Use and calibration of StepOnePlus™ Real-Time PCR System

1. INTRODUCTION

1.1 This method is based on the Applied Biosystems StepOnePlus™ Real-Time PCR manual. It is used to maintain the PCR machine by performing spatial and dye calibration. The method also describes the general steps for using the StepOnePlus™ Real-Time PCR System to set up a qPCR run.

2. EQUIPMENT AND SUPPLIES

2.1 StepOnePlusTM Real-Time PCR System (S/N 272006852) – PCR #1

2.2 StepOnePlusTM Real-Time PCR System (S/N 272008854) – PCR #3

2.3 CBS Optimizer PCR Workstation #1 (S/N 007631)

2.4 CBS Optimizer PCR Workstation #2 (S/N 007632)

2.5 Optical 96 well PCR reaction tray

2.6 Optical adhesive PCR reaction tray tape

2.7 Optical 96 well plate holder and stand

2.8 StepOnePlusTM Real-Rime PCR system Spectral Calibration kit (P/N 4371435)

2.9 Safety glasses

2.10 Fisher Mini Vortexes

2.11 Gloves, latex or nitriles

2.12 Tube racks

2.13 Micropipettes with sterile PCR grade filter tips

3. REAGENTS AND STANDARDS

3.1 Deionized water

3.2 PCR grade water – Autoclave deionized water for 15 min. Record in LIMS Autoclave log.

4 SETUP AND USE OF STEPONE PLUS INSTRUMENT FILE

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4.1 Select the corresponding test ID folder in \\DEAR(X:)\Labs\Lab3Tlhlab3\Biology\qPCR\Import files and open a recent text file. “Save as” the file with the current worksheet number.

4.2 Update the SPK and LFB serial numbers using the “Replace” tool if needed.

4.3 Use Edit/Replace from the menu to replace “Sample1” with the first sample ID. Do this with all the samples and refer to the plate layout so that the right sample IDs go into the correct wells.

4.4 Check the naming conventions for the DUP samples. For example, the spikes should be SampID_SPK_DUP_A_SPK#. When running purified DNA, samples should be named SampID_PUR. The spikes should be sampleID_SPK_PUR_SPKserial#. The extraction control should be named LFB_LFBPserial#.

4.5 Save the text file.

4.6 Open StepOne software and click “Advanced Setup” from the drop-down menu under New Experiment.

4.6.1 Under Experiment Properties, add the corresponding Experiment Name (XX-X-##-YYMMDD). Where XX corresponds to test ID (HF, BR, G2, GD, D3 or C2); X refers to the DNA extraction method (C for crude and P for purified); ## corresponds to the worksheet number and YYMMDD specifies the date of the run).

4.6.2 Type the user name initials and the instrument used (PCR1 or PCR3) in the comments box.

4.6.3 Make sure the experiment selections are correct (the defaults are 96 wells, Quantitation-Standard Curve, TaqMan® Reagents). For the ramp speed, click Standard-2 hours (Not Default).

4.7 Select File/Import from the menu, choose the corresponding text file from the Import files folder and click Import.

4.8 Click the Plate Setup tab and under Assign Target and Samples, make sure the samples and controls are in the correct wells. Delete the information from empty wells by un-clicking the assign target and the sample ID.

4.9 Save the file with the name assigned when setting up the experiment (See 4.6.1) in \\DEAR(X:)\Labs\Lab3 \\Tlhlab3\Biology\qPCR\Instrument Files.

4.10 The file is now ready to be used for the run. Load the plate into the instrument. Click on the Plate Set Up tab and start the run by clicking the green “Start Run” button in the upper right-hand corner of the screen.

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4.11 Run progress can be viewed from the touch screen of the instrument. At the beginning of a run do not leave the instrument or computer until you verify that the run has started.

4.12 When the run is complete, make sure to save the file on the thermocycler PC, so that the run information is saved correctly. Remove the reaction plate and discard.

5 QPCR RUN ANALYSIS AND EXPORTING FILE

5.1 In the instrument file, review the standard curve for both assays. Omit individual wells as needed. Efficiency should be 80-120% and R^2 should be ≥ 0.99. If the efficiency is below 70%, the R^2 is below 0.97, or the standard curves do not amplify properly the entire plate needs to be re-run. Compare the 3rd well spikes and SKETA assay controls with the samples to check for inhibition.

5.2 Select “Print Report” and print the Experiment Summary, Plate Layout, Standard Curves, and QC Summary for the data packet.

