

**PathoSEEK[®] *Pseudomonas aeruginosa* and *Staphylococcus aureus*
Assay with SenSATIVAx[®] Purification**

User Guide

Real-Time PCR (qPCR) Assay for the detection of *P. aeruginosa* and *S. aureus* in dried cannabis flower and MIP matrices

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Introduction

Some regulations require cannabis flower and cannabis products to be free of *Pseudomonas aeruginosa* and *Staphylococcus aureus*. There are three different PathoSEEK® qPCR Detection Assays that can be used to detect these species in cannabis flower, concentrates, infused edibles, and infused non-edibles, and this user guide will provide instructions for the following:

- PathoSEEK® *Staphylococcus* & *Pseudomonas* Multiplex Detection Assay
- PathoSEEK® *Staphylococcus aureus* Detection Assay
- PathoSEEK® *Pseudomonas aeruginosa* Detection Assay

Process Overview

The PathoSEEK® *Pseudomonas aeruginosa* and *Staphylococcus aureus* Multiplex Detection Assay uses a multiplexing strategy with an internal cannabis DNA control to ensure accurate detection of *Pseudomonas aeruginosa* and *Staphylococcus aureus* as well as cannabis DNA in a single reaction. The PathoSEEK® *Staphylococcus aureus* Detection Assay and PathoSEEK® *Pseudomonas aeruginosa* Detection Assay also use multiplexing to detect their respective microbial targets and the internal cannabis control. Unlike other techniques, this multiplexing strategy verifies the performance of the assay when detecting pathogens, resulting in the minimization of false negative results due to reaction set-up errors or failing experimental conditions. Below is a simplified depiction of the qPCR assays.

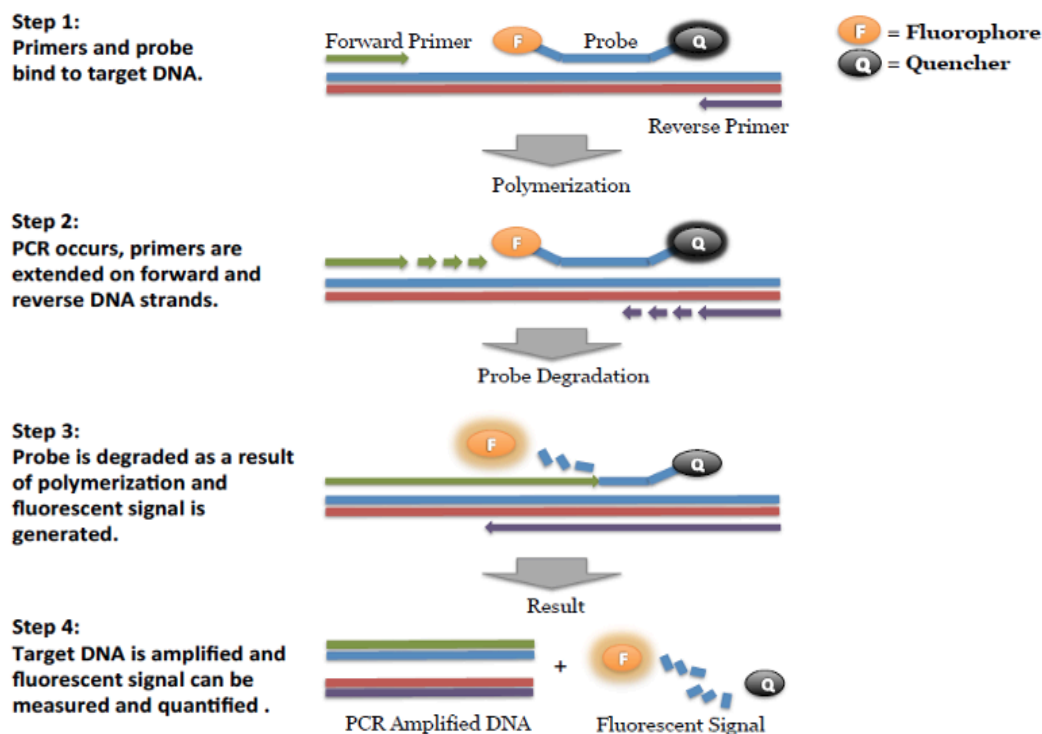


Figure 1: Overview of qPCR

The PathoSEEK Detection Assays can be used on the AriaMx Real-Time PCR Thermocycler (Agilent) or the CFX-96 (Bio-Rad). All three assays use the HEX Fluorophore for detection of cannabis DNA and the FAM Fluorophore for detection of *Pseudomonas aeruginosa* and/or *Staphylococcus aureus*.

SenSATIVAx is a proprietary DNA purification process that uses magnetic particles to isolate and purify both plant and microbial DNA from a raw, homogenized plant or MIP sample. The use of magnetic particles affords 8 or 96-tip automation, enabling high-throughput applications. DNA can be isolated from a single sample or a large batch in under 1 hour. Hands-on time is less than 45 minutes.

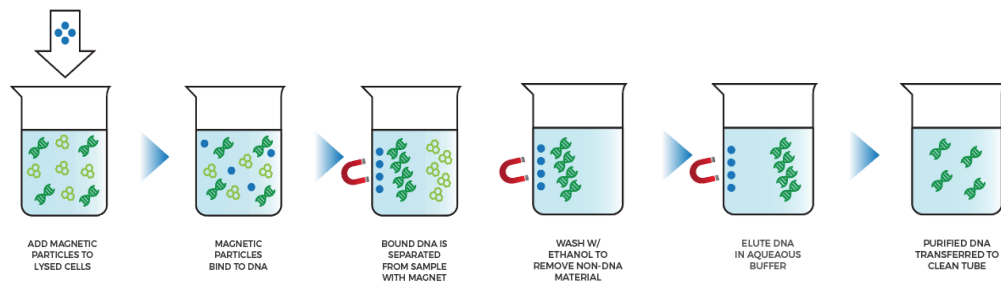


Figure 2: SenSATIVAx® Purification Protocol

Limit of Detection

The method is able to detect down to 1 CFU per test portion.

Materials and Methods

Test Kit Name: PathoSEEK® *Pseudomonas aeruginosa* and *Staphylococcus aureus* Assay with SenSATIVAx® DNA Purification

Test Kit Information

1. PathoSEEK *Staphylococcus* and *Pseudomonas* Multiplex Detection Assay - P/N 420138
2. PathoSEEK *Staphylococcus aureus* Detection Assay - P/N 420136
3. PathoSEEK *Pseudomonas aeruginosa* Detection Assay - P/N 420137
4. SenSATIVAx Flower/Leaf DNA Purification Kit - P/N 420001
5. SenSATIVAx MIP/Extract DNA Purification Kit - P/N 420004
6. PathoSEEK Staph and Pseudomonas Multiplex Positive Control - P/N 420338
7. PathoSEEK qPCR Master Kit v3 - P/N 420201

Test Kit Components

1. PathoSEEK *Staph* and *Pseudomonas* Multiplex Detection Assay — P/N 420138 (200 tests)
 - a. Assay — 1 tube. Store kit at -15 to -20°C. Expires 2 Years from Date of Manufacture
2. PathoSEEK *Staphylococcus aureus* Detection Assay — P/N 420136 (200 tests)
 - a. Assay — 1 tube. Store kit at -15 to -20°C. Expires 2 Years from Date of Manufacture
3. PathoSEEK *Pseudomonas aeruginosa* Detection Assay — P/N 420137 (200 tests)
 - a. Assay — 1 tube. Store kit at -15 to -20°C. Expires 2 Years from Date of Manufacture
4. SenSATIVAx Flower & Leaf DNA Purification Kit — P/N 420001 (200 tests)
 - a. MGC Lysis Buffer — 1 bottle. Store at 20-28°C. Expires 1 Year from Date of Manufacture
 - b. MGC Binding Buffer — 1 bottle. Store at 2-8°C. Expires 1 Year from Date of Manufacture
 - c. MGC Elution Buffer — 1 bottle. Store at 20-28°C. Expires 1 Year from Date of Manufacture
5. SenSATIVAx MIP DNA Purification Kit — P/N 420004 (200 tests).
 - a. Binding Buffer — 1 bottle. Store at 2-8°C. Expires 1 Year from Date of Manufacture
 - b. Elution Buffer — 1 bottle. Store at 20-28°C. Expires 1 Year from Date of Manufacture
 - c. Solution A — 1 bottle. Store at 20-28°C. Expires 1 Year from Date of Manufacture
 - d. Solution B — 1 bottle. Store at 20-28°C. Expires 1 Year from Date of Manufacture
6. PathoSEEK *Staph* and *Pseudomonas* Multiplex Control — P/N 420338 (50 reactions)
 - a. Control - 1 tube. Store at -15 to -20°C. Expires 2 Years from Date of Manufacture

7. PathoSEEK qPCR Master Kit v3— P/N 420201 Kit. (Store at -15 to -20°C) Expires 2 Years from

Date of Manufacture

- a. Reaction Buffer (10x).— 1 tube
- b. Nuclease Free Water.—2 tubes
- c. qPCR Master Mix - 1 tube

Supplies, Reagents, and Equipment

1. Agilent AriaMx Real-Time PCR System G8830A—Containing the following Optical Channels:
FAM and HEX

- a. Agilent HP Notebook PC option 650
- b. 96-Well Optical qPCR plate - Medicinal Genomics P/N 100164
- c. Adhesive optical seal for qPCR plates - Medicinal Genomics P/N 100177
- d. Optical Strip Caps.—Agilent #401425

Note: If using adhesive seals instead of strip caps, use Applied Biosystems MicroAmp Optical Film Compression Pad, Fisher Scientific, #43-126-39 to prevent evaporation and cross contamination between wells. Medicinal Genomics will provide at no charge if AriaMx is purchased through Medicinal Genomics.

