Individual Report	QCMD 2021 Epstein-Barr virus DNA EQA Programme	

Catalogue Code:	Ref Code:	Challenge:	Analysis Type:	Dataset:	Report UID:	Laboratory	
QAV024121	EBVDNA21	C2	Qualitative and Quantitative	495139	2677/495139/3867	CZ023	

Intended Results / Panel Composition

Sample Code	Sample Content	Matrix	Sample Relationships ^[1]	Detection Frequency ^[2]	Sample Status ^[3]	Consensus (Copies/ml) ^[4]		Range ^[5]
						(Log ₁₀)	(n)	
EBVDNA21C2-01	Epstein- Barr virus	Plasma	DS1_1	Frequently Detected	CORE	4.171	50	2.915 - 5.697
EBVDNA21C2-02	Epstein- Barr virus	Plasma	D1, DS1_3	Detected	CORE	3.167	46	2.407 - 5.752
EBVDNA21C2-03	Epstein- Barr virus	Plasma	D1, DS1_3	Frequently Detected	CORE	3.153	48	1.556 - 5.988
EBVDNA21C2-04	Epstein- Barr virus	Plasma	DS1_2	Frequently Detected	CORE	3.412	48	1.041 - 5.560
EBVDNA21C2-05	Negative	Plasma		Negative	CORE	N/A	N/A	N/A

[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] **Consensus (Copies/ml):** Mean consensus (Log₁₀) calculated from data returned by participants with outliers removed and number of quantitative results (n) returned for each panel member.

[5] Range: Maximum and minimum quantitative value (Copies/ml) reported by participants within this challenge without outliers removed.

For further details please refer to the current participant manual.

Your Summary Results

Units	Copies/ml
EQA Assessment Group ^[1]	Commercial
Core Panel Detection (Qualitative) Score ^[2]	0
Core Panel Estimation (Quantitative) Score ^[3]	0

Individual Report	QCMD 2021 Ep EQA Programn	stein-Barr virus DNA าe	Y	and the second se	

Catalogue Code:	Ref Code:	Challenge:	Analysis Type:	Dataset:	Report UID:	Laboratory	
QAV024121	EBVDNA21	C2	Qualitative and Quantitative	495139	2677/495139/3867	CZ023	

Core Panel Members Results

Sample Code	Unitage	EQA Assessment Group Consensus ^[4]	SD [5]	Quantitative Result		Qualitative Result		
				Your Result ^[6]	Estimation Score ^[7]	Percentage Correct (All) ^[8]	Your Result ^[9]	Detection Score ^[10]
EBVDNA21C2-01	Copies/ml	4.163	0.627	4.009	0	96.9	Positive	0
EBVDNA21C2-02	Copies/ml	3.146	0.490	3.012	0	92.7	Positive	0
EBVDNA21C2-03	Copies/ml	3.140	0.599	3.134	0	95.8	Positive	0
EBVDNA21C2-04	Copies/ml	3.419	0.732	3.324	0	99.0	Positive	0
EBVDNA21C2-05	Copies/ml	N/A	-	LOD/NR	N/A	94.8	Negative	0

All quantitative values above expressed in Log 10 Copies/ml.

[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] Core Panel Estimation (Quantitative) Score: An overall core panel estimation score provided per challenge / distribution.

[4] EQA Assessment Group Consensus: The mean value for all results within your EQA assessment group.

[5] SD: The standard deviation for results from your EQA assessment group.

[6] Your Quantitative Result: The quantitative result you returned for each sample within this EQA challenge. LOD/NR (limit of detection or not reported).

[7] Estimation Score: Your estimation (quantitative) scores are calculated based on your variation from the consensus for your EQA assessment group. With 0 (zero) scored if the quantitative value you reported is within one standard deviation (SD) from your EQA assessment group consensus, 1 (one) if your quantitative value is between one and two SDs, 2 (two) if your quantitative value is within two and three SDs and 3 (three) if your quantitative value is more than three SDs from the mean of your EQA assessment group.

