2019 CATALOGUE

EQA FOR MOLECULAR INFECTIOUS DISEASE TESTING





EQA FOR MOLECULAR INFECTIOUS DISEASE TESTING

QCMD (Quality Control for Molecular Diagnostics) is an independent External Quality Assessment (EQA) / Proficiency Testing (PT) scheme specialising in molecular testing of a wide range of infectious diseases.



IMPORTANCE OF EXTERNAL QUALITY ASSESSMENT

External Quality Assessment (EQA) or Proficiency Testing (PT) provides a means of periodically assessing a laboratory's performance in comparison with other laboratories using the same method and instrument.

Unlike Internal Quality Control (IQC), EQA provides an effective method of monitoring a laboratory's bias or accuracy through the analysis of 'blind samples'. Participation in an EQA scheme like QCMD will also support regulatory requirements and will assist in quality improvements.

EQA plays an essential role in assuring laboratory quality by supporting daily IQC. It facilitates interlaboratory performance comparison and encourages greater standardisation in testing. EQA has a number of functions:

- Helps maintain and improve the analytical quality of laboratory tests
- Provides an objective view of test system performance that IQC alone cannot provide
- Helps improve interlaboratory agreement
- Initiates corrective and preventative actions to resolve problems

Furthermore, participating in an EQA scheme is often a prerequisite to gaining accreditation, ISO 15189 states, "the laboratory shall participate in interlaboratory comparisons such as those organised by external quality assessment schemes".

In short, participation in an EQA scheme will give labs greater confidence and will provide evidence that the patient results they are reporting are reliable and accurate.

BENEFITS

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EXTENSIVE PROGRAMME OFFERING

Boasting the largest selection of molecular EQA programmes for infectious disease testing, you are sure to find what you're looking for.



FREQUENCY

Choose between one, two and four challenges* per year to suit your laboratory requirements. Reports are available within 2 weeks of the submission deadline (up to 4 weeks for the drug resistance / sequence based schemes), ensuring any corrective actions can be taken quickly.



HIGH QUALITY MATERIAL

The availability of whole pathogen samples in clinically relevant matrices mimics the performance of patient samples and ensures samples can be used to effectively monitor the performance of the entire testing process.



INTERNATIONAL ACCREDITATION

Where appropriate the EQA schemes are accredited to ISO 17043:2010 highlighting the superior quality and organisation of the QCMD scheme.



ONLINE EQA MANAGEMENT SYSTEM

IT EQA Management System (ITEMS) provides an online tool to easily manage all EQA activities from programme registration to submission of results and provision of EQA reports.

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HIGH LEVEL OF PARTICIPATION

With over 10,000 participant registrations in more than 100 countries, peer groups are maximised, increasing statistical validity.

COMPREHENSIVE REPORTS

Individual reports are provided with each EQA challenge. In line with the requirements of ISO17043, they provide the laboratories with their results and performance assessment in relation to their EQA assessment group (peer review group).

Supplementary reports which include scientific expert commentary may be provided at the end of the EQA cycle if appropriate.

HOW IT WORKS

The QCMD portfolio is extensive covering over 300 target organisms across more than 90 EQA programmes and pilot studies.

The following diagram provides an overview of the schemes operation.



BACTERIAL EQA PROGRAMMES

BACTERIAL 16S RIBOSOMAL RNA

B16SrRNA19

Designed to evaluate the ability to detect, identify and interpret which bacterial species are provided within each panel member using routine 16S rRNA molecular diagnostic procedures.

	Available Format(s)
Catalogue Number	QAB164183_1
Total Number of Challenges	1
Number of Samples	8 to 10
Distribution / Testing Period	Q4

Specifications

Target Pathogen – May include clinically relevant species of Serratia, Escherichia, Staphylococcus, Enterococcus and Klebsiella. Matrix – Physiological Buffer Sample Volume – 0.5 ml Analysis Type – Molecular typing Format – Liquid frozen Accreditation – Pending accreditation

BORDETELLA PERTUSSIS

BPDNA19

Designed to evaluate the ability to detect Bordetella pertussis using molecular methods.

	Available Format(s)
Catalogue Number	QAB094132_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q2

Specifications

Target Pathogen – Bordetella pertussis Matrix – Physiological Buffer Sample Volume – 1.0 ml Analysis Type – Qualitative Format – Liquid frozen Accreditation – ISO17043

BORRELIA BURGDORFERI SPP. (LYME DISEASE)

BbDNA19

Designed to assess the qualitative detection of *Borrelia burgdorferi* sensu stricto at different concentrations, and the qualitative detection of *B. burgdorferi* genospecies complex at different concentrations.

	Available Format(s)
Catalogue Number	QAB114147_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q3

Specifications

Target Pathogen – Borrelia burgdorferi spp. (Lyme Disease) Matrix – Microbiological Medium and/or Transport Medium Sample Volume – 1.0 ml Analysis Type – Qualitative Format – Liquid frozen Accreditation – ISO17043

CHLAMYDIA PSITTACI

CPS19

Designed to evaluate the ability to detect Chlamydia psittaci using molecular methods.

	Available Format(s)
Catalogue Number	QAB134165_1
Total Number of Challenges	1
Number of Samples	8 to 10
Distribution / Testing Period	Q2

Specifications

Target Pathogen – Chlamydia psittaci Matrix – Transport Medium Sample Volume – 1.0 ml Analysis Type – Qualitative Format – Liquid frozen Accreditation – Pending accreditation

CHLAMYDIA TRACHOMATIS

CTDNA19

Designed to assess the qualitative detection of *Chlamydia trachomatis* at various concentrations, and the ability to correctly identify different *C. trachomatis* strains using molecular methods.

	Available Format(s)	
Catalogue Number	QAB004101_1	QAB004101_2
Total Number of Challenges	1	2
Number of Samples	8 to 12	4 to 6
Distribution / Testing Period	Q3	Q1 and Q3

Specifications

Target Pathogen – Chlamydia trachomatis Matrix – Urine and/or Physiological Buffer Sample Volume – 4.0 ml Analysis Type – Qualitative Format – Liquid frozen Accreditation – ISO17043

CHLAMYDIA TRACHOMATIS AND NEISSERIA GONORRHOEAE

CTNg19

Designed to evaluate the ability to detect Chlamydia trachomatis and Neisseria gonorrhoeae using molecular methods.

	Available Format(s)	
Catalogue Number	QAB174191_1	QAB174191_2
Total Number of Challenges	1	2
Number of Samples	8 to 12	4 to 6
Distribution / Testing Period	Q3	Q1 and Q3

Specifications

Target Pathogen – Chlamydia trachomatis; Neisseria gonorrhoeae Matrix – Urine and/or Physiological Buffer Sample Volume – 4.0 ml Analysis Type – Qualitative Format – Liquid frozen Accreditation – ISO17043

CHLAMYDOPHILA PNEUMONIAE

CP19

Designed to evaluate the ability to detect *Chlamydophila pneumoniae* using molecular methods.

	Available Format(s)
Catalogue Number	QAB084107_1
Total Number of Challenges	1
Number of Samples	5 to 10
Distribution / Testing Period	Q2

Specifications

Target Pathogen – Chlamydophila pneumoniae Matrix – Bronchoalveolar Lavage (BAL) and/or Transport Medium Sample Volume – 0.5 ml Analysis Type – Qualitative Format – Liquid frozen Accreditation – ISO17043

CLOSTRIDIUM DIFFICILE (CD)

CDDNA19

Designed to evaluate the ability to detect Clostridium difficile using molecular methods.

	Available Format(s)	
Catalogue Number	QAB084125_1	QAB084125_2
Total Number of Challenges	1	2
Number of Samples	8 to 12	4 to 6
Distribution / Testing Period	Q4	Q2 and Q4

Specifications

Target Pathogen – Clostridium difficile (CD) Matrix – Microbiological Medium and/or Synthetic Faecal Matrix Sample Volume – 1.0 ml Analysis Type – Qualitative Format – Liquid frozen Accreditation – ISO17043

DIARRHEAGENIC ESCHERICHIA COLI

E.COLI19

Designed to evaluate the ability to detect diarrheagenic *Escherichia coli* strains using routine molecular methods.

	Available Format(s)
Catalogue Number	QAB154179_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q4

Specifications

Target Pathogen – Diarrheagenic Escherichia coli Matrix – Synthetic Faecal Matrix and/or Physiological Buffer Sample Volume – 1.0 ml Analysis Type – Qualitative Format – Liquid frozen Accreditation – Pending accreditation

EXTENDED SPECTRUM B-LACTAMASE AND CARBAPENEMASE

ESBL19

Designed to evaluate the ability to detect and determine different ESBL and Carbapenemases in a clinical setting.

	Available Format(s)
Catalogue Number	QAB134162_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q3

Specifications

Target Pathogen – Extended Spectrum β-lactamase and Carbapenemases Matrix – Physiological Buffer Sample Volume – 0.5 ml Analysis Type – Molecular typing Format – Liquid frozen Accreditation – Pending accreditation

GROUP B STREPTOCOCCUS

GBS19

Designed to assess in the qualitative detection of Group B Streptococcus using routine molecular methods.

	Available Format(s)
Catalogue Number	QAB174200_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q4

Specifications

Target Pathogen – Group B Streptococcus Matrix – Plasma, Synthetic CSF and/or Transport Medium Sample Volume – 1.0 ml Analysis Type – Qualitative Format – Liquid frozen Accreditation – Pending accreditation

HELICOBACTER PYLORI

H.PYLORI19

Designed to assess the qualitative detection of *H. pylori* and where appropriate, the identification of *H. pylori* antibiotic resistance status using routine molecular methods.

	Available Format(s)
Catalogue Number	QAB164190_1
Total Number of Challenges	1
Number of Samples	5 to 10
Distribution / Testing Period	Q3

Specifications

Target Pathogen – H. pylori Matrix – Synthetic Faecal Matrix and/or Physiological Buffer Sample Volume – 1.0 ml Analysis Type – Qualitative. Quantitative for information purposes only Format – Liquid frozen Accreditation – Pending accreditation

LEGIONELLA PNEUMOPHILA

LPDNA19

Designed to evaluate the ability to detect *Legionella pneumophila* using molecular methods.

