

EQA FOR MOLECULAR INFECTIOUS DISEASE TESTING

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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.



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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



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.



HIGH LEVEL OF PARTICIPATION

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



COMPREHENSIVE REPORTS

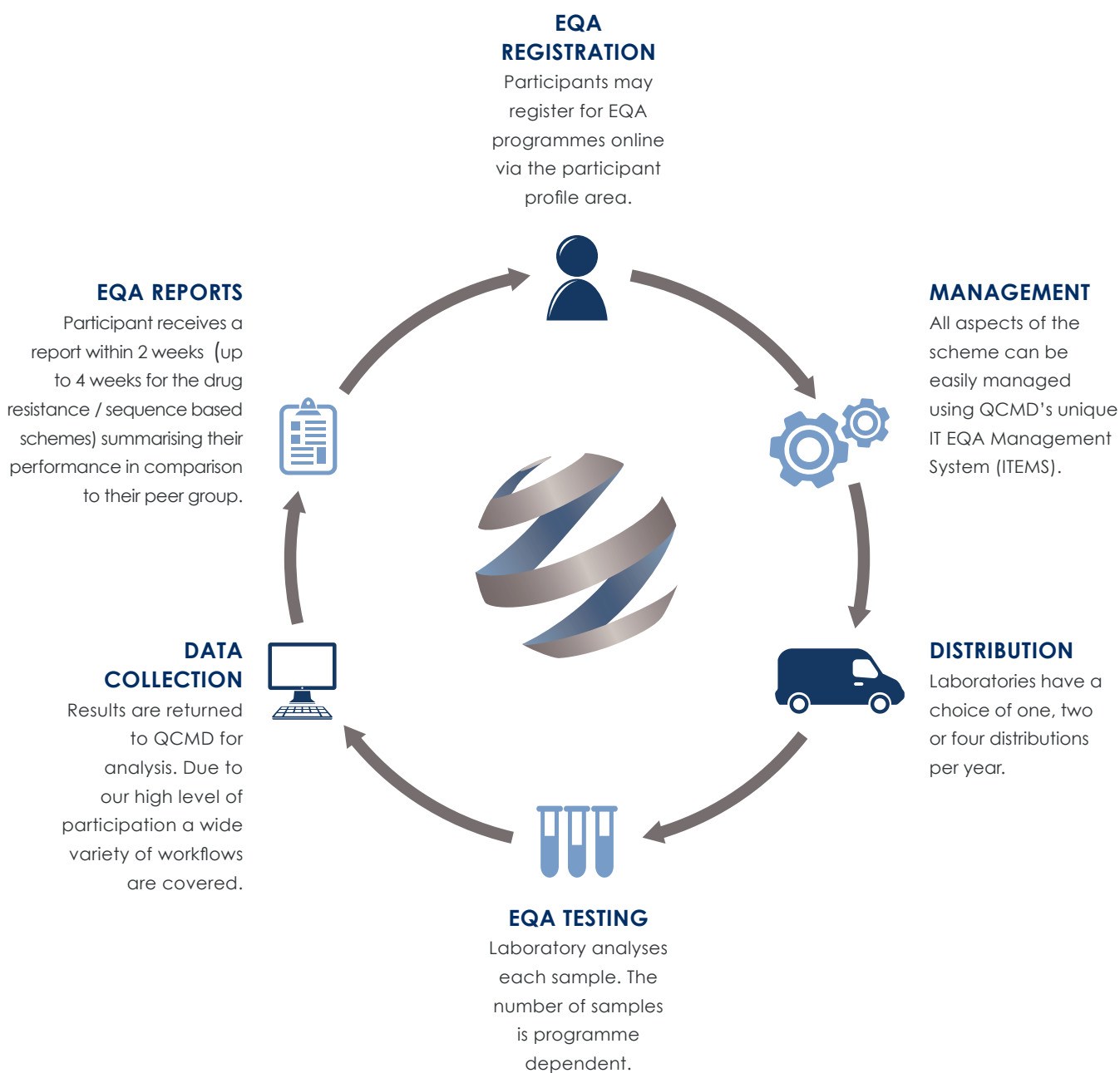
Individual reports are provided with each EQA challenge. In line with the requirements of ISO17043, they provide the laboratories with their results and performance assessment in relation to their EQA assessment group (peer review group).

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

B16SrRNA20

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

BPDNA20

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)

BbDNA20

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.

