Procedure for the Preparation and Analysis of Spiked Reference and Matrix Samples

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1. SCOPE AND APPLICATION

1.1. The analysis of a clean sample matrix, fortified with a known quantity of the target organism from a manufactured standard such as a Quality Control (QC) standard or a Certified Reference Material (CRM), permits the monitoring of initial and ongoing method performance, as long as manufacturer instructions for use are followed. It is also useful as a demonstration of capability for analysts. The laboratory will assess performance for accuracy with the established acceptance criteria in the COA.

2. SUMMARY OF METHOD

2.1. Four spiked reference samples of either *Enterococcus faecalis* (*E. faecalis*), Escherichia *coli* (*E. coli*), or *Klebsiella variicola,* are prepared using vials of phosphate buffer, or Autoclaved DI water, and analyzed according to the respective method.

2.2. Interferences

2.2.1. Temperature of the reference bacteria pellet can affect its activity. Analysis should begin when the capped pellet has acclimated to room temperature (15-30°C).

2.2.2. Time of inoculation can affect bacterial activity. Analyze spiked sample within 30 minutes of reconstitution.

2.2.3. Aseptic techniques must be used to ensure that there is no contamination of the sample via the media, reagents, supplies, or the analyst.

2.2.4. Mix contents thoroughly after each inoculation and again before processing by vigorously agitating the vials. This breaks up clumps of bacteria and homogenizes the sample.

2.3. Definitions

2.3.1. ***Enterococcus faecalis* –** Gram-positive cocci bacteria. One of several members of the genus *Enterococcus* that can be found in the feces of humans and other warm-blooded animals. Formerly of the Group D fecal streptococci group.

2.3.2. ***Escherichia coli* –** Gram-negative, non-spore forming, rod-shaped bacteria. One of several bacteria species known as fecal coliforms. Fecal coliforms are commonly found in the feces of humans and warm-blooded animals.

2.3.3. ***Klebsiella variicola* –** Gram-negative, non-spore forming, rod-shaped bacteria. One of several members of the family *Enterobacteriaceae.*

2.3.4. TSA - Tryptic Soy Agar. A non-selective soybean and casein enriched agar medium used for the growth of aerobic and facultative anaerobic bacteria.

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3. EQUIPMENT AND SUPPLIES

3.1. Sterile gloves, latex or nitrile

3.2. Inoculating loops, sterile

3.3. Glass serological disposable pipettes (1 mL), sterile and individually wrapped, with pipette pump; calibration checked on each new lot per SOP MB-03.3.

3.4. Plastic serological disposable pipettes (50 mL) sterile and individually wrapped, with pipette pump; calibration checked on each new lot per SOP MB-03.3.

3.5. Glass serological disposable pipettes (10 mL) sterile and individually wrapped, with pipette pump; calibration checked on each new lot per SOP MB-03.3.

3.6. Thermo Fisher Scientific Heratherm IMH 60 Model 51028066(S/N 42511812) or Fisher Scientific Isotemp Incubator Model 650F (S/N 303N0034) maintained at 35 ± 0.5 °C.

3.7. Water baths TSWB 4 (S/N 300471596) and TSWB 5 (S/N 300472033) maintained at 35.0 ± 0.5 °C. Water baths TSWB 3 (S/N 300311499) and TSWB 7 (S/N 300483022) maintained at 41.0 ± 0.5 °C, and TSWB 8 (S/N 300536247) and TSWB 9 (S/N 300561398) maintained at 44.5 ±0.2 °C.

3.8. Colilert-18 snap pack media (for the detection of E. coli and T. coli)

3.9. Enterolert snap pack media (for the detection of enterococci)

3.10. IDEXX Shrink-Banded Vessels without Sodium Thiosulfate

3.11. Antifoam dropper bottle

3.12. Quanti-Tray/2000 trays

3.13. UV light 365 nm, 6 watts

3.14. Colilert-18 Quanti-Tray/2000 comparators for E. coli and T. coli

3.15. Quanti-Tray rubber insert for Quanti-Tray/2000

3.16. Quanti-Tray Sealer Model Plus (S/N QTP 13215001064 or QTP 13181503550)

3.17. Vacuum Filtration system

3.18. Membrane filtration unit, sterile and wrapped in foil

3.19. UV Sterilizer unit

3.20. Forceps, sterile, straight or curved with smooth tip

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3.21. Membrane filters, sterile, white gridded, 47 mm diameter, 0.45±0.02 µm pore size