	Available Format(s)
Catalogue Number	QAB044122_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q1

Specifications

Target Pathogen – Legionella pneumophila Matrix – Bronchoalveolar lavage (BAL) and/or Transport Medium Sample Volume – 0.5 ml Analysis Type – Qualitative Format – Liquid frozen Accreditation – ISO17043

METHICILLIN RESISTANT STAPHYLOCOCCUS AUREUS (MRSA)

MRSADNA19

Designed to evaluate the ability to detect Methicillin Resistant *Staphylococcus aureus* using molecular methods.

	Available Format(s)
Catalogue Number	QAB064124_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q2

Specifications

Target Pathogen – Methicillin Resistant Staphylococcus aureus (MRSA) Matrix – Microbiological Medium and/or Transport Medium Sample Volume – 1.2 ml Analysis Type – Qualitative Format – Liquid ready-to-use Accreditation – ISO17043

METHICILLIN RESISTANT STAPHYLOCOCCUS AUREUS (MRSA) - TYPING

(epidemiology and outbreak studies)

MRSATP19

Designed to evaluate the ability to use molecular typing for outbreak analysis of Methicillin Resistant Staphylococcus aureus (MRSA).

	Available Format(s)
Catalogue Number	QAB074128_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q2

Specifications

Target Pathogen – Methicillin Resistant Staphylococcus aureus (MRSA) – Typing Matrix – Microbiological Medium and/or Transport Medium Sample Volume – 0.2 ml Analysis Type – Molecular typing Format – Liquid ready-to-use Accreditation – ISO17043

MYCOBACTERIUM TUBERCULOSIS (MTB)

MTBDNA19

Designed to evaluate the ability to detect Mycobacterium tuberculosis (M. bovis - BCG) using molecular methods.

	Available Format(s)	
Catalogue Number	QAB014129_1	QAB014129_2
Total Number of Challenges	1	2
Number of Samples	8 to 12	4 to 6
Distribution / Testing Period	Q4	Q2 and Q4

Specifications

Target Pathogen – Mycobacterium tuberculosis (M. bovis - BCG) Matrix – Sputum and/or Synthetic Sputum and/or Synthetic CSF Sample Volume – 1.0 ml Analysis Type – Qualitative Format – Liquid ready-to-use Accreditation – ISO17043

MYCOPLASMA PNEUMONIAE

MP19

Designed to evaluate the ability to detect Mycoplasma pneumoniae using molecular methods.

	Available Format(s)
Catalogue Number	QAB174192_1
Total Number of Challenges	1
Number of Samples	5 to 10
Distribution / Testing Period	Q2

Specifications

Target Pathogen – Mycoplasma pneumoniae Matrix – Bronchoalveolar lavage (BAL) and/or Transport Medium Sample Volume – 0.5 ml Analysis Type – Qualitative Format – Liquid frozen Accreditation – ISO17043

MYCOPLASMA SPP. (CELL CONTAMINATION)

MYCO19

Designed to evaluate the ability to detect and quantitate *Mycoplasma* species using molecular methods.

	Available Format(s)
Catalogue Number	QAB144168_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q4

Specifications

Target Pathogen – Mycoplasma spp. (cell contamination) Matrix – Physiological Buffer Sample Volume – 1.2 ml Analysis Type – Qualitative & Quantitative Format – Liquid frozen Accreditation – Pending accreditation

NEISSERIA GONORRHOEAE

NgDNA19

Designed to evaluate the ability to detect *Neisseria gonorrhoeae* using molecular technologies.

	Available Format(s)	
Catalogue Number	QAB034126_1	QAB034126_2
Total Number of Challenges	1	2
Number of Samples	8 to 12	4 to 6
Distribution / Testing Period	Q3	Q1 and Q3

Specifications

Target Pathogen – Neisseria gonorrhoeae Matrix – Urine and/or Physiological Buffer Sample Volume – 4.0 ml Analysis Type – Qualitative Format – Liquid frozen Accreditation – ISO17043

STAPHYLOCOCCUS AUREUS SPA

SASPA19

Designed to evaluate the ability to use molecular typing as a technique for identifying *Staphylococcus aureus*.

	Available Format(s)
Catalogue Number	QAB134164_1
Total Number of Challenges	1
Number of Samples	6 to 12
Distribution / Testing Period	Q2

Specifications

Target Pathogen – Staphylococcus aureus Matrix – Microbiological Medium and/or Transport Medium Sample Volume – 0.2 ml Analysis Type – Molecular typing Format – Liquid ready-to-use Accreditation – ISO17043

SYPHILIS

SYPH19

Designed to evaluate the ability to detect *Treponema pallidum* using molecular methods.

	Available Format(s)
Catalogue Number	QAB154180_1
Total Number of Challenges	1
Number of Samples	5 to 10
Distribution / Testing Period	Q4

Specifications

Target Pathogen – Treponema pallidum Matrix – Urine and/or Physiological Buffer Sample Volume – 1.0 ml Analysis Type – Qualitative Format – Liquid frozen Accreditation – Pending accreditation

VANCOMYCIN RESISTANT ENTEROCOCCI (VRE)

VRE19

Designed to evaluate the ability to detect and determine different VRE in clinically relevant sample types using molecular methods.

	Available Format(s)
Catalogue Number	QAB134163_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q3

Specifications

Target Pathogen – Vancomycin Resistant Enterococci Matrix – Microbiological medium and/or transport medium Sample Volume – 0.5 ml Analysis Type – Molecular Typing Format – Liquid frozen Accreditation – Pending accreditation

FUNGAL EQA PROGRAMMES

ASPERGILLUS SPP.

ASPDNA19

Designed to evaluate the ability to detect Aspergillus species using molecular methods.

	Available Format(s)
Catalogue Number	QAF104140_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q3

Specifications

Target Pathogen – Aspergillus species Matrix – Plasma and/or Physiological Buffer and/or Synthetic Sputum Sample Volume – 1.0 ml Analysis Type – Qualitative. Quantitative for information purposes only Format – Liquid frozen Accreditation – ISO17043

CANDIDA SPP.

CANDNA19

Designed to evaluate the ability to detect Candida species using molecular methods.

	Available Format(s)
Catalogue Number	QAF124151_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q3

Specifications

Target Pathogen – Candida species Matrix – Plasma and/or Physiological Buffer Sample Volume – 1.0 ml Analysis Type – Qualitative Format – Liquid frozen Accreditation – ISO17043

DERMATOPHYTOSIS

DERMA19

Designed to evaluate the ability to detect *dermatophytes* using routine molecular methods.

	Available Format(s)
Catalogue Number	QAF164187_1
Total Number of Challenges	1
Number of Samples	8 to 10
Distribution / Testing Period	Q3

Specifications

Target Pathogen – Dermatophytes and/or fungal infections Matrix – Physiological Buffer Sample Volume – 1.0 ml Analysis Type – Qualitative Format – Liquid frozen Accreditation – Pending accreditation

PNEUMOCYSTIS JIROVECII PNEUMONIA (PCP)

PCPDNA19

Designed to evaluate the ability to detect *Pneumocystis jirovecii* using molecular methods.

	Available Format(s)
Catalogue Number	QAF114144_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q3

Specifications

Target Pathogen – Pneumocystis jirovecii Matrix – Physiological Buffer Sample Volume – 1.0 ml Analysis Type – Qualitative and Quantitative Format – Liquid frozen Accreditation – ISO17043

MULTI-PATHOGEN/SYNDROMIC PROGRAMMES

BACTERIAL GASTROENTERITIS

GastroB19

Designed to evaluate the ability to detect a range of bacterial pathogens known to cause gastroenteritis using routine molecular diagnostic platforms and procedures. The panel members will resemble clinical samples and may include current clinically relevant strains of *Salmonella*, *Shigella*, *Yersinia*, *E.coli* 0157, *C. difficile* or *Campylobacter* species.

	Available Format(s)
Catalogue Number	QAB124153_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q4

Specifications

Target Pathogen – Salmonella, Shigella, Yersinia, E.coli 0157, C. difficile or Campylobacter species Matrix – Synthetic Faecal Matrix and/or Physiological Buffer

Sample Volume – 1.0 ml Analysis Type – Qualitative Format – Liquid frozen Accreditation – Pending accreditation

MALDI-TOF

MALDI19

Designed to evaluate the ability to detect and determine different clinically relevant isolates using MALDI-TOF and other similar mass spectrometry based technologies in the routine microbiology laboratory.

	Available Format(s)
Catalogue Number	QAB124155_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q3

Specifications

Target Pathogen – Clinically relevant isolates Matrix – Transport Medium Sample Volume – 1.0 ml Analysis Type – Typing Format – Liquid frozen Accreditation – Pending accreditation

PARASITIC GASTROENTERITIS

GastroP19

Designed to evaluate the ability to detect a range of parasitic pathogens known to cause gastroenteritis using routine molecular methods. The panel members will resemble clinical samples and may include current clinically relevant strains of *Giardia, Cryptosporidium* and *Entamoeba*.

	Available Format(s)
Catalogue Number	QAP124154_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q4

Specifications

Target Pathogen – Giardia, Cryptosporidium and Entamoeba Matrix – Synthetic Faecal Matrix and/or Physiological Buffer Sample Volume – 1.0 ml Analysis Type – Qualitative Format – Liquid frozen Accreditation – Pending accreditation

RESPIRATORY I

RESPI19

Designed to evaluate the ability to detect and determine various Influenza A & B and Respiratory syncytial virus strains. The panel is designed to represent various clinical scenarios.

	Available Format(s)
Catalogue Number	QAV164188_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q2

Specifications

Target Pathogen – Influenza A; Influenza B; Respiratory syncytial virus (RSV) Matrix – Transport Medium Sample Volume – 1.0 ml Analysis Type – Qualitative Format – Liquid frozen Accreditation – Pending accreditation

RESPIRATORY II

RESPII19

Designed to evaluate the ability to detect and determine human metapneumovirus, respiratory adenoviruses, rhinoviruses, coronaviruses, enterovirus and parainfluenza viruses. The panel is designed to represent various clinical scenarios.