Matrix – Microbiological Medium and/or Transport Medium

Sample Volume – 1.0 ml

Analysis Type – Qualitative

Format – Liquid frozen

Accreditation – ISO17043

CHLAMYDIA PSITTACI

CPS20

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

CTDNA20

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

CTNg20

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

CP20

Designed to evaluate the ability to detect *Chlamydomphila 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 – *Chlamydomphila 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)

CDDNA20

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.COLI20

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 β -LACTAMASE AND CARBAPENEMASE

ESBL20

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 – Various bacteria carrying ESBL and carbapenemase genes

Matrix – Physiological Buffer

Sample Volume – 0.5 ml

Analysis Type – Molecular typing

Format – Liquid frozen

Accreditation – Pending accreditation

GROUP B STREPTOCOCCUS

GBS20

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 – *Streptococcus agalactiae*

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.PYLORI20

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 – *Helicobacter 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

LPDNA20

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)

MRSADNA20

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)

MRSATP20

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)

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)

MTBDNA20

Designed to evaluate the ability to detect *Mycobacterium tuberculosis* 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*

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

MP20

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)

MYCO20

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* species

Matrix – Physiological Buffer

Sample Volume – 1.2 ml

Analysis Type – Qualitative & Quantitative

Format – Liquid frozen

Accreditation – Pending accreditation

NEISSERIA GONORRHOEAE

NgDNA20

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

SASPA20

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

SYPH20

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)

VRE20

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.

ASPDNA20

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.

CANDNA20

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

DERMA20

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*

Matrix – Physiological Buffer

Sample Volume – 1.0 ml

Analysis Type – Qualitative

Format – Liquid frozen

Accreditation – Pending accreditation

PNEUMOCYSTIS JIROVECII PNEUMONIA (PCP)

PCPDNA20

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

GastroB20

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	QAB124153_2
Total Number of Challenges	1	2
Number of Samples	8 to 12	4 to 6
Distribution / Testing Period	Q4	Q2 & 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

MALDI20

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

GastroP20

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*, *Entamoeba*, *Dientamoeba* and *Blastocystis*.

	Available Format(s)	
Catalogue Number	QAP124154_1	QAP124154_2
Total Number of Challenges	1	2
Number of Samples	8 to 12	4 to 6
Distribution / Testing Period	Q4	Q2 & Q4

Specifications

Target Pathogen – *Giardia*, *Cryptosporidium*, *Entamoeba*, *Dientamoeba* and *Blastocystis*

Matrix – Synthetic Faecal Matrix and/or Physiological Buffer

Sample Volume – 1.0 ml

Analysis Type – Qualitative

Format – Liquid frozen

Accreditation – Pending accreditation

RESPIRATORY I

RESPI20

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	QAV164188_2
Total Number of Challenges	1	2
Number of Samples	8 to 12	4 to 6
Distribution / Testing Period	Q3	Q1 & Q3

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

RESPII20

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	QAV164189_2
Total Number of Challenges	1	2
Number of Samples	8 to 12	4 to 6
Distribution / Testing Period	Q2	Q1 & Q3

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

RESPIII20

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	QAM174193_2
Total Number of Challenges	1	2
Number of Samples	8 to 12	4 to 6
Distribution / Testing Period	Q3	Q1 & Q3

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

SEXUALLY TRANSMITTED INFECTIONS I

STI_I20

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	QAB154177_2
Total Number of Challenges	1	2
Number of Samples	8 to 12	4 to 6
Distribution / Testing Period	Q4	Q2 & Q3

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_II20

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	QAM174201_2
Total Number of Challenges	1	2
Number of Samples	8 to 12	4 to 6
Distribution / Testing Period	Q4	Q2 & Q3

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

GastroV20

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	QAV124152_2
Total Number of Challenges	1	2
Number of Samples	8 to 12	4 to 6
Distribution / Testing Period	Q4	Q2 & 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

TGDNA20

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)

ADVDNA20

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

B19DNA20

Designed to evaluate the ability to detect and quantitate 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)

BKDNA20

Designed to evaluate the ability to detect and quantitate 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 – BK virus

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)

CHIKV20

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)

CVRNA20

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

CMVDBS20

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/ml however other units will be accepted

Sample Volume – 2 x 50µl

Analysis Type – Qualitative. Quantitative for information purposes only

Format – Dried blood spot

Accreditation – ISO17043

CYTOMEGALOVIRUS (CMV) DRUG RESISTANCE

CMVDR20

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)

CMVDNA20

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

CMVWB20

Designed to evaluate the ability to detect and quantitate 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)

DENVRNA20

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)

EVRNA20

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)

EVTP20

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	Q1

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)

EBVDNA20

Designed to evaluate the ability to detect and quantitate 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

EBVWB20

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/ml however other units will be accepted

Sample Volume – 1.0 ml

Analysis Type – Qualitative & Quantitative

Format – Liquid frozen

Accreditation – ISO17043

HEPATITIS A VIRUS (HAV)

HAVRNA20

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)

HBVDNA20

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

HBVDR20

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

HBVGT20

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)