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

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

[10] **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) 2021 Ep Programn	stein-Barr virus DN ne	Α		
Catalogue Code:	Ref Code:	Challenge:	Analysis Type:	Dataset:	Report UID:	Laboratory

Qualitative and Quantitative

495139

2677/495139/3867

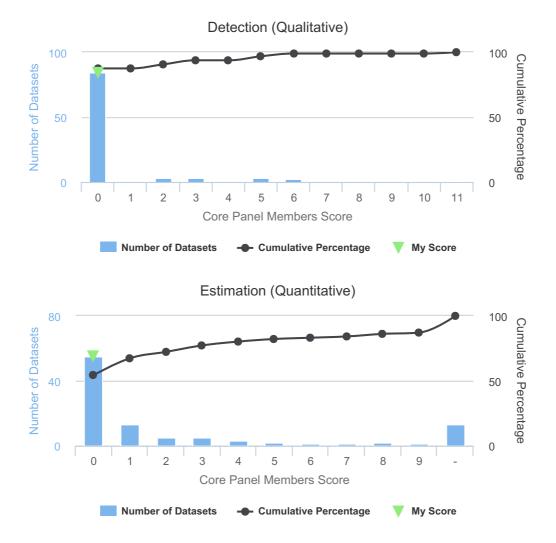
CZ023

Core Panel Member Score Breakdown

EBVDNA21

C2

QAV024121



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".

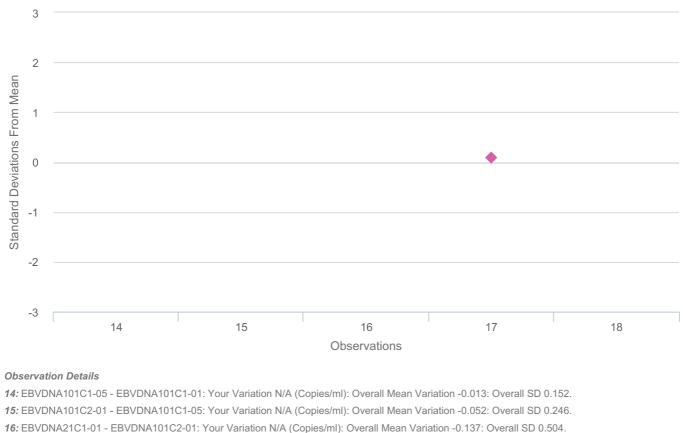
Core Panel Member Score Breakdown - Estimation: This figure gives you a breakdown of the quantitative estimation scores for all quantitative datasets returned within this EQA challenge / distribution independent of the EQA assessment group. Panel estimation scores are based on positive core panel members only. Those datasets that did not return quantitative values for all core samples are represented by '-'. *For further details please refer to the current participant manual.*

Individual	QCMD 2021 Ep	stein-Barr virus DN/	a 💋	QCAD
Report	EQA Programm	าe		Guality Control for Molecular Diagnostics

Catalogue Code:	Ref Code:	Challenge:	Analysis Type:	Dataset:	Report UID:	Laboratory
QAV024121	EBVDNA21	C2	Qualitative and Quantitative	495139	2677/495139/3867	CZ023

Duplicate Sample Performance Over Time





16: EBVDINA21C1-01 - EBVDINA101C2-01: Your Variation N/A (Copies/mil): Overall Mean Variation -0.137: Overall SD 0.504

17: EBVDNA21C1-05 - EBVDNA21C1-01: Your Variation 0.049 (Copies/ml): Overall Mean Variation 0.026: Overall SD 0.261.

18: EBVDNA21C2-01 - EBVDNA21C1-05: N/A

QCMD monitors your laboratory's performance over time based on the reported quantitative variation between duplicate panel members within the EQA challenge and, where appropriate, across EQA challenges.

The mean variation and standard deviation are calculated from the quantitative variation reported by each participant between duplicate panel members in the same unit of measurement once outliers have been removed. (See 'Observation Details')

Previous and current observations are plotted on the chart as the number of standard deviations your variation was from the mean variation for all participants who submitted corresponding results in the same unitage.

Any reported variation greater than ±3 SD will not be shown on the graph, but your variation value will be provided **in red** in the *Observation Details*. When "N/A" is displayed for an observation, either no valid quantitative results were provided or there was a change in reported unitage.

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Catalogue Code:	Ref Code:	Challenge:	Analysis Type:	Dataset:	Report UID:	Laboratory
QAV024121	EBVDNA21	C2	Qualitative and Quantitative	495139	2677/495139/3867	CZ023

My Workflow Details

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

Name	GeneProof Epstein-Barr Virus (EBV) PCR Kit + croBEE (v4)						
Description							
Targets	V Epstein-Barr virus						
Assays	 <i>Extraction</i> - GeneProof - GeneProof Commercial Kit Manufacturer: <i>GeneProof</i> Kit Type: <i>croBEE</i> <i>Amplification</i> - GeneProof - croBEE Real-Time PCR System Commercial Kit Manufacturer: <i>GeneProof</i> Kit Manufacturer: <i>GeneProof</i> Kit Type: <i>GeneProof</i> Epstein-Barr (EBV) PCR Kit Kit Version: ISEX 						

Further Programme Details

Number of Participants	107
Number of Countries	30
Number of Respondents	97
Number of Datasets Submitted	115
Quantitative Results Returned (All)	101 (87.8%)
- Quantitative Results Returned (Copies/ml)	51 (50.5%)
- Quantitative Results Returned (IU/mI)	50 (49.5%)

EQA Programme Aims

To assess the proficiency of laboratories in the detection and quantitation of Epstein-Barr virus (EBV).

