RIQAS

RANDOX INTERNATIONAL QUALITY ASSESSMENT SCHEME

METHOD QUESTIONNAIRE
SEROLOGY (SYPHILIS)
PROGRAMME
RQ9154

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.

This document must be retained by participant

REGISTRATION INSTRUCTIONS & RIQAS POLICIES

CRITERIA FOR PARTICIPATION

This programme is available to any laboratory running the Serology (SYPHILIS) parameters listed in this document. Quantitative results will be accepted on this programme. Titre results will also be accepted. **RATIO results will not be accepted.**

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/riqas-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/riqas-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 and you will be sent RIQAS literature, which will enable you to understand the RIQAS process and interpret your reports. Please quote this number on all correspondence with RIQAS.

B. ORDER NUMBER

If you are a UK or Irish participant, please state your official order number in the boxes provided. Other participants may order directly from their local Randox Laboratories representative.

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. 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.

H. 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.

I. 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 72 hours. 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.

J. 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 72 hrs.

NB Deletions of parameters cannot be made on RIQASNet. If you wish to delete a parameter please contact RIQAS directly on mail@riqas.com.

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.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

Results will normally be processed within 7-10 days of the FINAL DATE. PDF reports will be emailed the day after the results have been processed and for those registered for RIQASNet the PDF reports will be available on RIQASNet shortly after.

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 RIQAS Explained at www.randox.com/riqas-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

Randox Laboratories Ltd. complies with GDPR 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.

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 HAS NOT BEEN
ACCREDITED TO ISO/IEC 17043:2010

Please contact RIQAS at

Tel: +44 (0) 28 9445 4399

Fax: +44 (0) 28 9445 4398

E-Mail mail@riqas.com

RIQAS Scheme Co-ordinator: Stephen Doherty

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

RQ9154 - SEROLOGY (SYPHILIS) PROGRAMME

METHOD QUESTIONNAIRE

Syphilis - FTA-Abs U/ml

CODE		METHOD
VSFBSI	☐	Biocientifica Inmunofluor FTA-Abs
VSFBIM	☐	Biomerieux FTA-Abs
VSFDAS	☐	Dasit MarBlot Syphilis FTA-Abs
VSFEUR	☐	Euroimmun FTA-Abs
VSFHEG	☐	Hemagen FTA-Abs
VSFMDF	☐	Mastafluor FTA-Abs
VSFVIM	☐	Viro Immun FTA-Abs
VSFZEU	☐	Zeus Scientific FTA-ABS IFA