HCVRNA20

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

HCVDR20

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

HCVGT20

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)

HDV20

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)

HEVRNA20

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)

HSV DNA20

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

HSV DR20

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	Q1

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)

HHV6DNA20

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

HIVDNA20

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

HIVDR20

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)

HIVDRint20

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

HIVRNA20

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/ml however other units will be accepted

Sample Volume – 1.2 ml

Analysis Type – Qualitative & Quantitative

Format – Liquid frozen

Accreditation – ISO17043

HUMAN METAPNEUMOVIRUS (MPV)

MPV20

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

HPVPRES20

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)

INFRNA20

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 TYPING (HA)

INFTP20

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 Typing (HA)

Matrix – Transport Medium

Sample Volume – 1.0 ml

Analysis Type – Molecular typing

Format – Liquid frozen

Accreditation – ISO17043

JC VIRUS (JCV)

JCDNA20

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

MM20

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)

MERS20

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)

NVRNA20

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)

PINFRNA20

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)

PeVRNA20

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)

RSV20

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)

RVRNA20

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)

VZVDNA20

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)

WNVRNA20

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

ZIKA20

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

ARBO20

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 – 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

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

NTM20

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 MENINGITIS AND ENCEPHALITIS)

CNSI20

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	QAV174195_2
Total Number of Challenges	1	2
Number of Samples	8 to 12	4 to 6
Distribution / Testing Period	Q4	Q2 & 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 MENINGITIS AND ENCEPHALITIS)

CNSII20

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	QAM174196_2
Total Number of Challenges	1	2
Number of Samples	8 to 12	4 to 6
Distribution / Testing Period	Q4	Q2 & 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 IMMUNODEFICIENCY VIRUS TYPE 2 (HIV-2)

HIV2_20

This pilot study assesses the proficiency of laboratories in detection and quantitation of human immunodeficiency virus type 2 (HIV-2).

	Available Format(s)	
Catalogue Number	QAV204212_1	QAV204212_2
Total Number of Challenges	1	2
Number of Samples	8	4
Distribution / Testing Period	Q3	Q1 & Q3

Specifications

Target Pathogen – Human Immunodeficiency virus type 2

NA Target Source – Cultured and/or Clinical material

Matrix – Plasma

Analysis Type – Qualitative & Quantitative

Format – Liquid frozen

HUMAN PAPILLOMAVIRUS (SUREPATH)

HPVSURE20

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 – Human papillomavirus

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

MYCOBACTERIUM TUBERCULOSIS DRUG RESISTANCE

MTBDR20

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

MG20

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 – *Mycoplasma 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

SEPSIS

SEPSIS20

Designed to evaluate a range of pathogens associated with sepsis such as *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

TORQUE TENO VIRUS

TTV20

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 – Torque teno virus

NA Target Source – Cultured and/or Clinical material

Matrix – Transport Medium

Sample Volume – 1.0 ml

Analysis Type – Qualitative

Format – Liquid frozen

TRANSPLANTATION (VIRAL)

TRANS20

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	QAM174198_2
Total Number of Challenges	1	2
Number of Samples	8 to 12	4 to 6
Distribution / Testing Period	Q3	Q2 & Q4

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

TV20

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

VIRAL METAGENOMICS NGS

NGSmeta_20

This EQA pilot study aims to assess performance of existing metagenomics protocols as currently implemented by participating laboratories. Samples will be provided which will mimic cerebrospinal fluid samples containing known viral pathogens including enterovirus, herpes simplex virus and influenza virus.

Performance will be assessed based on the qualitative identification of viruses present in the samples, at the family, genus, species and subtype levels.

	Available Format(s)
Catalogue Number	QAV204213_1
Total Number of Challenges	1
Number of Samples	4 to 7
Distribution / Testing Period	Q4

Specifications

Target Pathogen – Enterovirus, herpes simplex virus and influenza virus.

