Individual Report	_	D 2021 Sy ramme	philis EQA	Z		

Catalogue Code:	Ref Code:	Challenge:	Analysis Type:	Dataset:	Report UID:	Laboratory
QAB154180	SYPH21	S	Qualitative	504142	2677/504142/3982	CZ023

Intended Results / Panel Composition

Sample Code	Sample Content	Matrix	Sample Relationships ^[1]	Detection Frequency ^[2]	Sample Status ^[3]	Percentag Correct (A	
						(%)	(n)
SYPH21S-01	T. pallidum	Transport Medium		Frequently Detected	CORE	100.0	65
SYPH21S-02	T. pallidum	Transport Medium		Frequently Detected	CORE	98.5	65
SYPH21S-03	T. pallidum	Transport Medium		Detected	EDUCATIONAL	76.9	65
SYPH21S-04	T. pallidum	Transport Medium		Frequently Detected	CORE	100.0	65
SYPH21S-05	T. pallidum	Transport Medium		Detected	CORE	92.3	65
SYPH21S-06	T. pallidum	Transport Medium		Frequently Detected	CORE	98.5	65
SYPH21S-07	T. pallidum	Transport Medium		Frequently Detected	CORE	100.0	65
SYPH21S-08	Negative	Transport Medium		Negative	CORE	98.5	65

[1] Sample Relationships: Indicates the relationships of the samples within this challenge. The highest titre member of dilution series DS1 is indicated by DS1_1 and further members of the series as DS1_2, DS1_3 etc. in order of reducing titre. Additional dilution series are indicated by DS2 (e.g DS2_1, DS2_2 etc.), DS3 (e.g. DS3_1, DS3_2 etc.). If one duplicate pair is present this is indicated by 'D1'. Further duplicate pairs are indicated by 'D2', 'D3' etc.
[2] Detection Frequency: To aid qualitative analysis each panel member is assigned a frequency of detection. This is based on the peer group consensus of all qualitative results returned from participants within the EQA challenge / distribution.

[3] **Sample Status:** EQA samples are defined as "CORE" or "EDUCATIONAL". Core proficiency samples are reviewed by the QCMD Scientific Expert(s). This is on the basis of scientific information, clinical relevance, current literature and, where appropriate, professional clinical guidelines. Participating laboratories are expected to report core proficiency samples correctly within the EQA challenge / distribution.

[4] **Percentage Correct (All):** Percentage of datasets (%) reporting the correct qualitative result and the total number of datasets (n) reported for each panel member.

For further details please refer to the current participant manual.

Your Summary Results

EQA Assessment Group ^[1]

Core Panel Detection (Qualitative) Score [2]

Commercial

0

QCMD, Technology Terrace, Todd Campus, West of Scotland Science Park, Glasgow, G20 0XA Tel: +44 (0) 141 945 6474, Fax: +44 (0) 141 945 5795 Web: www.qcmd.org

Individual	QCMD 2021 Syphilis EQA	QCAD
Report	Programme	Quality Control for Malecular Diagnostics

Catalogue Code:	Ref Code:	Challenge:	Analysis Type:	Dataset:	Report UID:	Laboratory
QAB154180	SYPH21	S	Qualitative	504142	2677/504142/3982	CZ023

Core Panel Members Results

Sample Code	Qualitative Results			Your Quantitative Data (for information only) [3]		
Coue	Percentage Correct (AII) ^[4]	Your Result ^[5]	Detection Score	Reported Value	Unitage	Cycle Threshold
SYPH21S-01	100.0	Positive	٥		N/A	31.19
SYPH21S-02	98.5	Positive	٥		N/A	32.56
SYPH21S-04	100.0	Positive	٥		N/A	31.75
SYPH21S-05	92.3	Positive	٥		N/A	34.67
SYPH21S-06	98.5	Positive	٥		N/A	32.03
SYPH21S-07	100.0	Positive	٥		N/A	32.37
SYPH21S-08	98.5	Negative	٥		N/A	-

[1] **EQA Assessment Group:** To aid data analysis, participant results are grouped according to the molecular amplification/detection method specified within their molecular workflow for this challenge / distribution. For further details refer to the *Additional Information: Individual Panel Member Analysis* section of this report.

