

Individual Report

QCMD 2020 Mycoplasma genitalium EQA Pilot Study



Catalogue Code: QAB184205	Ref Code: MG20	Challenge: S	Analysis Type: Qualitative	Dataset: 395362	Report UID: 2677/395362/3007	Laboratory CZ023
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Intended Results / Panel Composition

Sample Code	Sample Content	Matrix	Sample Relationships ^[1]	Detection Frequency ^[2]	Sample Status ^[3]	Percentage Correct (All) ^[4]	
						(%)	(n)
MG101S-01	M. genitalium G37	Urine	DS1_2	Detected	CORE	82.8	134
MG101S-02	M. genitalium G37	PBS	DS2_1	Frequently Detected	CORE	97.8	134
MG101S-03	Mycoplasma Negative	PBS		Negative	CORE	100.0	134
MG101S-04	M. genitalium M6303 (Macrolide Resistant)	PBS	DS3_1	Frequently Detected	CORE	99.3	134
MG101S-05	M. genitalium G37	PBS	DS2_3	Infrequently Detected	EDUCATIONAL	47.0	134
MG101S-06	M. genitalium M6593 (Macrolide Resistant)	PBS		Frequently Detected	CORE	95.5	134
MG101S-07	M. genitalium G37	PBS	DS2_2	Detected	CORE	86.6	134
MG101S-08	M. genitalium M6303 (Macrolide Resistant)	PBS	DS3_2	Detected	EDUCATIONAL	79.1	134
MG101S-09	M. pneumoniae	PBS		Negative	CORE	97.8	134
MG101S-10	M. genitalium G37	Urine	DS1_1	Frequently Detected	CORE	97.0	134

[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.

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Your Summary Results

EQA Assessment Group ^[1] Commercial

Core Panel Detection (Qualitative) Score ^[2] 0

Core Panel Members Results

Sample Code	Qualitative Results			Your Quantitative Data (for information only) ^[3]		
	Percentage Correct (All) ^[4]	Your Result ^[5]	Detection Score ^[6]	Reported Value	Unitage	Cycle Threshold
MG101S-01	82.8	Positive	0		N/A	36.52
MG101S-02	97.8	Positive	0		N/A	31.45
MG101S-03	100.0	Negative	0		N/A	-
MG101S-04	99.3	Positive	0		N/A	32.45
MG101S-06	95.5	Positive	0		N/A	34.53
MG101S-07	86.6	Positive	0		N/A	35.36
MG101S-09	97.8	Negative	0		N/A	-
MG101S-10	97.0	Positive	0		N/A	32.68

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

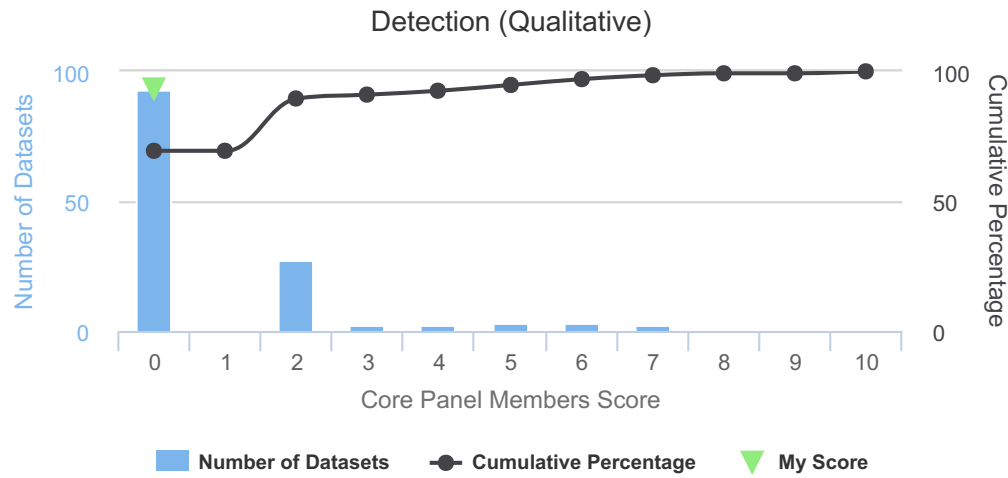
[5] Your Result: The qualitative result you reported for each sample within this EQA challenge / distribution.

[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.

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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	Mycoplasma genitalium/hominis PCR Kit-croBEE (v2)
Description	
Targets	<ul style="list-style-type: none"> B Mycoplasma genitalium B Mycoplasma hominis
Assays	<ul style="list-style-type: none"> A Extraction - GeneProof - croBEE NA16 Nucleic Acid Extraction System <ul style="list-style-type: none"> ● Commercial <ul style="list-style-type: none"> ○ Kit Manufacturer: <i>GeneProof</i> ○ Kit Type: <i>croBEE 201A Nucleic Acid Extraction Kit</i> R Amplification - GeneProof - croBEE Real-Time PCR System <ul style="list-style-type: none"> ● Multiplex ● Commercial <ul style="list-style-type: none"> ○ Kit Manufacturer: <i>GeneProof</i> ○ Kit Type: <i>Mycoplasma genitalium/hominis PCR Kit</i> ○ Kit Version: <i>ISEX</i>

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Educational Panel Members Results

Sample Code	Qualitative Results			Your Quantitative Data (for information only) ^[1]		
	Percentage Correct (All) ^[2]	Your Result ^[3]	Detection Score ^[4]	Reported Value	Unitage	Cycle Threshold
MG101S-05	47.0	Negative	1		N/A	-
MG101S-08	79.1	Positive	0		N/A	36.16

[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	144
Number of Countries	30
Number of Respondents	117
Number of Datasets Submitted	134
Qualitative Results Returned	134 (100.0%)


EQA Programme Aims

To assess laboratories ability to detect *Mycoplasma genitalium* using routine molecular diagnostic platforms 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.

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

Additional Core Samples Information

The following section has been categorised as shown below:


Core ► 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.