2. Bio-Rad CFX96 Touch™ Real-Time System.

- a. Bio-Rad Personal PC (Optional)
- b. 96-well Optical qPCR plate — Medicinal Genomics P/N 100164
- c. Adhesive optical seal for qPCR plates — Medicinal Genomics P/N 100177

3. Adjustable, variable volume pipettes (single or multichannel)—P10, P20, P50, P200 or P300 and P1000

4. Adjustable, variable volume filter pipettes tips—For P10, P20, P50, P200, P300 and P1000
5. Crushed ice
6. 96-well PCR Cryogenic Rack (optional) —VWR #89004-570
7. 1.5 mL Tube Benchtop Cryogenic Rack (if no ice)— VWR #89004-558 or equivalent
8. Freezer—Capable of maintaining -20°C
9. Tabletop Mini Plate Centrifuge—Fisher Scientific #14-100-143 or equivalent
10. Tabletop Mini Centrifuge—VWR #10067-588, #2631-0006 or equivalent
11. Vortex-Genie Pulse—Scientific Industries, SKU: SI-0236 or equivalent
12. High-Speed centrifuge— to accommodate 1.5mL tubes such as Eppendorf model 5414R or similar with ability to spin up to speeds of 14,000 - 19,300 RCF
13. Incubator—Capable of maintaining 37°C (VWR® Personal Size Incubator # 97025-630, or similar)
14. Filter Bags—Whirl Pak #B01385WA
15. Beaker or Solo Cup(optional)
16. Tryptic Soy Broth—Medicinal Genomics #420205 (Store at 2-8°C)
17. 1.5 mL Eppendorf Tubes
18. 15 mL or 50 mL conical tubes (for MIP)
19. 96-well Plate Magnet—Medicinal Genomics P/N 420202
20. 96-well Extraction Plate— Medicinal Genomics P/N 100298
21. Eppendorf Tube Rack
22. Scientific Scale—Capable of measuring to milligrams
23. Refrigerator—Capable of maintaining 2–8°C

24. 25 mL Serological Pipette —VWR 89130-890 or 89130-900 or equivalent.
25. 10% bleach.
26. 70% Ethanol — MGC P/N 420030
27. Chloroform (Ethanol as preservative/ Certified ACS) - Fisher Scientific, C298

Safety Precautions and Recommendations for Best Results

Safety Precautions

The PathoSEEK® *Pseudomonas aeruginosa* and *Staphylococcus aureus* Assay is a qPCR detection assay for detection of *Pseudomonas aeruginosa* and *Staphylococcus aureus* in cannabis matrices.

1. Assay users should observe standard microbiological practices and safety precautions when performing this assay. Wear protective gloves, lab coats, eye/face protection as indicated by your quality system.
2. It is the responsibility of each laboratory to handle waste and effluents processed according to their nature and degree of hazardousness. Waste and effluents processed must be treated and disposed of in accordance with all applicable local, state, and federal regulations.
3. Hazard Statement: 70% Ethanol
 - a. Highly flammable liquid and vapor. May cause respiratory irritation.
 - b. May cause drowsiness or dizziness. Causes damage to organs.
 - c. May cause damage to organs through prolonged or repeated exposure.
 - d. Please refer to the Safety Data Sheet (SDS) for more information and proper disposal.



4. Hazard Statement: Chloroform
 - a. Harmful if inhaled or swallowed.
 - b. Do not breathe vapor or mist. Do not ingest. Avoid contact with eyes, skin and clothing.Use only with adequate ventilation, which may require a chemical fume hood.

- c. Keep the container tightly closed and sealed until ready for use. Wash thoroughly after handling.
- d. Please refer to the Manufacturer Safety Data Sheet (SDS) for more information and proper disposal



Environment

1. The quality of results depends on the strict compliance with Good Laboratory Practices (for example, the EN ISO 7218 standard), especially concerning PCR.
 - a. Never circulate lab equipment from one workstation to another.
 - b. Always use a positive and negative control for each series of amplification reactions.
 - c. Periodically verify the accuracy and precision of pipettes, as well as correct functioning of the instruments.
 - d. Change gloves often, especially if you suspect contamination.
 - e. Clean workspaces periodically with 10% bleach and other decontaminating agents.
 - f. Use powder-free gloves.
 - g. If using qPCR reaction strip tubes instead of plates, avoid fingerprints and writing on caps because both can interfere with data acquisition.

Intended User

The PathoSEEK® *Pseudomonas aeruginosa* and *Staphylococcus aureus* Assays with SenSATIVAx® Purification and all components required to perform the method are intended for use by trained personnel familiar with laboratory techniques associated with pathogenic organism detection.

Sample Preparation

1. Aliquot Tryptic Soy Broth (TSB).

Note: TSB is a very good growth medium for microbes. It is best to pour the approximate amount of TSB into another sterile tube or container to avoid contaminating the whole bottle. Inspect stock of TSB for flocculants or signs of growth prior to aliquoting. Return it to the 2-8°C refrigerator immediately after use.

2. Wipe down the workspace with a 10% bleach solution, including the benchtop and all equipment being used.
3. Remove the MGC Binding Buffer and TSB from the 2-8°C refrigerator (it should come to room temperature, 20-28°C, before use).
4. Prepare consumables. Label all the filter bags with “[sample name] [date]”.
5. Prepare consumables. Label all the 1.5 mL centrifuge tubes needed with [sample name].
6. Label extraction plate with date, and if transferring eluted DNA to new plate label the destination plate also.
7. Before weighing out the sample to be tested, make sure that the entire sample is broken up and thoroughly homogenized. A well-homogenized sample will ensure more accurate testing.
8. *Cannabis flower, n grams* —Weigh flower sample material into one side of the mesh liner inside the Whirl-Pak bag. Add $9 \times n$ mL of TSB to each test portion. Close the Filter bag by folding the top over three times. Mix for 1 minute by hand. Incubate for 16-24 h at $37 \pm 1^\circ\text{C}$.
9. *MIP and concentrates, n grams*. — Weigh the MIP matrix into a 15 mL conical tube, 50 mL conical tube, or Whirl-Pak bag depending on MIP volume. Add $2.4 \times n$ mL of TSB to each test portion. Vortex or homogenize sample and TSB. Incubate for 16-24 h at $37 \pm 1^\circ\text{C}$.

10. If processing multiple samples, be sure to change gloves between each, to ensure there is no cross contamination between samples during the weighing process.

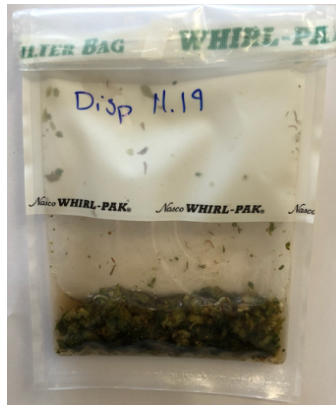


Figure 3: Homogenized cannabis flower and TSB.

DNA Purification

SenSATIVAx® for Flower/Leaf DNA Purification

1. Remove enriched samples from the incubator and hand homogenize again for 1 minute.
2. Aspirate **1 mL** from the side of the filter bag, free of plant debris, and dispense into the 1.5mL tube.
3. Add 50 μ L of MGC Lysis buffer and vortex for 10 seconds then let incubate on the bench for 2 minutes.
4. After 2 minute incubation spin for at least 1-3 minutes in a bench top mini centrifuge.

Note: The supernatant should be translucent at this point. If the sample is still opaque (cloudy) spin for longer. This is important for removing cellular debris. See Figure 4.

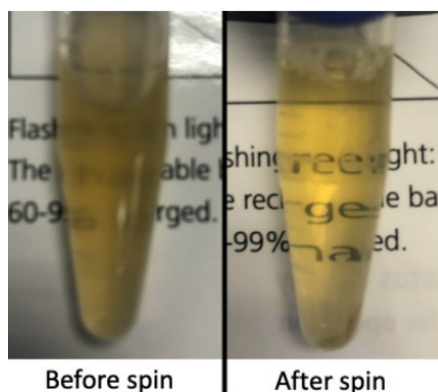


Figure 4: Example of translucent lysate after spinning.

5. Remove the 200 μ L of supernatant from the 1.5ml tube containing the centrifuged sample, being careful not to disturb the pellet at the bottom of the tube. Place the 200 μ L in a labeled 96 well extraction plate labeled with Extraction Plate Day1 [date]”

Note: Pellet size will vary depending on trichome density.

6. Vortex MGC Binding Buffer thoroughly before use, be sure that the magnetic particles are completely re-suspended in buffer for at least 30 seconds.

7. Add 200µL of MGC Binding Buffer to each sample, and pipette tip mix 15 times.

Note: Be careful to avoid adding too many bubbles by pipetting gently when tip mixing. This is extremely important as to not contaminate the wells in proximity.

8. Incubate the plate on the bench for at least 5 minutes.

9. Place the extraction plate onto the 96 well plate magnet plate for at least 5 minutes.

10. After the 5 min incubation, remove as much of the 400µL of the supernatant as possible. Be careful not to disturb or aspirate the beads. See Figure 5.

11. Add 400µL of 70% ethanol (EtOH) with the extraction plate still on the magnet plate. Wait at least 30 seconds and remove all the EtOH.

Note: Place the pipet tip at the bottom center of the well to remove all liquid.