Other methods, please specify on enrolment document

INSTRUMENT CODE	
REAGENT CODE	
OTHER UNITS, SPECIFY	

Syphilis - Immunoassay U/ml

CODE		METHOD
VSIARC	☐	Abbott Architect/ Alinity
VSIACL	☐	Acon Laboratories Foresight EIA
VSIADA	☐	Adaltis EIAgen Syphilis Ab Kit
VSIBSS	☐	Beckman Coulter / Sentinel Syphilis TP Latex
VSIBKE	☐	Biokit Bioelisa Syphilis
VSIBLN	☐	Bioland Nanosign Syphilis
VSIBMX	☐	Biomaxima Syphilis Test
VSIBIT	☐	Biomerieux Trepanostika
VSIBRB	☐	Biorad Bioplex Syphilis test
VSIBRS	☐	Bio-Rad Syphilis Total Anti-body test
VSICPE	☐	Captia ELISA
VSICCL	☐	Chemclin Immunoassay
VSIDAI	☐	Diagnostic Automation Inc
VSIDPE	☐	Diagnostic BioProbes ELISA
VSIDSE	☐	Diagnostics Systems ELISA
VSIDMT	☐	DiaMedix Trep-Sure ELISA
VSDIA	☐	DiaProph Med
VSILIA	☐	Diasorin Liaison
VSILIX	☐	Diasorin Liaison XL
VSIDTS	☐	Diasorin Treponema Screen
VSIDME	☐	Diasorin Murex
VSIDCH	☐	Diesse Chorus
VSIDEZ	☐	Diesse Enzywell
VSIEUE	☐	Euroimmun IFA
VSIFLG	☐	Fujirebio Lumipulse G Series
VSIGBE	☐	General Biologicals ELISA
VSIHUM	☐	Human ELISA
VSIIEA	☐	Iason EIASON ELISA
VSIL21	☐	Lab 21 EIA
VSIMIN	☐	Mindray Anti-TP
VSIODE	☐	Omega Diagnostics ELISA
VSIORE	☐	Orgenics ELISA
VSIPME	☐	Plasmatec ELISA
VSIRSS	☐	Radim Syphilis Screening
VSIC6	☐	Roche Cobas e601/602
VSIE8	☐	Roche Cobas e801
VSIEYS	☐	Roche Elecsys / Cobas e411
VSIRME	☐	Roche Modular
VSISDE	☐	SD Syphilis ELISA 3.0
VSICCP	☐	Siemens Centaur CP
VSICEN	☐	Siemens Centaur XP/XPT/Classic
VSISEZ	☐	Siemens Enzygnost
VSIDP2	☐	Siemens Immulite 2000
VSIVBE	☐	Vector-Best RecombiBest ELISA
VSIVE	☐	Vircell ELISA
VSIVIP	☐	Vitros Immunodiagnostic Syphilis kit
VSIVEC	☐	Vitros Syphilis TPA
VSIVTE	☐	Vitrotest ELISA
VSIWAE	☐	Wantai ELISA
VSIWLE	☐	Wiener Lab ELISA
VSIZAE	☐	ZAO Ecolab AgCL-RMP

Other methods, please specify on enrolment document

INSTRUMENT CODE	
REAGENT CODE	
OTHER UNITS, SPECIFY	

RQ9154 - SEROLOGY (SYPHILIS) PROGRAMME

METHOD QUESTIONNAIRE

Syphilis - RPR U/ml

CODE		METHOD
VSRASR	☐	Abbott Syfacard-R
VSRAG	☐	Agappe RPR
VSART	☐	Arkray RPR Test
VSRASI	☐	ASI RPR screening test
VSRAMR	☐	Atlas Medical RPR Card Test
VSRAXR	☐	Axis-Shield RPR test
VSRBDM	☐	BD Macro-Vue RPR card test
VSRBBS	☐	Bio-Bacter Serobacter
VSRBKR	☐	Biokit Reditest RPR
VSRBLO	☐	Biolabo RPR-Carbon
VSRBMX	☐	BioMaxima RPR test
VSRBIM	☐	Biomerieux Nosticon II RPR
VSRBRS	☐	Bio-Rad RPR test card
VSRBRX	☐	Biorex Syphilis RPR
VSRBSS	☐	Biosystems RPR kit
VSRBTR	☐	Biotec RPR test kit
VSRCEG	☐	Cenogenics RPR test
VSRCHL	☐	Chemelex RPR Carbon
VSRCMR	☐	Cormay RPR
VSRDAR	☐	Diagnostic Automation RPR kit
VSRDIA	☐	Dialab Syphilis RPR
VSRDDS	☐	DDS DiagnostIcs
VSREKO	☐	EKOlab Syphilis RPR
VSRELT	☐	ELITech RPR
VSREMR	☐	Emapol RPR kit
VSRFAD	☐	FAR Diagnostics RPR Carbon
VSRFDR	☐	Fortress Diagnostics/Biorex RPR Carbon
VSRFUR	☐	Fumouze RPR card test
VSRGID	☐	Giesse Diagnostics RPR Carbon
VSRGMT	☐	GM Test RPR
VSRHUM	☐	Human Syphilis RPR test
VSRIMR	☐	InterMedical RPR/Carbon
VSRIWI	☐	Inverness Medical Wampole Impact RPR
VSRLBR	☐	Lab 21 RPR test kit
VSRLCH	☐	Linear Chemicals RPR Carbon
VSRLLR	☐	Lorne Laboratories RPR
VSRLTA	☐	LTA Syphilis RPR
VSRMBR	☐	Mascia Brunelli RPR
VSRNMN	☐	Nal von Minden NADAL RPR Test
VSRNP	☐	Niarmedic Plus RPR
VSROMD	☐	Omega Diagnostics Immutrep (Semi-Quant)
VSRPST	☐	Plasmatec RPR test kit
VSRQCA	☐	Quimica Clinica Aplicada RPR
VSRRBX	☐	Ranbaxy Trepostat RPR
VSRRAN	☐	Randox RPR test
VSRRDR	☐	Reckon Diagnostics RPR test
VSRDGT	☐	RFCL Diagnova Trepostat
VSRRRP	☐	Roche/Sekisui RPR
VSRSBR	☐	Servibio RPR
VSRSGM	☐	SGM Italia RPR
VSRKHB	☐	Shanghai Kehua Bio-tech RPR kit
VSRSDR	☐	Span Diagnostics RPR test
VSRSRC	☐	Spinreact RPR Carbon
VSRSRQ	☐	Stanbio RPR Quicktest
VSRTDT	☐	Teco Diagnostics RPR test
VSRTUL	☐	Tulip Carbogen
VSRVBR	☐	Vector-Best RPR
VSRVDC	☐	Vedalab RPR Carbon
VSRWIE	☐	Wiener RPR slide test