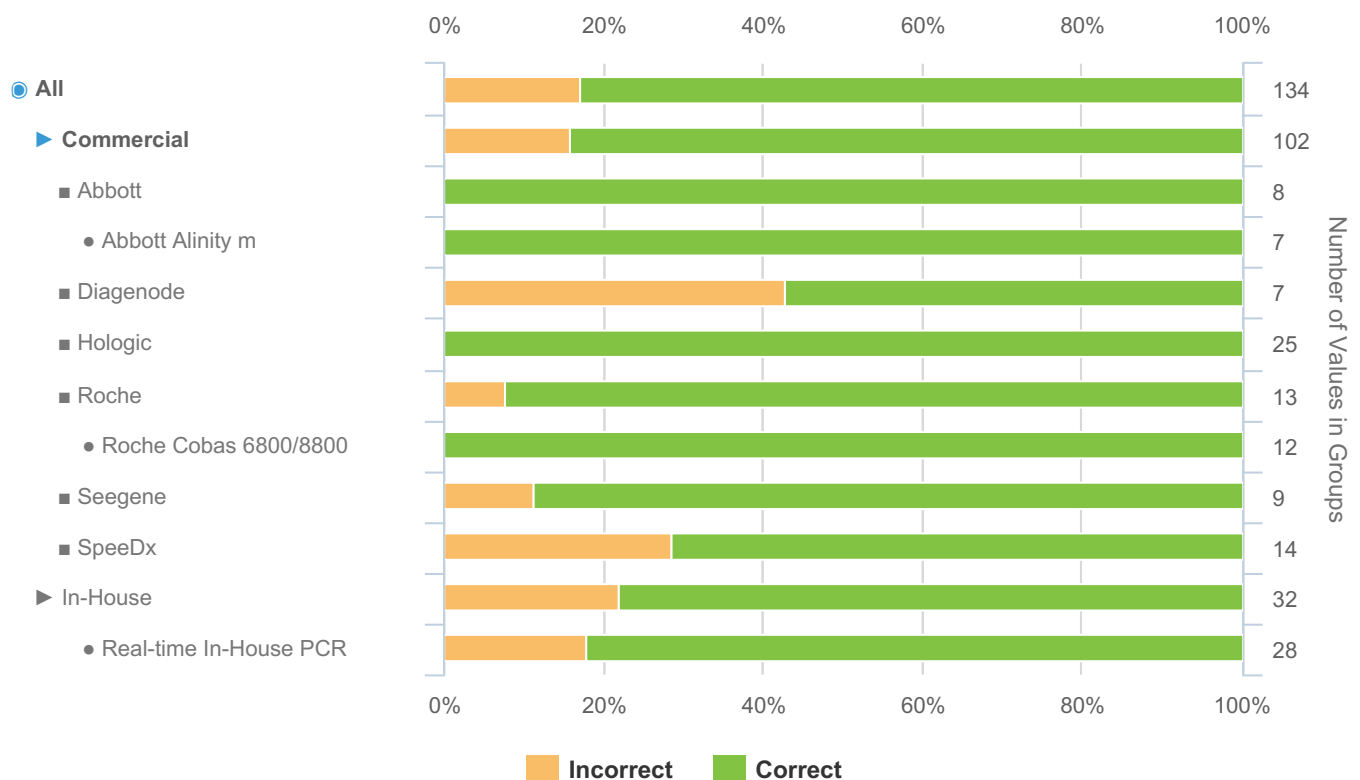
To allow meaningful assessment at the individual method level the EQA assessment group must consist of 5 or more datasets. If there are not sufficient datasets at the individual method level then your results will be included within a higher EQA assessment group based on whether it is a commercial or 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.

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MG101S-01 - Qualitative Results Breakdown

Sample Code	Sample Content	Matrix	Sample Relationships	Detection Frequency	Sample Status	Percentage Correct (All)	
						(%)	(n)
MG101S-01	M. genitalium G37	Urine	DS1_2	Detected	CORE	82.8	134



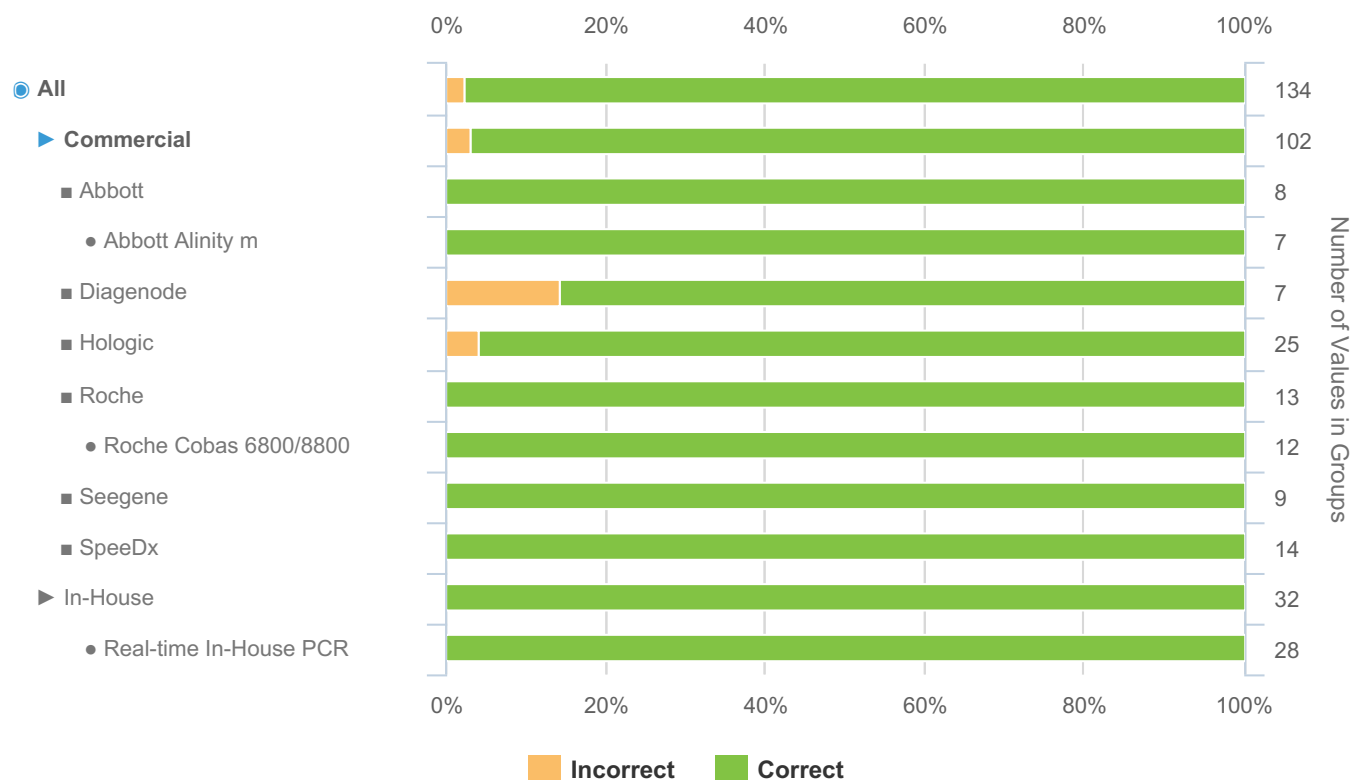
Groups below n=5: AB Analytica (n=2), AB Analytica - AB Analytica REALQUALITY RQ (n=2), Abbott - Abbott Real Time PCR (n=1), AusDiagnostics (n=1), AusDiagnostics - AusDiagnostics Easy-Plex (n=1), Bio-Rad (n=1), Bio-Rad - Bio-Rad Dx (n=1), Cepheid (n=1), Cepheid - Cepheid Xpert kit (n=1), Certest (n=1), Certest - Certest Real Time PCR (n=1), ELITech Group (n=3), ELITech Group - Elitech Elite Real Time kit (n=3), GeneProof (n=2), GeneProof - GeneProof Real Time PCR kit (n=2), InterLabService (n=2), InterLabService - InterLabService AmpliSens (n=2), PathoFinder (n=3), PathoFinder - PathoFinder Real Time PCR (n=3), Roche - Roche LightCycler (n=1), Sacace (n=3), Sacace - Sacace Real TM (n=3), TIB MOLBIOL (n=4), TIB MOLBIOL - TIB MOLBIOL LightMix (n=4), Vitassay (n=1), Vitassay - Vitassay Real-Time PCR (n=1), fast-track DIAGNOSTICS (n=2), fast-track DIAGNOSTICS - FTD real time PCR (n=2), In-House - Conventional In-House PCR (n=4)

Groups Rolled Up: Diagenode - Diagenode Real Time kit (n=7), Hologic - Hologic Aptima (n=25), Seegene - Seegene Real Time PCR (n=9), SpeedX - SpeedX Real Time PCR (n=14)

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MG101S-02 - Qualitative Results Breakdown

Sample Code	Sample Content	Matrix	Sample Relationships	Detection Frequency	Sample Status	Percentage Correct (All)	
						(%)	(n)
MG101S-02	M. genitalium G37	PBS	DS2_1	Frequently Detected	CORE	97.8	134



