

ThermoFisher
S C I E N T I F I C

Food and Water Proficiency Tests
Public Health England Schemes

***External quality assurance programme: the key
indicator of quality in the lab***

Overview: Some Definitions

- Food and Water **Proficiency Tests**
Public Health England Schemes
- *External quality assurance programme: the key indicator of quality in the lab*

- **Proficiency tests:**
- An exam which test how proficiency or skilled someone is in a particular activity, field of study, etc

- **External quality assessment (EQA)** is the challenge of the effectiveness of a laboratories quality management system.
- The term **external** refers to the fact that an organizer outside of the laboratories organisation provides a statement of quality to the laboratory.

Overview: Organization

- Typically, an EQA scheme consists of several rounds per year. In each round several (or many) participants receive test items, which are also called **samples**.
- The EQA organizer is also called the provider.
- The organizer ensures that the test items are sufficiently similar and homogeneous, typically according to ISO 13528.
- Often the test items' properties and analyte concentrations are known to the organizer, but not disclosed to participants before the final report.
- The participants' results are then compared to check if any participant had a bias towards e.g. higher values, or an unexpected imprecision.
- At the end of each round, the EQA organizer sends out reports and/or certificates to the participating laboratories.

https://en.wikipedia.org/wiki/External_quality_assessment

Overview - Why laboratories do PT?

- To demonstrate competence as part of accreditation requirement – ISO/IEC 17025 - *General requirements for the competence of testing and calibration laboratories*
- Helps to provide assurance of the results obtained provided they are treated and processed the same as other samples
- Helps improve laboratory processes and understanding of regulation/legislation
- To remain up to date with new and emerging organisms - educational
- To challenge processes/media/training with difficult or atypical organisms
- To allow Inter-laboratory comparison of performance
- To support work tendered for as an accredited laboratory
- Because you enjoy the challenge and the educational value that participating in PT brings!



Impact of incorrect results – food microbiology

FALSE NEGATIVE

Reporting incorrectly that a sample does not contain a pathogen

- Unsafe product released for sale
- People become ill/die
- Negative publicity
- Loss of reputation
- Financial cost

FALSE POSITIVE

Reporting incorrectly that a sample does contain a pathogen

- Product recall
- Incorrect product withdrawal
- Ban on export
- Loss of contracts
- Financial cost



Sources of error

- sample handling and/or processing errors
- inadequate staff training
- lack of understanding of legislation/guidelines
- incorrect methods or inappropriate media used
- equipment and culture media failures
- calculation and reporting errors



Why you benefit?

- **You**, the laboratory that examines food and water samples **need to produce test results that are reliable, accurate and clear**
- This helps to ensure that **the public is protected** from harm as you have assurance from PT your ability to confirm that the food or water released onto the market for sale is safe to consume or the water is safe to use
- Helps you to **identify gaps** in your processes, highlighting where quality improvements can be made
- It provides an opportunity to **improve staffs knowledge and experience** with organisms not frequently encountered
- Satisfactory performance with your PT provides assurance that your laboratory is **compliant with testing standards**, thereby meeting and maintaining accreditation requirements



Food and Water Proficiency Tests: Public Health England Schemes



Public Health
England

Proficiency testing
for food and
water microbiology

Public Health England's Proficiency Testing (PT) Schemes

- Public Health England (PHE) is an executive agency of the United Kingdom's Department of Health.
- PHE is the expert national public health agency which fulfils the Secretary of State for Health's statutory duty to protect health and address inequalities, and executes his power to promote the health and wellbeing of the nation.
- Our mission is to protect and improve the nation's health and to address inequalities, working with national and local government, the NHS, industry, academia, the public and the voluntary and community sector



Public Health England's Proficiency Testing (PT) Schemes

- Food and water examination laboratories play a vital role in protecting people's health by ensuring that food and waters are safe and do not pose a threat to health.
- One of PHE's goal is protecting the country from infectious diseases and environmental hazards, including the growing problem of infections that resist treatment with antibiotics.
- In support of this goal, PHE provides tools such as proficiency testing (PT) schemes and reference materials to support food and water microbiology laboratories in assuring their results.
- The PT schemes are used by laboratories that take quality seriously and understand the impact of the work they undertake.

Public Health England's Proficiency Testing (PT) Schemes

- **The vision** for the Food and Environmental Proficiency Testing Unit (FEPTU) aims to be: the leader in providing international PT schemes for food and water microbiology, supporting public health and raising awareness of:
 - the importance of producing accurate laboratory results
 - the impact of incorrect laboratory results on public health

- **The mission** for FEPTU is: 'To provide microbiology testing laboratories with reliable and robust proficiency testing samples to help to safeguard the public from potential harm that may be encountered from food and water sources. Our services will be exemplary, our resources will be used effectively and our experience and knowledge will be used to develop and foster success with our participants

Public Health England's Proficiency Testing (PT) Schemes

- Proficiency testing schemes are sometimes referred to as external quality assessment (EQA) schemes.
- FEPTU has been providing international PT schemes for food and water microbiology for more than 20 years.
- All the schemes are accredited by the United Kingdom Accreditation Service (UKAS) to the international standard ISO 17043: 2010 Competency assessment – General requirement for proficiency testing.
- PHE PT schemes are suitable for food and water microbiology laboratories worldwide in the food and water industries, private sector, and the public health and environmental health sectors.
- The schemes are provided by experts and driven by impact on public health and quality of service. These are uncompromised by commercial pressures.



Sample Type

- **Freeze-dried samples**

- Freeze-drying is a well-established process for preserving a wide range of micro-organisms and is used by many culture collections such as the United Kingdom's National Collection of Type Cultures (NCTC®). FEPTU use a freeze-drying process to prepare samples for a range of food EQA schemes.
- The samples are stable at ambient temperature although storage in refrigerated conditions is normally recommended

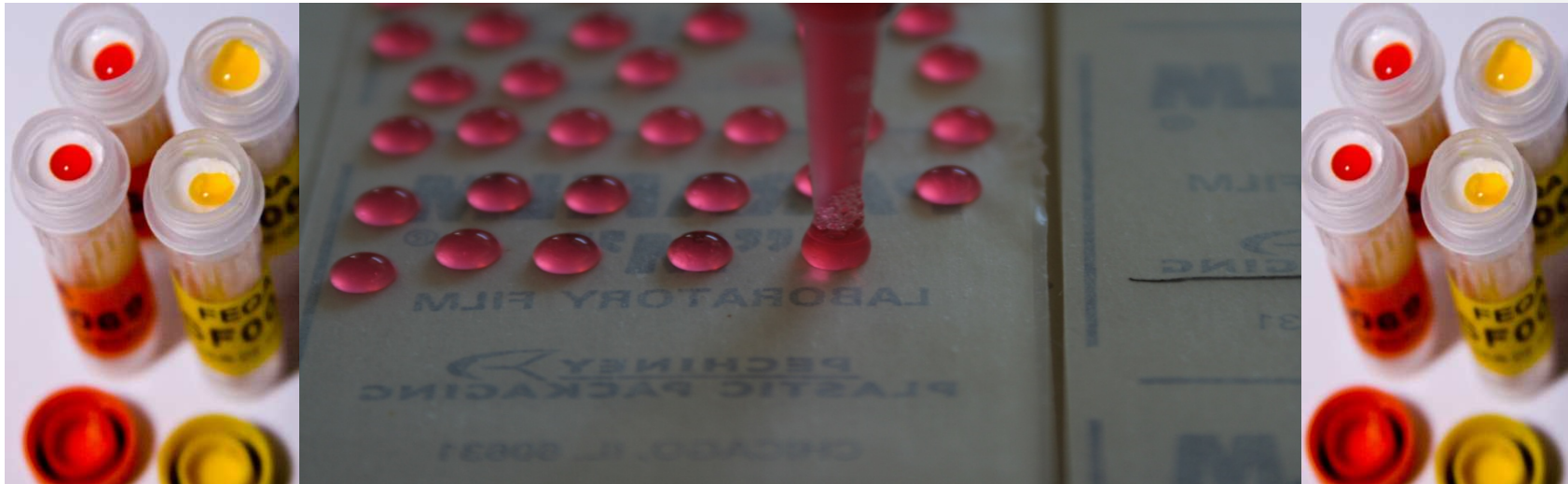
- **LENTICULE® discs**

- LENTICULE discs consist of control-dried cultures of viable micro-organisms that require storage at -20°C. They can be prepared to contain a wide range of micro-organisms, in pure or mixed cultures, at levels between 10 and 10⁸ cfu per LENTICULE disc. LENTICULE discs are used by PHE to prepare EQA samples for food and water microbiology schemes.



What Is a LENTICULE DISC?

