### Intended Results / Panel Composition

<table>
<thead>
<tr>
<th>Sample Code</th>
<th>Sample Content</th>
<th>Matrix</th>
<th>Sample Relationships</th>
<th>Detection Frequency</th>
<th>Sample Status</th>
<th>Consensus (IU/ml)</th>
<th>Range (Log)</th>
<th>(n)</th>
</tr>
</thead>
<tbody>
<tr>
<td>HBVDNA17C1-01</td>
<td>HBV Type A</td>
<td>Plasma</td>
<td>D1</td>
<td>Frequently Detected</td>
<td>CORE</td>
<td>3.073</td>
<td>220</td>
<td></td>
</tr>
<tr>
<td>HBVDNA17C1-02</td>
<td>HBV Type D</td>
<td>Plasma</td>
<td>D1</td>
<td>Frequently Detected</td>
<td>CORE</td>
<td>3.810</td>
<td>221</td>
<td></td>
</tr>
<tr>
<td>HBVDNA17C1-03</td>
<td>HBV Type A</td>
<td>Plasma</td>
<td>D1</td>
<td>Frequently Detected</td>
<td>CORE</td>
<td>3.070</td>
<td>220</td>
<td></td>
</tr>
<tr>
<td>HBVDNA17C1-04</td>
<td>Negative Plasma</td>
<td>Plasma</td>
<td></td>
<td></td>
<td>Negative CORE</td>
<td>N/A</td>
<td>N/A</td>
<td></td>
</tr>
</tbody>
</table>

**Sample Relationships:** Indicates the relationships of the samples within this challenge. Dilution series are indicated by 'DS1' with each panel member in the dilution series represented by a number in order of titre, where DS1_1 represents the highest titre within that dilution series. Further dilution series are indicated by 'DS2' 'DS3' etc. If one duplicate pair is present this is indicated by 'D1'. Further duplicate pairs are indicated by 'D2', 'D3' etc.

**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.

**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.

**Consensus (IU/ml):** Mean consensus (Log_{10}) calculated from data returned by participants with outliers removed and number of quantitative results (n) returned for each panel member.

**Range:** Maximum and minimum quantitative value (IU/ml) reported by participants within this challenge without outliers removed.

---

### Your Summary Results

<table>
<thead>
<tr>
<th>Units</th>
<th>IU/ml</th>
</tr>
</thead>
<tbody>
<tr>
<td>EQA Assessment Group</td>
<td>GeneProof Real Time PCR kit</td>
</tr>
<tr>
<td>Core Panel Detection (Qualitative) Score</td>
<td>0</td>
</tr>
<tr>
<td>Core Panel Estimation (Quantitative) Score</td>
<td>2</td>
</tr>
</tbody>
</table>
### Sample Code

<table>
<thead>
<tr>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>HBVDNA17C1-01</td>
<td>IU/ml</td>
<td></td>
<td>3.267</td>
<td>0.167</td>
<td>3.029</td>
<td>99.2</td>
<td>Positive</td>
<td>0</td>
<td></td>
<td>Positive</td>
</tr>
<tr>
<td>HBVDNA17C1-02</td>
<td>IU/ml</td>
<td></td>
<td>4.040</td>
<td>0.187</td>
<td>3.957</td>
<td>100.0</td>
<td>Positive</td>
<td>0</td>
<td></td>
<td>Positive</td>
</tr>
<tr>
<td>HBVDNA17C1-03</td>
<td>IU/ml</td>
<td></td>
<td>3.270</td>
<td>0.153</td>
<td>3.053</td>
<td>99.2</td>
<td>Positive</td>
<td>0</td>
<td></td>
<td>Positive</td>
</tr>
<tr>
<td>HBVDNA17C1-04</td>
<td>IU/ml</td>
<td></td>
<td>N/A</td>
<td>-</td>
<td>LOD/NR</td>
<td>98.4</td>
<td>Negative</td>
<td>0</td>
<td></td>
<td>Negative</td>
</tr>
</tbody>
</table>

All quantitative values above expressed in Log_{10} IU/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 (All)**: Percentage of datasets reporting the correct qualitative results for each panel members.