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.

Catalogue Code:	Ref Code:	Challenge:	Analysis Type:	Dataset:	Report UID:	Laboratory
QAV024121	EBVDNA21	C2	Qualitative and Quantitative	495139	2677/495139/3867	CZ023

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 ► Quantitative ► Copies/ml, IU/ml ► Qualitative

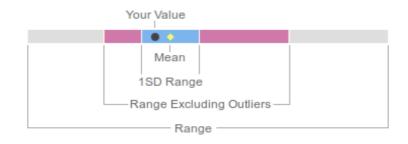
Individual Panel Member Analysis (Quantitative)

Quantitative 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 reported results using the same unit of measurement (i.e. Copies/ml or IU/ml).

The results below provide a breakdown of participant reported values on each of the panel members within this EQA challenge / distribution. Your result for each panel member is indicated by "your value". You can compare your value to the "mean" within your EQA assessment group and the overall consensus for each sample within this EQA challenge / distribution.

Key

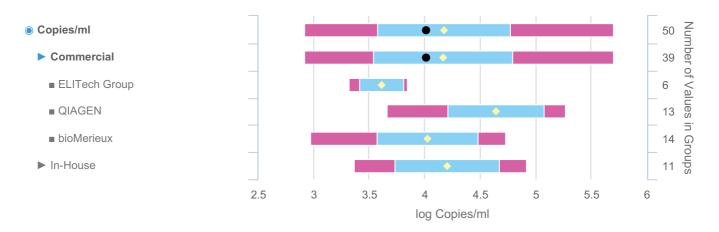


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Catalogue Code:	Ref Code:	Challenge:	Analysis Type:	Dataset:	Report UID:	Laboratory	
QAV024121	EBVDNA21	C2	Qualitative and Quantitative	495139	2677/495139/3867	CZ023	

EBVDNA21C2-01 - Quantitative Results Breakdown (Copies/ml)

Sample Code	Sample Content	Matrix	Sample Relationships	Detection Frequency	Sample Status	Consensus (Copies/ml)		Range
						(Log ₁₀)	(n)	
EBVDNA21C2-01	Epstein-Barr virus	Plasma	DS1_1	Frequently Detected	CORE	4.171	50	2.915 - 5.697



Groups below n=5: AB Analitica (n=1), AB Analitica - AB Analitica REALQUALITY RQ (n=1), Altona Diagnostics (n=1), Altona Diagnostics - Altona Diagnostics RealStar (n=1), GeneProof (n=4), GeneProof - GeneProof Real Time PCR kit (n=4)

Groups Rolled Up: ELITech Group - Elitech Elite Real Time kit (n=6), QIAGEN - QIAGEN Artus Real Time (n=13), bioMerieux - bioMerieux R-gene Quant Kit (n=14), In-House - Real-time In-House PCR (n=11)

Individual	QCMD 2021 Epstein-Barr virus DNA	QCMD
Report	EQA Programme	Quality Control for Malecular Diagnostics

Catalogue Code:	Ref Code:	Challenge:	Analysis Type:	Dataset:	Report UID:	Laboratory	
QAV024121	EBVDNA21	C2	Qualitative and Quantitative	495139	2677/495139/3867	CZ023	

EBVDNA21C2-02 - Quantitative Results Breakdown (Copies/ml)

Sample Code	Sample Content	Matrix	Sample Relationships	Detection Frequency	Sample Status	Consensus (Copies/ml)		Range
						(Log ₁₀)	(n)	
EBVDNA21C2-02	2 Epstein-Barr virus	Plasma	D1, DS1_3	Detected	CORE	3.167	46	2.407 - 5.752



Groups below n=5: Altona Diagnostics (n=1), Altona Diagnostics - Altona Diagnostics RealStar (n=1), GeneProof (n=4), GeneProof - GeneProof Real Time PCR kit (n=4)

Groups Rolled Up: ELITech Group - Elitech Elite Real Time kit (n=6), QIAGEN - QIAGEN Artus Real Time (n=12), bioMerieux - bioMerieux R-gene Quant Kit (n=12), In-House - Real-time In-House PCR (n=11)