NA Target Source – Cultured material

Matrix – Synthetic CSF + human cell lines

Sample Volume – 1.0ml

Analysis Type – Sequence Analysis

Format – Liquid frozen

YELLOW FEVER VIRUS

YFV20

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

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APPENDIX

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Adenovirus							Pg 22
ADVDNA20	QAV054133_1 QAV054133_2	1 2	8 to 12 4 to 6	Q3 Q2, Q3	Dry-ice	Qualitative & Quantitative	Viral EQA
Arthropod-borne viruses							Pg 45
ARBO20	QAM194206_1	1	8 to 12	Q2	Ambient	Qualitative	Pilot Study
Aspergillus spp.							Pg 15
ASPDNA20	QAF104140_1	1	8 to 12	Q3	Dry-ice	Qualitative	Fungal EQA
Atypical mycobacterium							Pg 45
NTM20	QAB194208_1	1	8 to 12	Q2	Ambient	Qualitative	Pilot Study
B19 virus							Pg 23
B19DNA20	QAV034116_1 QAV034116_2	1 2	8 4	Q3 Q1, Q3	Dry-ice	Qualitative & Quantitative	Viral EQA
Bacterial 16S Ribosomal RNA							Pg 4
B16SrRNA20	QAB164183_1	1	8 to 10	Q4	Dry-ice	Typing	Bacterial EQA
Bacterial Gastroenteritis							Pg 17
GastroB20	QAB124153_1 QAB124153_2	1 2	8 to 12 4 to 6	Q4 Q2 & Q4	Dry-ice	Qualitative	Multi-Pathogen / Syndromic EQA
BK virus (BKV)							Pg 23
BKDNA20	QAV144166_1 QAV144166_2	1 2	8 to 12 4 to 6	Q3 Q2, Q3	Dry-ice	Qualitative & Quantitative	Viral EQA
Bordetella pertussis							Pg 4
BPDNA20	QAB094132_1	1	8 to 12	Q2	Dry-ice	Qualitative	Bacterial EQA
Borrelia burgdorferi spp. (Lyme Disease)							Pg 5
BbDNA20	QAB114147_1	1	8 to 12	Q3	Dry-ice	Qualitative	Bacterial EQA
Candida spp.							Pg 15
CANDNA20	QAF124151_1	1	8 to 12	Q3	Dry-ice	Qualitative	Fungal EQA
Central Nervous System I (Viral Meningitis and Encephalitis)							Pg 46
CNSI20	QAV174195_1 QAV174195_2	1 2	8 to 12 4 to 6	Q4 Q2 & Q4	Dry-ice	Qualitative	Pilot Study
Central Nervous System II (Non-viral Meningitis and Encephalitis)							Pg 46
CNSII20	QAM174196_1 QAM174196_2	1 2	8 to 12 4 to 6	Q4 Q2 & Q4	Dry-ice	Qualitative	Pilot Study
Chikungunya virus (CHIKV)							Pg 24
CHIKV20	QAV154175_1	1	8 to 12	Q2	Ambient	Qualitative	Viral EQA
Chlamydia psittaci							Pg 5
CPS20	QAB134165_1	1	8 to 10	Q2	Dry-ice	Qualitative	Bacterial EQA
Chlamydia trachomatis							Pg 6
CTDNA20	QAB004101_1 QAB004101_2	1 2	8 to 12 4 to 6	Q3 Q1, Q3	Dry-ice	Qualitative	Bacterial EQA