	Available Format(s)
Catalogue Number	QAV164189_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q2

Specifications

Target Pathogen – Human metapneumovirus; respiratory adenoviruses; rhinoviruses; coronaviruses; enterovirus; parainfluenza viruses Matrix – Transport Medium Sample Volume – 1.0 ml Analysis Type – Qualitative Format – Liquid frozen Accreditation – Pending accreditation

RESPIRATORY III

RESPIII19

Designed to evaluate the ability to detect and determine various Bordetella pertussis, Legionella pneumoniae, Mycoplasma pneumoniae, Streptococcus pneumoniae or Haemophilus influenzae strains using molecular methods. The panel is designed to represent various clinical scenarios.

	Available Format(s)
Catalogue Number	QAM174193_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q4

Specifications

Target Pathogen – Bordetella pertussis, Legionella pneumoniae, Mycoplasma pneumoniae, Streptococcus pneumoniae or Haemophilus influenzae strains.

Matrix – Transport Medium Sample Volume – 1.0 ml Analysis Type – Qualitative Format – Liquid frozen Accreditation – Pending accreditation

SEPSIS

SEPSIS19

Designed to evaluate the ability to detect *Staphylococcus*, *Serratia*, *Escherichia* coli, *Enterococcus*, *Streptococcus*, *Klebsiella*, coagulase-negative *Staphylococcus*, *Pseudomonas* and *Candida* spp. using molecular methods.

	Available Format(s)
Catalogue Number	QAB164178_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q4

Specifications

Target Pathogen – Staphylococcus, Serratia, Escherichia coli, Enterococcus, Streptococcus, Klebsiella, coagulase-negative Staphylococcus, Pseudomonas and Candida spp. Matrix – Whole Blood and/or Plasma Sample Volume – 1.0 ml Analysis Type – Qualitative Format – Liquid frozen Accreditation – Pending accreditation

SEXUALLY TRANSMITTED INFECTIONS I

STI_119

Designed to evaluate the ability to detect a range of sexually transmitted infections known to cause disease using routine molecular methods. The panel members will resemble clinical samples and may include current clinically relevant strains of Mycoplasma genitalium, Mycoplasma hominis, Trichomonas vaginalis, Ureaplasma urealyticum and Gardnerella vaginalis.

	Available Format(s)
Catalogue Number	QAB154177_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q4

Specifications

Target Pathogen – Mycoplasma genitalium, Mycoplasma hominis, Trichomonas vaginalis, Ureaplasma urealyticum and Gardnerella vaginalis Matrix – Urine and/or Physiological Buffer Sample Volume – 4.0 ml Analysis Type – Qualitative Format – Liquid frozen Accreditation – Pending accreditation

SEXUALLY TRANSMITTED INFECTIONS II

STI_II19

Designed to evaluate the ability to detect a range of sexually transmitted infections known to cause disease using routine molecular methods. The panel members will resemble clinical samples and may include current clinically relevant strains of *Chlamydia trachomatis, Niesseria gonorrhoea, Treponema pallidum* and Herpes Simplex Virus (HSV) strains.

	Available Format(s)
Catalogue Number	QAM174201_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q4

Specifications

Target Pathogen – Chlamydia trachomatis, Niesseria gonorrhoea, Treponema pallidum and HSV Matrix – Urine and/or Physiological Buffer Sample Volume – 4.0 ml Analysis Type – Qualitative Format – Liquid frozen Accreditation – Pending accreditation

VIRAL GASTROENTERITIS

GastroV19

Designed to evaluate the ability to detect a range of viral pathogens known to cause gastroenteritis using routine molecular methods. The panel members will resemble clinical samples and may include current clinically relevant strains of norovirus, rotavirus, astrovirus, sapovirus and adenovirus.

	Available Format(s)
Catalogue Number	QAV124152_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q4

Specifications

Target Pathogen – norovirus, rotavirus, astrovirus, sapovirus and adenovirus Matrix – Synthetic Faecal Matrix and/or Physiological Buffer Sample Volume – 1.0 ml Analysis Type – Qualitative Format – Liquid frozen Accreditation – Pending accreditation

PARASITIC EQA PROGRAMMES

TOXOPLASMA GONDII

TGDNA19

Designed to evaluate the ability to detect Toxoplasma gondii using molecular methods.

	Available Format(s)	
Catalogue Number	QAP044123_1	QAP044123_2
Total Number of Challenges	1	2
Number of Samples	8 to 12	4 to 6
Distribution / Testing Period	Q4	Q2 and Q4

Specifications

Target Pathogen – Toxoplasma gondii Matrix – Amniotic Fluid and/or Plasma Sample Volume – 2.0 ml Analysis Type – Qualitative Format – Lyophilised Accreditation – ISO17043

VIRAL EQA PROGRAMMES

ADENOVIRUS (ADV)

ADVDNA19

Designed to evaluate the ability to detect Adenovirus using molecular methods.

	Available Format(s)	
Catalogue Number	QAV054133_1	QAV054133_2
Total Number of Challenges	1	2
Number of Samples	8 to 12	4 to 6
Distribution / Testing Period	Q3	Q2 and Q3

Specifications

Target Pathogen – Adenovirus Matrix – Transport Medium and/or Plasma Sample Volume – 1.0 ml Analysis Type – Qualitative & Quantitative Format – Liquid frozen Accreditation – ISO17043

B19 VIRUS

B19DNA19

Designed to evaluate the ability to detect and quantitatate B19 virus using molecular methods.

	Available Format(s)	
Catalogue Number	QAV034116_1	QAV034116_2
Total Number of Challenges	1	2
Number of Samples	8	4
Distribution / Testing Period	Q3	Q1 and Q3

Specifications

Target Pathogen – B19 virus Matrix – Plasma Units of Measurement – The primary unit is IU/ml however other units will be accepted Sample Volume – 1.2 ml Analysis Type – Qualitative & Quantitative Format – Liquid frozen Accreditation – ISO17043

BK VIRUS (BKV)

BKDNA19

Designed to evaluate the ability to detect and quantitatate various types of BK virus (BKV) and ensure the reliable quantification of BKV viral load using molecular methods.

	Available Format(s)	
Catalogue Number	QAV144166_1	QAV144166_2
Total Number of Challenges	1	2
Number of Samples	8 to 12	4 to 6
Distribution / Testing Period	Q3	Q2 and Q3

Specifications

Target Pathogen – BKV Matrix – Transport Medium and/or Plasma and/or Urine Units of Measurement – The primary unit is IU/ml however other units will be accepted Sample Volume – 1.0 ml Analysis Type – Qualitative & Quantitative Format – Liquid frozen Accreditation – ISO17043

CHIKUNGUNYA VIRUS (CHIKV)

CHIKV19

Designed to evaluate the ability to detect chikungunya virus using molecular methods.

	Available Format(s)
Catalogue Number	QAV154175_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q2

Specifications

Target Pathogen – Chikungunya virus Matrix – Transport Medium Sample Volume – 1.0 ml Analysis Type – Qualitative. Quantitative for information purposes only Format – Lyophilised Accreditation – Pending accreditation

CORONAVIRUS (CoV)

CVRNA19

Designed to evaluate the ability to detect coronavirus and different coronavirus genotypes using molecular methods.

	Available Format(s)
Catalogue Number	QAV064137_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q2

Specifications

Target Pathogen – Coronavirus Matrix – Transport Medium Sample Volume – 1.0 ml Analysis Type – Qualitative Format – Liquid frozen Accreditation – ISO17043

CYTOMEGALOVIRUS (CMV) DRIED BLOOD SPOTS

CMVDBS19

Designed to evaluate the ability to detect human cytomegalovirus (CMV) from dried blood spots using molecular methods.

	Available Format(s)
Catalogue Number	QAV064127_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q2

Specifications

Target Pathogen – Cytomegalovirus (CMV) Matrix – Dried Blood Spots Units of Measurement – The primary unit is IU/mI however other units will be accepted Sample Volume – 2 x 50µI Analysis Type – Qualitative. Quantitative for information purposes only Format – Dried blood spot Accreditation – ISO17043

CYTOMEGALOVIRUS (CMV) DRUG RESISTANCE

CMVDR19

Designed to evaluate the ability to detect CMV drug resistant mutations in the kinase UL97 and polymerase UL54 genes using molecular sequencing techniques.

	Available Format(s)
Catalogue Number	QAV144169_1
Total Number of Challenges	1
Number of Samples	4 to 7
Distribution / Testing Period	Q2

Specifications

Target Pathogen – Cytomegalovirus (CMV) Drug Resistance Matrix – Plasma and/or Physiological Buffer Sample Volume – 1.0 ml Analysis Type – Sequence analysis Format – Liquid frozen Accreditation – ISO17043

CYTOMEGALOVIRUS (CMV)

CMVDNA19

Designed to evaluate the ability to detect and quantitate human cytomegalovirus (CMV) using molecular methods.

	Available Format(s)	
Catalogue Number	QAV014120_1	QAV014120_2
Total Number of Challenges	1	2
Number of Samples	8 to 12	4 to 6
Distribution / Testing Period	Q3	Q2 and Q3

Specifications

Target Pathogen – Cytomegalovirus (CMV) Matrix – Plasma Units of Measurement – The primary unit is IU/ml however other units will be accepted Sample Volume – 1.0 ml Analysis Type – Qualitative & Quantitative Format – Liquid frozen Accreditation – ISO17043

CYTOMEGALOVIRUS (CMV) WHOLE BLOOD

CMVWB19

Designed to evaluate the ability to detect and quantitatate CMV from whole blood samples using molecular methods.

	Available Format(s)	
Catalogue Number	QAV124150_1	QAV124150_2
Total Number of Challenges	1	2
Number of Samples	8 to 12	4 to 6
Distribution / Testing Period	Q3	Q2 and Q3

Specifications

Target Pathogen – Cytomegalovirus (CMV) Matrix – Whole Blood Units of Measurement – The primary unit is IU/ml however other units will be accepted Sample Volume – 1.0 ml Analysis Type – Qualitative & Quantitative Format – Liquid frozen Accreditation – ISO17043

DENGUE VIRUS (DENV)

DENVRNA19

Designed to evaluate the ability to detect Dengue virus and ability to distinguish dengue virus from other flaviviruses using molecular methods.