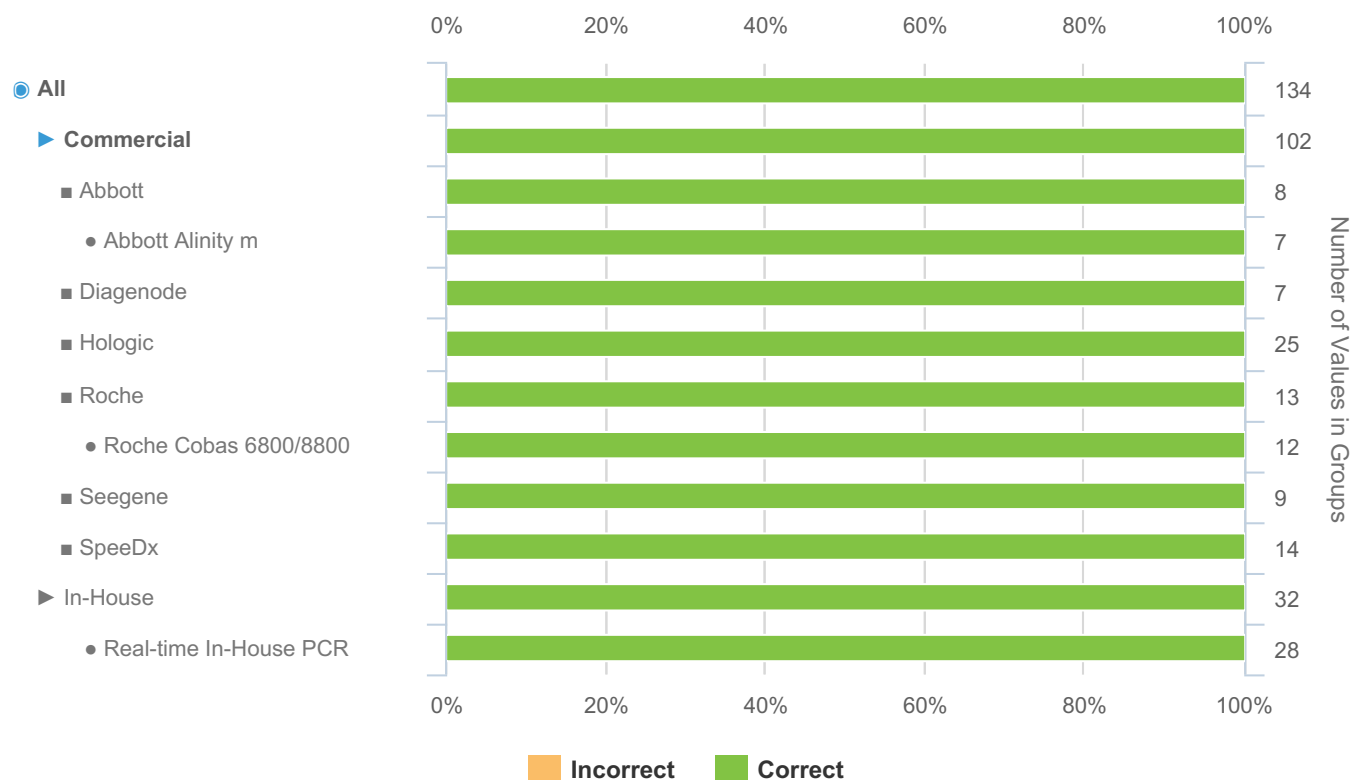
Groups below n=5: AB Analytica (n=2), AB Analytica - AB Analytica REALQUALITY RQ (n=2), Abbott - Abbott Real Time PCR (n=1), AusDiagnostics (n=1), AusDiagnostics - AusDiagnostics Easy-Plex (n=1), Bio-Rad (n=1), Bio-Rad - Bio-Rad Dx (n=1), Cepheid (n=1), Cepheid - Cepheid Xpert kit (n=1), Certest (n=1), Certest - Certest Real Time PCR (n=1), ELITech Group (n=3), ELITech Group - Elitech Elite Real Time kit (n=3), GeneProof (n=2), GeneProof - GeneProof Real Time PCR kit (n=2), InterLabService (n=2), InterLabService - InterLabService AmpliSens (n=2), PathoFinder (n=3), PathoFinder - PathoFinder Real Time PCR (n=3), Roche - Roche LightCycler (n=1), Sacace (n=3), Sacace - Sacace Real TM (n=3), TIB MOLBIOL (n=4), TIB MOLBIOL - TIB MOLBIOL LightMix (n=4), Vitassay (n=1), Vitassay - Vitassay Real-Time PCR (n=1), fast-track DIAGNOSTICS (n=2), fast-track DIAGNOSTICS - FTD real time PCR (n=2), In-House - Conventional In-House PCR (n=4)

Groups Rolled Up: Diagenode - Diagenode Real Time kit (n=7), Hologic - Hologic Aptima (n=25), Seegene - Seegene Real Time PCR (n=9), SpeedX - SpeedX Real Time PCR (n=14)

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MG101S-03 - Qualitative Results Breakdown

Sample Code	Sample Content	Matrix	Sample Relationships	Detection Frequency	Sample Status	Percentage Correct (All)	
						(%)	(n)
MG101S-03	Mycoplasma Negative	PBS		Negative	CORE	100.0	134