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.*
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.
Duplicate Sample Performance Over Time

Observation Details
1: HBVDNA15C1-04 - HBVDNA15C1-03: Your Variation -0.034 (IU/ml): Overall Mean Variation 0.002: Overall SD 0.110.
2: HBVDNA15C3-02 - HBVDNA15C1-04: Your Variation 0.292 (IU/ml): Overall Mean Variation 0.013: Overall SD 0.210.
3: HBVDNA16C1-01 - HBVDNA15C3-02: Your Variation 0.774 (IU/ml): Overall Mean Variation 0.121: Overall SD 0.221.
4: HBVDNA16C1-03 - HBVDNA16C1-01: Your Variation 0.023 (IU/ml): Overall Mean Variation -0.001: Overall SD 0.105.
5: HBVDNA16C3-01 - HBVDNA16C1-03: Your Variation 0.118 (IU/ml): Overall Mean Variation 0.022: Overall SD 0.194.
6: HBVDNA17C1-01 - HBVDNA16C3-01: Your Variation -0.466 (IU/ml): Overall Mean Variation 0.013: Overall SD 0.179.
7: HBVDNA17C1-03 - HBVDNA17C1-01: Your Variation 0.024 (IU/ml): Overall Mean Variation -0.004: Overall SD 0.079.

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.

Individual Report
Catalogue Code: QAV994110
Ref Code: HBVDNA17
Challenge: C1
Analysis Type: Qualitative and Quantitative
Dataset: 130350
Report UID: 2677/130350/824
Participant: CZ023-01

QCMD 2017 Hepatitis B virus DNA EQA Programme

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

Page 4 of 14
Issue Date: 01 May 2017
Report authorised by the QCMD Executive (1)
A UKAS accredited proficiency testing provider No.4385
My Workflow Details

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

<table>
<thead>
<tr>
<th>Name</th>
<th>Hepatitis B Virus (HBV) PCR Kit (v2)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Description</td>
<td></td>
</tr>
<tr>
<td>Targets</td>
<td>hepatitis B virus</td>
</tr>
<tr>
<td>Assays</td>
<td></td>
</tr>
<tr>
<td>Extraction - Manual Extraction Process</td>
<td>Commercial</td>
</tr>
<tr>
<td></td>
<td>Kit Manufacturer: GeneProof</td>
</tr>
<tr>
<td></td>
<td>Kit Type: PathogenFree DNA Isolation Kit</td>
</tr>
<tr>
<td>Amplification - Shanghai Hongshi Medical Technology - SLAN</td>
<td>Commercial</td>
</tr>
<tr>
<td></td>
<td>Kit Manufacturer: GeneProof</td>
</tr>
<tr>
<td></td>
<td>Kit Type: Hepatitis B Virus (HBV) PCR Kit</td>
</tr>
</tbody>
</table>

Further Programme Details

Number of Participants: 264
Number of Countries: 33
Number of Respondents: 251
Number of Datasets Submitted: 266
Quantitative Results Returned (All): 229 (86.1%)
- Quantitative Results Returned (Copies/ml): 8 (3.5%)
- Quantitative Results Returned (IU/ml): 221 (96.5%)
Qualitative Results Returned: 128 (48.1%)

EQA Programme Aims

To assess the proficiency of laboratories in the detection and quantitation of hepatitis B virus (HBV).
To assess the proficiency of laboratories in the detection and quantitation in different HBV genotypes.

Feedback and Enquiries

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

Any queries about this report should be addressed to the QCMD Neutral Office (neutraloffice@qcmd.org).
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
<table>
<thead>
<tr>
<th>Sample Code</th>
<th>Sample Content</th>
<th>Matrix</th>
<th>Sample Relationships</th>
<th>Detection Frequency</th>
<th>Sample Status</th>
<th>Consensus (IU/ml)</th>
<th>Range</th>
</tr>
</thead>
<tbody>
<tr>
<td>HBVDNA17C1-01</td>
<td>HBV Type A</td>
<td>Plasma</td>
<td>D1</td>
<td>Frequently Detected</td>
<td>CORE</td>
<td>3.073</td>
<td>220</td>
</tr>
</tbody>
</table>

**Groups below n=5:** Roche - Roche Cobas Ampliscreen (n=1), Beckman Coulter (n=2), Beckman Coulter - Beckman Coulter Veris (n=2), LG Life Science (n=1), LG Life Science - LG Life Science AdvanSure (n=1), Sacace (n=1), Sacace - Sacace Real TM (n=1), RTA Laboratories (n=1), RTA Laboratories - RTA Laboratories Real time (n=1), Hong Kong CH Gene (n=1), Hong Kong CH Gene - HK CH Gene Real Time PCR (n=1), Iontek (n=1), Iontek - Iontek Fluorion (n=1), In-House - Conventional In-House PCR (n=1), In-House - Real-time In-House PCR (n=4)