	Available Format(s)
Catalogue Number	QAV114148_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q2

Specifications

Target Pathogen – Dengue virus (DENV) Matrix – Transport Medium Sample Volume – 1.0 ml Analysis Type – Qualitative. Quantitative for information purposes only Format – Lyophilised Accreditation – ISO17043

ENTEROVIRUS (EV)

EVRNA19

Designed to evaluate the ability to detect and quantitate different types of enterovirus (EV) using molecular methods.

	Available Format(s)	
Catalogue Number	QAV984104_1	QAV984104_2
Total Number of Challenges	1	2
Number of Samples	8 to 12	4 to 6
Distribution / Testing Period	Q3	Q1 and Q3

Specifications

Target Pathogen – Enterovirus (EV) Matrix – Transport Medium Sample Volume – 1.0 ml Analysis Type – Qualitative. Quantitative for information purposes only Format – Liquid frozen Accreditation – ISO17043

ENTEROVIRUS TYPING (EV)

EVTP19

Designed to evaluate the ability to correctly identify specific enterovirus (EV) types using routine molecular method and procedures.

	Available Format(s)
Catalogue Number	QAV164185_1
Total Number of Challenges	1
Number of Samples	5 to 10
Distribution / Testing Period	Ql

Specifications

Target Pathogen – Enterovirus (EV) Matrix – Transport Medium Sample Volume – 1.0 ml Analysis Type – Molecular typing Format – Liquid frozen Accreditation – Pending accreditation

EPSTEIN-BARR VIRUS (EBV)

EBVDNA19

Designed to evaluate the ability to detect and quantitatate Epstein-Barr virus (EBV) in plasma samples using molecular methods.

	Available Format(s)	
Catalogue Number	QAV024121_1	QAV024121_2
Total Number of Challenges	1	2
Number of Samples	8 to 12	4 to 6
Distribution / Testing Period	Q3	Q2 and Q3

Specifications

Target Pathogen – Epstein-Barr virus (EBV) Matrix – Transport Medium and/or Plasma Units of Measurement – The primary unit is IU/ml however other units will be accepted Sample Volume – 1.0 ml Analysis Type – Qualitative & Quantitative Format – Liquid frozen Accreditation – ISO17043

EPSTEIN-BARR VIRUS (EBV) WHOLE BLOOD

EBVWB19

Designed to evaluate the ability to detect and quantitatate Epstein-Barr virus (EBV) in whole blood samples using molecular methods.

	Available Format(s)	
Catalogue Number	QAV134161_1	QAV134161_2
Total Number of Challenges	1	2
Number of Samples	8 to 12	4 to 6
Distribution / Testing Period	Q3	Q2 and Q3

Specifications

Target Pathogen – Epstein-Barr virus (EBV) Matrix – Whole Blood Units of Measurement – The primary unit is IU/mI however other units will be accepted Sample Volume – 1.0 mI Analysis Type – Qualitative & Quantitative Format – Liquid frozen Accreditation – ISO17043

HEPATITIS A VIRUS (HAV)

HAVRNA19

Designed to evaluate the ability to detect Hepatitis A virus (HAV) using molecular methods.

	Available Format(s)	
Catalogue Number	QAV124156_1	QAV124156_2
Total Number of Challenges	1	2
Number of Samples	8 to 10	4
Distribution / Testing Period	Q3	Q1 and Q3

Specifications

Target Pathogen – Hepatitis A virus (HAV) Matrix – Plasma Sample Volume – 1.2 ml Analysis Type – Qualitative. Quantitative for information purposes only Format – Liquid frozen Accreditation – ISO17043

HEPATITIS B VIRUS (HBV)

HBVDNA19

Designed to evaluate the ability to detect and quantitate Hepatitis B virus (HBV) and different HBV genotypes using molecular methods.

	Available Format(s)		
Catalogue Number	QAV994110_1	QAV994110_2	QAV994110_4
Total Number of Challenges	1	2	4
Number of Samples	8	4	4
Distribution / Testing Period	Q3	Q1 and Q3	Q1, Q2, Q3 and Q4

Specifications

Target Pathogen – Hepatitis B virus (HBV) Matrix – Plasma Units of Measurement – The primary unit is IU/ml however other units will be accepted Sample Volume – 1.2 ml Analysis Type – Qualitative & Quantitative Format – Liquid frozen Accreditation – ISO17043

HEPATITIS B VIRUS (HBV) DRUG RESISTANCE

HBVDR19

Designed to evaluate the ability to detect drug resistant mutations in the Hepatitis B virus (HBV) DNA polymerase gene using sequencing techniques and/or LiPA technology.

	Available Format(s)
Catalogue Number	QAV124160_1
Total Number of Challenges	1
Number of Samples	4 to 7
Distribution / Testing Period	Q3

Specifications

Target Pathogen – Hepatitis B virus (HBV) Drug Resistance Mutations Matrix – Plasma Sample Volume – 1.0 ml Analysis Type – Sequence Analysis Format – Liquid frozen Accreditation – ISO17043

HEPATITIS B VIRUS (HBV) GENOTYPING

HBVGT19

Designed to evaluate the ability to correctly identify Hepatitis B virus (HBV) genotypes using molecular methods.

	Available Format(s)
Catalogue Number	QAV064118_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q1

Specifications

Target Pathogen – Hepatitis B virus (HBV) Genotyping Matrix – Plasma Sample Volume – 1.2 ml Analysis Type – Molecular typing Format – Liquid frozen Accreditation – ISO17043

HEPATITIS C VIRUS (HCV)

HCVRNA19

Designed to evaluate the ability to detect and quantitate Hepatitis C virus (HCV) RNA and different HCV genotypes using molecular methods.

	Available Format(s)		
Catalogue Number	QAV994112_1	QAV994112_2	QAV994112_4
Total Number of Challenges	1	2	4
Number of Samples	8	4	4
Distribution / Testing Period	Q3	Q1 and Q3	Q1, Q2, Q3 and Q4

Specifications

Target Pathogen – Hepatitis C virus (HCV) Matrix – Plasma Units of Measurement – The primary unit is IU/ml however other units will be accepted Sample Volume – 1.2 ml Analysis Type – Qualitative & Quantitative Format – Liquid frozen Accreditation – ISO17043

HEPATITIS C VIRUS (HCV) DRUG RESISTANCE

HCVDR19

Designed to evaluate the ability to detect drug resistant mutations in the Hepatitis C virus (HCV) genotypes 1 and 3 (NS3 and NS5a regions) using sequencing techniques.

	Available Format(s)
Catalogue Number	QAV134167_1
Total Number of Challenges	1
Number of Samples	4 to 7
Distribution / Testing Period	Q3

Specifications

Target Pathogen – Hepatitis C virus (HCV) Drug Resistance Mutations Matrix – Plasma Sample Volume – 1.0 ml Analysis Type – Sequence Analysis Format – Liquid frozen Accreditation – ISO17043

HEPATITIS C VIRUS (HCV) GENOTYPING

HCVGT19

Designed to evaluate the ability to correctly genotype Hepatitis C virus (HCV) RNA using molecular methods.

	Available Format(s)
Catalogue Number	QAV034117_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q1

Specifications

Target Pathogen – Hepatitis C virus (HCV) RNA Matrix – Plasma Sample Volume – 1.2 ml Analysis Type – Molecular typing Format – Liquid frozen Accreditation – ISO17043

HEPATITIS D VIRUS (HDV)

HDV19

Designed to evaluate the ability to detect Hepatitis D virus (HDV) using molecular methods.

	Available Format(s)
Catalogue Number	QAV144170_1
Total Number of Challenges	1
Number of Samples	8 to 10
Distribution / Testing Period	Q3

Specifications

Target Pathogen – Hepatitis D virus (HDV) Matrix – Plasma Sample Volume – 1.0 ml Analysis Type – Qualitative & Quantitative Format – Liquid frozen Accreditation – Pending accreditation

HEPATITIS E VIRUS (HEV)

HEVRNA19

Designed to evaluate the ability to detect Hepatitis E virus (HEV) using molecular methods.

	Available Format(s)
Catalogue Number	QAV124157_1
Total Number of Challenges	1
Number of Samples	8 to 10
Distribution / Testing Period	Q3

Specifications

Target Pathogen – Hepatitis E virus (HEV) Matrix – Plasma Sample Volume – 0.6 ml Analysis Type – Qualitative. Quantitative for information purposes only Format – Liquid frozen Accreditation – ISO17043

HERPES SIMPLEX VIRUS 1 & 2 (HSV)

HSVDNA19

Designed to evaluate the ability to detect different types and concentrations of herpes simplex virus (HSV) using molecular methods.

	Available Format(s)	
Catalogue Number	QAV994105_1	QAV994105_2
Total Number of Challenges	1	2
Number of Samples	8 to 12	4 to 6
Distribution / Testing Period	Q3	Q1 and Q3

Specifications

Target Pathogen – Herpes simplex virus 1 & 2 (HSV) Matrix – Transport Medium Sample Volume – 1.0 ml Analysis Type – Qualitative. Quantitative for information purposes only Format – Liquid frozen Accreditation – ISO17043

HERPES SIMPLEX VIRUS DRUG RESISTANCE

HSVDR19

Designed to evaluate the ability to detect HSV drug resistance mutations in the HSV thymidine kinase (UL23) and DNA polymerase (UL30) genes using routine molecular methods.

	Available Format(s)
Catalogue Number	QAV164184_1
Total Number of Challenges	1
Number of Samples	4 to 7
Distribution / Testing Period	Ql

Specifications

Target Pathogen – HSV drug resistance mutations Matrix – Transport Medium Sample Volume – 1.0 ml Analysis Type – Sequence Analysis Format – Liquid frozen Accreditation – Pending accreditation

HUMAN HERPES VIRUS 6 (HHV6)

HHV6DNA19

Designed to evaluate the ability to detect various types of Human herpes virus 6 (HHV6) and quantitate HHV6 viral load using molecular methods.