[2] Core Panel Detection (Qualitative) Score: An overall core panel detection score provided per challenge / distribution.

[3] **Quantitative Data (for information only):** This is the quantitative value, unitage and cycle threshold you provided when you submitted your results. For qualitative programmes this information is not used as part of your formal EQA assessment.

[4] Percentage Correct (AII): Percentage of datasets (%) reporting the correct qualitative results for each panel member.

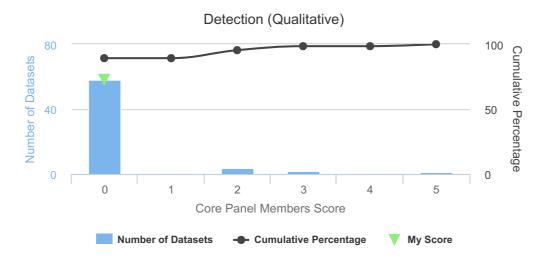
[5] Your Result: The qualitative result you reported for each sample within this EQA challenge / distribution.

[6] **Detection Score:** Your detection (qualitative) scores are based on the assigned detection frequency of each panel members, where 0 (zero) is "highly satisfactory" and 3 (three) is "highly unsatisfactory". Scores are provided for individual panel members.

For further details please refer to the current participant manual.

Individual Report	_	/ID 2021 S gramme	yphilis EQA			
Catalogue Code:	Ref Code:	Challenge:	Analysis Type:	Dataset:	Report UID:	Laboratory
QAB154180	SYPH21	S	Qualitative	504142	2677/504142/3982	CZ023

Core Panel Member Score Breakdown



Core Panel Member Score Breakdown - Detection: This figure gives you a breakdown of the qualitative detection scores for all qualitative datasets returned within this EQA challenge / distribution independent of the EQA assessment group. Panel detection scores are generated from only those panel members that are defined as "CORE".

For further details please refer to the current participant manual.

My Workflow Details

The details of the workflow(s) used to submit your results for this challenge.

Name	GeneProof Treponema pallidum PCR Kit-croBEE (v2)					
Description						
Targets	B Treponema pallidum					
Assays	 <i>Extraction</i> - GeneProof - croBEE NA16 Nucleic Acid Extraction System Commercial Kit Manufacturer: GeneProof Kit Type: croBEE 201A Nucleic Acid Extraction Kit 					
	 Amplification - GeneProof - croBEE Real-Time PCR System Commercial Kit Manufacturer: GeneProof Kit Type: GeneProof Treponema pallidum PCR Kit Kit Version: ISEX 					

Individual	QCMD 2021 Syphilis EQA	QCAD
Report	Programme	Quality Control for Molecular Diagnostics

Catalogue Code:	Ref Code:	Challenge:	Analysis Type:	Dataset:	Report UID:	Laboratory
QAB154180	SYPH21	S	Qualitative	504142	2677/504142/3982	CZ023

Educational Panel Members Results

Sample Code	Qualitative Results				e Data (for ii	nformation only) ^[1]
	Percentage Correct (AII) ^[2]	Your Result ^[3]	Detection Score [4]	Reported Value	Unitage	Cycle Threshold
SYPH21S-03	76.9	Positive	0		N/A	35.27

[1] **Quantitative Data (for information only):** This is the quantitative value, unitage and cycle threshold you provided when you submitted your results. For qualitative programmes this information is not used as part of your formal EQA assessment.

[2] Percentage Correct (All): Percentage of datasets (%) reporting the correct qualitative results for each panel member.

[3] Your Result: The qualitative result you reported for each sample within this EQA challenge / distribution.

[4] Detection Score: Your detection (qualitative) scores are based on the assigned detection frequency of each panel members, where 0 (zero) is "highly satisfactory" and 3 (three) is "highly unsatisfactory". Scores are provided for individual panel members.

For further details please refer to the current participant manual.

Further Programme Details

Number of Participants	71
Number of Countries	21
Number of Respondents	60
Number of Datasets Submitted	65
Qualitative Results Returned	65 (100.0%)

EQA Programme Aims

To assess laboratories ability to detect *Treponema pallidum* using their routine molecular diagnostic platform and procedures.