**Groups Rolled Up:** Hologic - Hologic Aptima (n=7), Abbott - Abbott Realtime (n=50), Anatolia Geneworks - Anatolia Geneworks Bosphore (n=14), Siemens - Siemens Versant (n=7), Qiagen - Qiagen Real Time (n=15)
**QCMD 2017 Hepatitis B virus DNA EQA Programme**

**Catalogue Code:** QAV994110  
**Ref Code:** HBVDNA17  
**Challenge:** C1  
**Analysis Type:** Qualitative and Quantitative  
**Dataset:** 130350  
**Report UID:** 2677/130350/824  
**Participant:** CZ023-01

**HBVDNA17C1-02 - Quantitative Results Breakdown (IU/ml)**

<table>
<thead>
<tr>
<th>Sample Code</th>
<th>Sample Content</th>
<th>Matrix</th>
<th>Sample Relationships</th>
<th>Detection Frequency</th>
<th>Sample Status</th>
<th>Consensus (IU/ml)</th>
<th>Range</th>
</tr>
</thead>
<tbody>
<tr>
<td>HBVDNA17C1-02</td>
<td>HBV Type D</td>
<td>Plasma</td>
<td></td>
<td>Frequently Detected</td>
<td>CORE</td>
<td>3.810</td>
<td>2.598 - 4.413</td>
</tr>
</tbody>
</table>

- **IU/ml**
  - **Commercial**
    - Hologic
    - Abbott
    - Roche
  - **GeneProof**
    - GeneProof Real Time PCR kit
  - **QIAGEN**
  - **In-House**

**Groups below n=5:** Roche - Roche Cobas Ampliscreen (n=1), Beckman Coulter (n=2), Beckman Coulter - Beckman Coulter Veris (n=2), LG Life Science (n=1), LG Life Science - LG Life Science AdvanSure (n=1), Sacace (n=1), Sacace - Sacace Real TM (n=1), RTA Laboratories (n=1), RTA Laboratories - RTA Laboratories Real time (n=1), Hong Kong CH Gene (n=1), Hong Kong CH Gene - HK CH Gene Real Time PCR (n=1), Iontek (n=1), Iontek - Iontek Fluorion (n=1), In-House - Conventional In-House PCR (n=1), In-House - Real-time In-House PCR (n=4)

**Groups Rolled Up:** Hologic - Hologic Aptima (n=8), Abbott - Abbott Realtime (n=50), Anatolia Geneworks - Anatolia Geneworks Bosphore (n=14), Siemens - Siemens Versant (n=7), QIAGEN - Qiagen Artus Real Time (n=15)
### HBVDNA17C1-03 - Quantitative Results Breakdown (IU/ml)

<table>
<thead>
<tr>
<th>Sample Code</th>
<th>Sample Content</th>
<th>Matrix</th>
<th>Sample Relationships</th>
<th>Detection Frequency</th>
<th>Sample Status</th>
<th>Consensus (IU/ml)</th>
<th>Range</th>
</tr>
</thead>
<tbody>
<tr>
<td>HBVDNA17C1-03</td>
<td>HBV Type A</td>
<td>Plasma</td>
<td>D1</td>
<td>Frequently Detected</td>
<td>CORE</td>
<td>3.070</td>
<td>220</td>
</tr>
</tbody>
</table>

#### Groups below n=5:
- Roche - Roche Cobas Ampliscreen (n=1), Beckman Coulter - Beckman Coulter Veris (n=2), LG Life Science (n=1), LG Life Science - LG Life Science AdvanSure (n=1), Sacace (n=1), Sacace - Sacace Real TM (n=1), RTA Laboratories (n=1), RTA Laboratories - RTA Laboratories Real time (n=1), Hong Kong CH Gene (n=1), Hong Kong CH Gene - HK CH Gene Real Time PCR (n=1), Iontek (n=1), Iontek - Iontek Fluorion (n=1), In-House - Conventional In-House PCR (n=1), In-House - Real-time In-House PCR (n=4)

#### Groups Rolled Up:
- Hologic - Hologic Aptima (n=8), Abbott - Abbott Realtime (n=49), Anatolia Geneworks - Anatolia Geneworks Bosphore (n=14), Siemens - Siemens Versant (n=7), QIAGEN - Qiagen Artus Real Time (n=15)
Additional Information: 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.