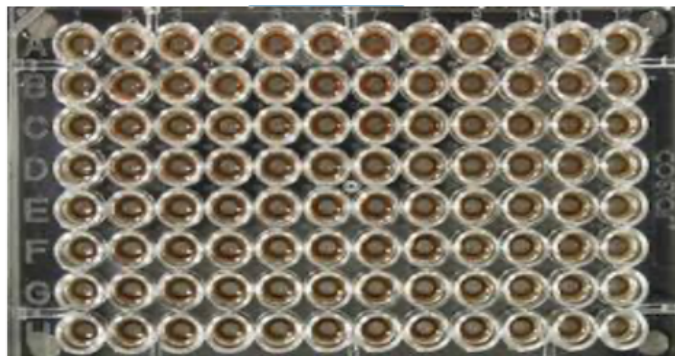


Figure 5: Extraction plate during wash step on magnetic plate.

12. Repeat 400 µL 70% EtOH wash with the extraction plate still on the magnet plate. Wait at least 30 seconds and remove all the EtOH.

Note: If EtOH still remains in the wells, go back in with a smaller pipet tip to remove the excess. Leftover EtOH can inhibit qPCR efficiency.

13. After all the EtOH has been removed, let the beads dry at room temperature. Allow the beads to dry for up to 15 minutes. Necessary drying time will vary based on complete removal of the second ethanol wash, as well as lab environment. Visually inspect beads for residual ethanol before the elution step.

Note: It is important to NOT allow the beads to dry for an extended period of time. Over-drying can cause a reduction in DNA yield.

14. Remove the extraction plate from the magnet plate and add 50 μ L of MGC Elution Buffer.

- a. Tip mix approximately 15 times or until the beads are completely re-suspended.

Note: The re-suspensions may appear varied in their appearance, but the result will be the same. See Figure 6.

- b. Incubate the plate for at least 1 minute on the bench, before returning the plate to the magnetic plate.
- c. Let the plate sit on the magnet for at least 1 minute before transferring the eluant to a new extraction plate labeled with “Final Extract [date]”.

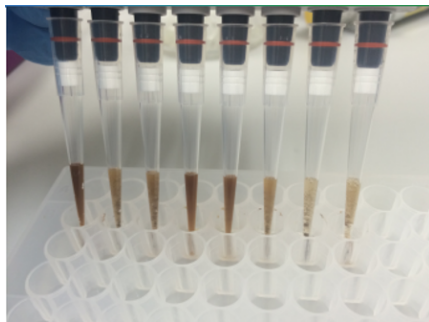


Figure 6: Multichannel pipette tips showing magnetic beads resuspended in elution buffer.

15. Seal the plate with the adhesive seal, making sure to completely seal the plate wells using a pen or flat object to slide back and forth along the seal. Store at -20°C until ready to perform the qPCR protocol.

SenSATIVAx® for MIP/Extracts DNA Purification

1. Prepare a SCCG positive control dilution of 1:5,000 (internal control)
 - a. Label a new 1.5mL Eppendorf tube (SCCG 1:50), add 1µl of SCCG positive control into 49µl of dH₂O. Vortex to mix thoroughly and quick spin tube. Label another 1.5mL Eppendorf tube (SCCG 1:5,000), add 99µl of dH₂O, then add 1µl of the SCCG 1:50 dilution. Vortex to mix thoroughly and quick spin tube. This will result in a 1:5,000 dilution of SCCG.

Note: It's easy to mis-pipette when trying to pipette only 1 µL of liquid. Visually check your pipette tip after aspirating 1 µL to ensure it is in the tip before adding it to the tube for dilutions 1 and 2.

- b. Place on ice until use.

Note: The 100 µL dilution prepared above is enough diluted SCCG for approximately 10 Purifications. If more purifications are going to be prepared at the same time, the initial 1:50 dilution can be used to make multiple 1:5,000 dilutions of SCCG.

2. Add initial sample weight x 4.6 mL SenSATIVAx Solution A to conical tube with enriched sample/TSB. Vortex the sample vigorously until homogenized.
3. Transfer 1 mL of the homogenized sample into a 1.5 mL tube.
4. Add 10µL of the SCCG internal control (1:5,000) to 1.5 mL tube and vortex to mix well.

5. Centrifuge for 10 minutes at 19,300 RCF using a high-speed bench top centrifuge.
 - a. If no bench top centrifuge is available, centrifuge for 15 minutes using a mini centrifuge.

Note: Some matrices will require the use of a high-speed centrifuge due to the presence of certain substances such as gelatin that hinder phase separation.

6. Transfer 600 μ L of the solution to a new tube. Push pipet tip through the top solid layer (if one exists), without disturbing the pellet on the bottom to aspirate the sample.
7. Add 600 μ L chloroform and vortex vigorously until solution turns a milky white color throughout

Note: This may require longer vortexing for some matrices

Caution: ALWAYS WEAR GLOVES WHEN HANDLING CHLOROFORM

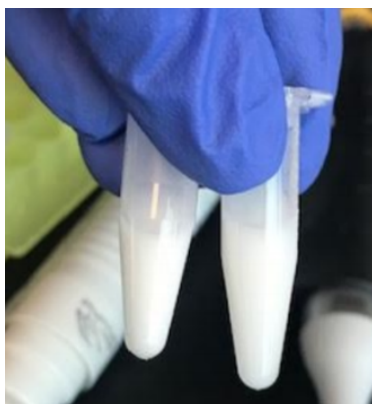


Fig 7. Example of a milky sample with chloroform in it.

8. Centrifuge for 5 minutes at 14,000 - 19,300 RCF using a bench top centrifuge
9. If no bench top centrifuge is available, centrifuge for 15 minutes using a mini centrifuge.

NOTE: If there is still any color in your aqueous layer (top layer) after centrifugation another chloroform wash may help give you a strong internal control signal (HEX) for every assay. Transfer 300 μ L of the top layer to a new 1.5 mL tube and add 300 μ L chloroform, vortex and centrifuge again.

10. Transfer 100µL of aqueous layer (TOP LAYER) from Step 7 to a well of the labeled 96 well extraction plate. Be careful not to disturb the lower chloroform layer.
 11. Add 100µL of SenSATIVAx Solution B to the 100 ul sample in the 96 well extraction plate and mix by pipetting up and down 3 times..
 12. Vortex MGC Binding Buffer thoroughly before use, be sure that the magnetic particles are completely re-suspended in buffer for at least 30 seconds.
 13. Add 200µL of MGC Binding Buffer to each sample, and pipette tip mix 15 times.
- Note: Be careful to avoid adding too many bubbles by pipetting gently when tip mixing. This is extremely important as to not contaminate the wells in proximity.*
14. Incubate the plate on the bench for at least 5 minutes.
 15. Place the extraction plate onto the 96 well plate magnet plate for at least 5 minutes.
 16. After the 5 min incubation, remove as much of the 400µL of the supernatant as possible. Be careful not to disturb or aspirate the beads.
 17. Add 400µL of 70% ethanol (EtOH) with the extraction plate still on the magnet plate. Wait at least 30 seconds and remove all the EtOH.

Note: Place the pipet tip at the bottom center of the well to remove all liquid.

18. Repeat 400 µL 70% EtOH wash with the extraction plate still on the magnet plate. Wait at least 30 seconds and remove all the EtOH.

Note: If EtOH still remains in the wells, go back in with a smaller pipet tip to remove the excess. Leftover EtOH can inhibit qPCR efficiency.

19. After all the EtOH has been removed, let the beads dry at room temperature. Allow the beads to dry for up to 15 minutes. Necessary drying time will vary based on complete removal of the

second ethanol wash, as well as lab environment. Visually inspect beads for residual ethanol before the elution step.

Note: It is important to NOT allow the beads to dry for an extended period of time. Over-drying can cause a reduction in DNA yield.

20. Remove the extraction plate from the magnet plate and add 50 µL of MGC Elution Buffer.

- a. Tip mix approximately 15 times or until the beads are completely re-suspended.

Note: The re-suspensions may appear varied in their appearance, but the result will be the same.

- b. Incubate the plate for at least 1 minute on the bench, before returning the plate to the magnetic plate.

- c. Let the plate sit on the magnet for at least 1 minute before transferring the eluant to a new extraction plate labeled with “Final Extract [date]”.

21. Seal the plate with the adhesive seal, making sure to completely seal the plate wells using a pen or flat object to slide back and forth along the seal. Store at -20°C until ready to perform the qPCR protocol.

Real-Time Quantitative PCR (qPCR) Setup Protocol

1. Remove qPCR reagents including qPCR Master Mix, water, reaction buffer and assay probe mixes to be used from the -20°C freezer. Place qPCR master mix on ice or leave at -20°C until ready to use. Allow remaining tubes to thaw at room temperature. Once thawed, immediately place tubes on ice.
2. Before preparing the reaction, invert or vortex and spin-down the reagents.
 - a. Assay probe mix tubes, reaction buffer, positive controls and water – Vortex quickly followed by a pulse spin-down in a microcentrifuge.
 - b. qPCR Master Mix – Invert the tube 5 times (do not vortex), followed by a pulse spin-down in a microcentrifuge.
 - c. Return all reagents to the ice.

Note: Do not vortex the qPCR Master Mix at any point during the protocol.

3. Make a separate master mix in a 1.5mL tube for each assay type being run. All probe mixes contain the internal plant control, SCCG probe mix, and the probe for the microbial targets. Label each tube with [Assay Name] MM. Always prepare enough master mix for 1 or 2 additional reactions over the total number of tests to account for pipetting and dead volumes.

Note: It is best to add the largest volume reagent first, in this case water.