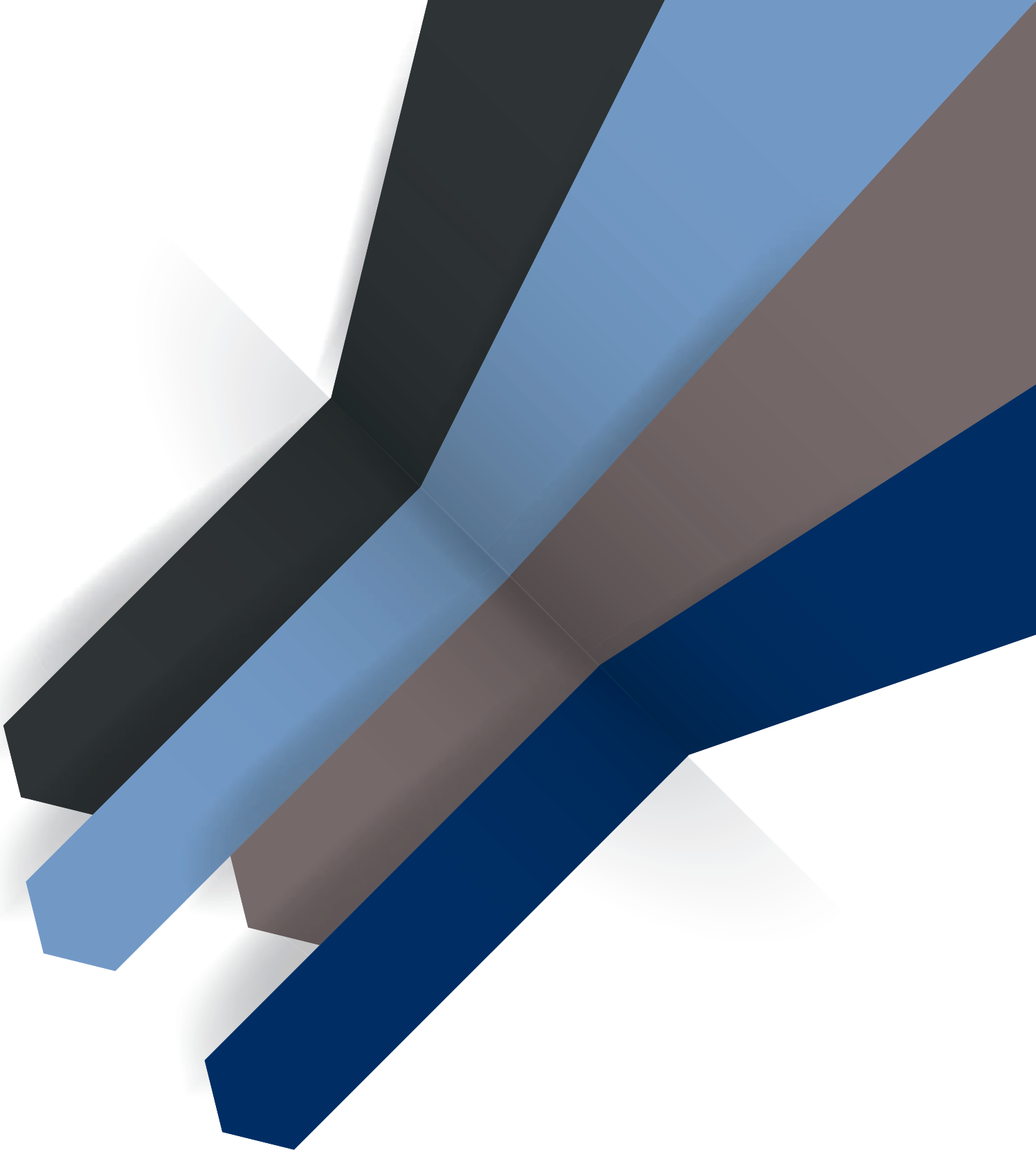
TARGET PATHOGEN							PAGE NUMBER
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Chlamydia trachomatis and Neisseria gonorrhoeae							Pg 6
CTNg20	QAB174191_1	1	8 to 12	Q3	Dry-ice	Qualitative	Multi-Pathogen / Syndromic EQA
	QAB174191_2	2	4 to 6	Q1, Q3			
Chlamydophila pneumoniae							Pg 7
CP20	QAB084107_1	1	5 to 10	Q2	Dry-ice	Qualitative	Bacterial EQA
Clostridium difficile (CD)							Pg 7
CDDNA20	QAB084125_1	1	8 to 12	Q4	Dry-ice	Qualitative	Bacterial EQA
	QAB084125_2	2	4 to 6	Q2, Q4			
Coronavirus (CoV)							Pg 24
CVRNA20	QAV064137_1	1	8 to 12	Q2	Dry-ice	Qualitative	Viral EQA
Cytomegalovirus (CMV) Dried Blood Spots							Pg 25
CMVDBS20	QAV064127_1	1	8 to 12	Q2	Ambient	Qualitative	Viral EQA
Cytomegalovirus (CMV) Drug Resistance							Pg 25
CMVDR20	QAV144169_1	1	4 to 7	Q2	Dry-ice	Drug Resistance / Sequencing	Viral EQA
Cytomegalovirus (CMV)							Pg 26
CMVDNA20	QAV014120_1	1	8 to 12	Q3	Dry-ice	Qualitative & Quantitative	Viral EQA
	QAV014120_2	2	4 to 6	Q2, Q3			
Cytomegalovirus (CMV) Whole Blood							Pg 26
CMVWB20	QAV124150_1	1	8 to 12	Q3	Dry-ice	Qualitative & Quantitative	Viral EQA
	QAV124150_2	2	4 to 6	Q2, Q3			
Dengue virus (DENV)							Pg 27
DENVRNA20	QAV114148_1	1	8 to 12	Q2	Ambient	Qualitative	Viral EQA
Dermatophytosis							Pg 16
DERMA20	QAF164187_1	1	8 to 10	Q3	Dry-ice	Qualitative	Fungal EQA
Diarrheagenic Escherichia coli							Pg 8
E.COLI20	QAB154179_1	1	8 to 12	Q4	Dry-ice	Qualitative	Bacterial EQA
Enterovirus (EV)							Pg 27
EVRNA20	QAV984104_1	1	8 to 12	Q3	Dry-ice	Qualitative	Viral EQA
	QAV984104_2	2	4 to 6	Q1, Q3			
Enterovirus Typing (EV)							Pg 28
EVTP20	QAV164185_1	1	5 to 10	Q1	Dry-ice	Typing	Viral EQA
Epstein-Barr virus (EBV)							Pg 28
EBVDNA20	QAV024121_1	1	8 to 12	Q3	Dry-ice	Qualitative & Quantitative	Viral EQA
	QAV024121_2	2	4 to 6	Q2, Q3			
Epstein-Barr virus (EBV) Whole Blood							Pg 29
EBVWB20	QAV134161_1	1	8 to 12	Q3	Dry-ice	Qualitative & Quantitative	Viral EQA
	QAV134161_2	2	4 to 6	Q2, Q3			
Extended Spectrum β-lactamase and Carbapenemase							Pg 8
ESBL20	QAB134162_1	1	8 to 12	Q3	Dry-ice	Typing	Bacterial EQA
Group B Streptococcus							Pg 9
GBS20	QAB174200_1	1	8 to 12	Q4	Dry-ice	Qualitative	Bacterial EQA

