSAN	H	Beckman Access/LXi725	
INDXI	H	Beckman Dxl 600/800	
INBIG	П	Biosource Gamma counter	
INCBE		Calbiotech ELISA	
INCDG	Ш	CDG Q-Strip	
INCIS	Ш	CIS IRMA	
INCIR INCLI	\vdash	CIS RIA coated blue	
INDAE	Н	Clinipro ELISA Diagnostic Automation ELISA	
INLIA	\vdash	Diasorin Liaison	
INLIX		Diasorin Liaison XL	
INDIA		Diasource IRMA	
INDRC		DIRUI CM Series	
INELI		ELISA	
INFJL		Fujirebio Lumipulse G Series	
INIMI	Ш	Immunotech IRMA	
INIVL INIZG	Н	Invitron Luminescence Izotop Gamma Counter	
INSLT	Н	Lifotronic eCL	
INLIR	H	Linco RIA	
INMC2	H	Mindray CL 8000i/6000i/2000i/1200i/1000	Di
INMC3		Mindray CL 900i	
INMOE		Monobind Inc. ELISA	
INNOV		Novatec ELISA	
INVEC	Ш	Ortho Vitros 3600/5600/ECi/XT 7600	
INDEL	_	Perkin Elmer DELFIA Perkin Elmer Wizard	
INWW INRAA	\vdash	Radim Alisei	
INC6	\vdash	Roche Cobas 6000 / 8000	
INRCE	\vdash	Roche Cobas 4000 / e411	
INEYS		Roche Elecsys	
INRME	H	Roche Modular E170	
INSAI		Siemens Atellica IM	
INCC	Ш	Siemens/Bayer ACS 180	
INCEN	Ш	Siemens Centaur	
INDPC	Н	Siemens/DPC Coat-a-count Siemens/DPC Immulite 1000	
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OTHER UNITS, SPECIFY				
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METHOD QUESTIONNAIRE

METHOD QUESTIONNAIRE				
Parathyroid Hormone (PTH) pmol/l				
CODE	METHOD			
PTHARC	Abbott Architect/ Alinity			
PTHABC	Autobio CLIA			
PTHSAN	Beckman Access/LXi725			
PTHDXI	Beckman Dxl 600/800			
PTHBLE	Bioline ELISA			
PTHBME	Biomerica ELISA			
PTHBIV	bioMerieux VIDAS PTH (1-84)			
PTHCIS	CIS IRMA			
PTHBYK	DiaSorin IRMA			
PTHLIA	DiaSorin Liaison 1-84 PTH			
PTHLIAN	DiaSorin Liaison N-TACT PTH II			
PTHLIX	DiaSorin Liaison XL 1-84 PTH			
PTHLIXN	DiaSorin Liaison XL N-TACT PTH II			
PTHDRG	DRG ELISA			
PTHDSI	DSL IRMA			
PTHFJL	Fujirebio Lumipulse G Series			
PTHIDS	IDS-iSYS PTH			
PTHSLT	Lifotronic eCL			
PTHMAI	Maccura I Series			
PTHMC2	Mindray CL 8000i/6000i/2000i/1200i/1000i			
PTHMC3	Mindray CL9001			
PTHVEC	Ortho Vitros 3600/5600/ECi/XT 7600			
PTHRCEB	Roche Cobas 4000 / e411 PTH (1-84)			
PTHRCE	Roche Cobas 4000 / e411 PTH			
PTHRCES	Roche Cobas 4000 / e411 PTH STAT			
PTHC6B	Roche Cobas e601/602 PTH (1-84)			
PTHC6	Roche Cobas e601/602 PTH			
PTHC6S	Roche Cobas e601/602 PTH STAT			
PTE8B	Roche Cobas e801 PTH (1-84)			
PTE8	Roche Cobas e801 PTH			
PTE8S	Roche Cobas e801 PTH STAT			
PTHEYSB	Roche Elecsys PTH (1-84)			
PTHEYS	Roche Elecsys PTH			
PTHEYSS	Roche Elecsys PTH STAT			
PTHRMEB	Roche Modular E170 PTH (1-84)			
PTHRME	Roche Modular E170 PTH			
PTHRMES	Roche Modular E170 PTH STAT			
PTHSCR	Scantibodies RIA			
PTHSAI	Siemens Atellica Solution			
PTHCEN	Siemens Centaur			
PTHDPI	Siemens/DPC Immulite 1000			
PTHDP2	Siemens/DPC Immulite 2000/2500			
PTHSNM	SNIBE Maglumi Analysers			
PTHTOS	Tosoh AIA Series			
PTHTOC	Tosoh AIA-CL Series			
Other method	ds, please specify on enrolment document			
INSTRUMENT (
REAGENT COD				
	<u> </u>			
OTHER UNITS,	SPECIFY			