	Available Format(s)	
Catalogue Number	QAV084119_1	QAV084119_2
Total Number of Challenges	1	2
Number of Samples	8 to 12	4 to 6
Distribution / Testing Period	Q3	Q2 and Q3

Specifications

Target Pathogen – Human herpes virus 6 (HHV6) Matrix – Transport Medium and/or Plasma Units of Measurement – The primary unit is IU/ml however other units will be accepted Sample Volume – 1.0 ml Analysis Type – Qualitative & Quantitative Format – Liquid frozen Accreditation – ISO17043

HUMAN IMMUNODEFICIENCY VIRUS TYPE 1 (HIV-1) - DNA

HIVDNA19

Designed to evaluate the ability to detect Human Immunodeficiency virus type 1 (HIV-1) pro-viral DNA using molecular methods.

	Available Format(s)	
Catalogue Number	QAV034114_1	QAV034114_2
Total Number of Challenges	1	2
Number of Samples	8	4
Distribution / Testing Period	Q3	Q1 and Q3

Specifications

Target Pathogen – Human Immunodeficiency virus type 1 (HIV-1) – DNA Matrix – Physiological Buffer Sample Volume – 0.1 ml Analysis Type – Qualitative. Quantitative for information purposes only Format – Liquid frozen Accreditation – ISO17043

HUMAN IMMUNODEFICIENCY VIRUS TYPE 1 (HIV-1) – DRUG RESISTANCE

HIVDR19

Designed to evaluate the ability to detect drug resistant mutations in the HIV-1 protease and reverse transcriptase genes using molecular sequencing techniques.

	Available Format(s)
Catalogue Number	QAV024131_1
Total Number of Challenges	1
Number of Samples	4 to 7
Distribution / Testing Period	Q3

Specifications

Target Pathogen – Human Immunodeficiency virus type 1 (HIV-1) – Drug Resistance Mutations Matrix – Plasma Sample Volume – 1.0 ml Analysis Type – Sequence Analysis Format – Liquid frozen Accreditation – ISO17043

HUMAN IMMUNODEFICIENCY VIRUS TYPE 1 (HIV-1) – DRUG RESISTANCE (INTEGRASE)

HIVDRint19

Designed to evaluate the ability to detect drug resistant mutations in the HIV-1 integrase gene using molecular sequencing techniques.

	Available Format(s)
Catalogue Number	QAV114146_1
Total Number of Challenges	1
Number of Samples	4 to 7
Distribution / Testing Period	Q3

Specifications

Target Pathogen – Human Immunodeficiency virus type 1 (HIV-1) – Drug Resistance (Integrase) Mutations Matrix – Plasma Sample Volume – 1.0 ml Analysis Type – Sequence Analysis Format – Liquid frozen Accreditation – ISO17043

HUMAN IMMUNODEFICIENCY VIRUS TYPE 1 (HIV-1) - RNA

HIVRNA19

Designed to evaluate the ability to detect and quantitate human immunodeficiency virus (HIV) RNA and different HIV genotypes using molecular methods.

	Available Format(s)		
Catalogue Number	QAV994108_1	QAV994108_2	QAV994108_4
Total Number of Challenges	1	2	4
Number of Samples	8	4	4
Distribution / Testing Period	Q3	Q1 and Q3	Q1, Q2, Q3 and Q4

Specifications

Target Pathogen – Human Immunodeficiency virus type 1 (HIV-1) – RNA Matrix – Plasma Units of Measurement – The primary unit is IU/mI however other units will be accepted Sample Volume – 1.2 ml Analysis Type – Qualitative & Quantitative Format – Liquid frozen Accreditation – ISO17043

HUMAN METAPNEUMOVIRUS (MPV)

MPV19

Designed to evaluate the ability to detect human metapneumovirus (MPV) and different human MPV types using molecular methods.

	Available Format(s)
Catalogue Number	QAV054135_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q2

Specifications

Target Pathogen – Human metapneumovirus (MPV) Matrix – Transport Medium Sample Volume – 1.0 ml Analysis Type – Qualitative Format – Liquid frozen Accreditation – ISO17043

HUMAN PAPILLOMAVIRUS (HPV) - PreservCyt

HPVPRES19

Designed to evaluate the ability to detect different high risk Human Papillomavirus (HPV) types within a PreservCyt[®] matrix using molecular methods.

	Available Format(s)	
Catalogue Number	QAV094130_1	QAV094130_2
Total Number of Challenges	1	2
Number of Samples	8 to 12	4 to 6
Distribution / Testing Period	Q4	Q2 and Q4

Specifications

Target Pathogen – Human Papillomavirus (HPV) – PreservCyt® Matrix – Transport Medium (PreservCyt®) Sample Volume – 4.0 ml Analysis Type – Qualitative Format – Liquid ready-to-use Accreditation – ISO17043

INFLUENZA A & B VIRUS (FLU)

INFRNA19

Designed to evaluate the ability to detect influenza virus RNA and distinguish Influenza virus types A and B using molecular methods.

	Available Format(s)	
Catalogue Number	QAV054134_1	QAV054134_2
Total Number of Challenges	1	2
Number of Samples	8 to 12	4 to 6
Distribution / Testing Period	Q4	Q2 and Q4

Specifications

Target Pathogen – Influenza A & B virus Matrix – Transport Medium Sample Volume – 1.0 ml Analysis Type – Qualitative Format – Liquid frozen Accreditation – ISO17043

INFLUENZA HAEMAGGLUTININ TYPING (HA)

INFHT19

Designed to evaluate the ability to detect different influenza virus subtypes in addition to the typing and subtyping of influenza viruses using molecular methods.

	Available Format(s)
Catalogue Number	QAV064138_1
Total Number of Challenges	1
Number of Samples	5 to 10
Distribution / Testing Period	Q4

Specifications

Target Pathogen – Influenza Haemagglutinin Typing (HA) Matrix – Transport Medium Sample Volume – 1.0 ml Analysis Type – Molecular typing Format – Liquid frozen Accreditation – ISO17043

JC VIRUS (JCV)

JCDNA19

Designed to evaluate the ability to detect and quantitate various types of JC virus (JCV) using molecular methods.

	Available Format(s)	
Catalogue Number	QAV074106_1	QAV074106_2
Total Number of Challenges	1	2
Number of Samples	8 to 12	4 to 6
Distribution / Testing Period	Q3	Q2 and Q3

Specifications

Target Pathogen – JC virus (JCV) Matrix – Transport Medium and/or Plasma Units of Measurement – The primary unit is IU/ml however other units will be accepted Sample Volume – 1.0 ml Analysis Type – Qualitative & Quantitative Format – Liquid frozen Accreditation – ISO17043

MEASLES / MUMPS

MM19

Designed to evaluate the ability to detect mumps and/or measles using routine molecular methods.

	Available Format(s)
Catalogue Number	QAV144171_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q3

Specifications

Target Pathogen – Mumps and/or Measles Matrix – Transport Medium Sample Volume – 1.0 ml Analysis Type – Qualitative Format – Liquid frozen Accreditation – Pending accreditation

MERS CORONAVIRUS (MERS-CoV)

MERS19

Designed to evaluate the ability to detect and determine MERS-CoV from other coronaviruses.

	Available Format(s)
Catalogue Number	QAV154181_1
Total Number of Challenges	1
Number of Samples	6 to 10
Distribution / Testing Period	Q2

Specifications

Target Pathogen – MERS coronavirus (MERS-CoV) Matrix – Transport Medium Sample Volume – 1.0 ml Analysis Type – Qualitative Format – Liquid frozen Accreditation – ISO17043

NOROVIRUS (NV)

NVRNA19

Designed to evaluate the ability to detect norovirus and different norovirus (NV) genogroups using molecular methods.

	Available Format(s)	
Catalogue Number	QAV084139_1	QAV084139_2
Total Number of Challenges	1	2
Number of Samples	8 to 12	4 to 6
Distribution / Testing Period	Q4	Q2 and Q4

Specifications

Target Pathogen – Norovirus (NV) Matrix – Transport Medium and/or Physiological Buffer and/or Synthetic Faecal Matrix Sample Volume – 1.0 ml VTM, 0.1ml Buffer Analysis Type – Qualitative Format – Liquid frozen Accreditation – ISO17043

PARAINFLUENZA VIRUS (PIV)

PINFRNA19

Designed to evaluate the ability to detect Parainfluenza virus and different Parainfluenza virus (PIV) types using molecular methods.

	Available Format(s)
Catalogue Number	QAV064136_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q2

Specifications

Target Pathogen – Parainfluenza virus (PIV) Matrix – Transport Medium Sample Volume – 1.0 ml Analysis Type – Qualitative Format – Liquid frozen Accreditation – ISO17043

PARECHOVIRUS (HPeV)

PeVRNA19

Designed to evaluate the ability to detect Parainfluenza virus and different Parainfluenza virus types using molecular methods.

	Available Format(s)	
Catalogue Number	QAV114145_1	QAV114145_2
Total Number of Challenges	1	2
Number of Samples	8 to 12	4 to 6
Distribution / Testing Period	Q3	Q1 and Q3

Specifications

Target Pathogen – Parechovirus (HPeV) Matrix – Transport Medium Sample Volume – 1.0 ml Analysis Type – Qualitative Format – Liquid frozen Accreditation – ISO17043

RESPIRATORY SYNCYTIAL VIRUS (RSV)

RSV19

Designed to evaluate the ability to detect different types of Respiratory syncytial virus (RSV) using molecular methods.

	Available Format(s)	
Catalogue Number	QAV054142_1	QAV054142_2
Total Number of Challenges	1	2
Number of Samples	8 to 12	4 to 6
Distribution / Testing Period	Q4	Q2 and Q4

Specifications

Target Pathogen – Respiratory syncytial virus (RSV) Matrix – Transport Medium Sample Volume – 1.0 ml Analysis Type – Qualitative Format – Liquid frozen Accreditation – ISO17043

RHINOVIRUS (RV)

RVRNA19

Designed to evaluate the ability to detect rhinovirus and different rhinovirus (RV) genotypes using molecular methods.

	Available Format(s)
Catalogue Number	QAV064143_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q2

Specifications

Target Pathogen – Rhinovirus (RV) Matrix – Transport Medium Sample Volume – 1.0 ml Analysis Type – Qualitative Format – Liquid frozen Accreditation – ISO17043

VARICELLA-ZOSTER VIRUS (VZV)

VZVDNA19

Designed to evaluate the ability to detect different types and concentrations of Varicella-Zoster virus (VZV) using molecular methods.