TARGET PATHOGEN							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.PYLORI20	QAB164190_1	1	5 to 10	Q3	Dry-ice	Qualitative	Bacterial EQA
Hepatitis A virus (HAV)							Pg 29
HAVRNA20	QAV124156_1 QAV124156_2	1 2	8 to 10 4 to 5	Q3 Q1, Q3	Dry-ice	Qualitative	Viral EQA
Hepatitis B virus (HBV)							Pg 30
HBVDNA20	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 virus (HBV) Drug Resistance							Pg 30
HBVDR20	QAV124160_1	1	4 to 7	Q3	Dry-ice	Drug Resistance / Sequencing	Viral EQA
Hepatitis B virus (HBV) Genotyping							Pg 31
HBVGT20	QAV064118_1	1	8 to 12	Q1	Dry-ice	Typing	Viral EQA
Hepatitis C virus (HCV)							Pg 31
HCVRNA20	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 virus (HCV) Drug Resistance							Pg 32
HCVDR20	QAV134167_1	1	4 to 7	Q3	Dry-ice	Drug Resistance / Sequencing	Viral EQA
Hepatitis C virus (HCV) Genotyping							Pg 32
HCVGT20	QAV034117_1	1	8 to 12	Q1	Dry-ice	Typing	Viral EQA
Hepatitis D virus (HDV)							Pg 33
HDV20	QAV144170_1	1	8 to 10	Q3	Dry-ice	Qualitative & Quantitative	Viral EQA
Hepatitis E virus (HEV)							Pg 33
HEVRNA20	QAV124157_1	1	8 to 10	Q3	Dry-ice	Qualitative	Viral EQA
Herpes simplex virus 1 & 2 (HSV)							Pg 34
HSVDNA20	QAV994105_1 QAV994105_2	1 2	8 to 12 4 to 6	Q3 Q1, Q3	Dry-ice	Qualitative	Viral EQA
Herpes simplex virus Drug Resistance							Pg 34
HSVDR20	QAV164184_1	1	4 to 7	Q1	Dry-ice	Sequence Analysis	Viral EQA
Human Immunodeficiency virus type 2 (HIV-2)							Pg 47
HIV2_20	QAV204212_1 QAV204212_2	1 2	8 4	Q1 Q1, Q3	Dry-ice	Qualitative & Quantitative	Pilot Study
Human herpes virus 6 (HHV6)							Pg 35
HHV6DNA20	QAV084119_1 QAV084119_2	1 2	8 to 12 4 to 6	Q3 Q2, Q3	Dry-ice	Qualitative & Quantitative	Viral EQA
Human Immunodeficiency virus type 1 (HIV-1) – DNA							Pg 35
HIVDNA20	QAV034114_1 QAV034114_2	1 2	8 4	Q3 Q1, Q3	Dry-ice	Qualitative	Viral EQA
Human Immunodeficiency virus type 1 (HIV-1) – Drug Resistance							Pg 36
HIVDR20	QAV024131_1	1	4 to 7	Q3	Dry-ice	Drug Resistance / Sequencing	Viral EQA
Human Immunodeficiency virus type 1 (HIV-1) – Drug Resistance (Integrase)							Pg 36
HIVDRint20	QAV114146_1	1	4 to 7	Q3	Dry-ice	Drug Resistance / Sequencing	Viral EQA