Feedback and Enquiries

Participants are encouraged to read the QCMD Participants' Manual, which can be downloaded from the QCMD website.

Any enquiries should be submitted through the 'Contact Us' form that you can find in the 'Help' section of your QCMD (ITEMS) Participant Profile Area.

	Individual Report	_	MD 2021 Sy gramme	yphilis EQA			
Catalogue Code: Ref C		Ref Code:	Challenge:	Analysis Type:	Dataset:	Report UID:	Laboratory

504142

2677/504142/3982

CZ023

Qualitative

Panel member analysis is separated into CORE samples followed by EDUCATIONAL samples.

S

Additional Core Samples Information

The following section has been categorised as shown below:

SYPH21

Core ► Qualitative

QAB154180

Individual Panel Member Analysis (Qualitative)

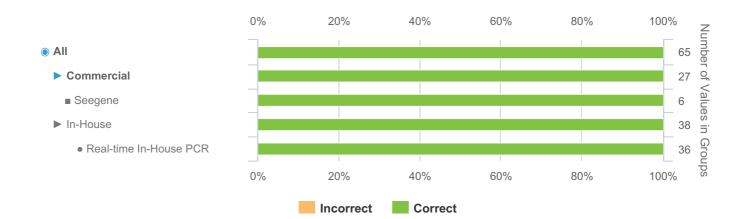
Qualitative analysis for each panel member is provided in relation to your EQA assessment group. EQA assessment groups are established using the molecular workflow information reported by all participants within this EQA challenge / distribution. The principal level of assessment is at the individual method level which is defined based on your reported "amplification/detection method" and other laboratories using the same or similar amplification/detection methods.

To allow meaningful assessment at the individual method level the EQA assessment group must consist of 5 or more datasets. If there are not sufficient datasets at the individual method level then your results will be included within a higher EQA assessment group based on whether it is a commercial or in house technology/method. The highest level assessment grouping is "All" participant reported qualitative results.

A breakdown of qualitative results reported by participants on each of the panel members within this EQA challenge / distribution is provided below. You can compare your results to those within your EQA assessment group and those obtained within other EQA assessment groups or to the overall consensus for each sample within this EQA challenge / distribution.

SYPH21S-01 - Qualitative Results Breakdown

Sample Code	Sample Content	Matrix	Sample Relationships	Detection Frequency	Sample Status	Percentage Correct (All)		
						(%)	(n)	
SYPH21S-01	T. pallidum	Transport Medium		Frequently Detected	CORE	100.0	65	



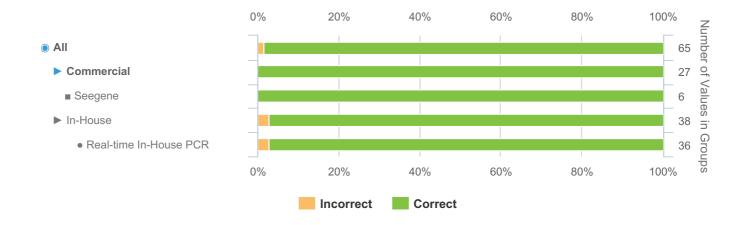
Individual Report		ID 2021 S gramme	yphilis EQA		Report UID: 2677/50/142/3082		
Catalogue Code: QAB154180	Ref Code: SYPH21	Challenge: S	Analysis Type: Qualitative	Dataset: 504142	Report UID: 2677/504142/3982	Laboratory CZ023	

Groups below n=5: AmpliSens (n=2), AmpliSens - AmpliSens Real Time PCR (n=2), AusDiagnostics (n=1), AusDiagnostics - AusDiagnostics High-Plex (n=1), Certest (n=1), Certest - Certest Real Time PCR (n=1), GeneProof (n=2), GeneProof - GeneProof Real Time PCR kit (n=2), PathoFinder (n=1), PathoFinder - PathoFinder Real Time PCR (n=1), Progenie Molecular (n=3), Progenie Molecular - Progenie Molecular RealCycler (n=3), Randox (n=1), Randox - Randox Multiplex Array (n=1), Sacace (n=3), Sacace - Sacace Real TM (n=3), TIB MOLBIOL (n=4), TIB MOLBIOL - TIB-MolBiol LightMix (n=4), fast-track DIAGNOSTICS (n=3), fast-track DIAGNOSTICS - FTD real time PCR (n=3), In-House - Conventional In-House PCR (n=2)