5.3 Select “Export”. In the Export Properties tab, select the following: Check “Results” only (default setting). Select “One File” from the drop-down menu (default setting).

5.6 Make sure the Export File Name is correct and delete _data from the file name. If samples from more than one worksheet are run together, the data need to be exported independently. Select File Type: “.txt” and Export File Location: \\Fldep1\dear\Labs\Lab3\Biology\qPCR\Export Files. Select the folder with the appropriate year and test ID.

5.7 Select “Start Export”, then select “Close Export Tool”.

6 MAINTENANCE OF STEP ONE PLUS™ REAL-TIME PCR SYSTEM

6.1 Perform a Spatial Calibration

6.1.1 The spatial calibration maps the positions of the wells on the sample block so that the StepOne™ software can associate increases in fluorescence during a run with specific wells of the reaction plate. The spatial calibration is performed every 18 months, or as often as necessary, depending on instrument use.

6.1.2 Make sure that the yellow Ethernet port of the instrument is connected to the Ethernet port of the computer running the software.

6.1.3 Get calibration plate 1 from the spectral calibration kit in the freezer.

6.1.4 Allow calibration plate 1 to warm to room temperature (approximately 5 min).

NOTE: Do not remove the calibration plate 1 from its packaging until ready to run. The fluorescent dyes in the wells of spectral calibration plates are photosensitive. Prolonged exposure to light can diminish the fluorescence signal strength of the dyes.

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6.1.5 Remove the calibration plate 1 from its packaging.

6.1.6 Vortex calibration plate 1 for 5 sec.

6.1.7 Centrifuge calibration plate 1 for 2 min at less than 1500 rpm.

NOTE: The calibration plate must be well mixed and centrifuged.

6.1.8 Check that all the liquid is at the bottom of each well. If not, centrifuge the calibration plate again at a higher rpm for a longer time.

NOTE: Ensure that the bottom of the calibration plate does not become dirty. This can contaminate the sample block and cause an abnormally high background signal.

6.1.9 Load calibration plate 1 into the instrument.

6.1.9.1 On the main screen of the StepOnePlus™ software, select Instrument —> Instrument Maintenance Manager.

6.1.9.2 In the navigation pane of the Instrument Maintenance Manager, click Spatial.

6.1.9.3 In the Instrument Maintenance Manager, click Start Calibration.

6.1.9.4 In the setup screen of the Spatial Calibration dialog box, select Start Calibration.

6.1.9.5 In the setup screen of the Spatial Calibration dialog box, select “The calibration plate is loaded into the instrument”, then click Next.

6.1.9.6 In the run screen, click Start Run.

6.1.10 To analyze the calibration data, in the analysis screen of the spatial calibration dialog box confirm the status of the calibration:

6.1.10.1. Passed: The instrument passed the calibration. Click Finish and save the calibration run by clicking Yes.

6.1.10.2. Failed: The instrument failed the calibration. Contact Applied Biosystems.

6.2 Perform a Dye Calibration

6.2.1 The dye calibration collects spectral data from a series of dye standards and distinguishes the individual contributions of each dye in the collective fluorescence collected by the instrument during a run. The dye calibration is performed every 18 months, or as often as necessary, depending on instrument use.

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6.2.2 Make sure that the yellow Ethernet port of the instrument is connected to the Ethernet port of the computer running the software.

6.2.3 Get calibration plate 1 from the spectral calibration kit in the freezer.

6.2.4 Allow calibration plate 1 to warm to room temperature.

NOTE: Do not remove the calibration plate 1 from its packaging until ready to run. The fluorescent dyes in the wells of spectral calibration plates are photosensitive. Prolonged exposure to light can diminish the fluorescence signal strength of the dyes.

6.2.5 Remove the calibration plate 1 from its packaging.

6.2.6 Vortex calibration plate 1 for 5 sec.

6.2.7 Centrifuge calibration plate 1 for 2 min at less than 1500 rpm.

NOTE: The calibration plate must be well mixed and centrifuged.

6.2.8 Check that all the liquid is at the bottom of each well of calibration plate 1. If not, centrifuge the calibration plate again at a higher rpm for a longer time.

NOTE: Ensure that the bottom of calibration plate does not become dirty. This can contaminate the sample block and cause an abnormally high background signal.