TARGET PATHOGEN							PAGE NUMBER
PROGRAMME CODE	CATALOGUE NUMBER	NO. OF CHALLENGES	PANEL MEMBERS PER CHALLENGE	DISTRIBUTION DATE(S)/ TESTING PERIOD	SHIPPING CONDITIONS	ANALYSIS TYPE	PROGRAMME TYPE
Human Immunodeficiency virus type 1 (HIV-1) – RNA							Pg 37
HIVRNA20	QAV994108_1	1	8	Q3	Dry-ice	Qualitative & Quantitative	Viral EQA
	QAV994108_2	2	4	Q1, Q3			
	QAV994108_4	4	8	Q1, Q2, Q3, Q4			
Human metapneumovirus (MPV)							Pg 37
MPV20	QAV054135_1	1	8 to 12	Q2	Dry-ice	Qualitative	Viral EQA
Human Papillomavirus (HPV) – PreservCyt							Pg 38
HPVPRES20	QAV094130_1	1	8 to 12	Q4	Ambient / Specialist	Qualitative	Viral EQA
	QAV094130_2	2	4 to 6	Q2, Q4			
Human Papillomavirus (Surepath)							Pg 47
HPVSURE20	QAV184204_1	1	8 to 12	Q4	Ambient	Qualitative	Pilot Study
Influenza A & B virus (FLU)							Pg 38
INFRNA20	QAV054134_1	1	8 to 12	Q4	Dry-ice	Qualitative	Viral EQA
	QAV054134_2	2	4 to 6	Q2, Q4			
Influenza Haemagglutinin Typing							Pg 39
INTP20	QAV064138_1	1	5 to 10	Q4	Dry-ice	Typing	Viral EQA
JC virus (JCV)							Pg 39
JCDNA20	QAV074106_1	1	8 to 12	Q3	Dry-ice	Qualitative & Quantitative	Viral EQA
	QAV074106_2	2	4 to 6	Q2, Q3			
Legionella pneumophila							Pg 10
LPDNA20	QAB044122_1	1	8 to 12	Q1	Dry-ice	Qualitative	Bacterial EQA
MALDI-TOF							Pg 17
MALDI20	QAB124155_1	1	8 to 12	Q3	Dry-ice	Typing	Multi-Pathogen / Syndromic EQA
Measles / Mumps							Pg 40
MM20	QAV144171_1	1	8 to 12	Q3	Dry-ice	Qualitative	Viral EQA
MERS coronavirus (MERS-CoV)							Pg 40
MERS20	QAV154181_1	1	6 to 10	Q2	Dry-ice	Qualitative	Viral EQA
Methicillin Resistant Staphylococcus aureus (MRSA)							Pg 10
MRSADNA20	QAB064124_1	1	8 to 12	Q2	Ambient	Qualitative	Bacterial EQA
Methicillin Resistant Staphylococcus aureus (MRSA) – Typing							Pg 11
MRSATP20	QAB074128_1	1	8 to 12	Q2	Ambient	Typing	Bacterial EQA
Mycobacterium tuberculosis (MTB)							Pg 11
MTBDNA20	QAB014129_1	1	8 to 12	Q4	Ambient	Qualitative	Bacterial EQA
	QAB014129_2	2	4 to 6	Q2, Q4			
Mycobacterium tuberculosis Drug Resistance							Pg 48
MTBDR20	QAB194209_1	1	6 to 10	Q4	Ambient	Typing	Pilot Study
Mycoplasma genitalium							Pg 48
MG20	QAB184205_1	1	6 to 10	Q3	Dry-ice	Qualitative	Pilot Study
Mycoplasma pneumoniae							Pg 12
MP20	QAB174192_1	1	5 to 10	Q2	Dry-ice	Qualitative	Bacterial EQA