Table 1: PCR Reagent Volumes

Reagents	1 Reaction	24 Reactions (Plus 1 excess rxn)	48 Reactions (Plus 2 excess rxn)
qPCR Master Mix	3.75 µL	93.75 µL	187.5 µL
Assay Probe Mix	1 µL	25 µL	50 µL
Reaction Buffer	0.8 µL	20 µL	40 µL
Water	8.2 µL	205 µL	410 µL
Total Assay Probe MM	13.75 µL	343.75 µL	687.5 µL

4. Once combined, gently tip mix or invert the tube 5 times to combine the assay master mix.
 - a. Pulse spin-down tube in microcentrifuge.
 - b. Place qPCR Master Mix tubes on ice until used.
 - c. For the positive control, make a 1:10 dilution
 - i. Add 1 µL of Positive Control to 9 µL nuclease free water (found in the kit), vortex to mix well and spin down the tube.
 1. For the negative control, use water (found in the kit)
- Note: It is best to add the largest volume reagent first, in this case the 9 µL water then the 1 µL of positive control, pipette mix or vortex control dilution to ensure control DNA is in solution.*
5. Place the Extraction Plate on the magnet. This is to ensure no magnetic beads are transferred into the qPCR reactions if there are some left over from the purification elution process.
 6. Use a 96-well optical qPCR plate and label the plate “qPCR Plate_ [date]”.

7. Carefully remove the seal from the Extraction Plate. If frozen, let the DNA thaw completely and spin the plate to avoid cross contamination between samples. Transfer 5 μ L of each sample into the corresponding well on the qPCR plate. Keep the extraction plate on the magnet when aspirating the 5 μ L.
 - a. Add 5 μ L of the diluted Positive Control to the corresponding positive control well. Then add 5 μ L of water to the corresponding negative control well.

Note: ALWAYS use a fresh tip for every liquid transfer into the qPCR plate

8. Add 13.75 μ L of specific Assay Probe MM to each corresponding sample well, positive control well, and negative control well in the qPCR plate. Gently tip mix a few times after each addition of qPCR master mix. Be careful not to introduce bubbles during this mix.
9. Seal the plate with strip caps or an adhesive seal.
10. Spin-down for at least 1 minute in plate microcentrifuge to bring well contents to the bottom of wells and help to get rid of reaction bubbles.

Note: Check for bubbles at the bottom of the wells (minimal bubbles on the surface of the liquid is acceptable). If bubbles remain in the bottom of the wells, spin-down for another minute.

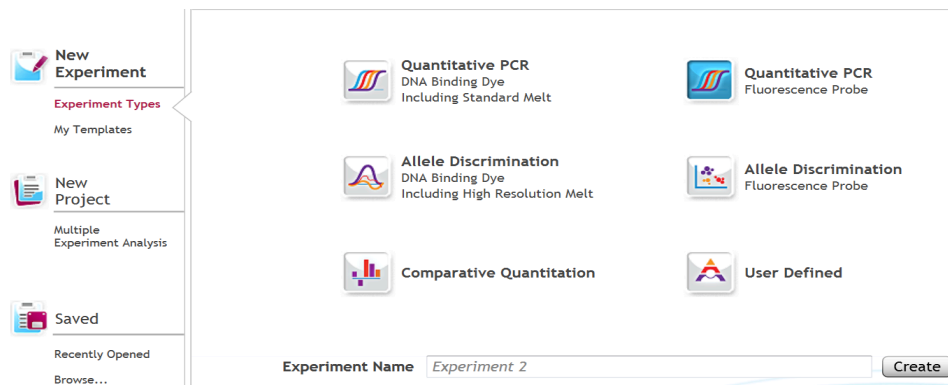
11. For the Agilent Aria: If using an adhesive seal; place the reusable compression pad (gray side down) on the plate directly lining up the holes in the pad with the holes in the plate.
12. Place the sealed plate onto the PCR instrument, positioning the A1 well in the top left corner.
13. Follow the software specific instructions to initiate the run.

Running the Agilent AriaMX

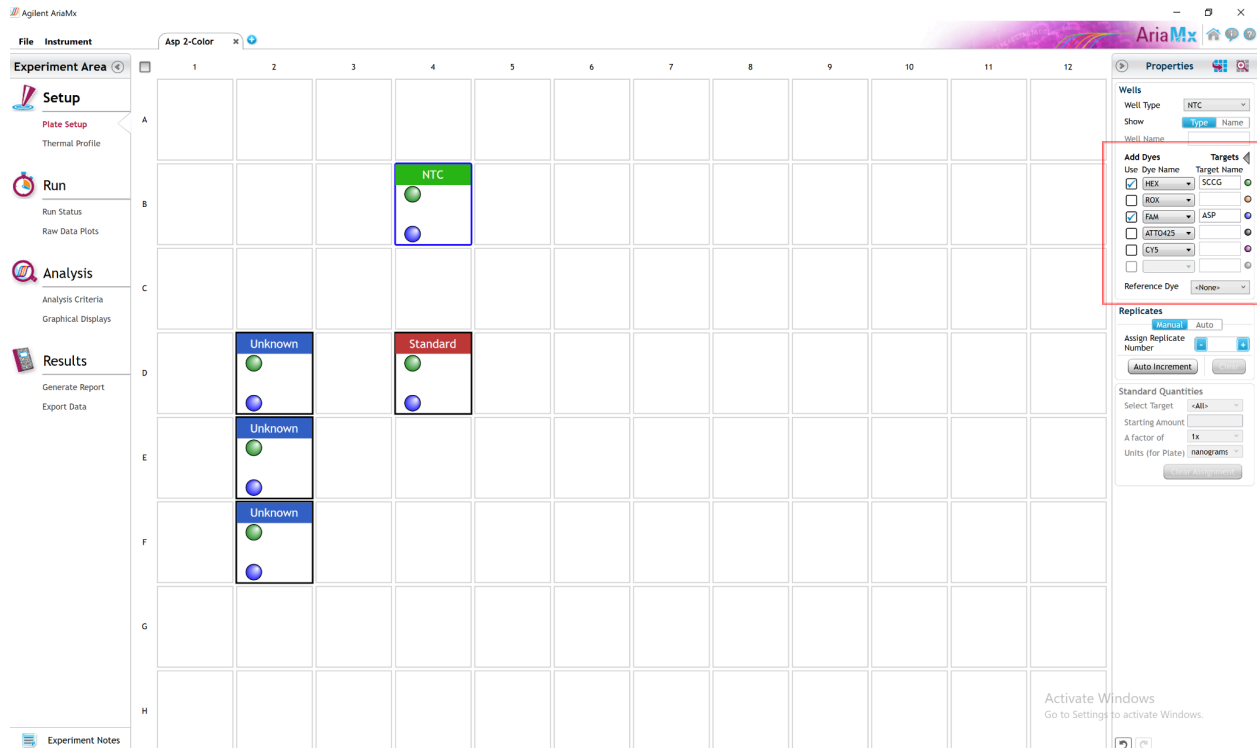
The following species will be detected on the following Fluorophores:

- P. aeruginosa and/or S. aureus: FAM
- Cannabis DNA: HEX

1. Create a New Experiment on the Agilent qPCR instrument. Select “Quantitative PCR: Fluorescence Probe” from Experiment Types.

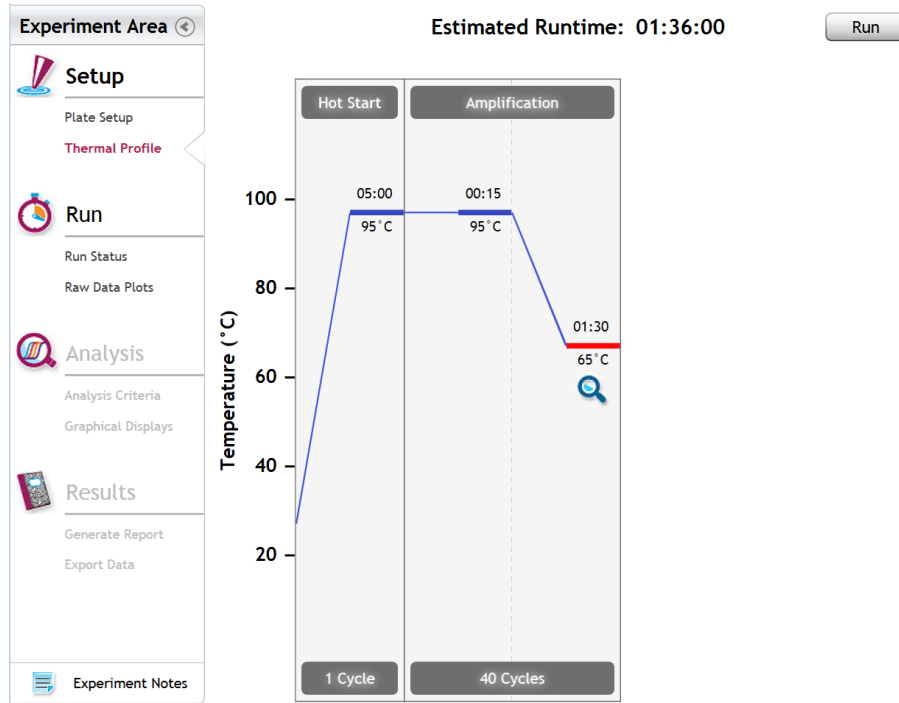


2. Under Setup > Plate Setup, highlight **only** wells that contain reactions and select FAM and HEX under **Add Dyes**. Having fluorescence detection turned on in empty wells can affect the calculations which the software makes to obtain the corrected data, ΔR log.