**HBVDNA17C1-01 - Qualitative Results Breakdown**

<table>
<thead>
<tr>
<th>Sample Code</th>
<th>Sample Content</th>
<th>Matrix</th>
<th>Sample Relationships</th>
<th>Detection Frequency</th>
<th>Sample Status</th>
<th>Percentage Correct All (%)</th>
<th>n</th>
</tr>
</thead>
<tbody>
<tr>
<td>HBVDNA17C1-01</td>
<td>HBV Type A</td>
<td>Plasma</td>
<td>D1</td>
<td>Frequently Detected</td>
<td>CORE</td>
<td>99.2</td>
<td>128</td>
</tr>
</tbody>
</table>

- Commercial
  - Abbott
  - Roche
    - Roche Cobas Taqscreen
    - Roche Cobas Taqman
  - Grifols
  - Anatolia Geneworks
  - QIAGEN
- In-House
  - Real-time In-House PCR
**Groups below n=5:** Hologic (n=3), Hologic - Hologic Aptima (n=3), Sacace (n=1), Sacace - Sacace Real TM (n=1), GFE Blut (n=1), GFE Blut - GFE Blut VSPK (n=1), InterLabService (n=1), InterLabService - InterLabService AmpliSens (n=1), Hong Kong CH Gene (n=1), Hong Kong CH Gene - HK CH Gene Real Time PCR (n=1), Iontek (n=1), Iontek - Iontek Fluorion (n=1), Siemens (n=2), Siemens - Siemens Versant (n=2), GeneProof (n=4), GeneProof - GeneProof Real Time PCR kit (n=4), In-House - Conventional In-House PCR (n=2)

**Groups Rolled Up:** Abbott - Abbott Realtime (n=17), Grifols - Grifols Procleix Ultrio (n=14), Anatolia Geneworks - Anatolia Geneworks Bosphore (n=5), QIAGEN - Qiagen Artus Real Time (n=11)

### HBVDNA17C1-02 - Qualitative Results Breakdown

<table>
<thead>
<tr>
<th>Sample Code</th>
<th>Sample Content</th>
<th>Matrix</th>
<th>Sample Relationships</th>
<th>Detection Frequency</th>
<th>Sample Status</th>
<th>Percentage Correct All (%</th>
<th>(n)</th>
</tr>
</thead>
<tbody>
<tr>
<td>HBVDNA17C1-02</td>
<td>HBV Type D</td>
<td>Plasma</td>
<td>Frequently Detected</td>
<td>CORE</td>
<td>100.0</td>
<td>128</td>
<td></td>
</tr>
</tbody>
</table>

#### Number of Values in Groups

<table>
<thead>
<tr>
<th>Commercial</th>
<th>Incorrect</th>
<th>Correct</th>
</tr>
</thead>
<tbody>
<tr>
<td>Abbott</td>
<td>128</td>
<td>112</td>
</tr>
<tr>
<td>Roche</td>
<td>51</td>
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</tr>
<tr>
<td>Roche Cobas Taqscreen</td>
<td>14</td>
<td>14</td>
</tr>
<tr>
<td>Roche Cobas Taqman</td>
<td>5</td>
<td>5</td>
</tr>
<tr>
<td>Grifols</td>
<td>11</td>
<td>14</td>
</tr>
<tr>
<td>Anatolia Geneworks</td>
<td>16</td>
<td>16</td>
</tr>
<tr>
<td>QIAGEN</td>
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<td>14</td>
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<tr>
<td>In-House</td>
<td>114</td>
<td>0%</td>
</tr>
</tbody>
</table>

**Groups below n=5:** Hologic (n=3), Hologic - Hologic Aptima (n=3), Sacace (n=1), Sacace - Sacace Real TM (n=1), GFE Blut (n=1), GFE Blut - GFE Blut VSPK (n=1), InterLabService (n=1), InterLabService - InterLabService AmpliSens (n=1), Hong Kong CH Gene (n=1), Hong Kong CH Gene - HK CH Gene Real Time PCR (n=1), Iontek (n=1), Iontek - Iontek Fluorion (n=1), Siemens (n=2), Siemens - Siemens Versant (n=2), GeneProof (n=4), GeneProof - GeneProof Real Time PCR kit (n=4), In-House - Conventional In-House PCR (n=2)
### HBVDNA17C1-03 - Qualitative Results Breakdown