	Available Format(s)	
Catalogue Number	QAV034103_1	QAV034103_2
Total Number of Challenges	1	2
Number of Samples	8 to 12	4 to 6
Distribution / Testing Period	Q3	Q1 and Q3

Specifications

Target Pathogen – Varicella-Zoster virus (VZV) Matrix – Transport Medium Sample Volume – 1.0 ml Analysis Type – Qualitative. Quantitative for information purposes only Format – Liquid frozen Accreditation – ISO17043

WEST NILE VIRUS (WNV)

WNVRNA19

Designed to evaluate the ability to detect West Nile virus and distinguish West Nile virus from other flaviviruses using molecular methods.

	Available Format(s)
Catalogue Number	QAV104141_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q2

Specifications

Target Pathogen – West Nile virus (WNV) Matrix – Transport Medium Sample Volume – 1.0 ml Analysis Type – Qualitative. Quantitative for information purposes only Format – Lyophilised Accreditation – ISO17043

ZIKA VIRUS

ZIKA19

Designed to evaluate the ability to detect Zika virus and determine the proficiency of laboratories in distinguishing Zika virus from other flaviviruses.

	Available Format(s)
Catalogue Number	QAV164186_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q2

Specifications

Target Pathogen – Zika virus Matrix – Transport Medium Sample Volume – 1.0 ml Analysis Type – Qualitative. Quantitative for information purposes only Format – Lyophilised Accreditation – Pending accreditation

PILOT STUDIES

ARTHROPOD-BORNE VIRUSES

ARBO19

Designed to evaluate the ability to detect different Arthropod-borne viruses (including viruses from *Flavi-, Toga-, Bunya-,* and/or *Reoviridae* families) using routine molecular methods. The panel is designed to represent various clinical scenarios and may include medically important arboviruses such as Tick-borne encephalitis viruses, sandfly fever viruses, Japanese encephalitis viruses, rift valley fever viruses, Usutu virus, Murray Valley encephalitis virus and St. Louis encephalitis virus.

	Available Format(s)
Catalogue Number	QAM194206_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q2

Specifications

Target Pathogen – Various flavivirus, Togavirus, Bunyavirus and/or Reoviridae families NA Target Source – Cultured and/or Clinical material Matrix – Transport Medium Sample Volume – 1 ml Analysis Type – Qualitative. Quantitative for information purposes only Format – Lyophilised

ATYPICAL MYCOBACTERIUM

NTM19

Designed to evaluate the ability to detect and differentiate Atypical mycobacterium or non-tuberculous mycobacterium using routine molecular methods.

	Available Format(s)
Catalogue Number	QAB194208_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q2

Specifications

Target Pathogen – Atypical mycobacterium or non-tuberculous mycobacterium (NTM) NA Target Source – Cultured and/or Clinical material Matrix – Transport Medium and/or Physiological Buffer Sample Volume – 1 ml Analysis Type – Qualitative Format – Liquid ready-to-use

CENTRAL NERVOUS SYSTEM I (VIRAL)

CNSI19

Designed to evaluate the ability to detect and determine various enterovirus, parechovirus, Herpes simplex virus 1/2, Varicella-Zoster virus and JC virus strains using routine molecular methods. The panel is designed to represent various clinical scenarios.

	Available Format(s)
Catalogue Number	QAV174195_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q4

Specifications

Target Pathogen – Various enterovirus, parechovirus, HSV1, HSV2, VZV and JCV
NA Target Source – Cultured and/or Clinical material
Matrix – Transport Medium
Sample Volume – 1.0 ml
Analysis Type – Qualitative. Quantitative for information purposes only
Format – Liquid frozen

CENTRAL NERVOUS SYSTEM II (NON-VIRAL)

CNSII19

Designed to evaluate the ability to detect and determine various Listeria spp., Neisseria meningitidis, Streptococcus pneumoniae, Streptococcus agalactiae, Escherichia coli K1, Aspergillus spp. and Haemophilus influenzae strains using routine molecular methods. The panel is designed to represent various clinical scenarios.

	Available Format(s)
Catalogue Number	QAM174196_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q4

Specifications

Target Pathogen – Various Listeria spp., Neisseria meningitidis, Streptococcus pneumoniae, Streptococcus agalactiae, E coli K1, Aspergillus spp. or Haemophilus influenzae strains
NA Target Source – Cultured and/or Clinical material
Matrix – Transport Medium
Sample Volume – 1.0 ml
Analysis Type – Qualitative. Quantitative for information purposes only
Format – Liquid frozen

HUMAN PAPILLOMAVIRUS (SUREPATH)

HPVSURE19

Designed to evaluate the ability to detect different high-risk Human Papillomavirus (HPV) types within a SurePath™ matrix using routine molecular methods.

	Available Format(s)
Catalogue Number	QAV184204_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q4

Specifications

Target Pathogen – HPV NA Target Source – Clinical material and/or cell lines containing HPV Matrix – Transport Medium (SurePath) Sample Volume – 2.0 ml Analysis Type – Qualitative Format – Lyophilised

IMMUNOCOMPROMISED

IC19

Designed to evaluate the ability to detect and determine various Toxoplasma gondii, Pneumocystis jirovecii, Aspergillus spp., Candida albicans and JC virus strains using routine molecular methods. The panel is designed to represent various clinical scenarios.

	Available Format(s)
Catalogue Number	QAM174197_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q4

Specifications

Target Pathogen – Various Toxoplasma gondii, Pneumocystis jirovecii, Aspergillus spp., Candida albicans and JC virus strains

NA Target Source – Cultured and/or Clinical material Matrix – Plasma and/or Synthetic Sputum and/or Saline Sample Volume – 1.0 ml Analysis Type – Qualitative. Quantitative for information purposes only Format – Liquid frozen

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MYCOBACTERIUM TUBERCULOSIS DRUG RESISTANCE

MTBDR19

Designed to evaluate the ability to detect and differentiate *Mycobacterium tuberculosis* drug resistant strains using routine molecular methods.

	Available Format(s)
Catalogue Number	QAB194209_1
Total Number of Challenges	1
Number of Samples	6 to 10
Distribution / Testing Period	Q4

Specifications

Target Pathogen – Mycobacterium tuberculosis NA Target Source – Cultured and/or Clinical material Matrix – Sputum and/or Synthetic Sputum and/or Synthetic CSF Analysis Type – Molecular typing Format – Liquid ready-to-use

MYCOPLASMA GENITALIUM

MG19

Designed to evaluate the ability to detect *Mycoplasma genitalium* using routine molecular methods.

	Available Format(s)
Catalogue Number	QAB184205_1
Total Number of Challenges	1
Number of Samples	6 to 10
Distribution / Testing Period	Q3

Specifications

Target Pathogen – Mycobacterium genitalium NA Target Source – Cultured and/or Clinical material Matrix – Urine and/or Saline Sample Volume – 4.0 ml Analysis Type – Qualitative Format – Liquid frozen

NEONATAL / NEW-BORN INFECTIONS

NEO19

Designed to evaluate the ability to detect and determine various cytomegalovirus, enterovirus, parechovirus, Herpes simplex virus 1/2, *Toxoplasma gondii* and Group B *Streptococcus* strains using routine molecular methods. The panel is designed to represent various clinical scenarios.

	Available Format(s)
Catalogue Number	QAM174199_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q4

Specifications

Target Pathogen – Various cytomegalovirus, enterovirus, parechovirus, Herpes simplex virus 1/2, Toxoplasma gondii and Group B Streptococcus strains NA Target Source – Cultured and/or Clinical material Matrix – Plasma and/or Transport Medium Sample Volume – 1.0 ml Analysis Type – Qualitative Format – Liquid frozen

TORQUE TENO VIRUS

TTV19

Designed to evaluate the ability to detect Torque teno virus (TTV) using routine molecular diagnostic platforms and procedures.

	Available Format(s)
Catalogue Number	QAV184203_1
Total Number of Challenges	1
Number of Samples	6 to 10
Distribution / Testing Period	Q4

Specifications

Target Pathogen – TTV NA Target Source – Cultured and/or Clinical material Matrix – Transport Medium Sample Volume – 1.0 ml Analysis Type – Qualitative Format – Liquid frozen

TRANSPLANTATION (VIRAL)

TRANS19

Designed to evaluate the ability to detect and determine various cytomegalovirus, Epstein-Barr virus, Human herpes virus 6, BK virus, B19 virus and adenovirus strains. The panel is designed to represent various clinical scenarios. Participating laboratories will be expected to test each panel using their appropriate molecular methods and to report their individual test results to QCMD.

	Available Format(s)
Catalogue Number	QAM174198_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q3

Specifications

Target Pathogen – Various EBV, HHV6, BKV, B19 and ADV NA Target Source – Cultured and/or Clinical material Matrix – Plasma and/or Transport Medium Sample Volume – 1.0 ml Analysis Type – Qualitative & Quantitative Format – Liquid frozen

TRICHOMONAS VAGINALIS

TV19

Designed to evaluate the ability to detect *Trichomonas vaginalis* using routine molecular methods.

	Available Format(s)
Catalogue Number	QAP184202_1
Total Number of Challenges	1
Number of Samples	6 to 10
Distribution / Testing Period	Q3

Specifications

Target Pathogen – Trichomonas vaginalis NA Target Source – Cultured and/or Clinical material Matrix – Transport Medium Sample Volume – 1.0 ml Analysis Type – Qualitative Format – Liquid frozen

YELLOW FEVER VIRUS

YFV19

Designed to evaluate the ability to detect Yellow fever virus and to distinguish Yellow fever virus from other flaviviruses using routine molecular methods.

	Available Format(s)
Catalogue Number	QAM194207_1
Total Number of Challenges	1
Number of Samples	8 to 12
Distribution / Testing Period	Q2

Specifications

Target Pathogen – Yellow fever virus NA Target Source – Cultured and/or Clinical material Matrix – Transport Medium Sample Volume – 1.0 ml Analysis Type – Qualitative. Quantitative for information purposes only Format – Lyophilised

"DESIGNED TO EVALUATE A LABORATORY'S ABILITY TO DETECT A WIDE RANGE OF MOLECULAR INFECTIOUS DISEASES."

