Results Interpretation Guide: Roche LightCycler® 480 Instrument I

Refer to Section 11 of the Instructions for Use (IFU 0009) for guidance in evaluating test results for run validity, interpretation of patient test results for a single PCR and Section 13, and for interpretation of results for patient samples.

1 Run Validity: Control Results
A. Record the name of the Test Run in the space “Test Run”.
B. Analyze PCR run according to Analysis with the Roche LightCycler 480 Instrument I in Section 11 of the IFU 0009.
C. Record “Well No.” and “CP” value or “Undetermined (UD)” for each of the 3 POSITIVE and 3 NEGATIVE Control replicates for Septin9 and ACTB (Internal Control) corresponding with PCR1, PCR2 and PCR3, included in the Test Run. If no CP value is obtained, the Control result is “Undetermined” (meaning no curve was generated).
D. Evaluate CP value based on TABLE 11: Validity Limits. Check “Pass” or “Fail”.
E. Confirm if VALID results were obtained for both POSITIVE and NEGATIVE Controls before proceeding to patient test results interpretation. Epi proColon POSITIVE and NEGATIVE Controls are considered VALID when all criteria in TABLE 11 are met for all three replicates per Control and both detectors (ACTB and Septin9). If both Controls are VALID, continue to Step 2, Patient Test Results: Interpretation of Results for a Single PCR. If either or both Controls are INVALID, the data for patient samples processed together with the Controls cannot be interpreted. Testing must be repeated for all patient samples included in this run.

2 Patient Test Results: Interpretation of Results for a Single PCR
A. Record the “Patient Name or ID No.” in the space “Patient ID”.
B. Record “Well No.” and “CP” values, or “Undetermined (UD)” for each patient samples for Septin9 and ACTB (Internal Control) corresponding with PCR1, PCR2 and PCR3, included in the Test Run. If no CP value is obtained, the result is “Undetermined” (meaning no curve was generated).
C. Evaluate CP value based on TABLE 12: Interpretation of Results for a Single PCR, and check appropriate box, INVALID, POSITIVE or NEGATIVE.
D. Continue to Step 3, Interpretation of Results for a Patient Sample.

3 Interpretation of Results for a Patient Sample
Refer to Section 13, Interpretation of Results for a Patient Sample, TABLE 16: Interpretation of Epi proColon 2.0 CE Test Results, for interpretation of patient test results.

POSITIVE Result: At least 2 of 3 POSITIVE Septin9 PCR replicates
NEGATIVE Result: At least 2 of 3 NEGATIVE Septin9 PCR replicates
INVALID Result: The test is “INVALID” in all other cases

NOTE: When no curve is generated, “Undetermined” is reported. If the Septin9 channel is Undetermined or “UD”, that specific well is Septin9 NEGATIVE. If the ACTB channel is Undetermined or “UD”, that specific well is NEGATIVE for ACTB and therefore, INVALID.
The Epi proColon® 2.0 CE Test Result Interpretation Forms

For use with the LightCycler® 480 Instrument I

There are two forms for recording and interpreting the Epi proColon 2.0 CE test results. Form 1 (below) includes areas for recording Control Results and results for two patients. Every Epi proColon Real-Time PCR test run requires that POSITIVE and NEGATIVE Controls be performed with patient samples. At the top of each Results Form, a space is provided to record the Run Name, Run Date, and the initials of the Technician performing the testing. Form 2 (below) offers space for three additional patients, and this page may be duplicated as needed.

Control results must be VALID before patient results are interpreted; patient results must be POSITIVE or NEGATIVE before being reported; INVALID patient test results should not be reported.

### Form 1

<table>
<thead>
<tr>
<th>Well No.</th>
<th>PCR Replicate</th>
<th>Detector</th>
<th>Acceptable Results Criteria</th>
<th>Single PCR Result</th>
<th>Final Patient Result</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>PCR1</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>PCR2</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>PCR3</td>
<td>Oct</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

### Form 2

<table>
<thead>
<tr>
<th>Well No.</th>
<th>PCR Replicate</th>
<th>Detector</th>
<th>Acceptable Results Criteria</th>
<th>Single PCR Result</th>
<th>Final Patient Result</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>PCR1</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>PCR2</td>
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<td></td>
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<td></td>
</tr>
<tr>
<td></td>
<td>PCR3</td>
<td>Oct</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

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**Epi proColon® 2.0 CE Test**

FOR CLINICAL LABORATORIES
Not For Use in the United States

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