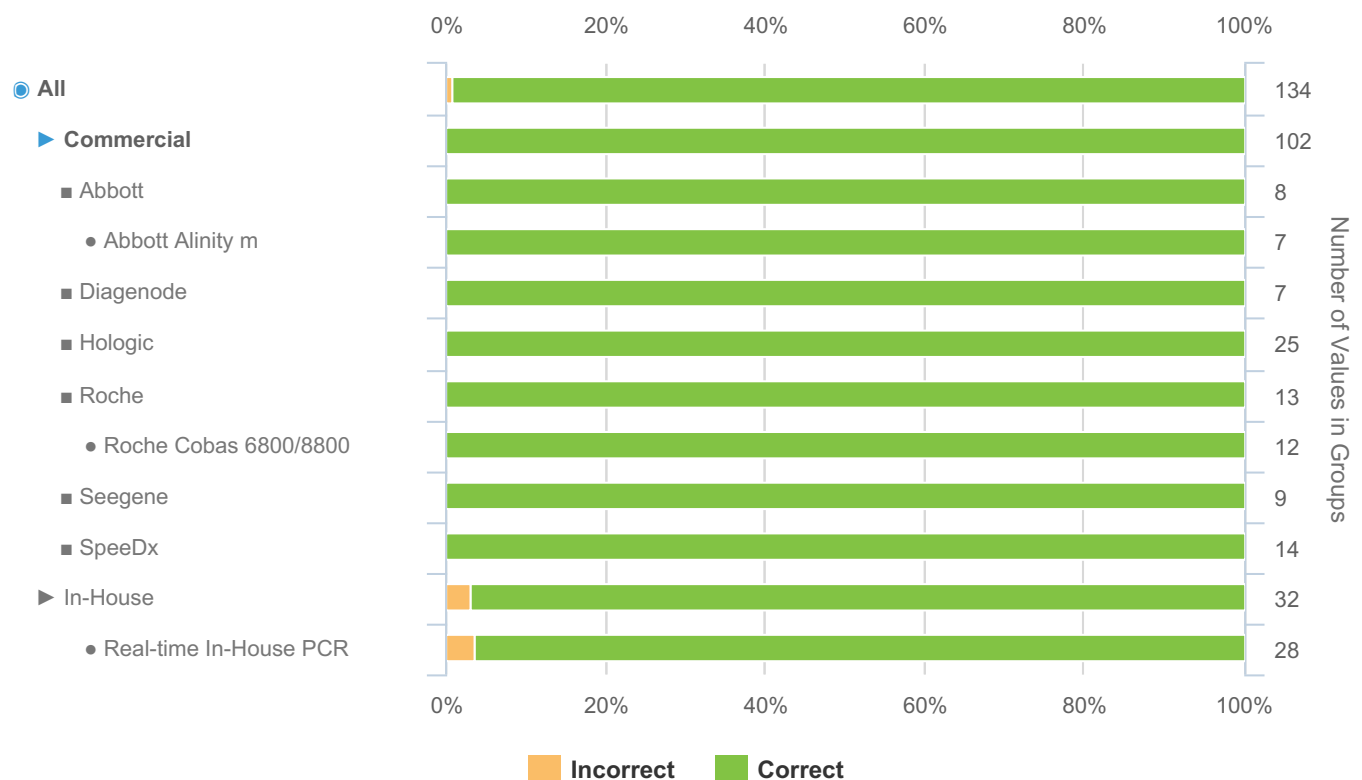
Groups below n=5: AB Analytica (n=2), AB Analytica - AB Analytica REALQUALITY RQ (n=2), Abbott - Abbott Real Time PCR (n=1), AusDiagnostics (n=1), AusDiagnostics - AusDiagnostics Easy-Plex (n=1), Bio-Rad (n=1), Bio-Rad - Bio-Rad Dx (n=1), Cepheid (n=1), Cepheid - Cepheid Xpert kit (n=1), Certest (n=1), Certest - Certest Real Time PCR (n=1), ELITech Group (n=3), ELITech Group - Elitech Elite Real Time kit (n=3), GeneProof (n=2), GeneProof - GeneProof Real Time PCR kit (n=2), InterLabService (n=2), InterLabService - InterLabService AmpliSens (n=2), PathoFinder (n=3), PathoFinder - PathoFinder Real Time PCR (n=3), Roche - Roche LightCycler (n=1), Sacace (n=3), Sacace - Sacace Real TM (n=3), TIB MOLBIOL (n=4), TIB MOLBIOL - TIB MOLBIOL LightMix (n=4), Vitassay (n=1), Vitassay - Vitassay Real-Time PCR (n=1), fast-track DIAGNOSTICS (n=2), fast-track DIAGNOSTICS - FTD real time PCR (n=2), In-House - Conventional In-House PCR (n=4)

Groups Rolled Up: Diagenode - Diagenode Real Time kit (n=7), Hologic - Hologic Aptima (n=25), Seegene - Seegene Real Time PCR (n=9), SpeedX - SpeedX Real Time PCR (n=14)

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MG101S-04 - Qualitative Results Breakdown

Sample Code	Sample Content	Matrix	Sample Relationships	Detection Frequency	Sample Status	Percentage Correct (All)	
						(%)	(n)
MG101S-04	M. genitalium M6303 (Macrolide Resistant)	PBS	DS3_1	Frequently Detected	CORE	99.3	134



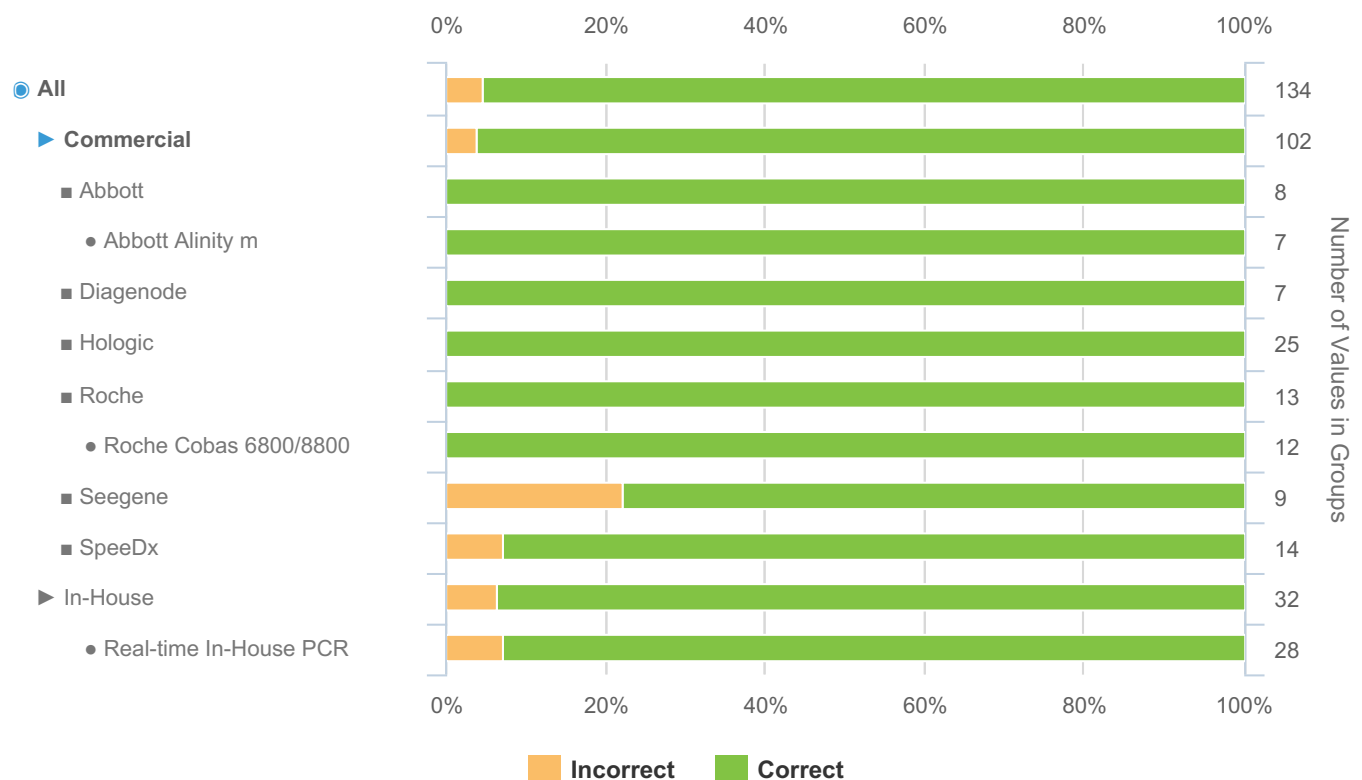
Groups below n=5: AB Analytica (n=2), AB Analytica - AB Analytica REALQUALITY RQ (n=2), Abbott - Abbott Real Time PCR (n=1), AusDiagnostics (n=1), AusDiagnostics - AusDiagnostics Easy-Plex (n=1), Bio-Rad (n=1), Bio-Rad - Bio-Rad Dx (n=1), Cepheid (n=1), Cepheid - Cepheid Xpert kit (n=1), Certest (n=1), Certest - Certest Real Time PCR (n=1), ELITech Group (n=3), ELITech Group - Elitech Elite Real Time kit (n=3), GeneProof (n=2), GeneProof - GeneProof Real Time PCR kit (n=2), InterLabService (n=2), InterLabService - InterLabService AmpliSens (n=2), PathoFinder (n=3), PathoFinder - PathoFinder Real Time PCR (n=3), Roche - Roche LightCycler (n=1), Sacace (n=3), Sacace - Sacace Real TM (n=3), TIB MOLBIOL (n=4), TIB MOLBIOL - TIB MOLBIOL LightMix (n=4), Vitassay (n=1), Vitassay - Vitassay Real-Time PCR (n=1), fast-track DIAGNOSTICS (n=2), fast-track DIAGNOSTICS - FTD real time PCR (n=2), In-House - Conventional In-House PCR (n=4)