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Catalogue Code:	Ref Code:	Challenge:	Analysis Type:	Dataset:	Report UID:	Laboratory	
QAV024121	EBVDNA21	C2	Qualitative and Quantitative	495139	2677/495139/3867	CZ023	

EBVDNA21C2-03 - Quantitative Results Breakdown (Copies/ml)

Sample Code	Sample Content	Matrix	Sample Relationships	Detection Frequency	Sample Status	Consensus (Copies/ml)		Range
						(Log ₁₀)	(n)	
EBVDNA21C2-03	Epstein-Barr virus	Plasma	D1, DS1_3	Frequently Detected	CORE	3.153	48	1.556 - 5.988



Groups below n=5: AB Analitica (n=1), AB Analitica - AB Analitica REALQUALITY RQ (n=1), Altona Diagnostics (n=1), Altona Diagnostics - Altona Diagnostics RealStar (n=1), GeneProof (n=4), GeneProof - GeneProof Real Time PCR kit (n=4)

Groups Rolled Up: ELITech Group - Elitech Elite Real Time kit (n=6), QIAGEN - QIAGEN Artus Real Time (n=12), bioMerieux - bioMerieux R-gene Quant Kit (n=13), In-House - Real-time In-House PCR (n=11)

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Catalogue Code:	Ref Code:	Challenge:	Analysis Type:	Dataset:	Report UID:	Laboratory	
QAV024121	EBVDNA21	C2	Qualitative and Quantitative	495139	2677/495139/3867	CZ023	

EBVDNA21C2-04 - Quantitative Results Breakdown (Copies/ml)

Sample Code	Sample Content	Matrix	Sample Relationships	Detection Frequency	Sample Status	Consensus (Copies/ml)		Range
						(Log ₁₀)	(n)	
EBVDNA21C2-04	Epstein-Barr virus	Plasma	DS1_2	Frequently Detected	CORE	3.412	48	1.041 - 5.560



Groups below n=5: AB Analitica (n=1), AB Analitica - AB Analitica REALQUALITY RQ (n=1), Altona Diagnostics (n=1), Altona Diagnostics - Altona Diagnostics RealStar (n=1), GeneProof (n=4), GeneProof - GeneProof Real Time PCR kit (n=4)

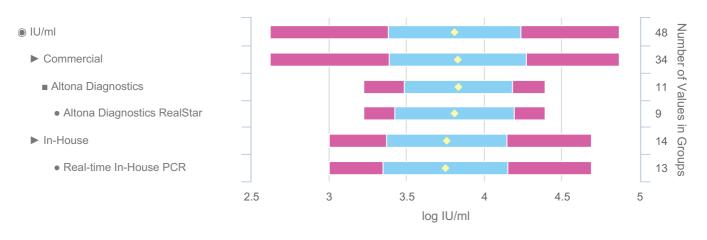
Groups Rolled Up: ELITech Group - Elitech Elite Real Time kit (n=6), QIAGEN - QIAGEN Artus Real Time (n=12), bioMerieux - bioMerieux R-gene Quant Kit (n=13), In-House - Real-time In-House PCR (n=11)

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Catalogue Code:	Ref Code:	Challenge:	Analysis Type:	Dataset:	Report UID:	Laboratory	
QAV024121	EBVDNA21	C2	Qualitative and Quantitative	495139	2677/495139/3867	CZ023	

EBVDNA21C2-01 - Quantitative Results Breakdown (IU/ml)

Sample Code	Sample Content	Matrix	Sample Relationships	Detection Frequency	Sample Status	Consens (IU/ml)	us	Range
						(Log ₁₀)	(n)	
EBVDNA21C2-01	Epstein-Barr virus	Plasma	DS1_1	Frequently Detected	CORE	3.806	48	2.620 - 4.866



Groups below n=5: AB Analitica (n=2), AB Analitica - AB Analitica REALQUALITY RQ (n=2), Abbott (n=4), Abbott - Abbott RealTime rn2000 (n=4), Altona Diagnostics - Altona Diagnostics AltoStar (n=2), Diagenode (n=1), Diagenode - Diagenode Real Time kit (n=1), ELITech Group (n=1), ELITech Group (n=1), ELITech Elite Real Time kit (n=1), GeneProof (n=1), GeneProof - GeneProof Real Time PCR kit (n=1), Hong Kong CH Gene (n=1), Hong Kong CH Gene (n=1), QIAGEN (n=4), QIAGEN - QIAGEN Artus Real Time (n=3), QIAGEN - Qiagen NeuMoDx (n=1), Roche (n=4), Roche - Roche Cobas (n=4), Sacace (n=1), Sacace - Sacace Real TM (n=1), bioMerieux (n=4), bioMerieux - bioMerieux R-gene Quant Kit (n=4), In-House - Conventional In-House PCR (n=1)