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Adenovirus							Pg 22	
ADVDNA19	QAV054133_1 QAV054133_2	1 2	8 to 12 4 to 6	Q3 Q2, Q3	Dry-ice	Qualitative & Quantitative	Viral EQA	
Arthropod-bo	rne viruses						Pg 45	
ARBO19	QAM194206_1	1	8 to 12	Q2	Ambient	Qualitative	Pilot Study	
Aspergillus sp	р.						Pg 15	
ASPDNA19	QAF104140_1	1	8 to 12	Q3	Dry-ice	Qualitative	Fungal EQA	
Atypical myc	obacterium						Pg 45	
NTM19	QAB194208_1	1	8 to 12	Q2	Ambient	Qualitative	Pilot Study	
B19 virus							Pg 23	
B19DNA19	QAV034116_1 QAV034116_2	1 2	8 4	Q3 Q1, Q3	Dry-ice	Qualitative & Quantitative	Viral EQA	
Bacterial 16S	Ribosomal RNA						Pg 4	
B16SrRNA19	QAB164183_1	1	8 to 10	Q4	Dry-ice	Typing	Bacterial EQA	
Bacterial Gastroenteritis Pg 17							Pg 17	
GastroB19	QAB124153_1	1	8 to 12	Q4	Dry-ice	Qualitative	Multi-Pathogen / Syndromic EQA	
BK virus (BKV)							Pg 23	
BKDNA19	QAV144166_1 QAV144166_2	1 2	8 to 12 4 to 6	Q3 Q2, Q3	Dry-ice	Qualitative & Quantitative	Viral EQA	
Bordetella per	rtussis						Pg 4	
BPDNA19	QAB094132_1	1	8 to 12	Q2	Dry-ice	Qualitative	Bacterial EQA	
Borrelia burgo	lorferi spp. (Lyme	e Disease)					Pg 5	
BbDNA19	QAB114147_1	1	8 to 12	Q3	Dry-ice	Qualitative	Bacterial EQA	
Candida spp.							Pg 15	
CANDNA19	QAF124151_1	1	8 to 12	Q3	Dry-ice	Qualitative	Fungal EQA	
Central Nervo	us System I (Vira	l)					Pg 46	
CNSI19	QAV174195_1	1	8 to 12	Q4	Dry-ice	Qualitative	Pilot Study	
Central Nervo	ous System II (Noi	n-viral)					Pg 46	
CNSII19	QAM174196_1	1	8 to 12	Q4	Dry-ice	Qualitative	Pilot Study	
Chikungunya	virus (CHIKV)						Pg 24	
CHIKV19	QAV154175_1	1	8 to 12	Q2	Ambient	Qualitative	Viral EQA	
Chlamydia ps	sittaci						Pg 5	
CPS19	QAB134165_1	1	8 to 10	Q2	Dry-ice	Qualitative	Bacterial EQA	
Chlamydia tro	achomatis						Pg 6	
CTDNA19	QAB004101_1 QAB004101_2	1 2	8 to 12 4 to 6	Q3 Q1, Q3	Dry-ice	Qualitative	Bacterial EQA	

TARGET PATHOGEN PA								
PROGRAMME CODE	CATALOGUE NUMBER	NO. OF CHALLENGES	PANEL MEMBERS PER CHALLENGE	DISTRIBUTION DATE(S)/ TESTING PERIOD	SHIPPING CONDITIONS	ANALYSIS TYPE	PROGRAMME TYPE	
Chlamydia trachomatis and Neisseria gonorrhoeae Pg 6								
CTNg19	QAB174191_1 QAB174191_2	1 2	8 to 12 4 to 6	Q3 Q1, Q3	Dry-ice	Qualitative	Multi-Pathogen / Syndromic EQA	
Chlamydophi	Chlamydophila pneumoniae							
CP19	QAB084107_1	1	5 to 10	Q2	Dry-ice	Qualitative	Bacterial EQA	
Clostridium di	ifficile (CD)						Pg 7	
CDDNA19	QAB084125_1 QAB084125_2	1 2	8 to 12 4 to 6	Q4 Q2, Q4	Dry-ice	Qualitative	Bacterial EQA	
Coronavirus (CoV)						Pg 24	
CVRNA19	QAV064137_1	1	8 to 12	Q2	Dry-ice	Qualitative	Viral EQA	
Cytomegalov	virus (CMV) Dried	Blood Spots					Pg 25	
CMVDBS19	QAV064127_1	1	8 to 12	Q2	Ambient	Qualitative	Viral EQA	
Cytomegalov	rirus (CMV) Drug	Resistance					Pg 25	
CMVDR19	QAV144169_1	1	4 to 7	Q2	Dry-ice	Drug Resistance / Sequencing	Viral EQA	
Cytomegalov	rirus (CMV)						Pg 26	
CMVDNA19	QAV014120_1 QAV014120_2	1 2	8 to 12 4 to 6	Q3 Q2, Q3	Dry-ice	Qualitative & Quantitative	Viral EQA	
Cytomegalov	rirus (CMV) Whole	e Blood					Pg 26	
CMVWB19	QAV124150_1 QAV124150_2	1 2	8 to 12 4 to 6	Q3 Q2, Q3	Dry-ice	Qualitative & Quantitative	Viral EQA	
Dengue virus	(DENV)						Pg 27	
DENVRNA19	QAV114148_1	1	8 to 12	Q2	Ambient	Qualitative	Viral EQA	
Dermatophyte	osis						Pg 16	
DERMA19	QAF164187_1	1	8 to 10	Q3	Dry-ice	Qualitative	Fungal EQA	
Diarrheagenia	c Escherichia col	i					Pg 8	
E.COLI19	QAB154179_1	1	8 to 12	Q4	Dry-ice	Qualitative	Bacterial EQA	
Enterovirus (E	V)						Pg 27	
EVRNA19	QAV984104_1 QAV984104_2	1 2	8 to 12 4 to 6	Q3 Q1, Q3	Dry-ice	Qualitative	Viral EQA	
Enterovirus Ty	ping (EV)						Pg 28	
EVTP19	QAV164185_1	1	5 to 10	Q1	Dry-ice	Typing	Viral EQA	
Epstein-Barr v	irus (EBV)						Pg 28	
EBVDNA19	QAV024121_1 QAV024121_2	1 2	8 to 12 4 to 6	Q3 Q2, Q3	Dry-ice	Qualitative & Quantitative	Viral EQA	
Epstein-Barr v	irus (EBV) Whole	Blood					Pg 29	
EBVWB19	QAV134161_1 QAV134161_2	1 2	8 to 12 4 to 6	Q3 Q2, Q3	Dry-ice	Qualitative & Quantitative	Viral EQA	
Extended Spe	ctrum ß-lactamo	ase and Carbo	penemase				Pg 8	
ESBL19	QAB134162_1	1	8 to 12	Q3	Dry-ice	Typing	Bacterial EQA	
Group B Strep	tococcus						Pg 9	
GBS19	QAB174200_1	1	8 to 12	Q4	Dry-ice	Qualitative	Bacterial EQA	

TARGET PATHO	OGEN						PAGE NUMBER
PROGRAMME CODE	CATALOGUE NUMBER	NO. OF CHALLENGES	PANEL MEMBERS PER CHALLENGE	DISTRIBUTION DATE(S)/ TESTING PERIOD	SHIPPING CONDITIONS	ANALYSIS TYPE	PROGRAMME TYPE
Helicobacter	pylori						Pg 9
H.PYLORI19	QAB164190_1	1	5 to 10	Q3	Dry-ice	Qualitative	Bacterial EQA
Hepatitis A vir	us (HAV)						Pg 29
HAVRNA19	QAV124156_1 QAV124156_2	1 2	8 to 10 4 to 5	Q3 Q1, Q3	Dry-ice	Qualitative	Viral EQA
Hepatitis B vire	us (HBV)						Pg 30
HBVDNA19	QAV994110_1 QAV994110_2 QAV994110_4	1 2 4	8 4 4	Q3 Q1, Q3 Q1, Q2, Q3, Q4	Dry-ice	Qualitative & Quantitative	Viral EQA
Hepatitis B vire	us (HBV) Drug Re	sistance					Pg 30
HBVDR19	QAV124160_1	1	4 to 7	Q3	Dry-ice	Drug Resistance / Sequencing	Viral EQA
Hepatitis B vire	us (HBV) Genotyj	ping					Pg 31
HBVGT19	QAV064118_1	1	8 to 12	Q1	Dry-ice	Typing	Viral EQA
Hepatitis C vir	us (HCV)						Pg 31
HCVRNA19	QAV994112_1 QAV994112_2 QAV994112_4	1 2 4	8 4 4	Q3 Q1, Q3 Q1, Q2, Q3, Q4	Dry-ice	Qualitative & Quantitative	Viral EQA
Hepatitis C vir	us (HCV) Drug Re	esistance					Pg 32
HCVDR19	QAV134167_1	1	4 to 7	Q3	Dry-ice	Drug Resistance / Sequencing	Viral EQA
Hepatitis C vir	us (HCV) Genoty	ping					Pg 32
HCVGT19	QAV034117_1	1	8 to 12	Q1	Dry-ice	Typing	Viral EQA
Hepatitis D vir	us (HDV)						Pg 33
HDV19	QAV144170_1	1	8 to 10	Q3	Dry-ice	Qualitative & Quantitative	Viral EQA
Hepatitis E viru	us (HEV)						Pg 33
HEVRNA19	QAV124157_1	1	8 to 10	Q3	Dry-ice	Qualitative	Viral EQA
Herpes simple	ex virus 1 & 2 (HS)	V)					Pg 34
HSVDNA19	QAV994105_1 QAV994105_2	1 2	8 to 12 4 to 6	Q3 Q1, Q3	Dry-ice	Qualitative	Viral EQA
Herpes simple	ex virus Drug Resi	stance					Pg 34
HSVDR19	QAV164184_1	1	4 to 7	Ql	Dry-ice	Sequence Analysis	Viral EQA
Human herpe	s virus 6 (HHV6)						Pg 35
HHV6DNA19	QAV084119_1 QAV084119_2	1 2	8 to 12 4 to 6	Q3 Q2, Q3	Dry-ice	Qualitative & Quantitative	Viral EQA
Human Immu	nodeficiency viru	us type 1 (HIV-	-1) – DNA				Pg 35
HIVDNA19	QAV034114_1 QAV034114_2	1 2	8 4	Q3 Q1, Q3	Dry-ice	Qualitative	Viral EQA
Human Immu	nodeficiency viru	us type 1 (HIV-	-1) – Drug Resisto	ance			Pg 36
HIVDR19	QAV024131_1	1	4 to 7	Q3	Dry-ice	Drug Resistance / Sequencing	Viral EQA
Human Immu	nodeficiency viru	us type 1 (HIV-	-1) – Drug Resisto	ance (Integrase)			Pg 36
HIVDRint19	QAV114146_1	1	4 to 7	Q3	Dry-ice	Drug Resistance / Sequencing	Viral EQA