3. Change the well types to reflect your plate set up. All wells should be set to Unknown except the negative control can be set to NTC well type. Add Target names to the dyes under **Targets**:
 - a. FAM : *S. aureus* and/or *P. aeruginosa*
 - b. HEX : Single Copy Cannabis Gene (SCCG) or Internal Control (IC)

4. Under Setup > Thermal Profile, create the following PCR thermal profile
 - a. Hot start at 95°C for 5 minutes, followed by 40 cycles of 95°C for 15 seconds and 65°C for 90 seconds.



5. Close the lid and click “Start Run”
6. Save the experiment with the [User] and [date]
7. When the qPCR run is complete, immediately dispose of the plate. Do not open the plate seal after the run to avoid contamination in the lab.

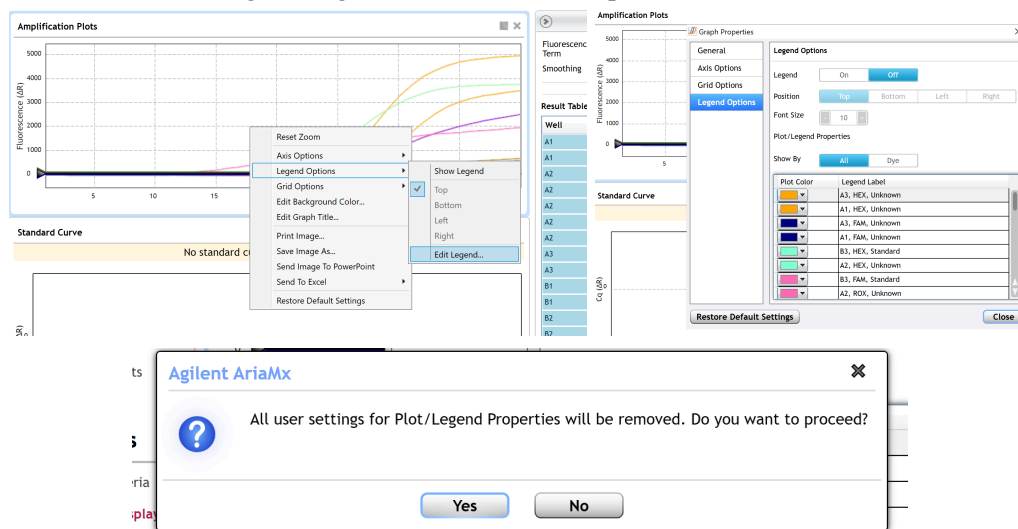
Data Analysis: Agilent AriaMX

PathoSEEK® P. aeruginosa and S. aureus - Data Analysis Quick Reference Table:

Table 3: All Matrices

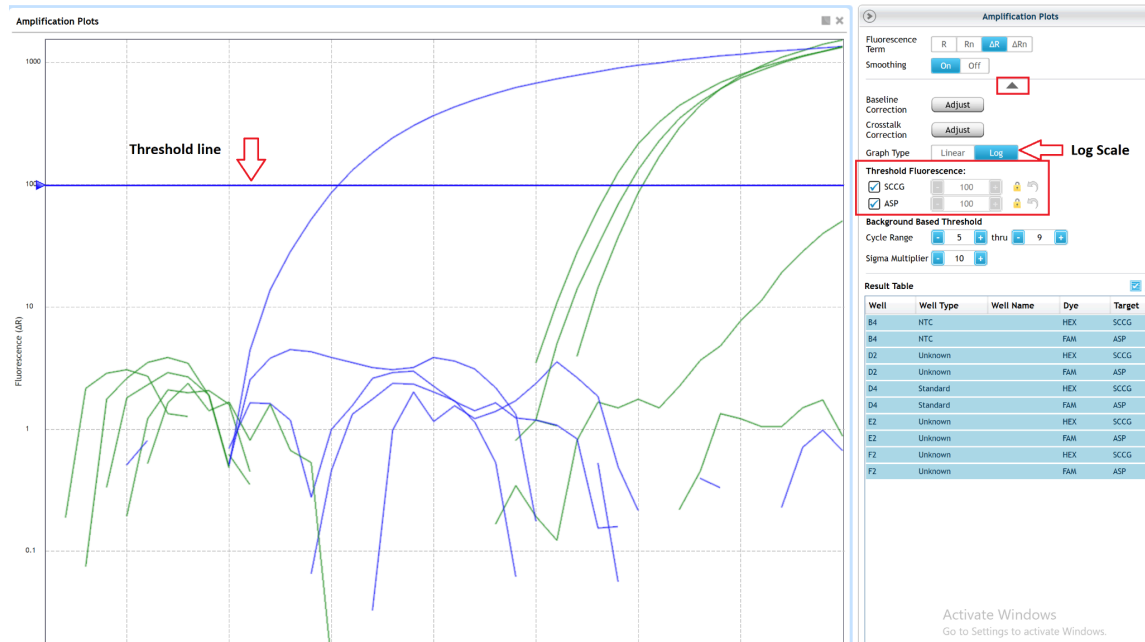
PathoSEEK® Assay	Cq Value	Fluor	Negative Control (Cq)	CFU threshold (CFU/g)
<i>P. aeruginosa</i> and/or <i>S. aureus</i>	≤ 40	FAM	No Cq	Presence/Absence
Internal Control*	≤ 35	HEX	*Internal control verifies the presence or absence of plant DNA	
Assay Positive Control	≤ 35	FAM		

1. Open the Data Analysis window when the run is complete.
2. Highlight the wells of interest in the Analysis Criteria under Analysis, then select Graphical Display
 - a. Amplification plots will be available for viewing
 - b. The Cq values will appear to the right in the table
 - c. Right click inside the graph, select Edit Legend under Legend Options
 - d. Change “All” to “Dye”
 - e. All user settings for Plot/Legend Properties will be removed. Do you want to proceed? Select “Yes”.
 - f. This will assign a single color to each fluorophore.



3. To analyze the results

- Start by turning the graph to Log Scale with a right click on the chart, select Axis options, enable y-axis log scale. Expand the amplification plots settings by clicking on the triangle
- Manually adjust thresholds to 100 RFU for the HEX and FAM fluorophores. (which will have the target name because you've added them in the plate setup.) Click the lock next to each target to prevent future changes.

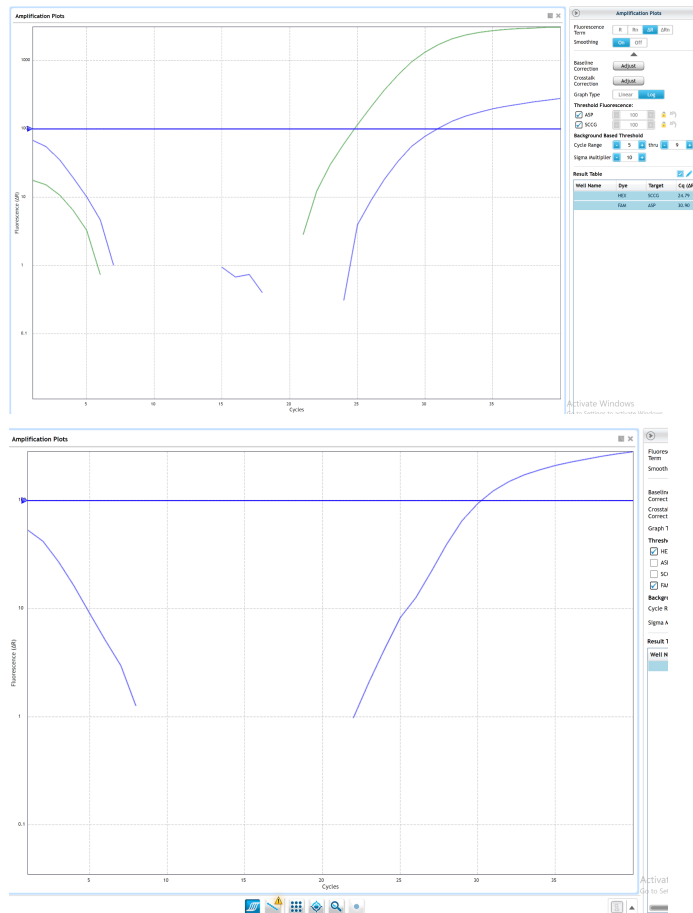


4. Controls

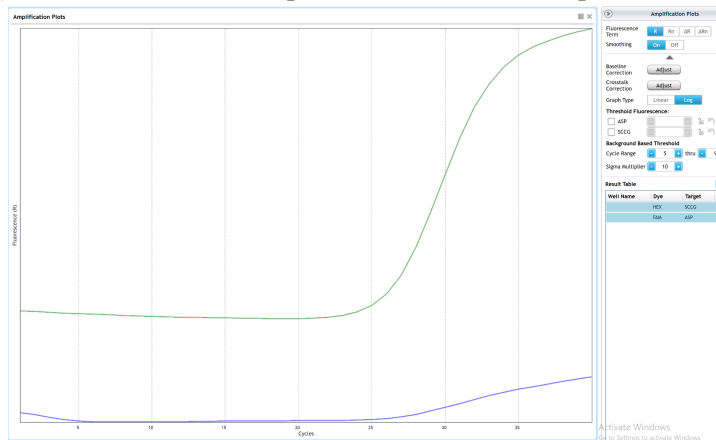
- Positive Control, on the FAM Fluorophore, has Cq value ≤ 35 .
 - Visually confirm with the curve on the graph.
- Negative Control, on the FAM Fluorophore, has no Cq Value.
 - Visually confirm with the curve on the graph.