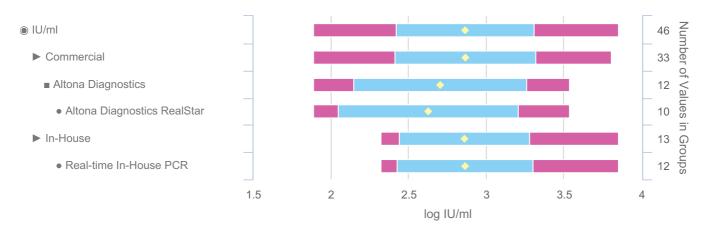
Groups Rolled Up:

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Catalogue Code:	Ref Code:	Challenge:	Analysis Type:	Dataset:	Report UID:	Laboratory	
QAV024121	EBVDNA21	C2	Qualitative and Quantitative	495139	2677/495139/3867	CZ023	

EBVDNA21C2-02 - Quantitative Results Breakdown (IU/ml)

Sample Code	Sample Content	Matrix	Sample Relationships	Detection Frequency	Sample Status	Consens (IU/ml)	us	Range
						(Log ₁₀)	(n)	
EBVDNA21C2-02	Epstein-Barr virus	Plasma	D1, DS1_3	Detected	CORE	2.861	46	1.886 - 3.848



Groups below n=5: Abbott (n=4), Abbott - Abbott RealTime rn2000 (n=4), Altona Diagnostics - Altona Diagnostics AltoStar (n=2), Diagenode (n=1), Diagenode - Diagenode Real Time kit (n=1), ELITech Group (n=1), ELITech Group - Elitech Elite Real Time kit (n=1), GeneProof - GeneProof Real Time PCR kit (n=1), Hong Kong CH Gene (n=1), Hong Kong CH Gene - HK CH Gene Real Time PCR (n=1), QIAGEN (n=4), QIAGEN - QIAGEN Artus Real Time (n=3), QIAGEN - Qiagen NeuMoDx (n=1), Roche (n=4), Roche - Roche Cobas (n=4), Sacace (n=1), Sacace - Sacace Real TM (n=1), bioMerieux (n=4), bioMerieux - bioMerieux R-gene Quant Kit (n=4), In-House - Conventional In-House PCR (n=1)

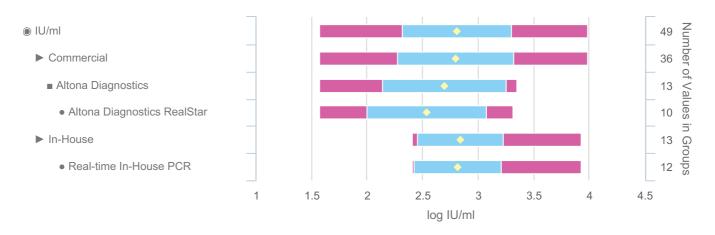
Groups Rolled Up:

Individual Report		D 2021 Ep Programi	ostein-Barr virus Dl me	NA		
Catalogua Cada	Pof Codo:	Challonger		Detect	Bonort IIID:	Laboratory

Catalogue Code:	Ref Code:	Challenge:	Analysis Type:	Dataset:	Report UID:	Laboratory
QAV024121	EBVDNA21	C2	Qualitative and Quantitative	495139	2677/495139/3867	CZ023

EBVDNA21C2-03 - Quantitative Results Breakdown (IU/ml)

Sample Code	Sample Content	Matrix	Sample Relationships	Detection Frequency	Sample Status	Consens (IU/ml)	us	Range
						(Log ₁₀)	(n)	
EBVDNA21C2-03	Epstein-Barr virus	Plasma	D1, DS1_3	Frequently Detected	CORE	2.804	49	1.568 - 3.982