TARGET PATHOGEN							PAGE NUMBER	
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Human Immunodeficiency virus type 1 (HIV-1) – RNA								
HIVRNA19	QAV994108_1 QAV994108_2 QAV994108_4	1 2 4	8 4 8	Q3 Q1, Q3 Q1, Q2, Q3, Q4	Dry-ice	Qualitative & Quantitative	Viral EQA	
Human metap	oneumovirus (MF	۷۷)					Pg 37	
MPV19	QAV054135_1	1	8 to 12	Q2	Dry-ice	Qualitative	Viral EQA	
Human Papillo	omavirus (HPV) -	- PreservCyt					Pg 38	
HPVPRES19	QAV094130_1 QAV094130_2	1 2	8 to 12 4 to 6	Q4 Q2, Q4	Ambient / Specialist	Qualitative	Viral EQA	
Human Papille	omavirus (Surepo	ath)					Pg 47	
HPVSURE19	QAV184204_1	1	8 to 12	Q4	Ambient	Qualitative	Pilot Study	
Immunocomp	promised						Pg 47	
IC19	QAM174197_1	1	8 to 12	Q4	Dry-ice	Qualitative	Pilot Study	
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INFRNA19	QAV054134_1 QAV054134_2	1 2	8 to 12 4 to 6	Q4 Q2, Q4	Dry-ice	Qualitative	Viral EQA	
Influenza Hae	magglutinin Typi	ing (HA)					Pg 39	
INFHT19	QAV064138_1	1	5 to 10	Q4	Dry-ice	Typing	Viral EQA	
JC virus (JCV)							Pg 39	
JCDNA19	QAV074106_1 QAV074106_2	1 2	8 to 12 4 to 6	Q3 Q2, Q3	Dry-ice	Qualitative & Quantitative	Viral EQA	
Legionella pr	eumophila						Pg 10	
LPDNA19	QAB044122_1	1	8 to 12	Q1	Dry-ice	Qualitative	Bacterial EQA	
MALDI-TOF			_				Pg 17	
MALDI19	QAB124155_1	1	8 to 12	Q3	Dry-ice	Typing	Multi-Pathogen / Syndromic EQA	
Measles / Mu	mps						Pg 40	
MM19	QAV144171_1	1	8 to 12	Q3	Dry-ice	Qualitative	Viral EQA	
MERS corona	virus (MERS-CoV)						Pg 40	
MERS19	QAV154181_1	1	6 to 10	Q2	Dry-ice	Qualitative	Viral EQA	
Methicillin Res	sistant Staphyloc	occus aureus	(MRSA)				Pg 10	
MRSADNA19	QAB064124_1	1	8 to 12	Q2	Ambient	Qualitative	Bacterial EQA	
Methicillin Res	sistant Staphyloc	occus aureus	(MRSA) – Typing				Pg 11	
MRSATP19	QAB074128_1	1	8 to 12	Q2	Ambient	Typing	Bacterial EQA	
Mycobacteriu	ım tuberculosis (MTB)					Pg 11	
MTBDNA19	QAB014129_1 QAB014129_2	1 2	8 to 12 4 to 6	Q4 Q2, Q4	Ambient	Qualitative	Bacterial EQA	
Mycobacteriu	um tuberculosis [Drug Resistanc	e				Pg 48	
MTBDR19	QAB194209_1	1	6 to 10	Q4	Ambient	Typing	Pilot Study	
Mycoplasma	genitalium						Pg 48	
MG19	QAB184205_1	1	6 to 10	Q3	Dry-ice	Qualitative	Pilot Study	
Mycoplasma	pneumoniae						Pg 12	
MP19	QAB174192_1	1	5 to 10	Q2	Dry-ice	Qualitative	Bacterial EQA	

TARGET PATHO	OGEN						PAGE NUMBER
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Mycoplasma spp. (cell contamination)							
MYCO19	QAB144168_1	1	8 to 12	Q4	Dry-ice	Qualitative & Quantitative	Bacterial EQA
Neisseria gon	orrhoeae						Pg 13
NgDNA19	QAB034126_1 QAB034126_2	1 2	8 to 12 4 to 6	Q3 Q1, Q3	Dry-ice	Qualitative	Bacterial EQA
Neonatal / Ne	ew-Born infection	IS					Pg 49
NEO19	QAM174199_1	1	8 to 12	Q4	Dry-ice	Qualitative	Pilot Study
Norovirus (NV)						Pg 41
NVRNA19	QAV084139_1 QAV084139_2	1 2	8 to 12 4 to 6	Q4 Q2, Q4	Dry-ice	Qualitative	Viral EQA
Parainfluenza	virus (PIV)						Pg 41
PINFRNA19	QAV064136_1	1	8 to 12	Q2	Dry-ice	Qualitative	Viral EQA
Parasitic Gast	roenteritis						Pg 18
GastroP19	QAP124154_1	1	8 to 12	Q4	Dry-ice	Qualitative	Multi-Pathogen / Syndromic EQA
Parechovirus	(HPeV)						Pg 42
PeVRNA19	QAV114145_1 QAV114145_2	1 2	8 to 12 4 to 6	Q3 Q1, Q3	Dry-ice	Qualitative	Viral EQA
Pneumocystis jirovecii pneumonia (PCP)						Pg 16	
PCPDNA19	QAF114144_1	1	8 to 12	Q3	Dry-ice	Qualitative & Quantitative	Fungal EQA
Respiratory I							Pg 18
RESPI19	QAV164188_1	1	8 to 12	Q2	Dry-ice	Qualitative	Multi-Pathogen / Syndromic EQA
Respiratory II							Pg 19
RESPII19	QAV164189_1	1	8 to 12	Q2	Dry-ice	Qualitative	Multi-Pathogen / Syndromic EQA
Respiratory III							Pg 19
RESPIII 19	QAM174193_1	1	8 to 12	Q4	Dry-ice	Qualitative	Multi-Pathogen / Syndromic EQA
Respiratory sy	vncytial virus (RS)	/)					Pg 42
RSV19	QAV054142_1 QAV054142_2	1 2	8 to 12 4 to 6	Q4 Q2, Q4	Dry-ice	Qualitative	Viral EQA
Rhinovirus (RV	()						Pg 43
RVRNA19	QAV064143_1	1	8 to 12	Q2	Dry-ice	Qualitative	Viral EQA
Sepsis							Pg 20
SEPSIS19	QAB164178_1	1	8 to 12	Q4	Dry-ice	Qualitative	Multi-Pathogen / Syndromic EQA
Sexually Trans	smitted Infections	s I					Pg 20
STI_119	QAB154177_1	1	8 to 12	Q4	Dry-ice	Qualitative	Multi-Pathogen / Syndromic EQA
Sexually Trans	smitted Infections	s II					Pg 21
STI_II19	QAM174201_1	1	8 to 12	Q4	Dry-ice	Qualitative	Multi-Pathogen / Syndromic EQA

TARGET PATHO	OGEN						PAGE NUMBER
PROGRAMME CODE	CATALOGUE NUMBER	NO. OF CHALLENGES	PANEL MEMBERS PER CHALLENGE	DISTRIBUTION DATE(S)/ TESTING PERIOD	SHIPPING CONDITIONS	ANALYSIS TYPE	PROGRAMME TYPE
Staphylococo	cus aureus spa						Pg 13
SASPA19	QAB134164_1	1	6 to 12	Q2	Ambient	Typing	Bacterial EQA
Syphilis							Pg 14
SYPH19	QAB154180_1	1	5 to 10	Q4	Dry-ice	Qualitative	Bacterial EQA
Torque teno v	irus (TTV)						Pg 49
TTV19	QAV184203_1	1	6 to 10	Q4	Dry-ice	Qualitative	Pilot Study
Toxoplasma g	gondii						Pg 22
TGDNA19	QAP044123_1 QAP044123_2	1 2	8 to 12 4 to 6	Q4 Q2, Q4	Ambient	Qualitative	Parasitic EQA
Transplantatio	on (viral)						Pg 50
TRANS19	QAM174198_1	1	8 to 12	Q3	Dry-ice	Qualitative & Quantitative	Pilot Study
Trichomonas	vaginalis						Pg 50
TV19	QAP184202_1	1	6 to 10	Q3	Dry-ice	Qualitative	Pilot Study
Vancomycin	Resistant Enteroc	cocci (VRE)					Pg 14
VRE19	QAB134163_1	1	8 to 12	Q3	Dry-ice	Qualitative	Bacterial EQA
Varicella-Zost	ter virus (VZV)						Pg 43
VZVDNA19	QAV034103_1 QAV034103_2	1 2	8 to 12 4 to 6	Q3 Q1, Q3	Dry-ice	Qualitative	Viral EQA
Viral Gastroe	nteritis						Pg 21
GastroV19	QAV124152_1	1	8 to 12	Q4	Dry-ice	Qualitative	Multi-Pathogen / Syndromic EQA
West Nile viru	s (WNV)						Pg 44
WNVRNA19	QAV104141_1	1	8 to 12	Q2	Ambient	Qualitative	Viral EQA
Yellow fever	/irus						Pg 51
YFV19	QAM194207_1	1	8 to 12	Q2	Ambient	Qualitative	Pilot Study
Zika virus							Pg 44
ZIKA19	QAV164186_1	1	8 to 12	Q2	Ambient	Qualitative	Viral EQA





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