Groups Rolled Up: Diagenode - Diagenode Real Time kit (n=7), Hologic - Hologic Aptima (n=25), Seegene - Seegene Real Time PCR (n=9), SpeedX - SpeedX Real Time PCR (n=14)

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MG101S-06 - Qualitative Results Breakdown

Sample Code	Sample Content	Matrix	Sample Relationships	Detection Frequency	Sample Status	Percentage Correct (All)	
						(%)	(n)
MG101S-06	M. genitalium M6593 (Macrolide Resistant)	PBS		Frequently Detected	CORE	95.5	134



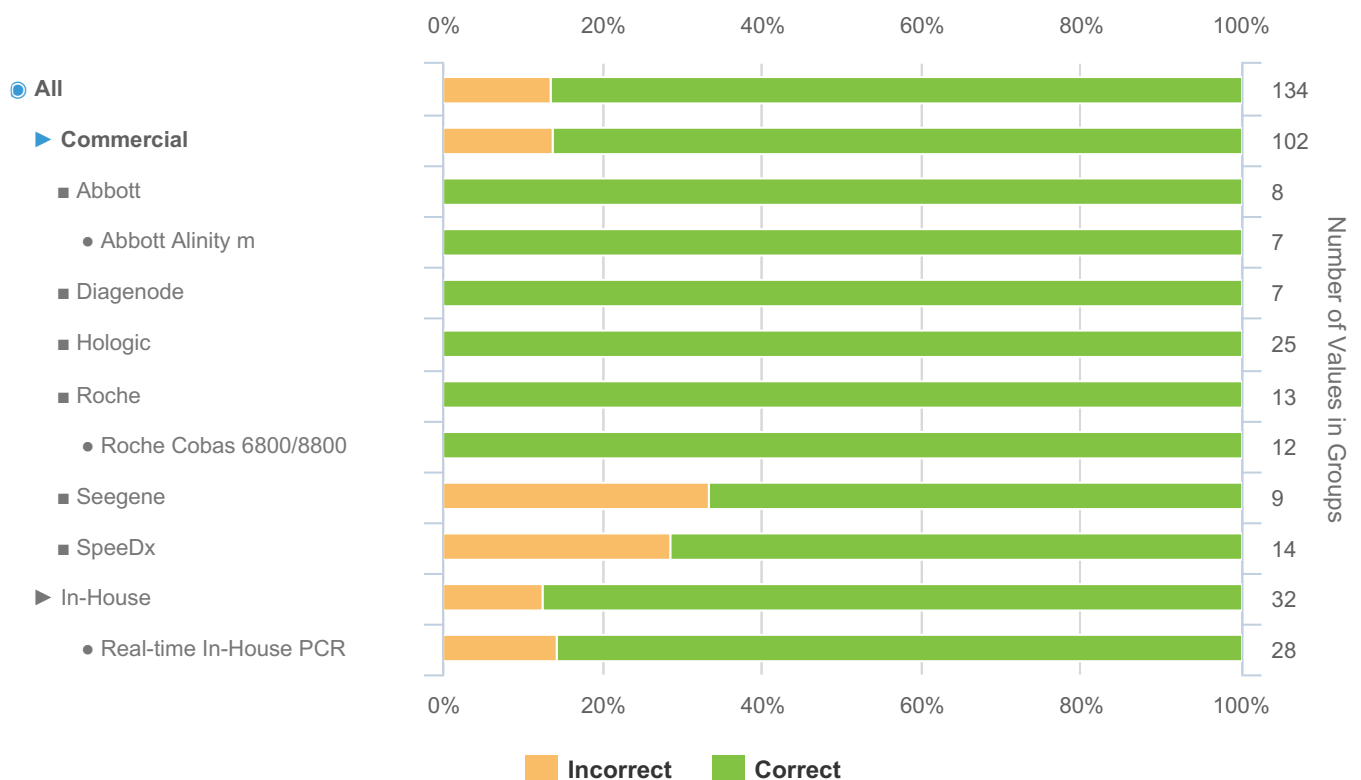
Groups below n=5: AB Analytica (n=2), AB Analytica - AB Analytica REALQUALITY RQ (n=2), Abbott - Abbott Real Time PCR (n=1), AusDiagnostics (n=1), AusDiagnostics - AusDiagnostics Easy-Plex (n=1), Bio-Rad (n=1), Bio-Rad - Bio-Rad Dx (n=1), Cepheid (n=1), Cepheid - Cepheid Xpert kit (n=1), Certest (n=1), Certest - Certest Real Time PCR (n=1), ELITech Group (n=3), ELITech Group - Elitech Elite Real Time kit (n=3), GeneProof (n=2), GeneProof - GeneProof Real Time PCR kit (n=2), InterLabService (n=2), InterLabService - InterLabService AmpliSens (n=2), PathoFinder (n=3), PathoFinder - PathoFinder Real Time PCR (n=3), Roche - Roche LightCycler (n=1), Sacace (n=3), Sacace - Sacace Real TM (n=3), TIB MOLBIOL (n=4), TIB MOLBIOL - TIB MOLBIOL LightMix (n=4), Vitassay (n=1), Vitassay - Vitassay Real-Time PCR (n=1), fast-track DIAGNOSTICS (n=2), fast-track DIAGNOSTICS - FTD real time PCR (n=2), In-House - Conventional In-House PCR (n=4)

Groups Rolled Up: Diagenode - Diagenode Real Time kit (n=7), Hologic - Hologic Aptima (n=25), Seegene - Seegene Real Time PCR (n=9), SpeedX - SpeedX Real Time PCR (n=14)

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MG101S-07 - Qualitative Results Breakdown

Sample Code	Sample Content	Matrix	Sample Relationships	Detection Frequency	Sample Status	Percentage Correct (All)	
						(%)	(n)
MG101S-07	M. genitalium G37	PBS	DS2_2	Detected	CORE	86.6	134