Other methods, please specify on enrolment document

INSTRUMENT CODE ☐

REAGENT CODE ☐

OTHER UNITS, SPECIFY ☐

RQ9154 - SEROLOGY (SYPHILIS) PROGRAMME

METHOD QUESTIONNAIRE

Syphilis - Rapid Tests U/ml

CODE		METHOD
VSPABO	☐	Abon Syphilis Ultra Rapid Test Device
VSPACS	☐	Acon Syphilis One Step
VSPDET	☐	Alere/Inv Medical Determine Syphilis TP
VSPADS	☐	All Diag Syphilitop Optima
VSPART	☐	Artron Syphilis Rapid Test
VSPWAR	☐	Beijing Wantai Syphilis Rapid test
VSPBBT	☐	Biofocus Biotracer Syphilis Rapid Card
VSPBLN	☐	Bioland Nanosign Syphilis test strip
VSPBSU	☐	Bioplastic Syphilis Ultra Rapid Device
VSPBIP	☐	Bio-Rad ID-PaGIA
VSPCTK	☐	CTK Biotech Rapid test
VSPDIP	☐	DiaMed ID PaGIA Test
VSPFDS	☐	Fortress Diagnostics Syphilis Strips
VSPHGS	☐	Healgen Scientific Syphilis Rapid Test
VSPHRT	☐	Human Hexagon Syphilis Rapid test
VSPHU	☐	Humasis Syphilis Card
VSPHSU	☐	Hydrex Syphilis Ultra
VSPIBS	☐	IBS Syphilis Rapid test
VSPING	☐	Innogenetics INNO-LIA test strips
VSPINO	☐	Innovacon Instalert Rapid tests
VSPITO	☐	InTec One Step Syphilis test
VSPKLQ	☐	Koroglu Laboquick Anti Syphilis
VSPL21	☐	Lab 21 Rapid tests
VSPMPA	☐	MP Diagnostics Assure Syphilis RT
VSPNMN	☐	Nal von Minden NADAL Syphilis test
VSPODI	☐	Omega Diagnostics Immutrep (Slide)
VSPOMV	☐	Omega Diagnostics Visitect
VSPORR	☐	Orgenics Rapid tests
VSPQPD	☐	Qualpro Diagnostics Rapid test
VSPRLS	☐	Rapid Labs Syphilis
VSPRDS	☐	Reckon Diagnostics Strip test
VSPSDB	☐	SD Bioline Syphilis Rapid Test
VSPSDT	☐	Span Diagnostics Trust test
VSPTDO	☐	Teco Diagnostics One Step Cassette
VSPTYD	☐	Toyo Diagnostics Cassette