- Small disc that contains control-dried cultures of viable microorganisms
- A preservation method unique to PHE and used in FEPTU for producing PT samples, reference materials and control strains



PT Sample Design

- Samples are designed to identify problems that may impact on people's health
- Wild-type strains used, **not** NCTC or ATCC cultures
- All strains fully characterised
- Samples provide a realistic challenge for routine examinations



Schemes Focus On:

- variations of methods used
- atypical organisms
- enumeration challenges associated with low levels
- limitation of confirmation tests



Statistic and Scoring

- Robust statistics used to evaluate enumeration results
- Scoring systems used for most schemes
- Provide Z-scores for some schemes
- Long-term evaluation of performance for all schemes with scoring systems
- Comparison of participants' results included
- Trend analyses available



PHE PT Participants

- Food and water industry
- Private testing laboratories
- Public health laboratories
- Government laboratories
- Hospital laboratories



Public Health
England



Participating Countries

| | | | | |
|-----------------|------------------|------------------|--------------|----------------|
| Albania | Cyprus | Jamaica | Peru | Thailand |
| Angola | Czech Republic | Japan | Philippines | Turkey |
| Argentina | Denmark | Korea | Poland | UAE |
| Austria | Estonia | Kuwait | Portugal | United Kingdom |
| Belgium | Falkland Islands | Latvia | Romania | USA |
| Bosnia | Finland | Liechtenstein | Saudi Arabia | Vietnam |
| Botswana | France | Lithuania | Serbia | |
| Brazil | Germany | Luxembourg | Singapore | |
| Bulgaria | Greece | Macedonia | Slovakia | |
| Canada | Hong Kong | Malta | Slovenia | |
| Cape Verde | Hungary | Namibia | South Africa | |
| Channel Islands | India | Netherlands | Spain | |
| China | Indonesia | Antilles | St. Helena | |
| Colombia | Ireland | Northern Ireland | Sweden | |
| Costa Rica | Israel | Norway | Switzerland | |
| Croatia | Italy | | Taiwan | |



General Schemes Features (1)

- All schemes have a yearly schedule
- Every set of samples (round) for a single scheme is referred to as a 'Distribution'
- Normally 2 to 6 distributions per year – depending on the scheme
- Every distribution has a unique distribution number
- Every distribution contains 2 or 3 samples each with a unique sample number
- Participants must receive all the samples from a distribution, but not all the distributions (minimum number required per year to give meaningful performance)

General Schemes Features (2)



- All schemes are accredited to 17043
- Samples that are stable and homogeneous
- All samples are designed, prepared and tested by in PHE microbiologists
- Quality control tests reflect commonly used methods
- Samples that challenge particular examinations e.g. isolation, identification, enumeration
- Micro-flora representative of real food and water samples

General Schemes Features (3)

- Significantly high proportion of positive samples
- Realistic levels of target organisms with background flora where appropriate
- Large numbers of participants (between 50 – 350 in each scheme) for robust data analyses
- High-level of international participation
- Efficient global sample delivery service
- Clear instructions and request forms

General Schemes Features (4)

- Easy-to-follow statistic and scoring systems
- Informative, educational and timely reports
- Continuous performance assessment reports
- Total confidentiality of performance
- Technical support from expert food and water microbiologists
- Informative web pages and on-line tools

Microbiology PT scheme - process

PHE - FEPTU

Prepare samples

Analyse results

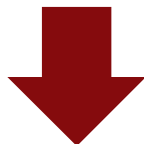
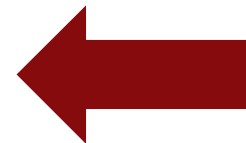
Prepare report

Participants

Examine samples

Report results

Evaluate



PHE Food and Water Schemes

- [Environmental swab scheme](#)
- [European food microbiology legislation scheme](#)
- [Non-pathogen scheme](#)
- [Norovirus and hepatitis A virus scheme](#)
- [Pathogenic vibrio scheme](#)
- [Shellfish scheme](#)
- [Shiga toxin Escherichia coli scheme](#)
- [Standard scheme](#)
- [Staphylococcus aureus enterotoxin \(SET\) detection scheme](#)

- [Bottled and mineral water scheme](#)
- [Drinking water scheme](#)
- [Legionella isolation scheme](#)
- [Legionella molecular scheme](#)
- [Recreational and surface water scheme](#)

- [Mycobacterium sp in water scheme](#)
- [Hospital tap water scheme](#)
- [Dialysis water scheme](#)
- [Endoscope rinse water scheme](#)

Food and Water Proficiency Tests: Public Health England Schemes



Public Health
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Protecting and improving the nation's health

Food schemes



Image by Zak Prior

Standard Scheme



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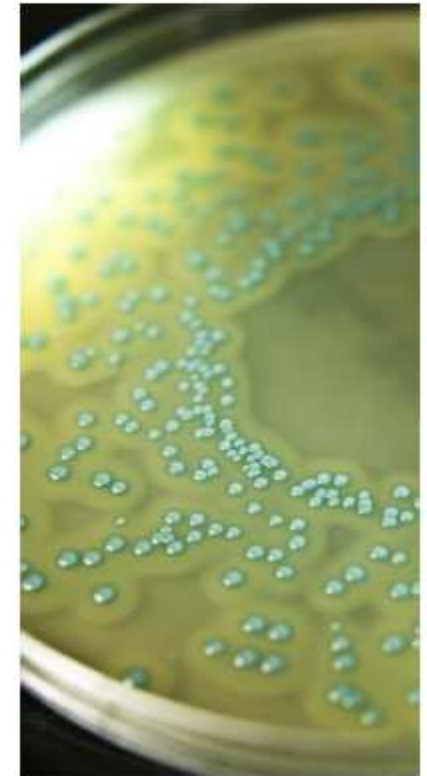
Standard Scheme

This microbiology scheme provides proficiency testing (PT) samples to food laboratories that routinely test for a range of food-borne pathogens and indicator organisms. To ensure that the public is protected from harm, food released onto the market for sale must be safe to be consumed.

This scheme helps you to identify gaps in your processes, highlighting where quality improvements can be made. It also provides an opportunity to improve staffs knowledge and experience with organisms not frequently encountered.

The scheme focuses on raising awareness of:

- the variations of methods used which can highlight differences in PT results
- atypical organisms circulating in the environment that may challenge a laboratory's interpretation of the PT sample
- the challenges associated with enumerating food samples and limitation of confirmation tests



Standard Scheme

STANDARD SCHEME

food microbiology examinations

Sample schedule for 2019

| Distribution number | Sample numbers | Dispatch date | Date results due by | Examinations and enumerations required Up to three sets of results can be reported |
|---------------------|------------------|---------------|---------------------|---|
| 312 | \$0661 \$0662 | 07/01/2019 | 15/02/2019 | <i>Campylobacter</i> spp. detection <i>Campylobacter</i> spp. enumeration <i>Escherichia coli</i> O157 detection <i>Salmonella</i> spp. detection Aerobic colony count <i>Enterobacteriaceae</i> enumeration |
| 314 | \$0665 \$0666 | 18/02/2019 | 29/03/2019 | Presumptive <i>Bacillus cereus</i> enumeration Coagulase-positive staphylococci enumeration <i>Listeria</i> spp. (including <i>L. mono</i>) enumeration <i>Listeria monocytogenes</i> enumeration Aerobic colony count Coliform enumeration |
| 316 | \$0669 \$0670 | 08/04/2019 | 17/05/2019 | <i>Cronobacter</i> spp. detection <i>Listeria</i> spp. (including <i>L. mono</i>) detection <i>Listeria monocytogenes</i> detection <i>Clostridium perfringens</i> enumeration Aerobic colony count <i>Escherichia coli</i> enumeration |
| 318 | \$0673 \$0674 | 03/06/2019 | 12/07/2019 | <i>Campylobacter</i> spp. detection <i>Campylobacter</i> spp. enumeration <i>Escherichia coli</i> O157 detection <i>Salmonella</i> spp. detection Aerobic colony count <i>Enterobacteriaceae</i> enumeration <i>Yersinia enterocolitica</i> |
| 320 | \$0677 \$0678 | 29/07/2019 | 06/09/2019 | Presumptive <i>Bacillus cereus</i> enumeration Coagulase-positive staphylococci enumeration <i>Listeria</i> spp. (including <i>L. mono</i>) enumeration <i>Listeria monocytogenes</i> enumeration Aerobic colony count Coliform enumeration |
| 322 | \$0681 \$0682 | 30/09/2019 | 08/11/2019 | <i>Listeria</i> spp. (including <i>L. mono</i>) detection <i>Listeria monocytogenes</i> detection <i>Salmonella</i> spp. detection <i>Clostridium perfringens</i> enumeration Sulfite-reducing anaerobic bacteria Aerobic colony count <i>Escherichia coli</i> enumeration |

Note for this scheme molecular methods can be used for pathogen examination or as a confirmation test

European Food Microbiology Legislation Scheme



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European Food Microbiology Legislation Scheme



This unique microbiological scheme provides proficiency testing samples to laboratories that examine food products in accordance with European legislation specified in Regulation (EC) 2073/2005 Microbiological Criteria for Foodstuffs associated with Regulation (EC) 852/2004 and subsequent amendments.

The scheme assesses participants' ability to test and interpret laboratory results in accordance with EU food safety and process hygiene criteria. This scheme is of particular importance to nominated national reference and official control laboratories as part of compliance to Regulation (EC) 882/2004.

This PT scheme helps you to identify gaps in your processes, highlighting where quality improvements can be made. It also provides an opportunity to improve staffs knowledge and experience with organisms not frequently encountered.