5. Unknown *P. aeruginosa* and/or *S. aureus* Targets

- A “presence” or failing result for the unknown *P. aeruginosa* and/or *S. aureus* target(s).
 - Any Cq value for the FAM Fluorophore ≤ 40 .



- ii. Visually confirm with the curve on the graph. It is very important to check the raw data view (R) to confirm with the amplification curve when a presence result occurs.



- iii. Review the raw data for each sample:

Select the wells of interest, use the plate view of the Raw Data Plots to scroll across the plate to confirm amplification in sample wells and positive control wells. This review also ensures that thermal cycling is uniform across the plate.

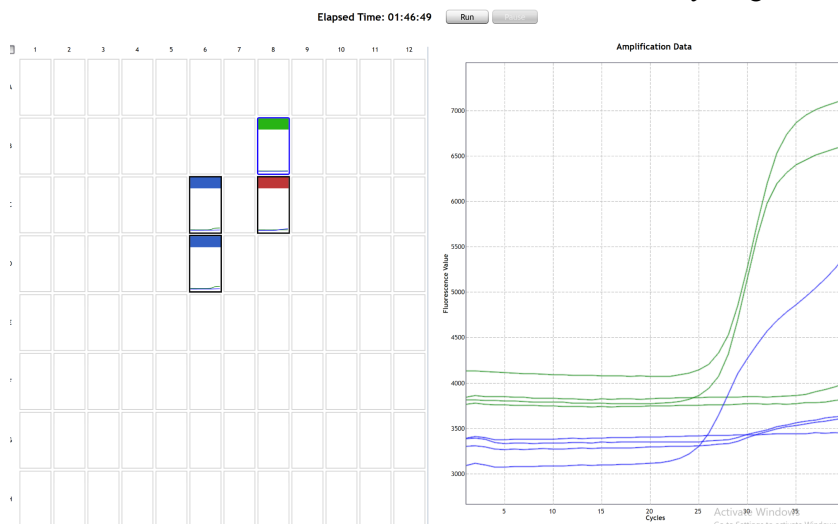


Figure 7: Raw data (R) plots for which the cycle quantification (Cq) will be generated.

- iv. Look at the raw data in the "Graphical Displays" view. For every dye that gives a Cq make sure the baseline is flat and the fluorescence signal grows rapidly over 5-7 cycles as seen below. (To view the raw data, select "R" next to "Fluorescence Term" at the top right-hand side of the screen.)

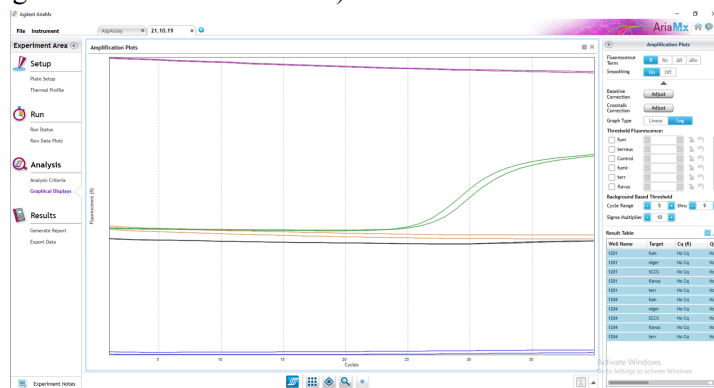
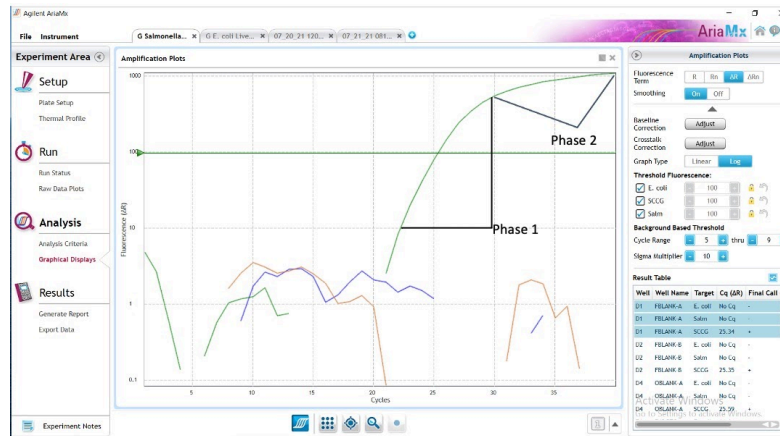


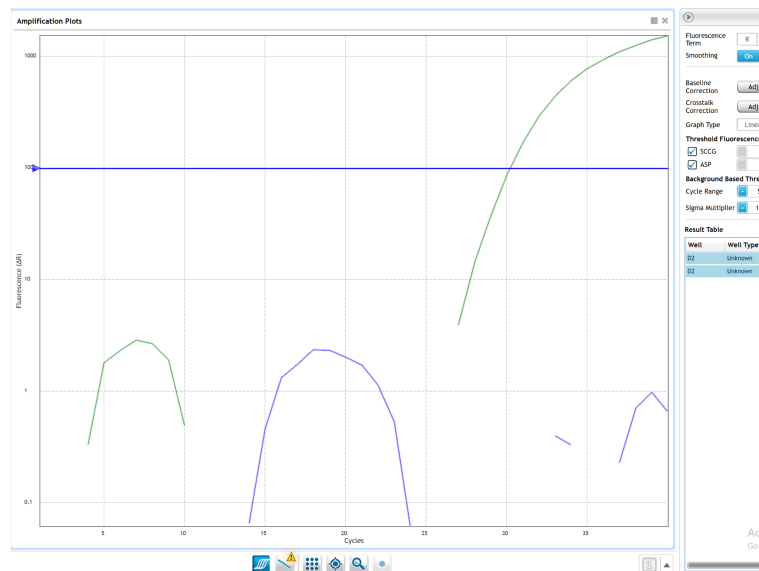
Figure: Example of good baseline with sigmoidal shaped curve increase in the internal control fluorescence.

- v. Check the amplification plots for a semi-logarithmic curve with two distinct phases that crosses the threshold:
 1. Select the ΔR Fluorescence Term and look at the shape of the amplification plot. The graph should have the classic semi-logarithmic shape with two visually distinct phases.
 2. Phase 1 shows exponential growth in the fluorescence, in a span of around 5 cycles.

3. Phase 2 is a plateau where the amplification signal growth ends but remains level.



- b. An “absence” or passing result for the unknown *P. aeruginosa* and/or *S. aureus* targets.
 - i. No Cq value for the FAM Fluorophore
 1. Visually confirm that there is no curve on the graph.
 2. Cq of <35 for the internal control on the HEX fluorophore.
 - a. Visually confirm the curve on the graph.

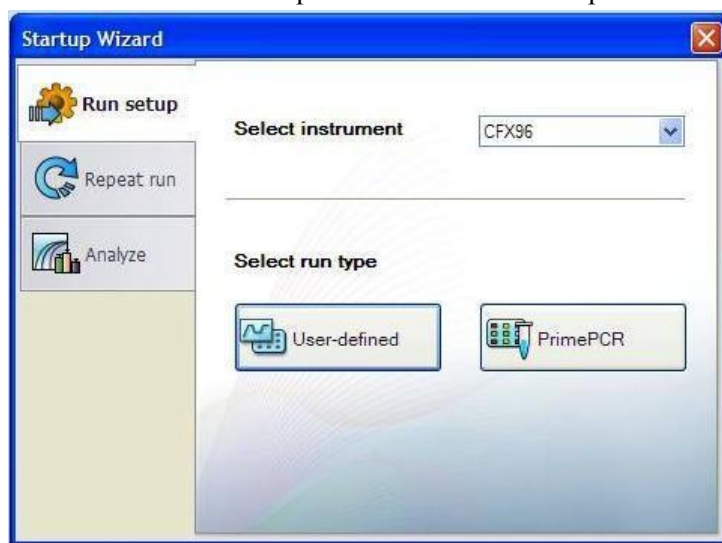


Running the BioRad CFX96

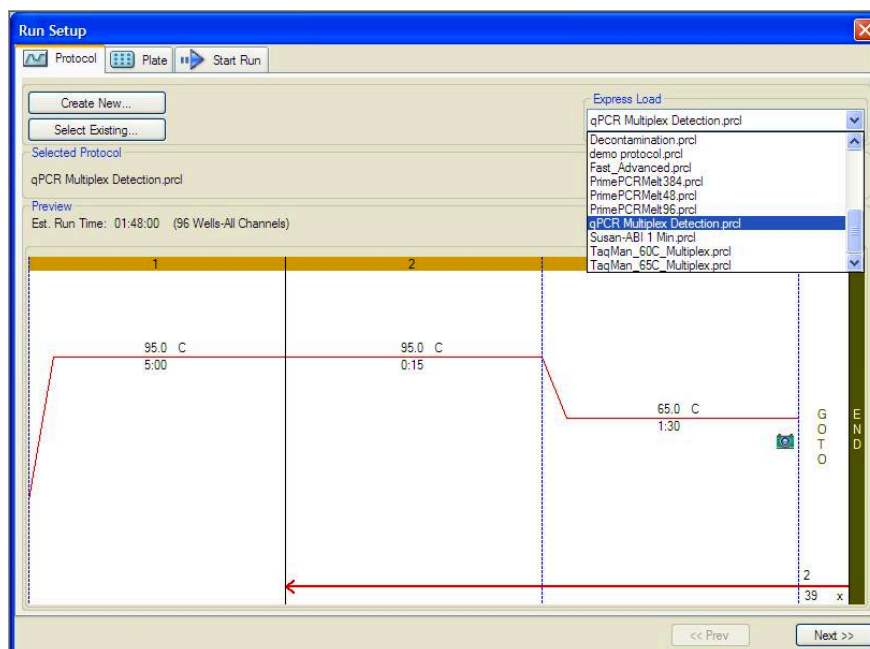
The following targets will be detected on the following fluorophores:

- P. aeruginosa and/or S. aureus: FAM
- Cannabis DNA: HEX

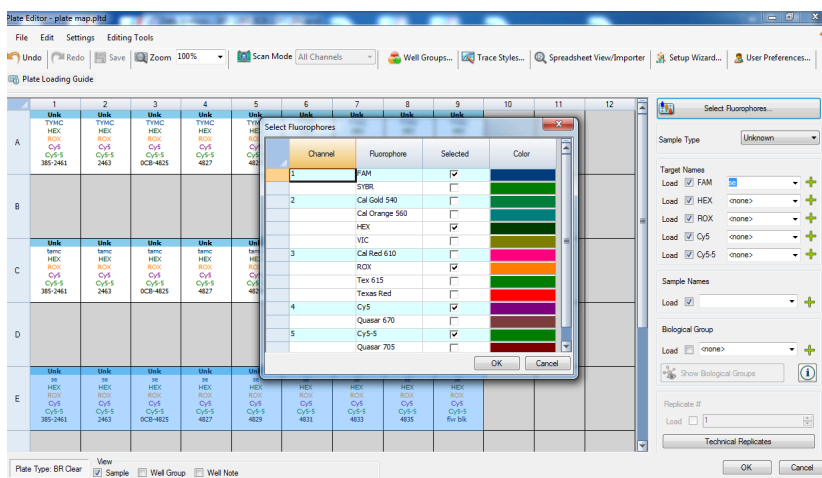
1. Start the qPCR Cycling program
 - a. Select User-Defined in the Startup Wizard under Run setup



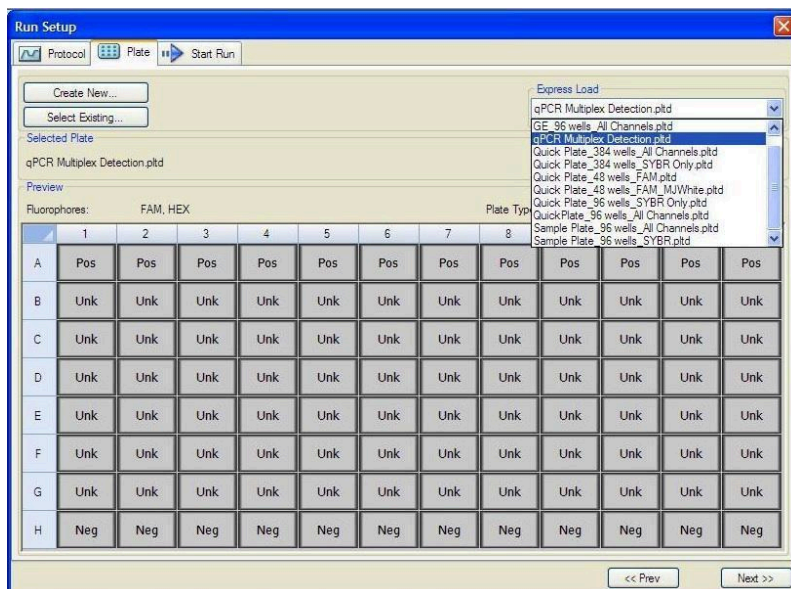
2. Use the Express Load dropdown menu to pick the qPCR Multiplex Detection Program and click “Next”.
 - a. If not already pre-programmed, create a cycling program with the following specifications and save as “qPCR Multiplex Detection”
 - i. Hot start at 95°C for 5 minutes, followed by 40 cycles of 95°C for 15 seconds and 65°C for 90 seconds.



3. Design your plate under the plate tab in the Run Setup
 - a. Select the qPCR Multiplex Detection from the dropdown menu. If not already present, click “Create New”
 - b. Plate editor window will appear. Choose FAM and HEX Fluorophores and click “OK”.



- c. If plate layout was previously saved, click “Edit Selected” to move to the Plate Editor Screen.

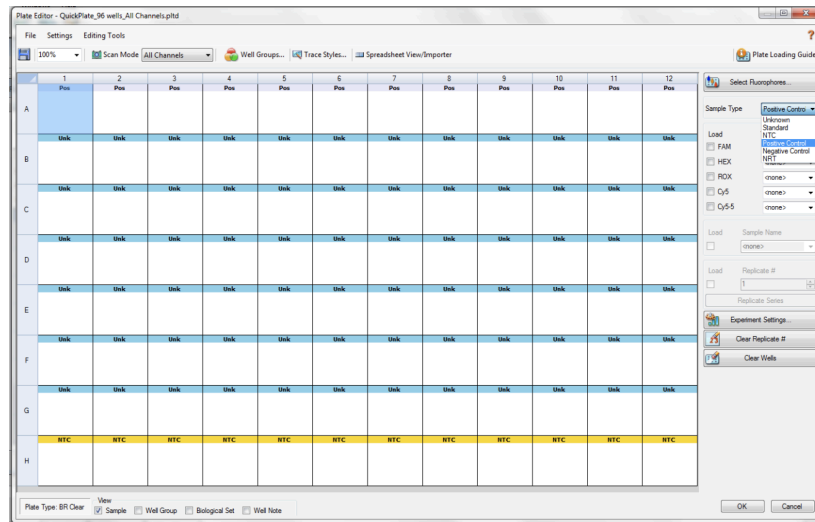


- d. On the Plate Editor Screen, change the Sample Type to correlate with your specific plate setup.

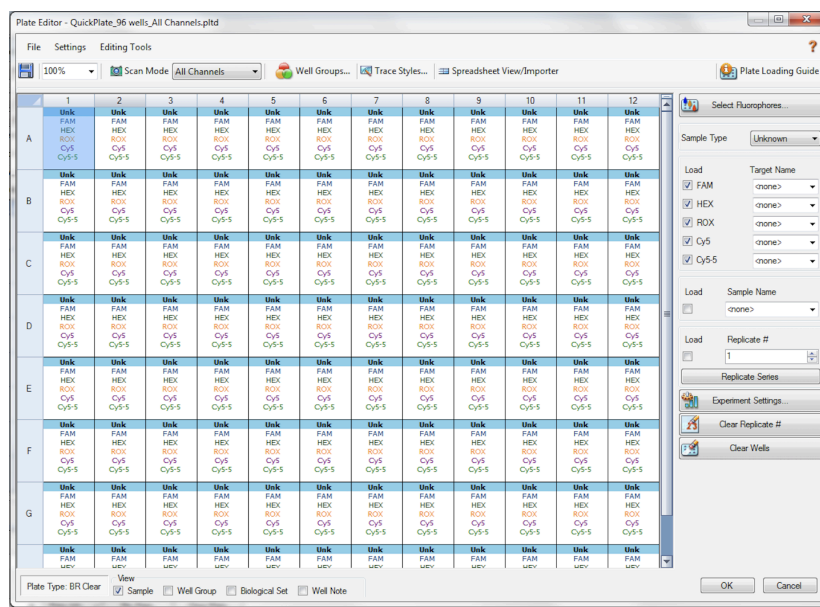
NOTE: To select the Sample Type, highlight the wells you would like to define, then choose from the dropdown menu one of three types:

*Unknown
Positive Control
Negative Control*

- e. Make sure All Channels is selected from the dropdown menu at the top.



- f. Attach the fluorophores to the wells being used.
- i. Highlight all the wells being used
 - ii. For *P. aeruginosa* and/or *S. aureus* Assay, **Highlight the well locations and click on FAM and HEX**
- g. When the plate is designed correctly, click OK.
- h. Click “yes” to save your plate. If creating plate layout for the first time, save as “qPCR Multiplex Detection”. If you do not save the plate, it will return to the default plate.



Note: Saving will override the template (that is fine).

- i. Close the lid and click Start Run.
- j. Save the experiment with the [User] and [date]
- k. When the run is complete, immediately dispose of the plate after qPCR. **Do not remove the plate seal after the run to avoid contamination in the lab.**

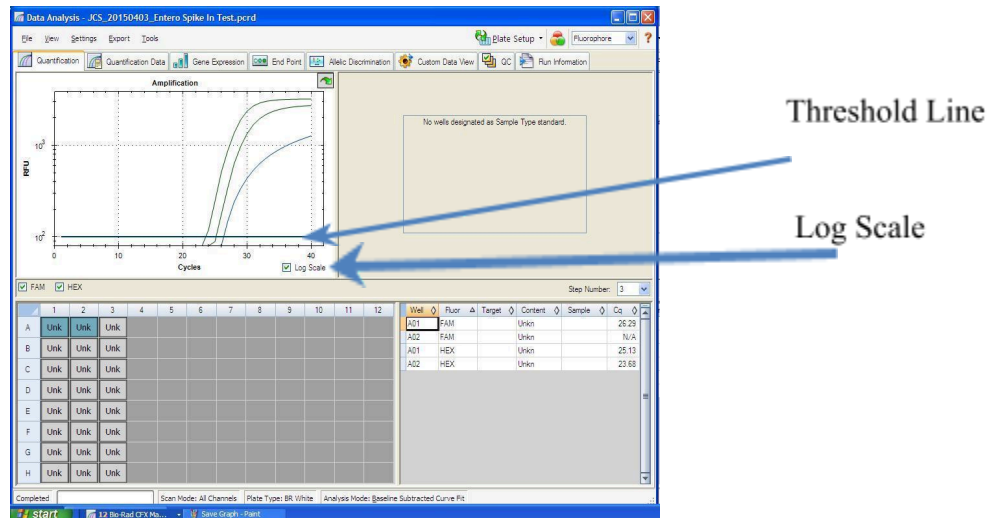
Data Analysis BioRad CFX96

PathoSEEK® P. aeruginosa and/or S. aureus Data Analysis Quick Reference Table:

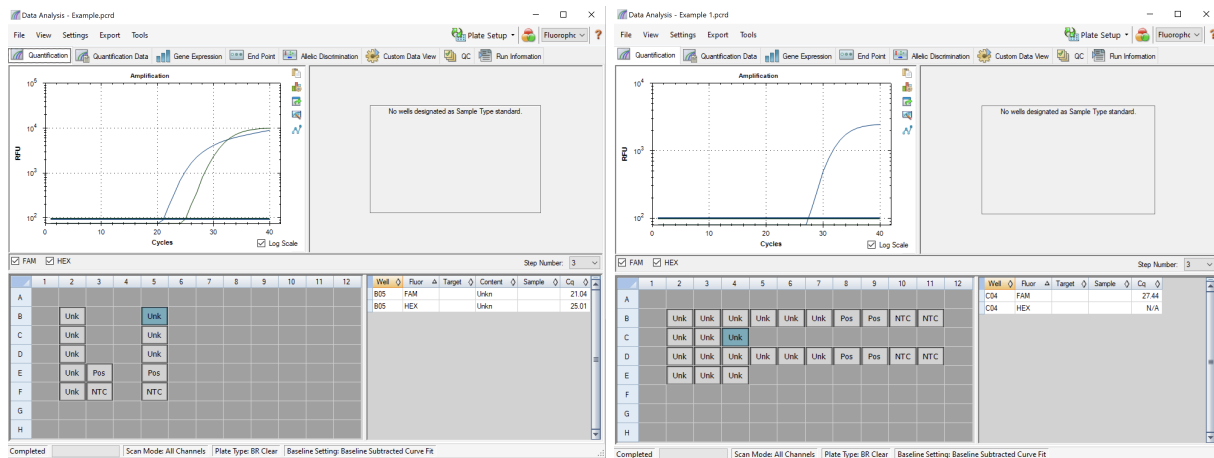
Table 4: All Matrices

PathoSEEK® Assay	Cq Value	Fluor	Negative Control (Cq)	
P. aeruginosa and/or S. aureus	≤ 40	FAM	No Cq	
Internal Control*	≤35	HEX	*Internal control verifies the presence or absence of plant DNA	
Assay Positive Control	≤35	FAM		

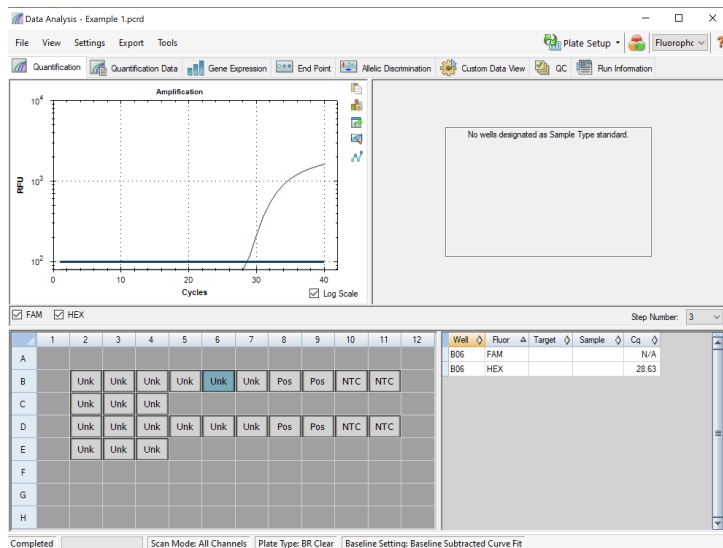
1. The Data Analysis window will open automatically when the run is complete.
2. Highlight the well of interest.
 - a. The graph will appear above.
 - b. The Cq values will appear to the right.
3. To analyze the results
 - a. Start by turning the graph to Log Scale and manually moving the threshold to 10² for all fluorophores.
 - b. To turn the graph to Log Scale, click on the box at the bottom right of the graph.
 - c. To adjust the threshold, click on the horizontal lines, and move them to the specified value mentioned above on the y-axis.



- d. Controls
 - i. Assay-specific Positive Control, on the FAM fluorophore, has a Cq value ≤ 35 .
 1. Visually confirm with the curve on the graph.
 - ii. Assay-specific Negative Control, on the FAM fluorophore, has no Cq value.
 1. Visually confirm with the curve on the graph.
- e. Unknown *P. aeruginosa* and/or *S. aureus* Targets: The following fluorophore detects *P. aeruginosa* and/or *S. aureus*:
 - i. *P. aeruginosa* and/or *S. aureus*: FAM
 - ii. A “presence” or failing result for the unknown *P. aeruginosa* and/or *S. aureus*.
 1. Any Cq value for the FAM fluorophore ≤ 40 .
 2. Visually confirm with the curve on the graph. It is very important to confirm with the amplification curve when a presence result occurred. Sometimes the background amplification will give a false positive reading.



- iii. An “absence” or passing result for the unknown *P. aeruginosa* and/or *S. aureus* targets.
1. No Cq value for the FAM fluorophore.
 2. Visually confirm no curve on the graph.
 3. Internal Control, on the HEX fluorophore, has a Cq value ≤ 35 for flower samples, < 40 for all other matrices.
 4. Visually confirm with the curve on the graph



Troubleshooting Guide

Symptom	Reason	Solution
Internal Control Failure	Extraction Failure	Repeat SenSATIVAx™ and PathoSEEK™ by following the protocol.
	Residual ethanol in elution	Ethanol is an inhibitor to PCR. Return to the SenSATIVAx™ protocol and repeat all steps.
	Mix up in Reaction Setup	Repeat the qPCR by following the protocol.
	Missing Fluorophore on plate set up	In the Data Analysis window click on View/Edit Plate Setup from the Settings drop down. All wells should have both FAM and HEX. Once completed and the window is closed the analysis should automatically update.
	Matrix related (ex: age, remediation, ground)	Repeat SenSATIVAx and PathoSEEK following the protocol. If unacceptable internal control results are obtained a second time, Internal Cannabis Control may be spiked into samples during extraction for the third run. Please visit our Help Center for more information.
Internal Control Positive result on positive or negative control samples or samples that do not contain plant DNA No Cq or a Cq of more than 35 is acceptable. Any Cq lower than 35 constitutes a rerun	Plant DNA contamination in a reagent	Troubleshoot which reagent was contaminated; use new reagents, thoroughly clean all pipettes and bench areas with 10% bleach solution.
	qPCR bench too close to extraction area	Designate separate benches, pipettes etc. for extractions and qPCR setup
Positive Negative Control	Small Cq value <15	Visually confirm that there is an amplification curve. If not, this is a low level background and is to be expected.
	Contamination	Repeat the qPCR by following the protocol.
	Insufficient pre-setup bleaching	Wipe down the lab workspace and all equipment with 10% Bleach. Repeat qPCR.
Negative Positive Control	Mix up in Reaction Setup	Repeat the qPCR by following the protocol.
Total run failure	Excessive vortex of the qPCR Master Mix	Repeat the qPCR by following the protocol.
Background Amplification	Unclear	This is usually seen with a very low Cq reading (<15), the curve is usually missing the exponential growth phase, but rather a gradual increase of fluorescence signal. This is usually a negative result, but should be repeated.

Glossary and Definitions

Deoxyribonucleic acid (DNA) is a molecule that encodes the genetic instructions used in the development and functioning of all known living organisms.

Polymerase Chain Reaction (PCR) is a technology in molecular biology used to amplify a single copy or a few copies of a piece of DNA across several orders of magnitude, generating thousands to millions of copies of a particular DNA sequence.

A **fluorophore** is a fluorescent chemical compound that can re-emit light upon light excitation.

The **Negative Controls** are the reactions where no Cq is expected. It helps to ensure that all Assay-specific reactions are clean of contaminants.

The assay-specific **Positive Controls** are the reactions where a Cq is expected. It helps ensure that all Assay-specific reactions are working correctly. The Assay specific Positive Control is targeting the pathogen using the FAM Fluorophore.

Amplification of the **Internal Control** is expected in every reaction containing cannabis DNA. It ensures the DNA isolation procedure was successful. The internal control targets plant DNA, or more specifically, a Single Copy Control Gene (SCCG), using the HEX Fluorophore.

MIP is short for Marijuana Infused Product. A MIP is cannabis plant material or concentrate mixed into a consumable.

Revision History

Version	Date	Description
v1	June 2023	Update to User Guide format
v2	June 2024	Removal of optional Grim Reefer Free DNA Removal. Added information to the disclaimer section concerning remediated samples. Updated troubleshooting guide steps for internal control failures.

DISCLAIMER

This test was developed, and its performance characteristics determined by Medicinal Genomics Company, for laboratory use. Any deviations from this protocol are not supported by MGC.

This test has not been validated on remediated (irradiated, ozone treated, acid treated, hydrogen peroxide treated, etc.) samples. Samples that have undergone remediation may cause discordant results between plating methods and PathoSEEK methods. When remediated samples produce a result above the action limit on qPCR, we recommend confirming viability with an approved plating method.

Results may vary based on laboratory conditions. Altitude and humidity are among factors known to affect the growth of bacterial and fungal species.

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