Groups Rolled Up: Seegene - Seegene Allplex (n=6)

SYPH21S-02 - Qualitative Results Breakdown

Sample Code	Sample Content	Matrix	Sample Relationships	Detection Frequency	Sample Status	Percentage Correct (All)	
						(%)	(n)
SYPH21S-02	T. pallidum	Transport Medium		Frequently Detected	CORE	98.5	65



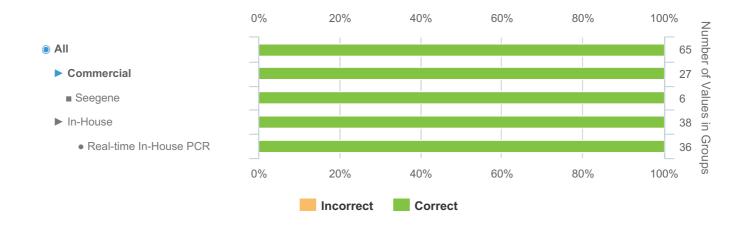
Groups below n=5: AmpliSens (n=2), AmpliSens - AmpliSens Real Time PCR (n=2), AusDiagnostics (n=1), AusDiagnostics - AusDiagnostics High-Plex (n=1), Certest (n=1), Certest - Certest Real Time PCR (n=1), GeneProof (n=2), GeneProof - GeneProof Real Time PCR kit (n=2), PathoFinder (n=1), PathoFinder - PathoFinder Real Time PCR (n=1), Progenie Molecular (n=3), Progenie Molecular - Progenie Molecular RealCycler (n=3), Randox (n=1), Randox - Randox Multiplex Array (n=1), Sacace (n=3), Sacace - Sacace Real TM (n=3), TIB MOLBIOL (n=4), TIB MOLBIOL - TIB-MolBiol LightMix (n=4), fast-track DIAGNOSTICS (n=3), fast-track DIAGNOSTICS - FTD real time PCR (n=3), In-House - Conventional In-House PCR (n=2)

ndividual	QCMD 2021 Syphilis EQA	QCMD
Report	Programme	Quality Cantrol for Malecular Diagnostics

Catalogue Code:	Ref Code:	Challenge:	Analysis Type:	Dataset:	Report UID:	Laboratory
QAB154180	SYPH21	S	Qualitative	504142	2677/504142/3982	CZ023

SYPH21S-04 - Qualitative Results Breakdown

Sample Code	Sample Content	Matrix	Sample Relationships	Detection Frequency	Sample Status	Percentage (All)	Correct
						(%)	(n)
SYPH21S-04	T. pallidum	Transport Medium		Frequently Detected	CORE	100.0	65



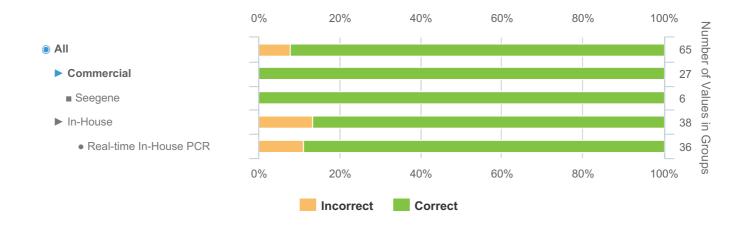
Groups below n=5: AmpliSens (n=2), AmpliSens - AmpliSens Real Time PCR (n=2), AusDiagnostics (n=1), AusDiagnostics - AusDiagnostics High-Plex (n=1), Certest (n=1), Certest - Certest Real Time PCR (n=1), GeneProof (n=2), GeneProof - GeneProof Real Time PCR kit (n=2), PathoFinder (n=1), PathoFinder - PathoFinder Real Time PCR (n=1), Progenie Molecular (n=3), Progenie Molecular - Progenie Molecular RealCycler (n=3), Randox (n=1), Randox - Randox Multiplex Array (n=1), Sacace (n=3), Sacace - Sacace Real TM (n=3), TIB MOLBIOL (n=4), TIB MOLBIOL - TIB-MolBiol LightMix (n=4), fast-track DIAGNOSTICS (n=3), fast-track DIAGNOSTICS - FTD real time PCR (n=3), In-House - Conventional In-House PCR (n=2)