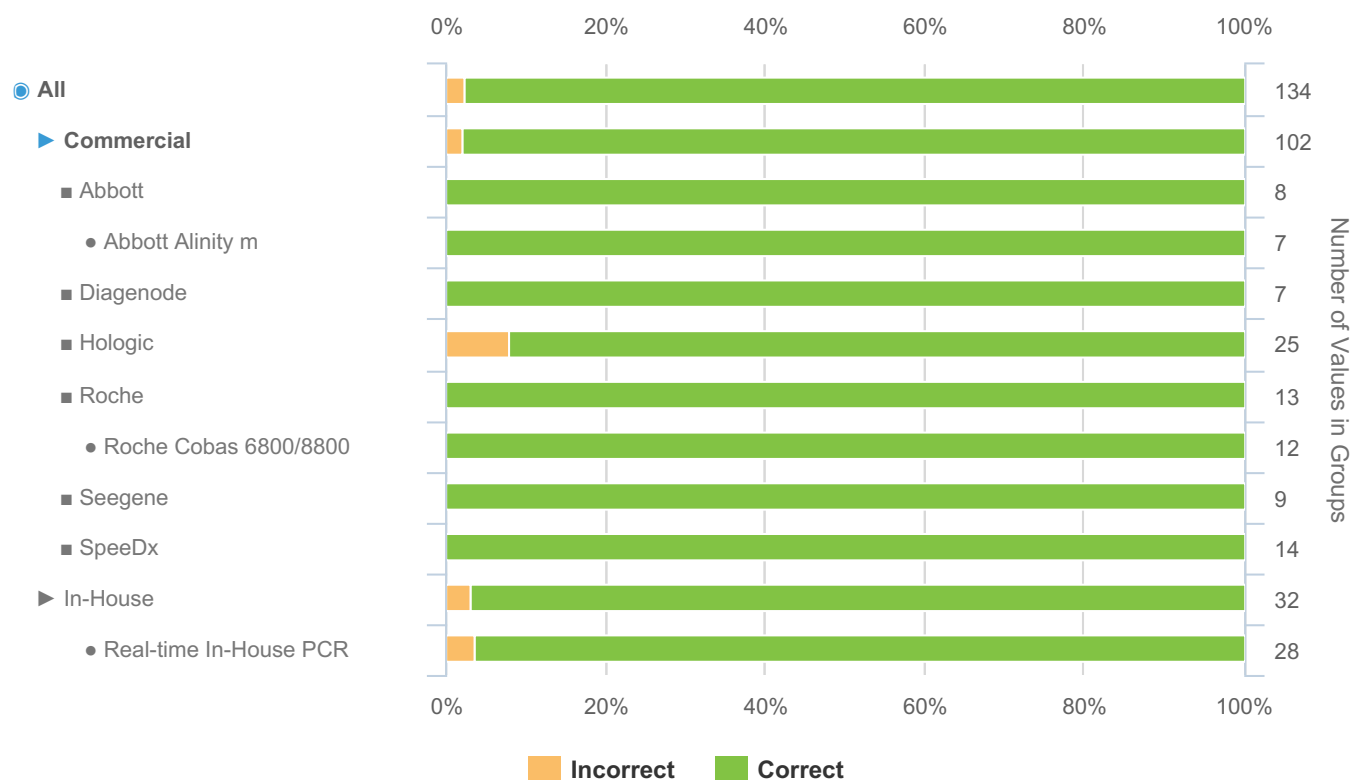
Groups below n=5: AB Analytica (n=2), AB Analytica - AB Analytica REALQUALITY RQ (n=2), Abbott - Abbott Real Time PCR (n=1), AusDiagnostics (n=1), AusDiagnostics - AusDiagnostics Easy-Plex (n=1), Bio-Rad (n=1), Bio-Rad - Bio-Rad Dx (n=1), Cepheid (n=1), Cepheid - Cepheid Xpert kit (n=1), Certest (n=1), Certest - Certest Real Time PCR (n=1), ELITech Group (n=3), ELITech Group - Elitech Elite Real Time kit (n=3), GeneProof (n=2), GeneProof - GeneProof Real Time PCR kit (n=2), InterLabService (n=2), InterLabService - InterLabService AmpliSens (n=2), PathoFinder (n=3), PathoFinder - PathoFinder Real Time PCR (n=3), Roche - Roche LightCycler (n=1), Sacace (n=3), Sacace - Sacace Real TM (n=3), TIB MOLBIOL (n=4), TIB MOLBIOL - TIB MOLBIOL LightMix (n=4), Vitassay (n=1), Vitassay - Vitassay Real-Time PCR (n=1), fast-track DIAGNOSTICS (n=2), fast-track DIAGNOSTICS - FTD real time PCR (n=2), In-House - Conventional In-House PCR (n=4)

Groups Rolled Up: Diagenode - Diagenode Real Time kit (n=7), Hologic - Hologic Aptima (n=25), Seegene - Seegene Real Time PCR (n=9), SpeedX - SpeedX Real Time PCR (n=14)

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MG101S-09 - Qualitative Results Breakdown

Sample Code	Sample Content	Matrix	Sample Relationships	Detection Frequency	Sample Status	Percentage Correct (All)	
						(%)	(n)
MG101S-09	M. pneumoniae	PBS		Negative	CORE	97.8	134



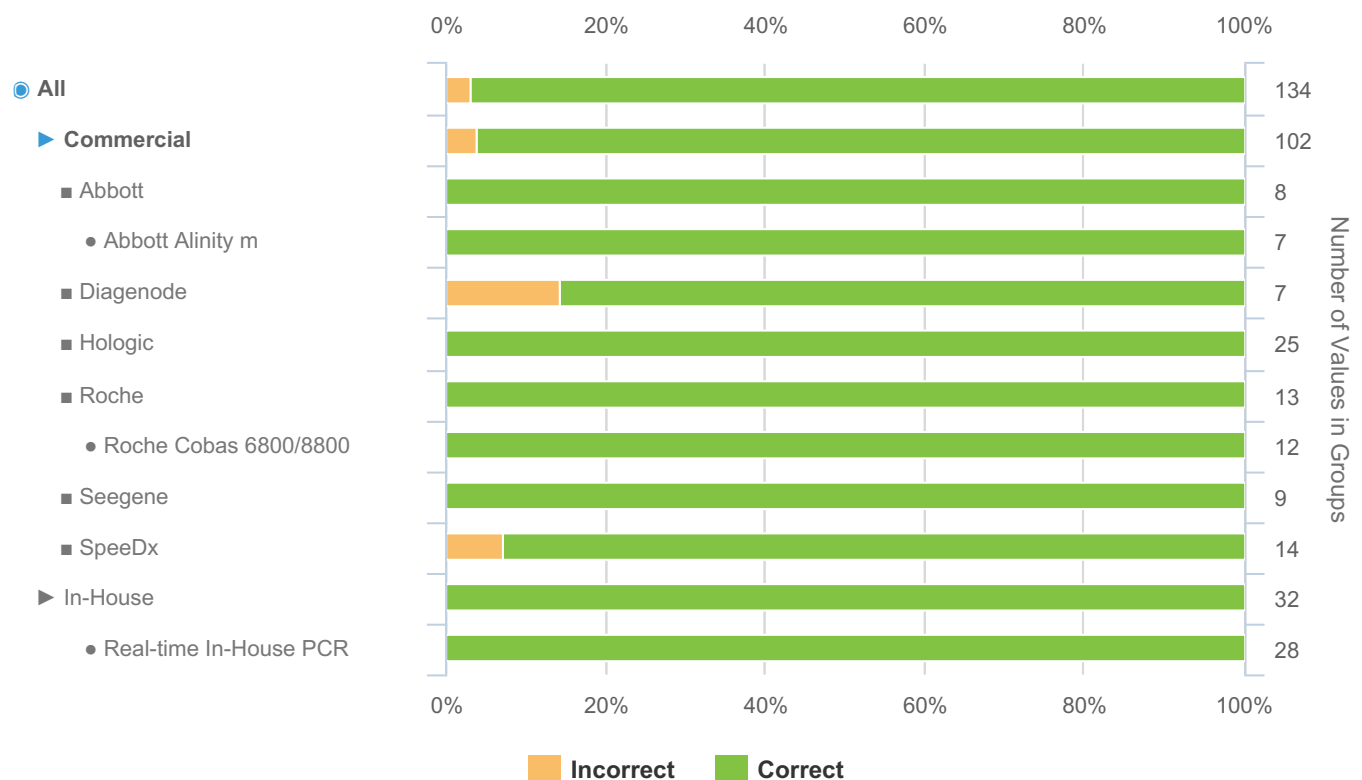
Groups below n=5: AB Analytica (n=2), AB Analytica - AB Analytica REALQUALITY RQ (n=2), Abbott - Abbott Real Time PCR (n=1), AusDiagnostics (n=1), AusDiagnostics - AusDiagnostics Easy-Plex (n=1), Bio-Rad (n=1), Bio-Rad - Bio-Rad Dx (n=1), Cepheid (n=1), Cepheid - Cepheid Xpert kit (n=1), Certest (n=1), Certest - Certest Real Time PCR (n=1), ELITech Group (n=3), ELITech Group - Elitech Elite Real Time kit (n=3), GeneProof (n=2), GeneProof - GeneProof Real Time PCR kit (n=2), InterLabService (n=2), InterLabService - InterLabService AmpliSens (n=2), PathoFinder (n=3), PathoFinder - PathoFinder Real Time PCR (n=3), Roche - Roche LightCycler (n=1), Sacace (n=3), Sacace - Sacace Real TM (n=3), TIB MOLBIOL (n=4), TIB MOLBIOL - TIB MOLBIOL LightMix (n=4), Vitassay (n=1), Vitassay - Vitassay Real-Time PCR (n=1), fast-track DIAGNOSTICS (n=2), fast-track DIAGNOSTICS - FTD real time PCR (n=2), In-House - Conventional In-House PCR (n=4)