TARGET PATHOGEN							PAGE NUMBER
PROGRAMME CODE	CATALOGUE NUMBER	NO. OF CHALLENGES	PANEL MEMBERS PER CHALLENGE	DISTRIBUTION DATE(S)/ TESTING PERIOD	SHIPPING CONDITIONS	ANALYSIS TYPE	PROGRAMME TYPE
Mycoplasma spp. (cell contamination)							Pg 12
MYCO20	QAB144168_1	1	8 to 12	Q4	Dry-ice	Qualitative & Quantitative	Bacterial EQA
Neisseria gonorrhoeae							Pg 13
NgDNA20	QAB034126_1	1	8 to 12	Q3	Dry-ice	Qualitative	Bacterial EQA
	QAB034126_2	2	4 to 6	Q1, Q3			
Norovirus (NV)							Pg 41
NVRNA20	QAV084139_1	1	8 to 12	Q4	Dry-ice	Qualitative	Viral EQA
	QAV084139_2	2	4 to 6	Q2, Q4			
Parainfluenza virus (PIV)							Pg 41
PINFRNA20	QAV064136_1	1	8 to 12	Q2	Dry-ice	Qualitative	Viral EQA
Parasitic Gastroenteritis							Pg 18
GastroP20	QAP124154_1	1	8 to 12	Q4	Dry-ice	Qualitative	Multi-Pathogen / Syndromic EQA
	QAP124154_2	2	4 to 6	Q2 & Q4			
Parechovirus (HPeV)							Pg 42
PeVRNA20	QAV114145_1	1	8 to 12	Q3	Dry-ice	Qualitative	Viral EQA
	QAV114145_2	2	4 to 6	Q1, Q3			
Pneumocystis jirovecii pneumonia (PCP)							Pg 16
PCPDNA20	QAF114144_1	1	8 to 12	Q3	Dry-ice	Qualitative & Quantitative	Fungal EQA
Respiratory I							Pg 18
RESPI20	QAV164188_1	1	8 to 12	Q3	Dry-ice	Qualitative	Multi-Pathogen / Syndromic EQA
	QAV164188_2	2	4 to 6	Q1, Q3			
Respiratory II							Pg 19
RESPII20	QAV164189_1	1	8 to 12	Q3	Dry-ice	Qualitative	Multi-Pathogen / Syndromic EQA
	QAV164189_2	2	4 to 6	Q1, Q3			
Respiratory III							Pg 19
RESPIII20	QAM174193_1	1	8 to 12	Q3	Dry-ice	Qualitative	Multi-Pathogen / Syndromic EQA
	QAM174193_2	2	4 to 6	Q1, Q3			
Respiratory syncytial virus (RSV)							Pg 42
RSV20	QAV054142_1	1	8 to 12	Q4	Dry-ice	Qualitative	Viral EQA
	QAV054142_2	2	4 to 6	Q2, Q4			
Rhinovirus (RV)							Pg 43
RVRNA20	QAV064143_1	1	8 to 12	Q2	Dry-ice	Qualitative	Viral EQA
Sepsis							Pg 49
SEPSIS20	QAB164178_1	1	8 to 12	Q4	Dry-ice	Qualitative	Multi-Pathogen / Syndromic EQA
Sexually Transmitted Infections I							Pg 20
STI_I20	QAB154177_1	1	8 to 12	Q3	Dry-ice	Qualitative	Multi-Pathogen / Syndromic EQA
	QAB154177_2	2	4 to 6	Q2, Q3			
Sexually Transmitted Infections II							Pg 20
STI_II20	QAM174201_1	1	8 to 12	Q3	Dry-ice	Qualitative	Multi-Pathogen / Syndromic EQA
	QAM174201_2	2	4 to 6	Q2, Q3			

TARGET PATHOGEN							PAGE NUMBER
PROGRAMME CODE	CATALOGUE NUMBER	NO. OF CHALLENGES	PANEL MEMBERS PER CHALLENGE	DISTRIBUTION DATE(S)/ TESTING PERIOD	SHIPPING CONDITIONS	ANALYSIS TYPE	PROGRAMME TYPE
<i>Staphylococcus aureus spa</i>							Pg 13
SASPA20	QAB134164_1	1	6 to 12	Q2	Ambient	Typing	Bacterial EQA
Syphilis							Pg 14
SYPH20	QAB154180_1	1	5 to 10	Q4	Dry-ice	Qualitative	Bacterial EQA
Torque teno virus (TTV)							Pg 49
TTV20	QAV184203_1	1	6 to 10	Q4	Dry-ice	Qualitative	Pilot Study
<i>Toxoplasma gondii</i>							Pg 22
TGDNA20	QAP044123_1	1	8 to 12	Q4	Ambient	Qualitative	Parasitic EQA
	QAP044123_2	2	4 to 6	Q2, Q4			
Transplantation (viral)							Pg 50
TRANS20	QAM174198_1	1	8 to 12	Q3	Dry-ice	Qualitative & Quantitative	Pilot Study
	QAM174198_2	2	4 to 6	Q2, Q4			
<i>Trichomonas vaginalis</i>							Pg 50
TV20	QAP184202_1	1	6 to 10	Q3	Dry-ice	Qualitative	Pilot Study
Vancomycin Resistant Enterococci (VRE)							Pg 14
VRE20	QAB134163_1	1	8 to 12	Q3	Dry-ice	Qualitative	Bacterial EQA
Varicella-Zoster virus (VZV)							Pg 43
VZVDNA20	QAV034103_1	1	8 to 12	Q3	Dry-ice	Qualitative	Viral EQA
	QAV034103_2	2	4 to 6	Q1, Q3			
Viral Gastroenteritis							Pg 21
GastroV20	QAV124152_1	1	8 to 12	Q4	Dry-ice	Qualitative	Multi-Pathogen / Syndromic EQA
	QAV124152_2	2	4 to 6	Q2, Q4			
Viral Metagenomics NGS							Pg 51
NGSmta_20	QAV204213_1	1	4 to 7	Q4	Dry-ice	Qualitative	Viral EQA
West Nile virus (WNV)							Pg 44
WNVRNA20	QAV104141_1	1	8 to 12	Q2	Ambient	Qualitative	Viral EQA
Yellow fever virus							Pg 51
YFV20	QAM194207_1	1	8 to 12	Q2	Ambient	Qualitative	Pilot Study
Zika virus							Pg 44
ZIKA20	QAV164186_1	1	8 to 12	Q2	Ambient	Qualitative	Viral EQA



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