3.22. Bunsen burner

3.23. Beaker, 30 mL

4. REAGENTS AND STANDARDS

4.1. Microbiological Reference Materials:

4.1.1. Certified Reference Material (CRM)

4.1.2. *Escherichia coli NCTC 9001*

4.1.3. *Klebsiella variicola* ATCC 31488

4.1.4. *Enterococcus faecalis* NCTC 775

4.1.5. *Pseudomonas aeruginosa* NCTC 12951

4.1.6. *Streptococcus bovis* NCTC 8177

4.2. Microbiologics Kwik-Stik™ Reference Culture Microorganisms

4.2.1. *Pseudomonas aeruginosa* ATCC® 10145

4.2.2. *Aerococcus viridans* ATCC 10400

4.2.3. *Serratia marcesens* ATCC 43862

Note: Kwik-Stik cultures are prepared according to SOP MB-03.1

4.3. Dilu-Lok bottle with 99 mL Phosphate Buffer with $MgCl_2$ for membrane filtration method.

4.4. Autoclaved DI Water for Quanti- Tray

4.5. Tryptic Soy agar (TSA) - Prepare per SOP MB-02.33. Dry plates at room temperature for 24 hours prior to use.

5. SAMPLE PREPARATION PROCEDURE

5.1. Retrieve four vials of the target organism from the freezer and allow the capped vials to equilibrate to room temperature (15-30°C). Do not open vial until equilibration is complete. This should take about 15 minutes.

5.2. Once the sample pellet is at room temperature, open the vial and aseptically transfer the pellet to the hydration fluid. Shake vigorously to dissolve.

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5.3. Analyze spiked samples according to method requirements.

5.4. POSITIVE CONTROLS

5.4.1. For the detection of E. coli and T. coli, use the Universal Microbe CRM or a single pure E. coli pellet, and a single pure *K. variicola,* respectively. The Universal Microbe can be used for T. coli detection if K. variicola is not readily available. For the detection of enterococci, use a single pure E. *faecalis* pellet. Membrane filtration analysis requires four Dilu-Lok bottles spiked with the target bacteria and analyzed according to SOP MB-01.1. Quanti-Tray requires four spiked IDEXX vessels, containing Autoclaved DI Water, with the target bacteria and analyzed according to SOP MB-01.15

5.5. NEGATIVE CONTROLS

5.5.1. For Colilert-18 Quanti-tray analysis of E. coli and T. coli, use *Pseudomonas aeruginosa*, either pellet or plated working cultures, and spike one IDEXX vessel containing 100 mL of Autoclaved DI Water and analyze.

5.5.2. For Enterolert Quanti-Tray analysis of enterococci, use *Streptococcus bovis* and *E. coli* pellets (or *Escherichia coli* plated working culture) to spike one IDEXX vessel containing 100 mL Autoclaved DI Water for each culture and analyze. Plated working cultures *Aerococcus viridans* and *Serratia marcescens,* can also be used as negative controls.

5.5.3. For membrane filtration, spike one Dilu-Lok bottle with *Pseudomonas aeruginosa* and filter 0.1 mL.

6. SAMPLE ANALYSIS

6.1. For initial method performance (IPR), ongoing method performance (OPR), and initial demonstration of capability (IDOC) for analysts, prepare and analyze four Reference Samples by fortifying the corresponding hydration fluid with the target organism.

6.1.1. For membrane filtration, spike four Dilu-Lok bottles with a positive control microbe and analyze according to SOP MB-01.1. Filter 6.0, 7.0 and 8.0-mL volume if using the Universal water microbe cocktail. If using a single pure E. coli culture, the suggested dilutions are 10, 15, and 20 mL.

6.1.2. For Quanti-Tray analysis, spike four IDEXX vessels containing 100 mL of Autoclaved DI water with a positive control microbe and analyze according to SOP MB-01.15.

6.1.3. Analyze blanks and negative controls in addition to the four reference samples.

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6.2. Analysts must also complete a continuing demonstration of capability (CDOC) at least once annually. CDOCs can be satisfied by completion of Proficiency Tests (PTs), or media QC including a method blank and duplicate to meet precision requirements. Otherwise, CDOCs can be satisfied as described below.

6.2.1. For Quanti-Tray analysis, spike one IDEXX vessel containing 100 ml of autoclaved DI water with a positive control microbe. Shake vessel until pellet is completely dissolved and run sample as a 40 mL dilution with a duplicate. Include a method blank and analyze according to SOP MD-01.15. (A lower dilution volume for the sample spike can be used).

6.2.2. For membrane filtration, spike one Dilu-Lok with a positive control and analyze according to SOP MB-01.1. Filter 6.0, 7.0 and 8.0-mL volume if using the Universal water microbe cocktail. If using a single pure E. coli culture, the suggested dilutions are 10, 15, and 20 mL. Duplicate the highest dilution for precision and include a method blank.

7. CALCULATIONS AND PRECISION

7.1 Calculations of each spiked sample (4 for IDOC; 1 for CDOC) are made based off SOP MB 01.1 for membrane filtration and using the IDEXX MPN Generator (only for calculations, not for creating a Quanti-Tray Excel file) explained in SOP MB 01.15 for Quanti-Tray. Averages are calculated from these results for COA comparison.

7.2 Retrieve the Certificate of Analysis (COA) for the reference culture located in the following folder: \\floridadep\data\DEAR\Labs\Lab5\Biology\BenchCommon\COA\Micro supplies\cultures and check that the average results fall within the appropriate Acceptable Range specified.

7.3 Check precision, log range value and mean range, by comparing results against the microbiology Control Charts All Tests 032514 excel file, found in the following folder: \\floridadep\data\DEAR\Labs\Lab5\Biology\BenchCommon\Micro.