6.2.9 Load calibration plate 1 into the instrument.

6.2.9.1 On the main screen of the StepOnePlus™ software, select Instrument —> Instrument Maintenance Manager.

6.2.9.2 In the navigation pane of the Instrument Maintenance Manager, click Dye.

6.2.9.3 In the Dye screen, select System Dye Calibration.

6.2.9.4 Select Start Calibration.

6.2.9.5 In the setup screen, select “Plate 1 is loaded into the instrument”, then click Next.

6.2.9.6 In the run screen, click Start Run.

6.2.9.7 While the instrument is running calibration plate 1, prepare calibration plate 2 by repeating steps 6.2.5–6.2.8.

6.2.9.8 When Calibration plate 1 is done, the wizard will prompt you to load and run calibration plate 2.

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6.2.9.9 In the setup screen, select “Plate 2 is loaded in the instrument”, then click Continue Calibration.

6.2.10 To analyze the calibration data, in the analysis screen of the dye calibration dialog box confirm the status of the calibration:

6.2.10.1 Passed: The instrument passed the calibration. Continue with step 6.2.10.3.

6.2.10.2 Failed: The instrument failed the calibration. Troubleshoot the error as described in the “Dye Calibration Failure” of the Applied Biosystems StepOnePlus™ Installation, Networking and Maintenance manual.

6.2.10.3 For each dye on the plate, confirm the grouping of the dye spectra (refer to the diagrams in the Maintain the System chapter of the Applied Biosystems StepOnePlus™ Installation, Networking and Maintenance manual).

6.2.10.3.1 In the plate layout, press Ctrl, and select the wells that contain the same dye.

6.2.10.3.2 Inspect the raw data for each spectrum, confirming that the peak is within the detectable range of the instrument, free of irregular spectral peaks and presents the correct channel for the dye.

6.3 Perform a Background Calibration

6.3.1 This calibration measures the level of background fluorescence in the instrument by continuously reading a background plate containing PCR buffer for 10 min at 60°C. The background calibration should be performed every three months or as often as necessary. The software averages the spectra recorded during the run and extracts it into a calibration file. This file is used by the StepOne software during subsequent runs to remove the background fluorescence from the run data.

6.3.2 Get the background plate from the spectral calibration kit in the freezer.

Note: A new background plate can be created following the manual instructions on page 103.

6.3.3 Allow the background plate to warm to room temperature (at least 5 min).

6.3.4 Remove the background plate from its packaging.

Note: Keep the packing sleeve. The background plate can be used up to three times if it is stored in its original packing sleeve.

6.3.5 Vortex the background plate for 5 seconds

6.3.6 Centrifuge the background plate for 2 min at less than 1500 rpm.

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6.3.7 Make sure the liquid is at the bottom of each well of the background plate. If not, centrifuge at a greater rpm and for a longer time.

Note: Ensure that the bottom of the calibration plate does not become dirty. This can contaminate the sample block and cause an abnormally high background signal.

6.3.8 Load the background plate into the instrument.

6.3.8.1 On the main screen of the StepOnePlus™ software, select Instrument —> Instrument Maintenance Manager.

6.3.8.2 In the navigation pane of the Instrument Maintenance Manager, click Background

6.3.8.3 Click Start Calibration.

6.3.8.4 In the Setup screen of the Background Calibration dialog box, select "The background plate is loaded into the instrument", then click Next.

6.3.8.5 In the Run screen, click START RUN.

6.3.9 To analyze the calibration data, in the Background Calibration dialog box confirm the status of the calibration:

6.3.9.1 Passed- The instrument passed the calibration. Go to step 6.3.9.3.

6.3.9.2 Failed- The instrument failed the calibration. Troubleshoot the errors following the manual's suggestions on page 138.

6.3.9.3 Check the raw data for spectral peaks that diverge significantly from that of the neighboring cells. This indicates fluorescent contamination and the irregular signal needs to be troubleshooted following the manufacturer manual suggestions on page 138.

6.3.10 If all the spectra are acceptable, click **Finish.**

6.3.11 When prompted to save the background calibration, click **Yes**.

7 QUALITY CONTROL

7.1 Update quality control laboratory manual to record qPCR instrument maintenance.

8 REFERENCES

8.1 DEP Laboratory Quality Manual
http://publicfiles.dep.state.fl.us/dear/labs/lab_qualitymanual_18.pdf

8.2 Health and Safety Plan Health and Safety Plan

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8.3 Applied Biosystems StepOne™ and StepOnePlus™, Real-Time PCR Systems, Installation, Networking, and Maintenance manual

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