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METHOD QUESTIONNAIRE

WIL IIIO	D QUESTIONNAIRE		
1-25-(O	H)2-Vitamin D pmol/I (PILOT)		
CODE	METHOD		
VDLIA	DiaSorin Liaison		
VDLIX	DiaSorin Liaison XL		
VDBYK	DiaSorin RIA		
VDDIE	DIAsource, ELISA		
VDDIR	DIAsource RIA		
VDHPLC	HPLC		
VDIDE	IDS ELISA		
VDIDS VDIDR	IDS iSYS IDS RIA		
VDEYS	Roche Elecsys		
Other m	ethods, please specify on enrolment document		
INSTRUM	ENT CODE		
REAGENT	CODE		
OTHER III	NITS, SPECIFY		
OTTLK	NITO, OF LOTE 1		
25-OH-\	Vitamin D nmol/l		
	- · · · · · · · · · · · · · · · · · · ·		
CODE	METHOD		
VDARC	Abbott Architect (3L52) Abbott Architect (5D02)/ Aligity (9D45)		
VDARC2 VDAPI	Abbott Architect (5P02)/ Alinity (8P45) Applied Biosystems API 4000		
VDAFI	Agappe Mispa i3		
VDAIC	Aptasys Indra CLIA		
VDABC	Autobio CLIA		
VDSAN	Beckman Access 25 OH Vitamin D Total		
VDDXI	Beckman Dxl 600 / 800		
VDBIO	Biohit Total 25 OH Vitamin D bioMerieux Vidas/mini Vidas/Vidas 3		
VDBIV VDCIR	Chongqing ISIA 25hydroxy vitD rapid test		
VDCDG	CDG Q-Strip		
VDLIA	DiaSorin Liaison		
VDLIX	DiaSorin Liaison XL		
VDBYK	DiaSorin RIA		
VDDIE	DIAsource ELISA		
VDDIR	DIAsource RIA		
VDDIA VDDRC	Diazyme Vitamin D DIRUI CM Series		
VDEUE	Euroimmune ELISA		
VDFJL	Fujirebio Lumipulse G Series		
VDHP	HPLC .		
VDIBL	IBL ELISA		
VDIDE	IDS ELISA		
VDIDS VDIDR	IDS iSYS IDS RIA		
VDLM	LC/MS		
VDSLT	Lifotronic eCL		
VDMOE	Monobind Inc. ELISA		
VDMC2	Mindray CL 8000i/6000i/2000i/1200i/1000i		
VDMC3	Mindray CL900i		
VDORA	Orgentec Alegria Elisa		
VDVEC	Ortho Vitros 3600/5600/ECi/XT 7600		
VDEVE	Randox Evolution		
VDC6 VDR2	Roche Vitamin D Total Roche Vitamin D Total II		
VDE82	Roche Vitariii D Total II e801		
VDR3	Roche Vitamin D Total III		
VDE83	Roche Vitmain D Total III e402/E803		
VDSYI	Shenzhen YHLO iFlash Series		
VDSAI	Siemens Atellica Solution		
VDCEN	Siemens Centaur		
VDSDE VDSNM	Siemens Dimension EXL Vitamin D Total SNIBE Maglumi Analyser		
VDSNW	Tosoh AIA Series		
VDTOC	Tosoh AIA-CL Series		
VDWXE	Waters Quattro Premier XE		
Other m	ethods, please specify on enrolment document		
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REAGENT	CODE		
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OTHER U	NITS, SPECIFY		

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