Groups below n=5: AB Analitica (n=2), AB Analitica - AB Analitica REALQUALITY RQ (n=2), Abbott (n=4), Abbott - Abbott RealTime rn2000 (n=4), Altona Diagnostics - Altona Diagnostics AltoStar (n=3), Diagenode (n=1), Diagenode - Diagenode Real Time kit (n=1), ELITech Group (n=1), ELITech Group - Elitech Elite Real Time kit (n=1), GeneProof (n=1), GeneProof - GeneProof Real Time PCR kit (n=1), Hong Kong CH Gene (n=1), Hong Kong CH Gene (n=1), QIAGEN (n=4), QIAGEN - QIAGEN Artus Real Time (n=3), QIAGEN - Qiagen NeuMoDx (n=1), Roche (n=4), Roche - Roche Cobas (n=4), Sacace (n=1), Sacace - Sacace Real TM (n=1), bioMerieux (n=4), bioMerieux - bioMerieux R-gene Quant Kit (n=4), In-House - Conventional In-House PCR (n=1)

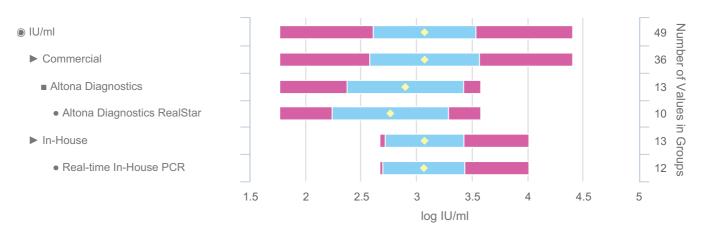
Groups Rolled Up:

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Catalogue Code:	Ref Code:	Challenge:	Analysis Type:	Dataset:	Report UID:	Laboratory	
QAV024121	EBVDNA21	C2	Qualitative and Quantitative	495139	2677/495139/3867	CZ023	

EBVDNA21C2-04 - Quantitative Results Breakdown (IU/ml)

Sample Code	Sample Content	Matrix	Sample Relationships	Detection Frequency	Sample Status	Consens (IU/ml)	us	Range
						(Log ₁₀)	(n)	
EBVDNA21C2-04	Epstein-Barr virus	Plasma	DS1_2	Frequently Detected	CORE	3.068	49	1.763 - 4.402



Groups below n=5: AB Analitica (n=2), AB Analitica - AB Analitica REALQUALITY RQ (n=2), Abbott (n=4), Abbott - Abbott RealTime rn2000 (n=4), Altona Diagnostics - Altona Diagnostics AltoStar (n=3), Diagenode (n=1), Diagenode - Diagenode Real Time kit (n=1), ELITech Group (n=1), ELITech Group - Elitech Elite Real Time kit (n=1), GeneProof (n=1), GeneProof - GeneProof Real Time PCR kit (n=1), Hong Kong CH Gene (n=1), Hong Kong CH Gene (n=1), QIAGEN (n=4), QIAGEN - QIAGEN Artus Real Time (n=3), QIAGEN - Qiagen NeuMoDx (n=1), Roche (n=4), Roche - Roche Cobas (n=4), Sacace (n=1), Sacace - Sacace Real TM (n=1), bioMerieux (n=4), bioMerieux - bioMerieux R-gene Quant Kit (n=4), In-House - Conventional In-House PCR (n=1)

Groups Rolled Up:

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		D 2021 Ep Programi	ostein-Barr virus DI ne	NA		
Catalogua Codo:	Pof Code	Challonger	Analysis Type	Datacati	Penert IIID:	Laboratory

Catalogue Code:	Ref Code:	Challenge:	Analysis Type:	Dataset:	Report UID:	Laboratory	
QAV024121	EBVDNA21	C2	Qualitative and Quantitative	495139	2677/495139/3867	CZ023	

EBVDNA21C2-01 - Qualitative Results Breakdown

Sample Code	Sample Content	Matrix	Sample Relationships	Detection Frequency	Sample Status	Percentage (All)	Correct
						(%)	(n)
EBVDNA21C2-01	Epstein-Barr virus	Plasma	DS1_1	Frequently Detected	CORE	96.9	96



Groups below n=5: AB Analitica (n=3), AB Analitica - AB Analitica REALQUALITY RQ (n=3), Abbott (n=2), Abbott - Abbott RealTime rn2000 (n=2), Altona Diagnostics - Altona Diagnostics AltoStar (n=3), AusDiagnostics (n=1), AusDiagnostics - AusDiagnostics Easy-Plex (n=1), Diagenode (n=1), Diagenode - Diagenode Real Time kit (n=1), Hong Kong CH Gene (n=1), Hong Kong CH Gene - HK CH Gene Real Time PCR (n=1), PrimerDesign (n=1), PrimerDesign Genesig (n=1), QIAGEN - Qiagen NeuMoDx (n=1), Roche (n=3), Roche - Roche Cobas (n=3), Sacace (n=1), Sacace - Sacace Real TM (n=1), Seegene (n=2), Seegene - Seegene Allplex (n=1), Seegene - Seegene Seeplex (n=1), fast-track DIAGNOSTICS (n=1), fast-track DIAGNOSTICS - FTD real time PCR (n=1), In-House - Conventional In-House PCR (n=2)