This scheme focuses on raising awareness:

- of the test requirements for a particular food category and how to interpret microbiological results for batches
- of updates in legislation and testing requirements
- of the challenges associated with enumerating food samples
- of atypical organisms circulating in the environment that may challenge a laboratory's interpretation of the PT sample due to methods, media used and confirmation tests done

European Food Microbiology Legislation Scheme

EUROPEAN MICROBIOLOGY LEGISLATION SCHEME

food microbiology examinations

Sample schedule for 2019

| Distribution number | Sample numbers | Dispatch date | Dates results due by | Food category | Examinations and enumerations required |
|---------------------|----------------------------|---------------|----------------------|----------------------|--|
| EFL47 | EFL139 EFL140 EFL141 | 07/01/2019 | 08/02/2019 | Meat | According to the Food Safety or Process Hygiene Criteria |
| EFL48 | EFL142 EFL143 EFL144 | 18/02/2019 | 22/03/2019 | Dairy | |
| EFL49 | EFL145 EFL146 EFL147 | 03/06/2019 | 05/07/2019 | Ready to eat product | |
| EFL50 | EFL148 EFL149 EFL150 | 30/09/2019 | 01/11/2019 | Miscellaneous | |

Non-Pathogen Scheme



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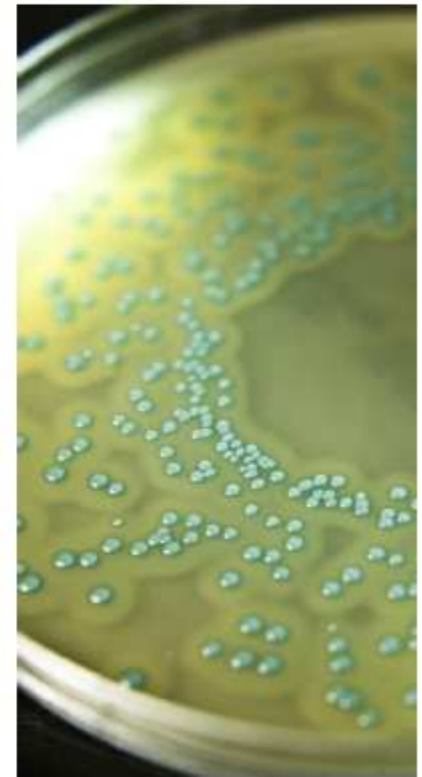
This microbiology scheme provides proficiency testing (PT) samples to laboratories that examine food samples for spoilage and indicator organisms. This scheme is suitable for laboratories on food production sites that do not want to introduce pathogens and ensure products are contamination free.

This scheme helps you to identify gaps in your processes, highlighting where quality improvements can be made. It also provides an opportunity to improve staffs knowledge and experience with organisms not frequently encountered.

The scheme focuses on raising awareness of:

- the variations of methods used which can highlight differences in PT results
- atypical organisms circulating in the environment that may challenge a laboratory's interpretation of the PT sample
- the challenges associated with enumerating food samples and limitation of confirmation tests

Non-Pathogen Scheme



Non-Pathogen Scheme

NON-PATHOGEN SCHEME

food microbiology examinations

Sample schedule for 2019

| Distribution number | Sample numbers | Dispatch date | Date results due by | Examinations and enumerations required <i>Up to three sets of results can be reported</i> |
|---------------------|------------------|---------------|---------------------|--|
| NP062 | NP0179 NP0180 | 07/01/2019 | 22/02/2019 | Pseudomonas spp. Yeasts Moulds Coliforms Enterobacteriaceae |
| NP063 | NP0181 NP0182 | 08/04/2019 | 24/05/2019 | Escherichia coli Enterococci Lactic acid bacteria Aerobic colony count at 30°C |
| NP064 | NP0183 NP0184 | 29/07/2019 | 13/09/2019 | All the above enumerations are included for all samples An option to register for PYM (Pseudomonas spp., yeasts and moulds only) is available |

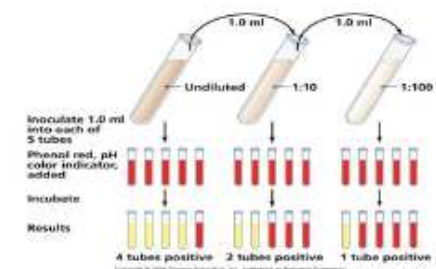
Shellfish Scheme



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Shellfish Scheme



This unique microbiology scheme provides proficiency testing (PT) samples to laboratories that examine raw bivalve molluscs from harvesting sites in accordance with Regulation (EC) No. 854/2004 and from the production chain between harvest and consumption, in accordance with Regulation (EC) 2073/2005 and subsequent amendments. This scheme is of particular importance to nominated national reference and official control laboratories as part of compliance to Regulation (EC) 882/2004.

The scheme is organised in collaboration with the Centre for Environment, Fisheries and Aquaculture Science (Cefas), Weymouth, United Kingdom (UK) and helps you to identify gaps in your processes, highlighting where quality improvements can be made. It also provides an opportunity to improve staffs knowledge and experience with organisms not frequently encountered.

PHE's Shellfish Scheme focuses on raising awareness of:

- interpreting tube combination results and the associated most probable numbers value per 100g for *Escherichia coli*
- updates in ISO methods

Shellfish Scheme

SHELLFISH SCHEME food microbiology examinations

Sample schedule for 2019

| Distribution number | Sample numbers | Dispatch date | Date results due by | Examinations and enumerations required |
|---------------------|------------------|---------------|---------------------|---|
| SF062 | SF0132 SF0133 | 18/02/2019 | 22/03/2019 | Escherichia coli MPN Salmonella spp. dection |
| SF063 | SF0134 SF0135 | 03/06/2019 | 05/07/2019 | |
| SF064 | SF0136 SF0137 | 30/09/2019 | 01/11/2019 | |

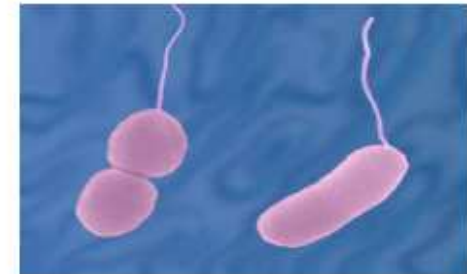
Pathogenic Vibrio Scheme



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Pathogenic *Vibrio* Scheme



This unique microbiology scheme provides proficiency testing (PT) samples to laboratories that examine *Vibrio* spp. in food and water samples. This scheme challenges the detection of predominant species of vibrio that are of public health concern. Also included as part of the scheme design is enumeration of *Vibrio parahaemolyticus*.

This PT scheme helps you to identify gaps in your processes, highlighting where quality improvements can be made. It also provides an opportunity to improve staffs knowledge and experience with organisms not frequently encountered.

PHE's Pathogenic *Vibrio* Scheme focuses on raising awareness of:

- the different *Vibrio* spp. that may be isolated from food or water samples
- the limitations of confirmation tests for *Vibrio* spp.
- culture media and batch to batch variations that may exist when analysing food or water samples
- the challenges associated with enumerating *V. parahaemolyticus* in food samples
- the increasing significance of other *Vibrio* spp. in food or water samples

Pathogenic Vibrio Scheme

PATHOGENIC VIBRIO SCHEME

food microbiology examinations

Sample schedule for 2019

| Distribution number | Sample numbers | Dispatch date | Date results due by | Examinations and enumerations required |
|---------------------|----------------|---------------|---------------------|--|
| V055 | V0150 V0151 | 18/02/2019 | 22/03/2019 | Detection: <i>Vibrio cholerae</i> non-O1 and non-O139 strains <i>Vibrio vulnificus</i> |
| V056 | V0152 V0153 | 03/06/2019 | 05/07/2019 | Detection and enumeration: <i>Vibrio parahaemolyticus</i> |
| V057 | V0154 V0155 | 30/09/2019 | 01/11/2019 | |

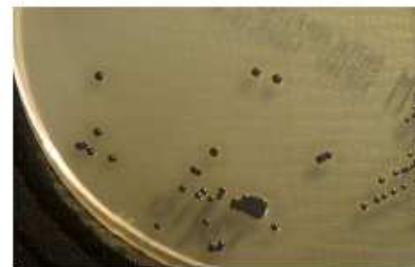
Staphylococcus aureus enterotoxin Scheme



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Staphylococcus aureus enterotoxin Scheme



This unique microbiology scheme provides proficiency testing (PT) samples to laboratories that examine for *Staphylococcus aureus* enterotoxin in food samples. Staphylococcal food poisoning is caused by consumption of food or beverages containing pre-formed enterotoxins. The scheme is suitable for laboratories that test food for *Staphylococcus aureus* enterotoxins (A-E) using a range of kits and methods.

EC Regulation 2073/2005 lays down microbiological criteria for various combinations of food commodities and microorganisms, their toxins or metabolites. *Staphylococcus aureus* enterotoxins are extremely difficult to eliminate from foods, they are resistant to heat, freezing and irradiation. They will survive commercial pasteurisation processes and may even survive processes used for the sterilisation of canned foods.

This PT scheme helps you to identify gaps in your processes, highlighting where quality improvements can be made. It also provides an opportunity to improve staffs knowledge and experience with organisms not frequently encountered.

This scheme focuses on raising awareness of:

- the different methods used to detect enterotoxins in food samples
- the limitation and issues with some of the kit methods available

Staphylococcus aureus enterotoxin Scheme

STAPHYLOCOCCUS AUREUS ENTEROTOXIN SCHEME

food microbiology examinations

Sample schedule for 2019

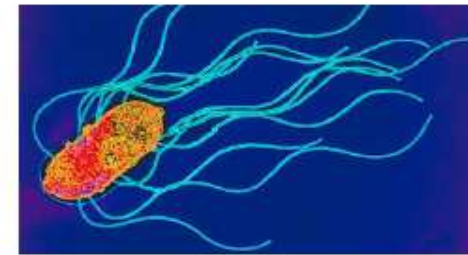
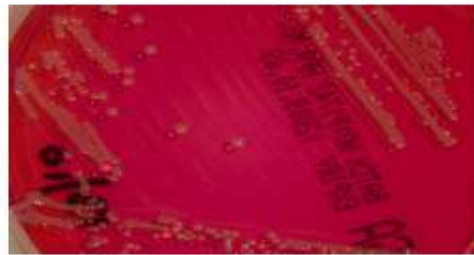
| Distribution number | Sample numbers | Dispatch date | Date results due by | Examinations required |
|---------------------|------------------|---------------|---------------------|--|
| STA040 | ST0079 ST0080 | 07/01/2019 | 08/02/2019 | <i>Staphylococcus aureus</i> enterotoxin |
| STA041 | ST0081 ST0082 | 08/04/2019 | 10/05/2019 | |
| STA042 | ST0083 ST0084 | 29/07/2019 | 30/08/2019 | |

Environmental Swab Scheme



Protecting and improving the nation's health

Environmental Swab Scheme



This microbiology scheme provides proficiency testing (PT) samples to laboratories that examine swabs for microbial pathogens and/or hygiene indicator organisms. The scheme focuses on raising awareness of pathogens implicated in foodborne outbreaks, the challenges associated with enumerating samples and the accurate reporting of results from random/template area swab.