Individual	QCMD 2021 Syphilis EQA	2	QCMD
Report	Programme		Quality Control for Molecular Diagnostics

Catalogue Code:	Ref Code:	Challenge:	Analysis Type:	Dataset:	Report UID:	Laboratory
QAB154180	SYPH21	S	Qualitative	504142	2677/504142/3982	CZ023

SYPH21S-05 - Qualitative Results Breakdown

Sample Code	Sample Content	Matrix	Sample Relationships	Detection Frequency	Sample Status	Percentage (All)	Correct
						(%)	(n)
SYPH21S-05	T. pallidum	Transport Medium		Detected	CORE	92.3	65



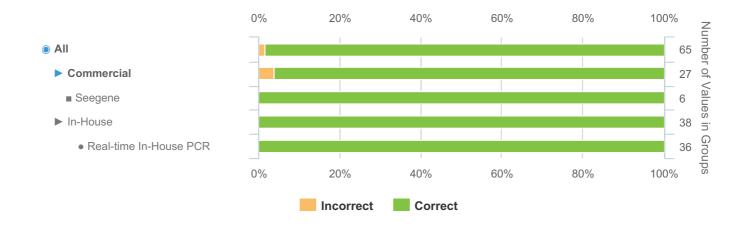
Groups below n=5: AmpliSens (n=2), AmpliSens - AmpliSens Real Time PCR (n=2), AusDiagnostics (n=1), AusDiagnostics - AusDiagnostics High-Plex (n=1), Certest (n=1), Certest - Certest Real Time PCR (n=1), GeneProof (n=2), GeneProof - GeneProof Real Time PCR kit (n=2), PathoFinder (n=1), PathoFinder - PathoFinder Real Time PCR (n=1), Progenie Molecular (n=3), Progenie Molecular - Progenie Molecular RealCycler (n=3), Randox (n=1), Randox - Randox Multiplex Array (n=1), Sacace (n=3), Sacace - Sacace Real TM (n=3), TIB MOLBIOL (n=4), TIB MOLBIOL - TIB-MolBiol LightMix (n=4), fast-track DIAGNOSTICS (n=3), fast-track DIAGNOSTICS - FTD real time PCR (n=3), In-House - Conventional In-House PCR (n=2)

Individual Report	QCMD 2021 Syphilis EQA Programme	Z	QCMD Quality Control for Molecular Diagnostics

Catalogue Code:	Ref Code:	Challenge:	Analysis Type:	Dataset:	Report UID:	Laboratory
QAB154180	SYPH21	S	Qualitative	504142	2677/504142/3982	CZ023

SYPH21S-06 - Qualitative Results Breakdown

Sample Code	Sample Content	Matrix	Sample Relationships	Detection Frequency	Sample Status	Percentage (All)	Correct
						(%)	(n)
SYPH21S-06	T. pallidum	Transport Medium		Frequently Detected	CORE	98.5	65