7.4 Enter the highest and lowest result out of the four IDOC spikes into Control Charts to determine precision and accuracy. For CDOC, enter the result of the spike and it's duplicate. Remove these numbers from control charts once precision results are collected, and enter them into the DOC excel file: IPR_OPR_data121823-w-form found in the following folder: X:\Labs\Lab5\Biology\BenchCommon\Micro\DOCs.

7.5 Record filter, incubation, read dates/times, analyst, buffered water and media used, QC record numbers for consumables used, and blanks and dups with their respective Test IDs for each sample spike in Prep Worksheet Manager per SOP BB-30. Use LIMS Prep Worksheet Manager to create a MICRO-QC worksheet. Laboratory → Technical → Prep Worksheet Manager→ Preparation ID: MICRO QC.

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8. SAFETY/HAZARDOUS WASTE MANAGEMENT

8.1. Always autoclave any expired or unused bacterial cultures for 45 minutes before discarding in the trash.

8.2. Expired or unused IDEXX media is collected by hazardous waste.

8.3. Discard any unused melted agar in hazardous waste container. Do not reuse media that has re-solidified.

8.4. Refer to Bureau of Laboratories Health and Safety Plan for safety equipment, use of laboratory hoods, and chemical storage policies.

8.5. Read SDS information for reagent handling and disposal.

9. REFERENCES

9.1. EPA Method 1600 (Sept 2014)

9.2. Standard Methods 9020 (2017)

9.3. Standard Methods 9222 (2006)

9.4. NELAC Quality Systems Manual

9.5. Difco™ & BBL™ Manual
\\floridadep\data\DEAR\Labs\Lab5\Biology\BenchCommon\Methods & Library Files

9.6. FDEP Laboratory Quality Manual https://floridadep.gov/dear/florida-dep-laboratory/content/dep-laboratory-quality-assurance-manual-and-sops

9.7. FDEP Laboratory Health and Safety Plan
https://floridadep.sharepoint.com/dear/Shared%20Documents/FDEP%20Health%20%20Safety%20Plan%20B%20-%20June%202021.pdf

9.8. DEP SOP MB-02.33, *Preparation of Dehydrated Culture Media*

9.9. DEP SOP MB-03.1, *Maintenance of Bacterial Cultures*

9.10. DEP SOP MB-03.3, *Calibration and Use of Mechanical and Disposable Pipettes*

9.11. IDEXX Quanti-Tray/2000 Colilert-18 method instructions

9.12. IDEXX Quanti-Tray/2000 Enterolert method instructions

9.13. IDEXX Quanti-Tray Sealer PLUS Manual

9.14. DEP SOP MB 01.1, *Membrane Filter Method for Fecal Coliforms*

9.15. DEP SOP MB 01.15, *Quanti-Tray/2000 Colilert-18 Method for Escherichia coli, Total coliforms, and Enterolert Method for Enterococci*

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APPENDIX OF CHANGES

02/19/2008 Added SOPs to references section and updated safety and hazardous waste management section.

02/11/2010 Added correct version of referenced method.

02/02/2011 Updated references to include new versions of Quality Manual and Health and Safety Plan.

01/05/2012 Removed procedure for ADB broth and added reference to procedure in media SOP.

06/20/2016 Added Summary of Method, including Interferences and Definitions. In Equipment and Supplies, included calibration SOPs and Incubation Model and Serial Number. In Reagents and Standards, detailed preparation and use of TSA and included *Escherichia coli*. (Also included *E. coli* throughout the SOP.) In Sub-section Sample Spiking, included Quanti-Tray procedure. Added Difco™ & BBL™ Manual and Pipette Calibration SOP to References.

12/18/2017 Modified sentence structure in the following areas: Summary of Method, Interferences, Undiluted spiking suspension (USS) and Serial dilution of USS. Added one significant figure to volume amounts and added the words Dilu-lok, species name, and Sample Spiking to figure 8.1.

01/03/2019 Updated references to match current versions.

12/19/2019 Added new equipment references used for incubation and storage purposes. Updated Flowchart of Spiking Suspension Procedure for specific DOC tests.

12/17/2020 Updated equipment references for incubators in use. Updated links for Quality Manual and Health and Safety Plan.

12/28/2021 Updated link for the Difco™ & BBL™ Manual. Added a note to section 5.4.1, and updated reference to the flowchart.

12/21/2022 Updated SOP by adding the new protocol to determine initial method performance (IPR), ongoing method performance (OPR), and demonstration of capability (DOC) for analysts. Former procedure remains in the SOP, unedited, to serve as an alternative method. Updated FDEP Laboratory Health and Safety Plan.

12/20/2023 Updated CDOC procedures for analysts. Updated hyperlinks. Added TSWB 9 to the equipment list. Updated Section 7 to include DOC calculations, location to LIMS modules for data entry and hyperlinks to data entry folders. Updated section 5 and 6 to clarify and remove redundancy on the procedure of analyzing negative controls. Section 10 was removed as it is no longer a procedure performed in the laboratory.

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