Determining that foodborne pathogens are absent in ready to eat food preparation area or retail setting is of particular importance to prevent public health incidents.

Determination of the number of aerobic viable micro-organisms, *Escherichia coli* and *Enterobacteriaceae* on a specified area of a surface can provide an indication of cleanliness/sanitation. This allows monitoring of cleaning procedures over time, providing useful information on general environmental conditions.

This scheme allows you to:

- assure your calculation ability for the different swab areas (random and template)
- have confidence in determining the pathogen/s likely to be implicated based on an outbreak scenario provided

Environmental Swab Scheme

| ENVIRONMENTAL SWAB SCHEME | | | | |
|--------------------------------|------------------|---------------|---------------------|--|
| food microbiology examinations | | | | |
| Sample schedule for 2019 | | | | |
| Distribution number | Sample numbers | Dispatch date | Date results due by | Examinations required |
| ES14 | ES0027 ES0028 | 18/02/2019 | 15/03/2019 | Relevant pathogen relating to outbreak details as provided on the request form Campylobacter spp. Coagulase-positive staphylococci Escherichia coli O157 Listeria monocytogenes Salmonella spp. |
| ES15 | ES0029 ES0030 | 08/04/2019 | 03/05/2019 | <u>Enumeration of hygiene indicator organisms</u> Aerobic colony count Bacillus cereus Escherichia coli Enterobacteriaceae Listeria spp. |
| ES16 | ES0031 ES0032 | 29/07/2019 | 23/08/2019 | Relevant pathogen relating to outbreak details as provided on the request form Campylobacter spp. Coagulase-positive staphylococci Escherichia coli O157 Listeria monocytogenes Salmonella spp. |
| ES17 | ES0033 ES0034 | 30/09/2019 | 25/10/2019 | <u>Enumeration of hygiene indicator organisms</u> Aerobic colony count Bacillus cereus Escherichia coli Enterobacteriaceae Listeria spp. |

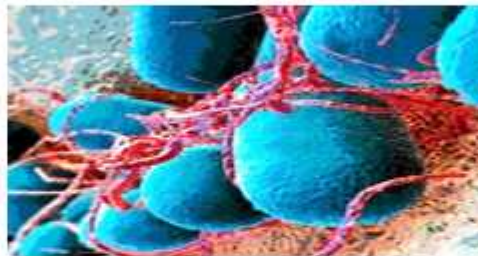
Shiga Toxin-Producing *Escherichia coli* Scheme



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Shiga toxin-producing *Escherichia coli* Scheme



This microbiology molecular scheme provides proficiency testing (PT) samples to laboratories that analyse food samples for Shiga toxin-producing *Escherichia coli* (STEC) in accordance with European legislation Commission Regulation (EU) No 209/2013 amending Regulation (EC) No 2073/2005 as regards microbiological criteria for sprouts and the sampling rules for poultry carcasses and fresh poultry meat.

The scheme focuses on raising awareness of:

- the variations in the molecular methods used which can highlight differences in PT results obtained
- the limitation of molecular methods available on the market
- updates and compliance with ISO methods or other standards
- the correct interpretation to apply on PCR results obtained

Shiga Toxin-Producing Escherichia coli Scheme

SHIGA TOXIN-PRODUCING *ESCHERICHIA COLI* SCHEME

food microbiology examinations

Sample schedule for 2019

| Distribution number | Sample numbers | Dispatch date | Date results due by | Examinations required Samples can only be examined using molecular methods |
|---------------------|------------------|---------------|---------------------|--|
| STX6 | STX011 STX012 | 07/01/2019 | 01/02/2019 | Detection of the major virulence genes associated with <i>Escherichia coli</i> serogroups O157, O111, O26, O103, O145 and O104:H4 (STEC) |
| STX7 | STX013 STX014 | 03/06/2019 | 28/06/2019 | Includes detection of <i>stx</i> -coding genes and the presence of the intimin-coding gene <i>eae</i> |

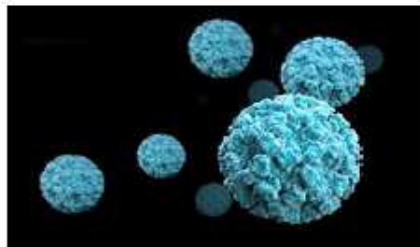
Norovirus & Hepatitis A Virus Scheme



Norovirus & Hepatitis A Virus Scheme



Protecting and improving the nation's health



This scheme provides proficiency testing (PT) samples to laboratories that examine food products or waters for hepatitis A virus and norovirus GI and GII using the reverse-transcription polymerase chain reaction (RT-PCR). This PT challenges laboratories in detection and quantification (copies per sample) of both these viruses. Implementing Regulation (EU) 2017/1142 (amending Regulation (EC) 669/2009) requires virus testing of frozen raspberries imported from Serbia, while Regulation (EC) 2073/2005 recommends adoption of virus testing for live bivalve molluscs once methods are developed.

This scheme is organised in collaboration with Cefas as the European Union Reference Laboratory (EURL) for monitoring bacteriological and viral contamination of bivalve molluscs. This scheme helps you to identify gaps in your processes, highlighting where quality improvements can be made. It also provides an opportunity to improve staffs knowledge and experience with organisms not frequently encountered.

The scheme focuses on raising awareness of:

- the variations of methods used which can highlight differences in PT results
- the limitation of methods available on the market
- updates in ISO methods

Norovirus & Hepatitis A Virus Scheme

NOVAVIRUS AND HEPATITIS A VIRUS SCHEME

food and water microbiology examinations

Sample schedule for 2019

| Distribution number | Sample numbers | Dispatch date | Date results due by | Examinations required Samples can only be examined using molecular methods |
|---------------------|--------------------|---------------|---------------------|---|
| NHV005 | NHV0009 NHV0010 | 18/02/2019 | 22/03/2019 | Detection of Norovirus GI and GII and Hepatitis A Virus |
| NHV006 | NHV0011 NHV0012 | 30/09/2019 | 01/11/2019 | Optional: Quantification of Norovirus GI and GII and Hepatitis A Virus |

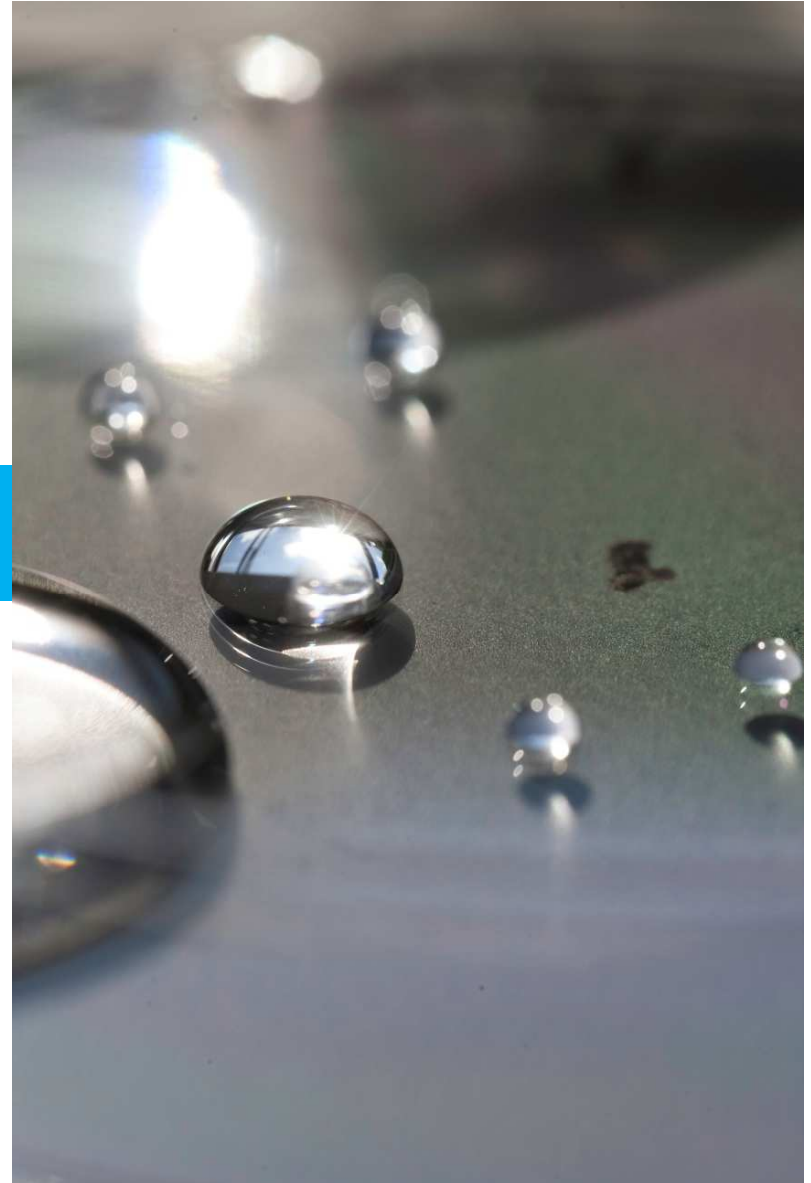


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Water schemes

Image by Zak Prior



Legionella Isolation Scheme



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Legionella Isolation Scheme



This microbiology scheme provides proficiency testing (PT) samples to laboratories that examine waters for legionellae. This scheme challenges the detection and accurate enumeration of different serogroups of *Legionella pneumophila*, and other *Legionella* spp..