Groups Rolled Up: Diagenode - Diagenode Real Time kit (n=7), Hologic - Hologic Aptima (n=25), Seegene - Seegene Real Time PCR (n=9), SpeedX - SpeedX Real Time PCR (n=14)

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MG101S-10 - Qualitative Results Breakdown

Sample Code	Sample Content	Matrix	Sample Relationships	Detection Frequency	Sample Status	Percentage Correct (All)	
						(%)	(n)
MG101S-10	M. genitalium G37	Urine	DS1_1	Frequently Detected	CORE	97.0	134



Groups below n=5: AB Analytica (n=2), AB Analytica - AB Analytica REALQUALITY RQ (n=2), Abbott - Abbott Real Time PCR (n=1), AusDiagnostics (n=1), AusDiagnostics - AusDiagnostics Easy-Plex (n=1), Bio-Rad (n=1), Bio-Rad - Bio-Rad Dx (n=1), Cepheid (n=1), Cepheid - Cepheid Xpert kit (n=1), Certest (n=1), Certest - Certest Real Time PCR (n=1), ELITech Group (n=3), ELITech Group - Elitech Elite Real Time kit (n=3), GeneProof (n=2), GeneProof - GeneProof Real Time PCR kit (n=2), InterLabService (n=2), InterLabService - InterLabService AmpliSens (n=2), PathoFinder (n=3), PathoFinder - PathoFinder Real Time PCR (n=3), Roche - Roche LightCycler (n=1), Sacace (n=3), Sacace - Sacace Real TM (n=3), TIB MOLBIOL (n=4), TIB MOLBIOL - TIB MOLBIOL LightMix (n=4), Vitassay (n=1), Vitassay - Vitassay Real-Time PCR (n=1), fast-track DIAGNOSTICS (n=2), fast-track DIAGNOSTICS - FTD real time PCR (n=2), In-House - Conventional In-House PCR (n=4)

Groups Rolled Up: Diagenode - Diagenode Real Time kit (n=7), Hologic - Hologic Aptima (n=25), Seegene - Seegene Real Time PCR (n=9), Speedx - Speedx Real Time PCR (n=14)

Individual Report		QCMD 2020 Mycoplasma genitalium EQA Pilot Study				
Catalogue Code: QAB184205	Ref Code: MG20	Challenge: S	Analysis Type: Qualitative	Dataset: 395362	Report UID: 2677/395362/3007	Laboratory 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.

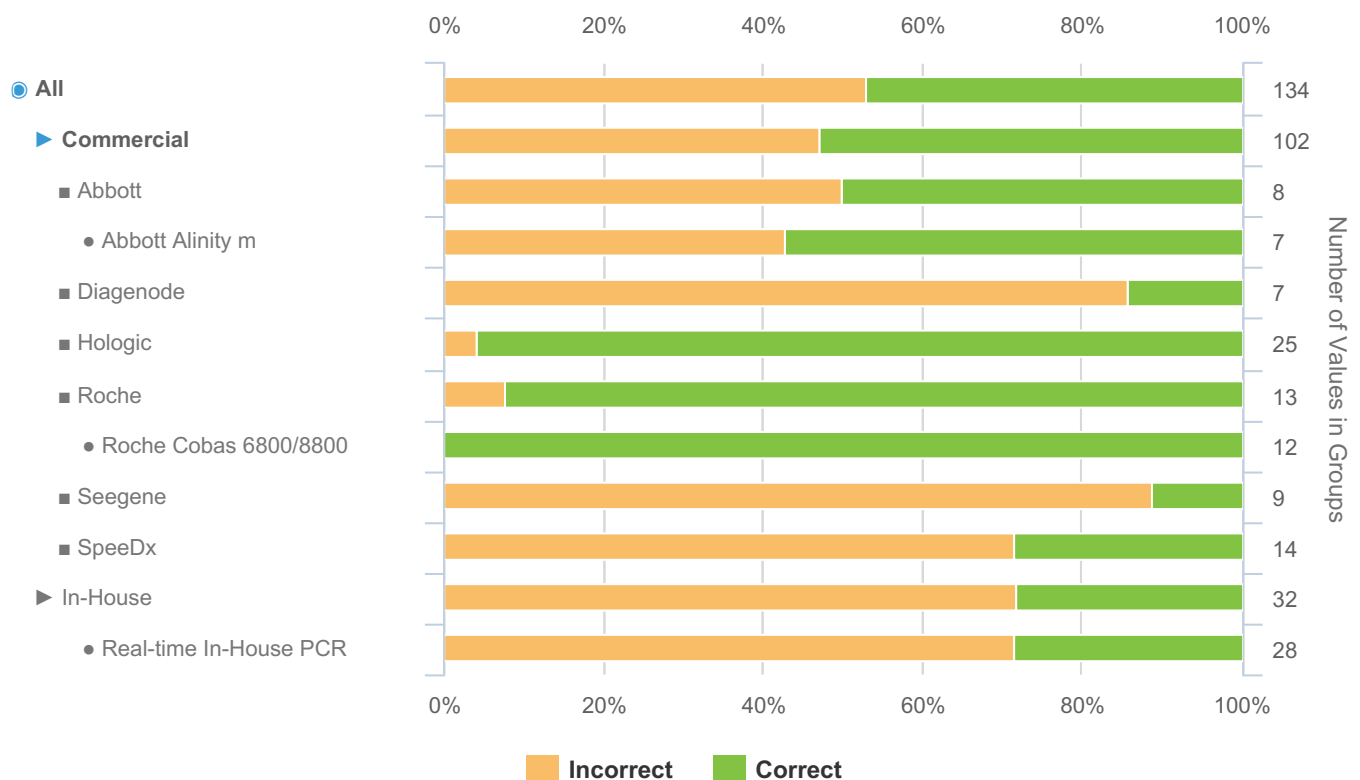
To allow meaningful assessment at the individual method level the EQA assessment group must consist of 5 or more datasets. If there are not sufficient datasets at the individual method level then your results will be included within a higher EQA assessment group based on whether it is a commercial or 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.