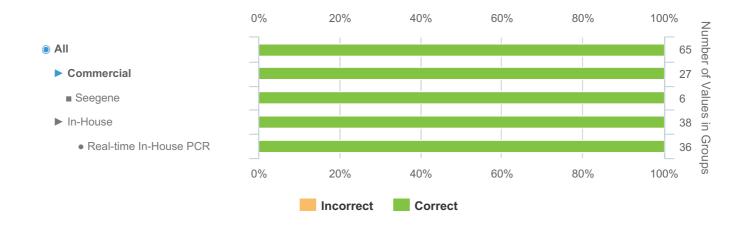
Groups below n=5: AmpliSens (n=2), AmpliSens - AmpliSens Real Time PCR (n=2), AusDiagnostics (n=1), AusDiagnostics - AusDiagnostics High-Plex (n=1), Certest (n=1), Certest - Certest Real Time PCR (n=1), GeneProof (n=2), GeneProof - GeneProof Real Time PCR kit (n=2), PathoFinder (n=1), PathoFinder - PathoFinder Real Time PCR (n=1), Progenie Molecular (n=3), Progenie Molecular - Progenie Molecular RealCycler (n=3), Randox (n=1), Randox - Randox Multiplex Array (n=1), Sacace (n=3), Sacace - Sacace Real TM (n=3), TIB MOLBIOL (n=4), TIB MOLBIOL - TIB-MolBiol LightMix (n=4), fast-track DIAGNOSTICS (n=3), fast-track DIAGNOSTICS - FTD real time PCR (n=3), In-House - Conventional In-House PCR (n=2)

Individual	QCMD 2021 Syphilis EQA	QCAD
Report	Programme	Quality Control for Malecular Diagnostics

Catalogue Code:	Ref Code:	Challenge:	Analysis Type:	Dataset:	Report UID:	Laboratory
QAB154180	SYPH21	S	Qualitative	504142	2677/504142/3982	CZ023

SYPH21S-07 - Qualitative Results Breakdown

Sample Code	Sample Content	Matrix	Sample Relationships	Detection Frequency	Sample Status	Percentage (All)	Correct
						(%)	(n)
SYPH21S-07	T. pallidum	Transport Medium		Frequently Detected	CORE	100.0	65



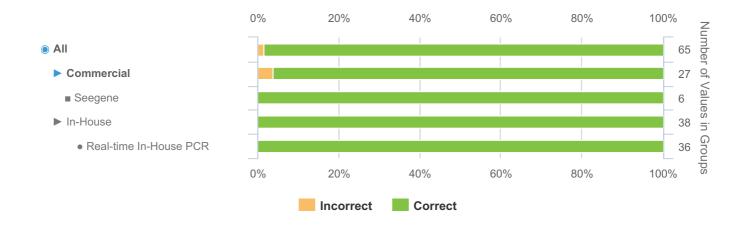
Groups below n=5: AmpliSens (n=2), AmpliSens - AmpliSens Real Time PCR (n=2), AusDiagnostics (n=1), AusDiagnostics - AusDiagnostics High-Plex (n=1), Certest (n=1), Certest - Certest Real Time PCR (n=1), GeneProof (n=2), GeneProof - GeneProof Real Time PCR kit (n=2), PathoFinder (n=1), PathoFinder - PathoFinder Real Time PCR (n=1), Progenie Molecular (n=3), Progenie Molecular - Progenie Molecular RealCycler (n=3), Randox (n=1), Randox - Randox Multiplex Array (n=1), Sacace (n=3), Sacace - Sacace Real TM (n=3), TIB MOLBIOL (n=4), TIB MOLBIOL - TIB-MolBiol LightMix (n=4), fast-track DIAGNOSTICS (n=3), fast-track DIAGNOSTICS - FTD real time PCR (n=3), In-House - Conventional In-House PCR (n=2)

Individual	QCMD 2021 Syphilis EQA	QCMD
Report	Programme	Quality Control for Molecular Diagnostics

Catalogue Code:	Ref Code:	Challenge:	Analysis Type:	Dataset:	Report UID:	Laboratory
QAB154180	SYPH21	S	Qualitative	504142	2677/504142/3982	CZ023

SYPH21S-08 - Qualitative Results Breakdown

Sample Code	Sample Content	Matrix	Sample Relationships		Sample Status	Percentage (All)	e Correct
						(%)	(n)
SYPH21S-08	8 Negative	Transport Medium		Negative	CORE	98.5	65