Legionella spp. are the causative agent of legionellosis infections, varying in severity from a mild self-limiting febrile illness (Pontiac fever) to a potentially fatal atypical pneumonia (Legionnaires' disease). *Legionella* is recognised as a significant cause of sporadic and epidemic community-acquired and nosocomial-acquired pneumonia with many cases being associated with travel making it difficult to identify the source of infection.

This PT scheme helps you to identify gaps in your processes, highlighting where quality improvements can be made. It also provides an opportunity to improve staffs knowledge and experience with organisms not frequently encountered.

Legionella Isolation Scheme

LEGIONELLA ISOLATION SCHEME

water microbiology examinations

Sample schedule for 2019

| Distribution number | Sample numbers | Dispatch date | Date results due by | Examinations and enumerations required |
|---------------------|----------------|---------------|---------------------|--|
| G114 | G114A G114B | 28/01/2019 | 08/03/2019 | Detection, enumeration and identification of Legionella spp. |
| G115 | G115A G115B | 13/05/2019 | 21/06/2019 | |
| G116 | G116A G116B | 01/07/2019 | 09/08/2019 | |
| G117 | G117A G117B | 04/11/2019 | 13/12/2019 | |

Legionella Molecular Scheme



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Legionella Molecular Scheme



This unique microbiology scheme provides proficiency testing samples to laboratories that examine waters for legionellae using molecular platforms. Both detection and genomic quantification results are assessed.

Legionella spp. are the causative agent of legionellosis infections, varying in severity from a mild self-limiting febrile illness (Pontiac fever), to a potentially fatal atypical pneumonia (Legionnaires' disease). *Legionella* is recognised as a significant cause of sporadic and epidemic community-acquired and nosocomial-acquired pneumonia with many cases being associated with travel making it difficult to identify the source of infection.

Legionella Molecular Scheme

LEGIONELLA MOLECULAR SCHEME

water microbiology examinations

Sample schedule for 2019

| Distribution number | Sample numbers | Dispatch date | Date results due by | Examinations and enumerations required Samples can only be examined using molecular methods |
|---------------------|----------------|---------------|---------------------|--|
| LM5 | LM5A LM5B | 11/03/2019 | 19/04/2019 | Detection and quantification of Legionella spp. |
| LM6 | LM6A LM6B | 02/09/2019 | 11/10/2019 | |

Recreational and Surface Water Scheme



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Recreational and Surface Water Scheme

This microbiology scheme provides proficiency testing (PT) samples to laboratories that examine recreational and surface waters routinely for microbial contents. This scheme challenges the accurate enumeration of micro-organisms and detection of pathogen that maybe present in these types of waters. Sample types included in this scheme design are river, lakes and streams, bathing beach (marine), swimming pool and hydrotherapy pool waters.

This PT scheme helps you to identify gaps in your processes, highlighting where quality improvements can be made. It also provides an opportunity to improve staffs knowledge and experience with organisms not frequently encountered.

PHE's Recreational and Surface Water Scheme focuses on raising awareness of:

- the challenges associated with enumerating samples containing low levels of organisms
- variations in the methods used which can highlight differences in results obtained
- issues with confirmatory tests carried out on isolates
- updates in local guidelines which maybe different to European ones



Recreational and Surface Water Scheme

| RECREATIONAL AND SURFACE WATER SCHEME | | | | |
|---------------------------------------|----------------|---------------|---------------------|---|
| water microbiology examinations | | | | |
| Sample schedule for 2019 | | | | |
| Distribution number | Sample numbers | Dispatch date | Date results due by | Examinations and enumerations required |
| S88 | S88A S88B | 28/01/2019 | 01/03/2019 | <u>swimming pool / hydrotherapy pool waters:</u> Coagulase-positive staphylococci Coliform bacteria <i>Escherichia coli</i> Enterococci <i>Pseudomonas aeruginosa</i> Total staphylococci Colony count (37°C/24 hours) |
| S89 | S89A S89B | 11/03/2019 | 12/04/2019 | <u>marine (bathing beach) waters:</u> <i>Escherichia coli</i> Enterococci <i>Salmonella</i> spp. |
| S90 | S90A S90B | 13/05/2019 | 14/06/2019 | <u>river, lake or stream waters:</u> Coliform bacteria <i>Escherichia coli</i> Enterococci Faecal coliforms <i>Clostridium perfringens</i> <i>Salmonella</i> spp. |
| S91 | S91A S91B | 01/07/2019 | 02/08/2019 | <u>swimming pool / hydrotherapy pool waters:</u> Coagulase-positive staphylococci Coliform bacteria <i>Escherichia coli</i> Enterococci <i>Pseudomonas aeruginosa</i> Total staphylococci Colony count (37°C/24 hours) |
| S92 | S92A S92B | 02/09/2019 | 04/10/2019 | <u>marine (bathing beach) waters:</u> <i>Escherichia coli</i> Enterococci <i>Salmonella</i> spp. |
| S93 | S93A S93B | 04/11/2019 | 06/12/2019 | <u>river, lake or stream waters:</u> Coliform bacteria <i>Escherichia coli</i> Enterococci Faecal coliforms <i>Clostridium perfringens</i> <i>Salmonella</i> spp. |

Drinking Water Scheme



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Drinking Water Scheme



This microbiology scheme provides proficiency testing (PT) samples to laboratories that examine drinking waters routinely. This scheme challenges the accurate enumeration of low levels of micro-organisms. Drinking water can contain many different bacteria that can be a risk to health, it is essential that all water intended for drinking is properly treated and routinely effectively monitored for safe consumption.

This PT scheme helps you to identify gaps in your processes, highlighting where quality improvements can be made. It also provides an opportunity to improve staffs knowledge and experience with organisms not frequently encountered.

PHE's Drinking Water Scheme focuses on raising awareness of:

- the challenges associated with enumerating samples containing low levels of organisms
- variations in the methods used which can highlight differences in results obtained
- issues with confirmatory tests carried out on isolates
- updates in local guidelines or policies

Drinking Water Scheme

DRINKING WATER SCHEME

water microbiology examinations

Sample schedule for 2019

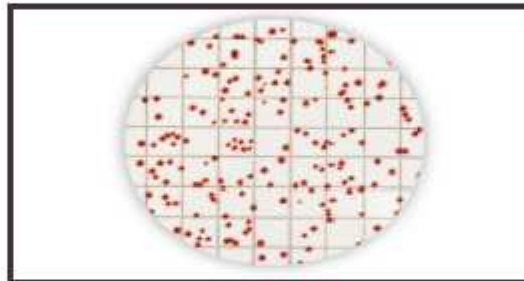
| Distribution number | Sample numbers | Dispatch date | Date results due by | Examinations and enumerations required Up to three sets of results can be reported |
|---------------------|-------------------------|---------------|---------------------|---|
| W185 | W185A W185B W185C | 28/01/2019 | 01/03/2019 | Coliform bacteria Escherichia coli Enterococci Clostridium perfringens Pseudomonas aeruginosa Colony count (37°C/48 hours) Colony count (22°C/72 hours) |
| W186 | W186A W186B W186C | 11/03/2019 | 12/04/2019 | |
| W187 | W187A W187B W187C | 13/05/2019 | 14/06/2019 | |
| W188 | W188A W188B W188C | 01/07/2019 | 02/08/2019 | |
| W189 | W189A W189B W189C | 02/09/2019 | 04/10/2019 | |
| W190 | W190A W190B W190C | 04/11/2019 | 06/12/2019 | |

Bottled and Mineral Water Scheme



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Bottled and Mineral Water Scheme



This microbiology scheme provides proficiency testing (PT) samples to laboratories that examine bottle and mineral waters routinely. Bottled and mineral water is a food product and therefore, must comply with strict safety requirements, as well as industry guides to good hygiene and manufacturing practices.

This PT scheme helps you to identify gaps in your processes, highlighting where quality improvements can be made. It also provides an opportunity to improve staffs knowledge and experience with organisms not frequently encountered.

PHE's Bottled and Mineral Water Scheme focuses on raising awareness of:

- the challenges associated with enumerating samples containing low levels of organisms
- issues with confirmatory tests carried out on isolates

Bottled and Mineral Water Scheme

BOTTLED AND MINERAL WATER SCHEME

water and food microbiology examinations

Sample schedule for 2019

| Distribution number | Sample numbers | Dispatch date | Date results due by | Examinations and enumerations required |
|---------------------|------------------|---------------|---------------------|--|
| BMW21 | BMW21A BMW21B | 28/01/2019 | 22/02/2019 | Coliform bacteria Escherichia coli Enterococci |
| BMW22 | BMW22A BMW22B | 13/05/2019 | 07/06/2019 | Pseudomonas aeruginosa Sporulated sulphite-reducing anaerobes Colony count (37°C/24 hours) |
| BMW23 | BMW23A BMW23B | 02/09/2019 | 27/09/2019 | Colony count (22°C/72 hours) |



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External Quality Assessment Schemes for 'Hospital Waters'



Hospital Water

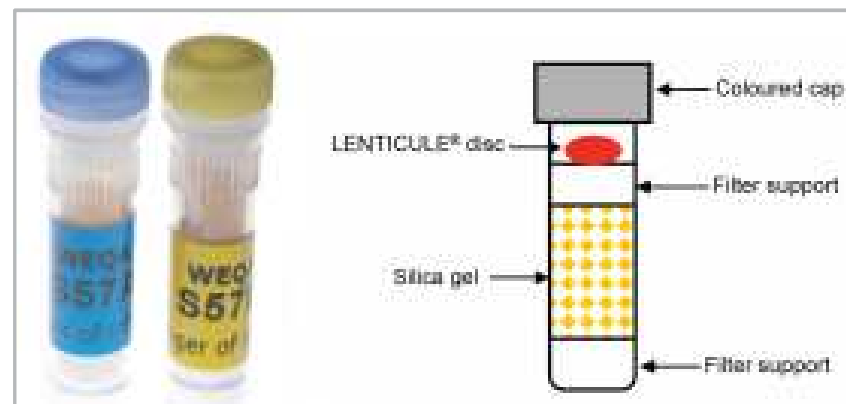
- Does your laboratory examine waters from a hospital environment, such as:
 - • water used to rinse endoscopes
 - • water used to prepare dialysis fluid
 - • hydrotherapy pool water
 - • tap water taken from augmented care units, such as neonatal and burns units?