Other methods, please specify on enrolment document

INSTRUMENT CODE ☐

REAGENT CODE ☐

OTHER UNITS, SPECIFY ☐

RQ9154 - SEROLOGY (SYPHILIS) PROGRAMME

METHOD QUESTIONNAIRE

Syphilis - TPHA U/ml

CODE		METHOD
VSTAMT	☐	Altlas Medical TPHA kit
VSTAXD	☐	Axiom Diagnostic TPHA
VSTAXT	☐	Axis-Shield Microsyph TPHA
VSTBIK	☐	Biokit Syphagen TPLH
VSTBLO	☐	Biolabo TPHA
VSTBIM	☐	Biomerieux TPHA
VSTBRS	☐	Bio-Rad TPHA kit
VSTBSS	☐	Biosystems TPHA kit
VSTBTC	☐	Biotec TPHA kit
VSTBPC	☐	BPC Biosed Syphilide TPHA
VSTCYD	☐	Cypress Diagnostics TPHA
VSTDIA	☐	Dialab Syphilis TPHA
VSTDSF	☐	Diesse Syphilis Fast
VSTEUR	☐	Eurobio TPHA test
VSTELT	☐	ELITech TPHA
VSTFAR	☐	FAR TPHA kit
VSTFDT	☐	Fortress Diagnostics TPHA
VSTFSP	☐	Fujirebio Serodia Particle Agg.
VSTFST	☐	Futura Systems TPHA
VSTHUM	☐	Human Syphilis TPHA liquid
VSTL21	☐	Lab 21 TPHA
VSTLTA	☐	LTA Syphilis TPHA
VSTLCH	☐	Linear Chemicals TPHA
VSTMBT	☐	Mascia Brunelli TPHA
VSTNMN	☐	Nal von Minden NADAL TPHA Test
VSTNBM	☐	Newbio TPHA kit
VSTOMD	☐	Omega Diagnostics Immutrep
VSTOXO	☐	Oxoid TPHA test
VSTPST	☐	Plasmatec THPA test kit
VSTRAN	☐	Randox TPHA kit
VSTSGM	☐	SGM Italia TPHA
VSTSCG	☐	Siemens Cellognost
VSTSOB	☐	Sobioda TPHA
VSTSRT	☐	Spinreact TPHA
VSTZAE	☐	ZAO Ecolab TPHA

Other methods, please specify on enrolment document

INSTRUMENT CODE ☐

REAGENT CODE ☐

OTHER UNITS, SPECIFY ☐

RQ9154 - SEROLOGY (SYPHILIS) PROGRAMME

METHOD QUESTIONNAIRE

Syphilis - VDRL U/ml

CODE		METHOD
VSVADV	☐	All.Diag VDRL check
VSVBCD	☐	Becton Dickinson VDRL
VSVBIO	☐	Bioclin Quibasa VDRL
VSVMD	☐	Biomed VDRL
VSVBIM	☐	Biomerieux VDRL
VSVBRS	☐	Bio-Rad VDRL Latex Pasteur
VSVBTC	☐	Biotec VDRL Kit
VSVBTV	☐	Britania VDRL - Sepssi
VSVCEG	☐	Cenogenics VDRL test
VSVDSY	☐	Diagast Sypal CB
VSVDIA	☐	Dialab Syphilis VDRL
VSVDMA	☐	Diasorin Murex VDRL
VSVICV	☐	INCDMI Cantacuzino Spl VDRL
VSVIDV	☐	Investigacion Diagnostics VDRL Test
VSVLBV	☐	Labtest VDRL
VSVLIC	☐	Licon VDRL-USR
VSVOMD	☐	Omega Diagnostics Immutrep
VSVOXO	☐	Oxoid VDRL test
VSVPST	☐	Plasmatec VDRL test kit
VSVROD	☐	Rodelg Laboratories VDRL
VSVSDM	☐	Sclavo Diagnostics Microgen VDRL
VSVSCA	☐	Siemens VDRL Cardiolipin
VSVSDT	☐	Span Diagnostics TRUST VDRL
VSVSRV	☐	Spinreact VDRL
VSVTUG	☐	Tulip Carbogen
VSVWIE	☐	Wiener Lab VDRL test

Other methods, please specify on enrolment document

INSTRUMENT CODE	☐
REAGENT CODE	☐
OTHER UNITS, SPECIFY	☐