Groups below n=5: AmpliSens (n=2), AmpliSens - AmpliSens Real Time PCR (n=2), AusDiagnostics (n=1), AusDiagnostics - AusDiagnostics High-Plex (n=1), Certest (n=1), Certest - Certest Real Time PCR (n=1), GeneProof (n=2), GeneProof - GeneProof Real Time PCR kit (n=2), PathoFinder (n=1), PathoFinder - PathoFinder Real Time PCR (n=1), Progenie Molecular (n=3), Progenie Molecular - Progenie Molecular RealCycler (n=3), Randox (n=1), Randox - Randox Multiplex Array (n=1), Sacace (n=3), Sacace - Sacace Real TM (n=3), TIB MOLBIOL (n=4), TIB MOLBIOL - TIB-MolBiol LightMix (n=4), fast-track DIAGNOSTICS (n=3), fast-track DIAGNOSTICS - FTD real time PCR (n=3), In-House - Conventional In-House PCR (n=2)

Individual Report	QCMD 2021 Syphilis EQA Programme	

Catalogue Code:	Ref Code:	Challenge:	Analysis Type:	Dataset:	Report UID:	Laboratory
QAB154180	SYPH21	S	Qualitative	504142	2677/504142/3982	CZ023

Additional Educational Samples Information

The following section has been categorised as shown below:

Educational
Qualitative

Individual Panel Member Analysis (Qualitative)

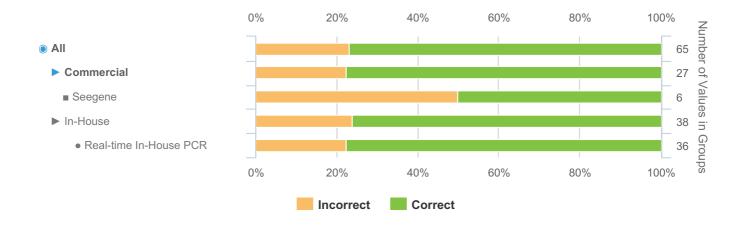
Qualitative analysis for each panel member is provided in relation to your EQA assessment group. EQA assessment groups are established using the molecular workflow information reported by all participants within this EQA challenge / distribution. The principal level of assessment is at the individual method level which is defined based on your reported "amplification/detection method" and other laboratories using the same or similar amplification/detection methods.

To allow meaningful assessment at the individual method level the EQA assessment group must consist of 5 or more datasets. If there are not sufficient datasets at the individual method level then your results will be included within a higher EQA assessment group based on whether it is a commercial or in house technology/method. The highest level assessment grouping is "All" participant reported qualitative results.

A breakdown of qualitative results reported by participants on each of the panel members within this EQA challenge / distribution is provided below. You can compare your results to those within your EQA assessment group and those obtained within other EQA assessment groups or to the overall consensus for each sample within this EQA challenge / distribution.

SYPH21S-03 - Qualitative Results Breakdown

Sam Code	•	Sample Content	Matrix	Sample Relationships	Detection Frequency	Sample Status	Percentag Correct (A	·
							(%)	(n)
SYPI	H21S-03	T. pallidum	Transport Medium		Detected	EDUCATIONAL	76.9	65



Individual	QCMD 2021 Syphilis EQA	QCMD
Report	Programme	Quality Control for Molecular Diagnostics

Catalogue Code: R	Ref Code:	Challenge:	Analysis Type:	Dataset:	Report UID:	Laboratory
QAB154180 S	SYPH21	S	Qualitative	504142	2677/504142/3982	CZ023

Groups below n=5: AmpliSens (n=2), AmpliSens - AmpliSens Real Time PCR (n=2), AusDiagnostics (n=1), AusDiagnostics - AusDiagnostics High-Plex (n=1), Certest (n=1), Certest - Certest Real Time PCR (n=1), GeneProof (n=2), GeneProof - GeneProof Real Time PCR kit (n=2), PathoFinder (n=1), PathoFinder - PathoFinder Real Time PCR (n=1), Progenie Molecular (n=3), Progenie Molecular - Progenie Molecular RealCycler (n=3), Randox (n=1), Randox - Randox Multiplex Array (n=1), Sacace (n=3), Sacace - Sacace Real TM (n=3), TIB MOLBIOL (n=4), TIB MOLBIOL - TIB-MolBiol LightMix (n=4), fast-track DIAGNOSTICS (n=3), fast-track DIAGNOSTICS - FTD real time PCR (n=3), In-House - Conventional In-House PCR (n=2)

Groups Rolled Up: Seegene - Seegene Allplex (n=6)

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