- If so, participation in Public Health England's (PHE) 'Hospital Waters' external quality assessment (EQA) schemes will allow you to:
 - • demonstrate competence with the microbiological examination of these samples
 - • improve understanding of the interpretation of the results obtained.

- PHE provides EQA schemes specifically designed for laboratories that examine water samples. The schemes are very similar to those provided by the United Kingdom National External Quality Assessment Service (UK NEQAS). UK NEQAS specialises in schemes for laboratory medicine, whereas the PHE schemes focus on competence in testing water samples from a hospital environment; they are operated to the same rigorous quality standards as UK NEQAS.

Hospital Water

- The EQA test samples are provided in LENTICULE® disc format. The discs consist of a certified quantity of microorganisms in a water soluble matrix.
- They are easy to use and ideally suited for tests where accurate enumeration is paramount. On receipt of your EQA samples you can test immediately or store the discs at -20° C.



Endoscope Rinse Water Scheme



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Endoscope Rinse Water Scheme



This unique microbiology scheme provides proficiency testing (PT) samples to laboratories that examine endoscope rinse waters routinely for microbial contents. This scheme challenges the accurate enumeration of low levels of micro-organisms that may be present in this type of water.

Polices and guidance recommend that total viable counts should be made on the final rinse water. This is because the most significant problem associated with the use of automatic re-processors is contaminated rinse water that comes into contact with the endoscope after the disinfection process, increasing the risk of infection to patients.

This PT scheme helps you to identify gaps in your processes, highlighting where quality improvements can be made. It also provides an opportunity to improve staffs knowledge and experience with organisms not frequently encountered.

The scheme focuses on raising awareness of:

- the variation of different methods and media used and highlighting subsequent impact on PT results
- interpreting the microbiological results obtained
- the importance of testing *Mycobacterium* spp. and highlighting the difficulties associated with isolating this organism

Endoscope Rinse Water Scheme

ENDOSCOPE RINSE WATER SCHEME

water microbiology examinations

Sample schedule for 2019

| Distribution number | Sample numbers | Dispatch date | Date results due by | Examinations and enumerations required |
|---------------------|----------------|---------------|---------------------|---|
| EW21 | EW21A EW21B | 28/01/2019 | 22/02/2019 | Total viable counts (28°C - 32°C for 5 days) <i>Pseudomonas aeruginosa</i> |
| EW22 | EW22A EW22B | 13/05/2019 | 07/06/2019 | |
| EW23 | EW23A EW23B | 02/09/2019 | 27/09/2019 | Selected distribution: Yeasts/moulds (EW21) |



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Dialysis Water Scheme



This unique microbiology scheme provides proficiency testing (PT) samples to laboratories that examine water used to prepare dialysis fluid to confirm its safe use. This scheme challenges the accurate enumeration of low levels of micro-organisms which is critical for patient safety.

The European Renal Best Practice Guidelines recommends that the quality of water produced by the water treatment facility should meet the concentration limits for microbiological contaminants detailed in ISO 13959:2014: *Water for haemodialysis and related therapies*. This states that dialysis water shall contain a total viable microbial count of less than 100 colony forming units (cfu)/mL.

This PT scheme helps you to identify gaps in your processes, highlighting where quality improvements can be made. It also provides an opportunity to improve staffs knowledge and experience with organisms not frequently encountered.

The scheme focuses on raising awareness of:

- the variation of different methods and media used and highlighting subsequent impact on PT results
- interpreting the microbiological results obtained
- the requirement of microbiological testing as stated in the 'European Best Practice Guidelines for Haemodialysis'

Dialysis Water

DIALYSIS WATER SCHEME

water microbiology examinations

Sample schedule for 2019

| Distribution number | Sample numbers | Dispatch date | Date results due by | Examinations and enumerations required |
|---------------------|----------------|---------------|---------------------|--|
| DW18 | DW18A DW18B | 11/03/2019 | 05/04/2019 | Total viable counts (17°C - 23°C for 7 days) |
| DW19 | DW19A DW19B | 01/07/2019 | 26/07/2019 | |
| DW20 | DW20A DW20B | 04/11/2019 | 29/11/2019 | |

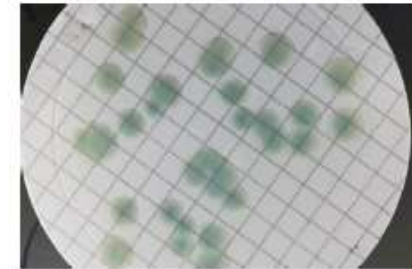
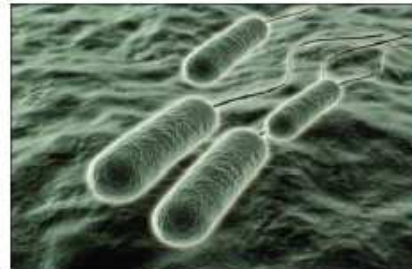
Hospital Tap Water Scheme



Public Health
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Hospital Tap Water Scheme



This unique microbiology scheme is suitable for laboratories in the water testing and other microbiology sectors that examine hospital tap waters from augmented care units for *Pseudomonas aeruginosa* at low levels.

Hospital water is a recognised potential source of *Pseudomonas aeruginosa*, which is a microorganism that can act as an opportunistic pathogen and colonise and infect vulnerable patients. Several outbreaks of *P. aeruginosa* have been attributed to contaminated water systems in hospitals.

This proficiency testing scheme helps you to identify gaps in your processes, highlighting where quality improvements can be made. It also provides an opportunity to improve staffs knowledge and experience with organisms not frequently encountered.

The scheme focuses on raising awareness of:

- the challenges associated with enumerating samples containing low levels of organisms
- interpreting the results obtained
- media issues (especially when non *P. aeruginosa* organisms are in the sample)

Hospital Tap Water Scheme

HOSPITAL TAP WATER SCHEME

water microbiology examinations

Sample schedule for 2019

| Distribution number | Sample numbers | Dispatch date | Date results due by | Examinations and enumerations required |
|---------------------|------------------|---------------|---------------------|--|
| HTW15 | HTW15A HTW15B | 11/03/2019 | 05/04/2019 | <i>Pseudomonas aeruginosa</i> |
| HTW16 | HTW16A HTW16B | 01/07/2019 | 26/07/2019 | |
| HTW17 | HTW17A HTW17B | 04/11/2019 | 29/11/2019 | |

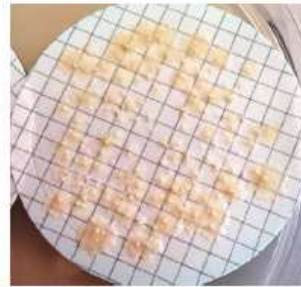
Mycobacterium sp in Water Scheme



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Mycobacterium spp. in Water Scheme



This unique microbiology scheme provides proficiency testing (PT) samples to laboratories that examine endoscope rinse and heater cooler unit (HCU) waters for *Mycobacterium* spp. This scheme challenges the detection, accurate enumeration and identification of this organism from these hospital water samples.

HCUs are used during open heart surgeries to warm or cool a patient as part of their care. It has recently been recognised that there is the potential for *Mycobacterium chimaera* or other species to grow in a water tank in the HCU. When the water evaporates, the mycobacteria may become dispersed into the environment as aerosols and may infect a patient during certain types of open heart surgery.

Flexible endoscopes are complex reusable instruments that require unique consideration with respect to decontamination. Their external surfaces and internal channels for air, water, aspiration and accessories are all potentially exposed to body fluids and other contaminants. Environmental non-pathogenic mycobacteria present a particular problem when they occur in the final rinse-water of some instruments used for diagnosis.

This PT scheme helps you to identify gaps in your processes, highlighting where quality improvements can be made. It also provides an opportunity to improve staffs knowledge and experience with organisms not frequently encountered.