<table>
<thead>
<tr>
<th>Sample Code</th>
<th>Sample Content</th>
<th>Matrix</th>
<th>Sample Relationships</th>
<th>Detection Frequency</th>
<th>Sample Status</th>
<th>Percentage Correct All (%)</th>
<th>(n)</th>
</tr>
</thead>
<tbody>
<tr>
<td>HBVDNA17C1-03</td>
<td>HBV Type A</td>
<td>Plasma</td>
<td>D1</td>
<td>Frequently Detected</td>
<td>CORE</td>
<td>99.2</td>
<td>128</td>
</tr>
</tbody>
</table>

#### Number of Values in Groups

- Incorrect: 128
- Correct: 112

#### Percentage Correct All (%)

- All: 128
- Commercial: 17
- Abbott: 55
- Roche: 37
- Roche Cobas Taqscreen: 14
- Roche Cobas Taqman: 5
- Grifols: 14
- Anatolia Geneworks: 11
- QiAGEN: 16
- In-House: 14
- Real-time In-House PCR: 14

### Groups below n=5:

- Hologic (n=3), Hologic - Hologic Aptima (n=3), Sacace (n=1), Sacace - Sacace Real TM (n=1), GFE Blut (n=1), GFE Blut - GFE Blut VSPK (n=1), InterLabService (n=1), InterLabService - InterLabService AmpliSens (n=1), Hong Kong CH Gene (n=1), Hong Kong CH Gene - HK CH Gene Real Time PCR (n=1), Iontek (n=1), Iontek - Iontek Fluorion (n=1), Siemens (n=2), Siemens - Siemens Versant (n=2), GeneProof (n=4), GeneProof - GeneProof Real Time PCR kit (n=4), In-House - Conventional In-House PCR (n=2)

### Groups Rolled Up:

- Abbott - Abbott Realtime (n=17), Grifols - Grifols Procleix Ultrio (n=14), Anatolia Geneworks - Anatolia Geneworks Bosphore (n=5), QiAGEN - Qiagen Artus Real Time (n=11)

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**QCMD 2017 Hepatitis B virus DNA EQA Programme**

**Catalogue Code:** QAV994110  
**Ref Code:** HBVDNA17  
**Challenge:** C1  
**Analysis Type:** Qualitative and Quantitative  
**Dataset:** 130350  
**Report UID:** 2677/130350/824  
**Participant:** CZ023-01

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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  
Issue Date: 01 May 2017
HBVDNA17C1-04 - Qualitative Results Breakdown

<table>
<thead>
<tr>
<th>Sample Code</th>
<th>Sample Content</th>
<th>Matrix</th>
<th>Sample Relationships</th>
<th>Detection Frequency</th>
<th>Sample Status</th>
<th>Percentage Correct All (%</th>
<th>(n)</th>
</tr>
</thead>
<tbody>
<tr>
<td>HBVDNA17C1-04</td>
<td>Negative Plasma</td>
<td>Plasma</td>
<td></td>
<td>Negative</td>
<td>CORE</td>
<td>98.4</td>
<td>128</td>
</tr>
</tbody>
</table>

Groups below n=5: Hologic (n=3), Hologic - Hologic Aptima (n=3), Sacace (n=1), Sacace - Sacace Real TM (n=1), GFE Blut (n=1), GFE Blut - GFE Blut VSPK (n=1), InterLabService (n=1), InterLabService - InterLabService AmpliSens (n=1), Hong Kong CH Gene (n=1), Hong Kong CH Gene - HK CH Gene Real Time PCR (n=1), Iontek (n=1), Iontek - Iontek Fluorion (n=1), Siemens (n=2), Siemens - Siemens Versant (n=2), GeneProof (n=4), GeneProof - GeneProof Real Time PCR kit (n=4), In-House - Conventional In-House PCR (n=2)

Groups Rolled Up: Abbott - Abbott Realtime (n=17), Grifols - Grifols Procleix Utrio (n=14), Anatolia Geneworks - Anatolia Geneworks Bosphore (n=5), QIAGEN - QIAGEN Artus Real Time (n=11)
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