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Catalogue Code:	Ref Code:	Challenge:	Analysis Type:	Dataset:	Report UID:	Laboratory	
QAV024121	EBVDNA21	C2	Qualitative and Quantitative	495139	2677/495139/3867	CZ023	

EBVDNA21C2-02 - Qualitative Results Breakdown

Sample Code	Sample Content	Matrix	Sample Relationships	Detection Frequency	Sample Status	Percentage (All)	Correct
						(%)	(n)
EBVDNA21C2-02	Epstein-Barr virus	Plasma	D1, DS1_3	Detected	CORE	92.7	96



Groups below n=5: AB Analitica (n=3), AB Analitica - AB Analitica REALQUALITY RQ (n=3), Abbott (n=2), Abbott - Abbott RealTime rn2000 (n=2), Altona Diagnostics - Altona Diagnostics AltoStar (n=3), AusDiagnostics (n=1), AusDiagnostics - AusDiagnostics Easy-Plex (n=1), Diagenode (n=1), Diagenode - Diagenode Real Time kit (n=1), Hong Kong CH Gene (n=1), Hong Kong CH Gene - HK CH Gene Real Time PCR (n=1), PrimerDesign (n=1), PrimerDesign Genesig (n=1), QIAGEN - Qiagen NeuMoDx (n=1), Roche (n=3), Roche - Roche Cobas (n=3), Sacace (n=1), Sacace - Sacace Real TM (n=1), Seegene (n=2), Seegene - Seegene Allplex (n=1), Seegene - Seegene Seeplex (n=1), fast-track DIAGNOSTICS (n=1), fast-track DIAGNOSTICS - FTD real time PCR (n=1), In-House - Conventional In-House PCR (n=2)

Individual Report		D 2021 Ep Programr	ostein-Barr virus DN ne	Α		
Cotologua Codo	Def Cada	Challanger	Analysia Tyrnay	Detect	Benert UID	Loboratory

Catalogue Code:	Ref Code:	Challenge:	Analysis Type:	Dataset:	Report UID:	Laboratory	
QAV024121	EBVDNA21	C2	Qualitative and Quantitative	495139	2677/495139/3867	CZ023	

EBVDNA21C2-03 - Qualitative Results Breakdown

Sample Code	Sample Content	Matrix	Sample Relationships	Detection Frequency	Sample Status	Percentage (All)	Correct
						(%)	(n)
EBVDNA21C2-03	Epstein-Barr virus	Plasma	D1, DS1_3	Frequently Detected	CORE	95.8	96



Groups below n=5: AB Analitica (n=3), AB Analitica - AB Analitica REALQUALITY RQ (n=3), Abbott (n=2), Abbott - Abbott RealTime rn2000 (n=2), Altona Diagnostics - Altona Diagnostics AltoStar (n=3), AusDiagnostics (n=1), AusDiagnostics - AusDiagnostics Easy-Plex (n=1), Diagenode (n=1), Diagenode - Diagenode Real Time kit (n=1), Hong Kong CH Gene (n=1), Hong Kong CH Gene - HK CH Gene Real Time PCR (n=1), PrimerDesign (n=1), PrimerDesign Genesig (n=1), QIAGEN - Qiagen NeuMoDx (n=1), Roche (n=3), Roche - Roche Cobas (n=3), Sacace (n=1), Sacace - Sacace Real TM (n=1), Seegene (n=2), Seegene - Seegene Allplex (n=1), Seegene - Seegene Seeplex (n=1), fast-track DIAGNOSTICS (n=1), fast-track DIAGNOSTICS - FTD real time PCR (n=1), In-House - Conventional In-House PCR (n=2)

Individual Report		D 2021 Ep Programr	ostein-Barr virus DN ne	Α		
Catalogua Cada	Def Ceder	Challanger	Analysia Tyme	Detect	Depart LUD:	Laboratory

Catalogue Code:	Ref Code:	Challenge:	Analysis Type:	Dataset:	Report UID:	Laboratory	
QAV024121	EBVDNA21	C2	Qualitative and Quantitative	495139	2677/495139/3867	CZ023	