The scheme focuses on raising awareness of:

- the variation of different methods and media used and highlighting subsequent impact on PT results
- interpreting the microbiological results obtained
- the importance of testing *Mycobacterium* spp. and highlighting the difficulties associated with isolating this organism

Mycobacterium sp in Water Scheme

MYCOBACTERIUM SPP. IN WATER SCHEME

water microbiology examinations

Sample schedule for 2019

| Distribution number | Sample numbers | Dispatch date | Date results due by | Examinations and enumerations required |
|---------------------|------------------|---------------|---------------------|--|
| MY001 | MY001A MY001B | 13/05/2019 | 02/08/2019 | Endoscope rinse water |
| MY002 | MY002A MY002B | 01/07/2019 | 20/09/2019 | Heater cooler water |
| MY003 | MY003A MY003B | 04/11/2019 | 24/01/2020 | Endoscope rinse water |

Learning from the Challenge



What can be learnt from challenging samples

- Exposure to **new organisms** of public health concern – raising awareness of their existence and allowing you to assess suitability of your current method/s or validating new ones
- Raising awareness of **atypical organisms** that exist in the environment and a greater understanding of the impact on laboratory testing and results
- Helps you to understand the **limitations of methods/media** used
- Helps you to understand the **limitations of confirmation tests**
- Allows you to understand **gaps in your procedures** – especially if an approved method is not followed
- Helps your laboratory understand how accurate your test results are



Water schemes



Image by Zak Prior

Legionella Isolation Scheme

- To provide external quality assessment samples to challenge the detection and enumeration of legionellae.
- The scheme focuses on raising awareness:
 - of the different *Legionella* spp. that maybe isolated from water samples
 - of the issues with batch to batch variations of the media used
 - of the confirmatory tests done and their limitations
 - of the importance of following standardised methods that are internationally recognised
 - updates in local guidelines or polices
- Four distributions a year with two samples in each



Distribution G107A – May 2017

- Contents:
 - *Legionella pneumophila* sg1 (1.80×10^4) (wild strain)
 - *Acinetobacter junii* (1.60×10^5) (wild strain)
 - *Pseudomonas fluorescens* (4.00×10^6) (wild strain)

- *Acinetobacter junii* and *Pseudomonas fluorescens* formed colonies on glycine vancomycin polymyxin B cycloheximide (GVPC) medium after processing

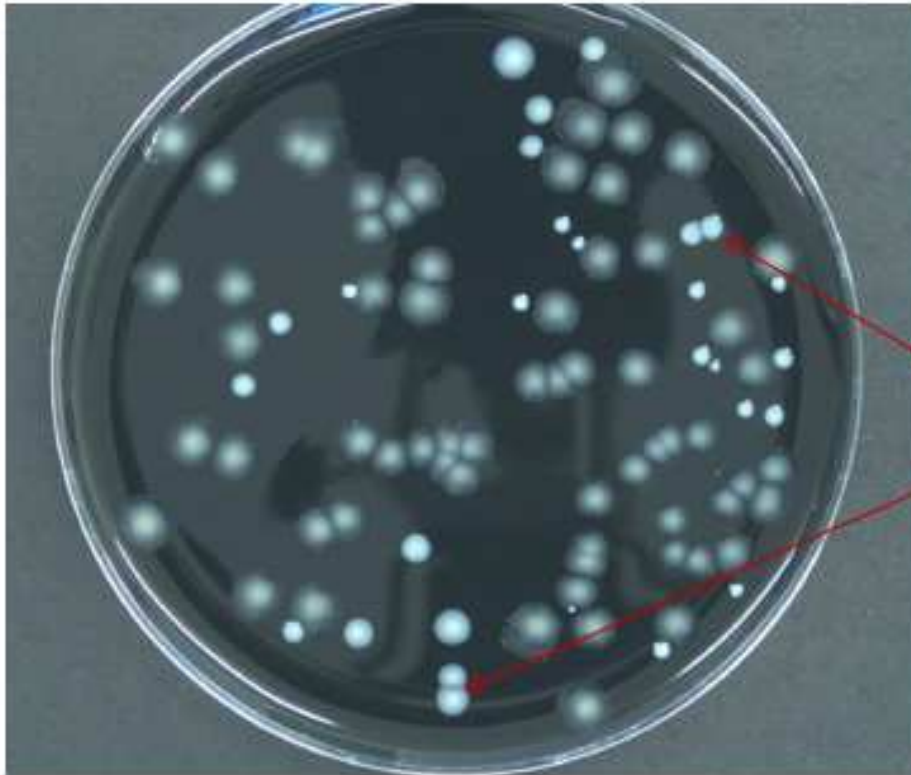
- All levels are colony forming unit per disc

G107A - findings

- Only 103/156 (66%) of the participants correctly reported the presence of *Legionella pneumophila* in this sample
- Participants clearly experienced difficulties with isolating this organism amongst the high level of background flora included in this sample
- The participants' median was 7.00×10^3 cfu L⁻¹ and the expected range (statistically calculated*) was 4.34×10^2 – 3.94×10^4 cfu L⁻¹
- The standard deviation was $0.47 \log_{10}$ cfu L⁻¹
- This sample was not scored

•* Median \pm 0.75 log₁₀ or counts within 11th to 89th percentiles (whichever is greater)

G107A: FEPTU's results



GVPC agar: 400 μL of 1:10 acid treated sample following 10 days incubation at 36 $^{\circ}\text{C}$

Legionella pneumophila colonies

FEPTU's median based on nine samples: $1.80 \times 10^4 \text{ L}^{-1}$

1 litre of the sample using a Nylon membrane filter with a pore size of 0.22 μm
Membrane suspended in 5 mL of Page's saline, vortexed to disperse any *Legionella* into the saline
Dilutions of 1:10 and 1:100 were done
For the acid treatment 400 μL of the sample was treated with 400 μL Buffer acid at pH of 2.2 for 5 minutes

G107A – strain information

- *L. pneumophila* colonies were clearly visible on acid treated samples at both 1:10 and 1:100 dilutions on GVPC agar
- Colonies on GVPC were grey, circular, entire and smooth with a ground glass appearance
- Were between 1-3 mm
- As expected the strain did not grow following a sub-culture onto buffered charcoal yeast extract (BCYE) without cysteine agar
- Gave a positive reaction with Oxoid's *Legionella* latex test



Interesting or a nuisance?

- Interesting:
 - To note the wide variations of methods followed – no two methods were the same
 - The high level of background flora obviously challenged a laboratory process
 - 50/119 (42%) still use ISO 11731:1998 (newer version available)
 - 27/93 (29%) did not check the pH of acid prior to use
- Nuisance:
 - It means if your quality policy states that failures with PT will be investigated:
 - then this is an action for you
 - resources such as time and consumables cost money
 - raise a non-conformance
 - explain to your accreditation body or service users



Drinking Water Scheme

- To provide external quality assessment samples for general routine examinations undertaken by routine water microbiology laboratories
- The scheme focuses on raising awareness:
 - of the challenges associated with enumerating samples containing low levels of organisms
 - of variations in the methods used which can highlight the differences in results obtained
 - of issues with confirmatory tests carried out on isolates
 - updates in local and national guidelines and policies
- Six distributions a year with three samples in each



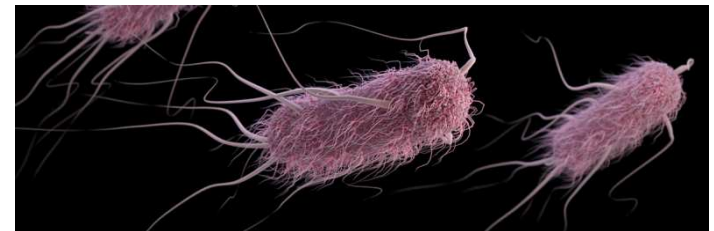
- Contents

- *Escherichia coli* (14 per 100 mL) (wild strain)
- *Enterobacter cloacae* (8.0×10^2 per disc) (wild strain)
- *Klebsiella oxytoca* (1.9×10^3 per disc) (wild strain)
- *Enterococcus faecium* (43 per 100 mL) (NCTC 7171)
- *Staphylococcus hyicus* (47 per mL) (NCTC 10350)

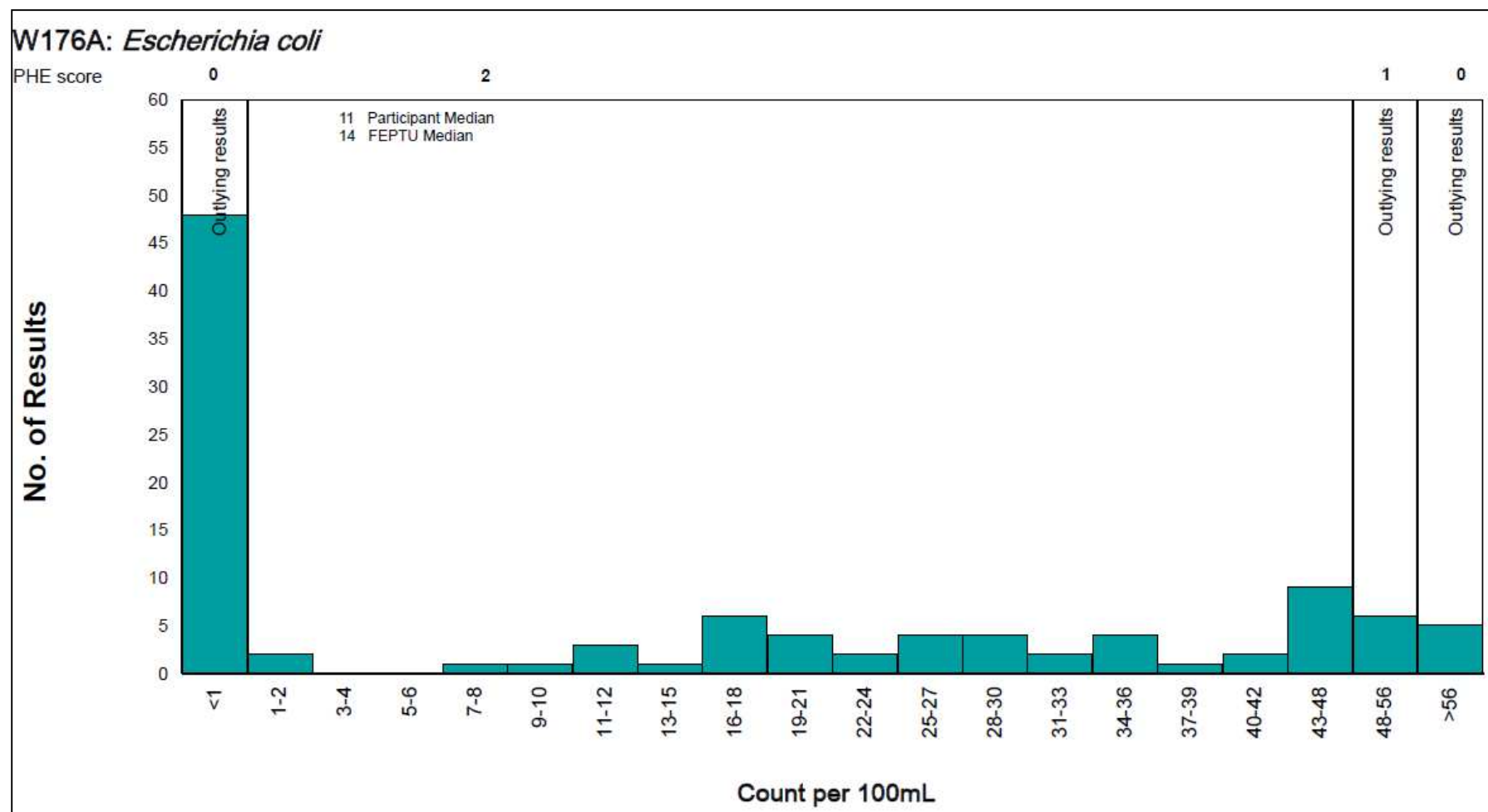
- *Escherichia coli* examination was not scored