Individual Report		QCMD 2020 Mycoplasma genitalium EQA Pilot Study			 <small>Quality Control for Molecular Diagnostics</small>	
Catalogue Code: QAB184205	Ref Code: MG20	Challenge: S	Analysis Type: Qualitative	Dataset: 395362	Report UID: 2677/395362/3007	Laboratory CZ023

MG101S-05 - Qualitative Results Breakdown

Sample Code	Sample Content	Matrix	Sample Relationships	Detection Frequency	Sample Status	Percentage Correct (All)	
						(%)	(n)
MG101S-05	M. genitalium G37	PBS	DS2_3	Infrequently Detected	EDUCATIONAL	47.0	134



Groups below n=5: AB Analytica (n=2), AB Analytica - AB Analytica REALQUALITY RQ (n=2), Abbott - Abbott Real Time PCR (n=1), AusDiagnostics (n=1), AusDiagnostics - AusDiagnostics Easy-Plex (n=1), Bio-Rad (n=1), Bio-Rad - Bio-Rad Dx (n=1), Cepheid (n=1), Cepheid - Cepheid Xpert kit (n=1), Certest (n=1), Certest - Certest Real Time PCR (n=1), ELITech Group (n=3), ELITech Group - Elitech Elite Real Time kit (n=3), GeneProof (n=2), GeneProof - GeneProof Real Time PCR kit (n=2), InterLabService (n=2), InterLabService - InterLabService AmpliSens (n=2), PathoFinder (n=3), PathoFinder - PathoFinder Real Time PCR (n=3), Roche - Roche LightCycler (n=1), Sacace (n=3), Sacace - Sacace Real TM (n=3), TIB MOLBIOL (n=4), TIB MOLBIOL - TIB MOLBIOL LightMix (n=4), Vitassay (n=1), Vitassay - Vitassay Real-Time PCR (n=1), fast-track DIAGNOSTICS (n=2), fast-track DIAGNOSTICS - FTD real time PCR (n=2), In-House - Conventional In-House PCR (n=4)

Groups Rolled Up: Diagenode - Diagenode Real Time kit (n=7), Hologic - Hologic Aptima (n=25), Seegene - Seegene Real Time PCR (n=9), SpeedX - SpeedX Real Time PCR (n=14)

Individual Report

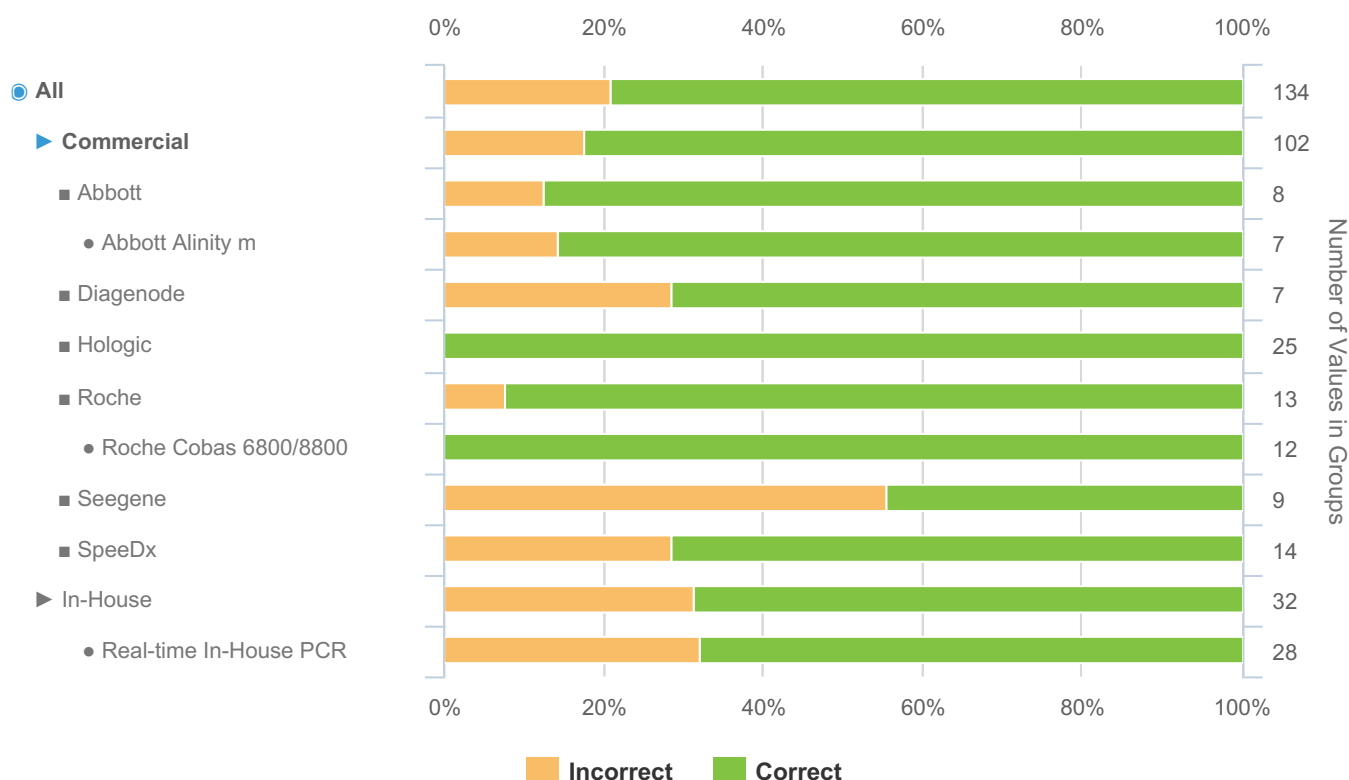
QCMD 2020 Mycoplasma genitalium EQA Pilot Study



Catalogue Code: QAB184205	Ref Code: MG20	Challenge: S	Analysis Type: Qualitative	Dataset: 395362	Report UID: 2677/395362/3007	Laboratory: CZ023
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MG101S-08 - Qualitative Results Breakdown

Sample Code	Sample Content	Matrix	Sample Relationships	Detection Frequency	Sample Status	Percentage Correct (All)	
						(%)	(n)
MG101S-08	M. genitalium M6303 (Macrolide Resistant)	PBS	DS3_2	Detected	EDUCATIONAL	79.1	134



Groups below n=5: AB Analytica (n=2), AB Analytica - AB Analytica REALQUALITY RQ (n=2), Abbott - Abbott Real Time PCR (n=1), AusDiagnostics (n=1), AusDiagnostics - AusDiagnostics Easy-Plex (n=1), Bio-Rad (n=1), Bio-Rad - Bio-Rad Dx (n=1), Cepheid (n=1), Cepheid - Cepheid Xpert kit (n=1), Certest (n=1), Certest - Certest Real Time PCR (n=1), ELITech Group (n=3), ELITech Group - Elitech Elite Real Time kit (n=3), GeneProof (n=2), GeneProof - GeneProof Real Time PCR kit (n=2), InterLabService (n=2), InterLabService - InterLabService AmpliSens (n=2), PathoFinder (n=3), PathoFinder - PathoFinder Real Time PCR (n=3), Roche - Roche LightCycler (n=1), Sacace (n=3), Sacace - Sacace Real TM (n=3), TIB MOLBIOL (n=4), TIB MOLBIOL - TIB MOLBIOL LightMix (n=4), Vitassay (n=1), Vitassay - Vitassay Real-Time PCR (n=1), fast-track DIAGNOSTICS (n=2), fast-track DIAGNOSTICS - FTD real time PCR (n=2), In-House - Conventional In-House PCR (n=4)

Groups Rolled Up: Diagenode - Diagenode Real Time kit (n=7), Hologic - Hologic Aptima (n=25), Seegene - Seegene Real Time PCR (n=9), SpeedX - SpeedX Real Time PCR (n=14)

Individual Report		QCMD 2020 Mycoplasma genitalium EQA Pilot Study				
Catalogue Code: QAB184205	Ref Code: MG20	Challenge: S	Analysis Type: Qualitative	Dataset: 395362	Report UID: 2677/395362/3007	Laboratory CZ023

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