EBVDNA21C2-04 - Qualitative Results Breakdown

Sample Code	Sample Content	Matrix	Sample Relationships	Detection Frequency	Sample Status	Percentage (All)	Correct
						(%)	(n)
EBVDNA21C2-04	Epstein-Barr virus	Plasma	DS1_2	Frequently Detected	CORE	99.0	96



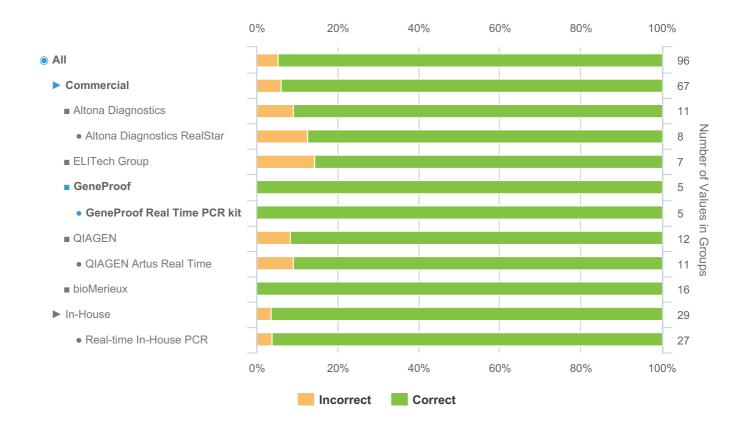
Groups below n=5: AB Analitica (n=3), AB Analitica - AB Analitica REALQUALITY RQ (n=3), Abbott (n=2), Abbott - Abbott RealTime rn2000 (n=2), Altona Diagnostics - Altona Diagnostics AltoStar (n=3), AusDiagnostics (n=1), AusDiagnostics - AusDiagnostics Easy-Plex (n=1), Diagenode (n=1), Diagenode - Diagenode Real Time kit (n=1), Hong Kong CH Gene (n=1), Hong Kong CH Gene - HK CH Gene Real Time PCR (n=1), PrimerDesign (n=1), PrimerDesign Genesig (n=1), QIAGEN - Qiagen NeuMoDx (n=1), Roche (n=3), Roche - Roche Cobas (n=3), Sacace (n=1), Sacace - Sacace Real TM (n=1), Seegene (n=2), Seegene - Seegene Allplex (n=1), Seegene - Seegene Seeplex (n=1), fast-track DIAGNOSTICS (n=1), fast-track DIAGNOSTICS - FTD real time PCR (n=1), In-House - Conventional In-House PCR (n=2)

Individual Report		CMD 2021 Epstein-Barr virus DNA QA Programme					
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Catalogue Code:	Ref Code:	Challenge:	Analysis Type:	Dataset:	Report UID:	Laboratory	
QAV024121	EBVDNA21	C2	Qualitative and Quantitative	495139	2677/495139/3867	CZ023	

EBVDNA21C2-05 - Qualitative Results Breakdown

Sample Code	Sample Content	Matrix	Sample Relationships	Detection Frequency	Sample Status	Percentage (All)	Correct
						(%)	(n)
EBVDNA21C2-05	Negative	Plasma		Negative	CORE	94.8	96



Groups below n=5: AB Analitica (n=3), AB Analitica - AB Analitica REALQUALITY RQ (n=3), Abbott (n=2), Abbott - Abbott RealTime rn2000 (n=2), Altona Diagnostics - Altona Diagnostics AltoStar (n=3), AusDiagnostics (n=1), AusDiagnostics - AusDiagnostics Easy-Plex (n=1), Diagenode (n=1), Diagenode - Diagenode Real Time kit (n=1), Hong Kong CH Gene (n=1), Hong Kong CH Gene - HK CH Gene Real Time PCR (n=1), PrimerDesign (n=1), PrimerDesign Genesig (n=1), QIAGEN - Qiagen NeuMoDx (n=1), Roche (n=3), Roche - Roche Cobas (n=3), Sacace (n=1), Sacace - Sacace Real TM (n=1), Seegene (n=2), Seegene - Seegene Allplex (n=1), Seegene - Seegene Seeplex (n=1), fast-track DIAGNOSTICS (n=1), fast-track DIAGNOSTICS - FTD real time PCR (n=1), In-House - Conventional In-House PCR (n=2)

Individual Report	QCMD 2021 Epstein-Barr virus DNA EQA Programme	QUALITY CONTrol for Malecular Diagnostic

Catalogue Code:	Ref Code:	Challenge:	Analysis Type:	Dataset:	Report UID:	Laboratory
QAV024121	EBVDNA21	C2	Qualitative and Quantitative	495139	2677/495139/3867	CZ023

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