W176A – *Escherichia coli*

- 55/112 (49%) of the participants reported a false negative result for this examination
- Sample contained an *E. coli* with participants' median being 11 colony forming units per 100mL
- The *E. coli* was a weak β -glucuronidase strain and therefore will have accounted for the false negative results by participants using a test that requires this enzyme to be metabolised to produce a positive result



W176A – Participants results



W176A – method breakdown

| Method | Number of laboratories reporting a false negative result by method (%) | Range of counts reported per 100mL |
|--|--|------------------------------------|
| Colilert (18/250) | 37/37 (100) | - |
| Chromogenic coliform agar (CCA) | 9/26 (35) | 1 - 94 |
| MLGA | 2/14 (14) | 11-154 |
| MLSA | 1/3 (33) | 28 - 56 |
| MLSB | 1/17 (6) | 9 - 78 |
| <u>Tergitol</u> | 1/1 (100) | - |
| TBX | 2/8 (25) | 8 - 39 |
| Other or no method provided | 2/6 (33) | 25 - 55 |

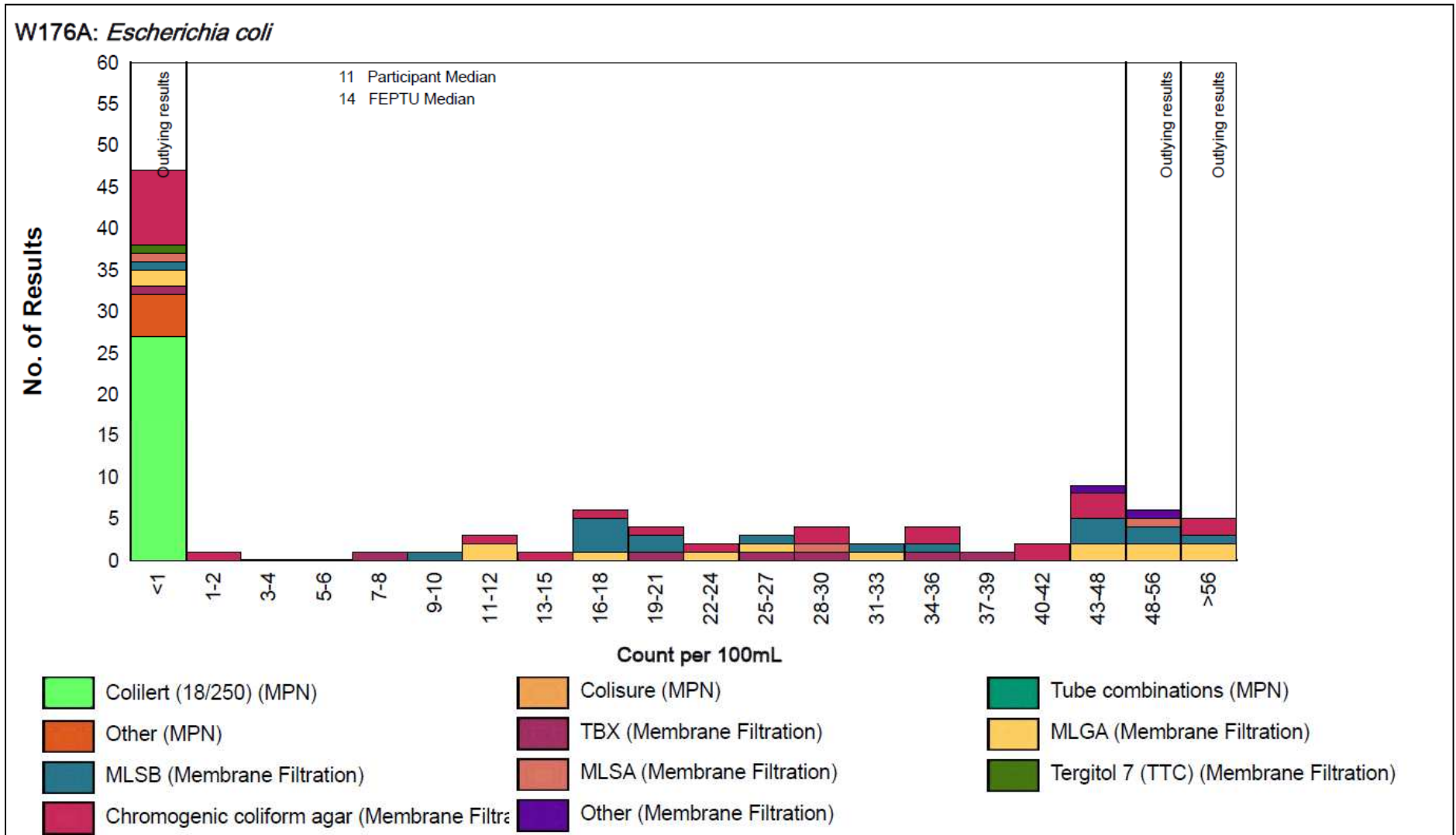
W176A – method breakdown

| Method | Number of laboratories reporting a false negative result by method (%) | Range of counts reported per 100mL |
|--|--|------------------------------------|
| Colilert (18/250) | 37/37 (100) | - |
| Chromogenic coliform agar (CCA) | 9/26 (35) | 1 - 94 |
| MLGA | 2/14 (14) | 11-154 |
| MLSA | 1/3 (33) | 28 - 56 |
| MLSB | 1/17 (6) | 9 - 78 |
| <u>Tergitol</u> | 1/1 (100) | - |
| TBX | 2/8 (25) | 8 - 39 |
| Other or no method provided | 2/6 (33) | 25 - 55 |

W176A – method breakdown

| Method | Number of laboratories reporting a false negative result by method (%) | Range of counts reported per 100mL |
|--|--|------------------------------------|
| Colilert (18/250) | 37/37 (100) | - |
| Chromogenic coliform agar (CCA) | 9/26 (35) | 1 - 94 |
| MLGA | 2/14 (14) | 11-154 |
| MLSA | 1/3 (33) | 28 - 56 |
| MLSB | 1/17 (6) | 9 - 78 |
| <u>Tergitol</u> | 1/1 (100) | - |
| TBX | 2/8 (25) | 8 - 39 |
| Other or no method provided | 2/6 (33) | 25 - 55 |

W176A – method breakdown



Interesting or a nuisance?

- Interesting:
 - To note that 17/26 (65%) participants that used CCA reported the presence of *E. coli* – unknown if they did a confirmation test
 - All Colilert® users failed to detect the *E. coli*
 - Media or methods using substrates for targeting specific organisms are not always 100% sensitive – be aware
- Nuisance:
 - It means if your quality policy states that failures with PT will be investigated:
 - then this is an action for you
 - resources such as staffs' time and consumables cost money
 - raise a non-conformance
 - explain to your accreditation body or service users



Food schemes



Image by Zak Prior



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Standard Scheme

Standard Scheme

- Laboratories that routinely test for a range of food-borne pathogens and indicator organisms
- Participants are often private laboratories that test foods for clients in the food industry who may submit products routinely for microbial assessment, end product testing and customer complaints

Pathogens

Salmonella spp.

Listeria monocytogenes

Listeria spp.

Escherichia coli O157 (non-toxigenic strains)

Campylobacter spp.

Cronobacter spp.

Enumerations

Presumptive *Bacillus cereus*

Campylobacter spp.

Coliforms

Clostridium perfringens

Coagulase-positive staphylococci

Listeria monocytogenes

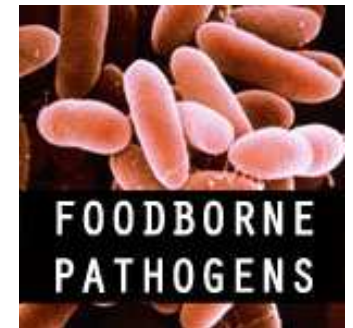
Listeria spp.

Aerobic colony count

Escherichia coli

Enterobacteriaceae

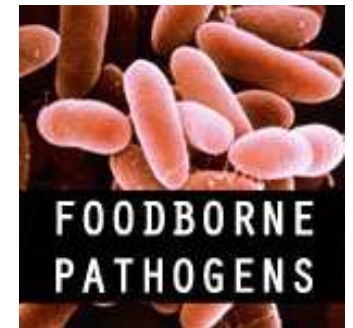
Six distributions a year with two samples in each



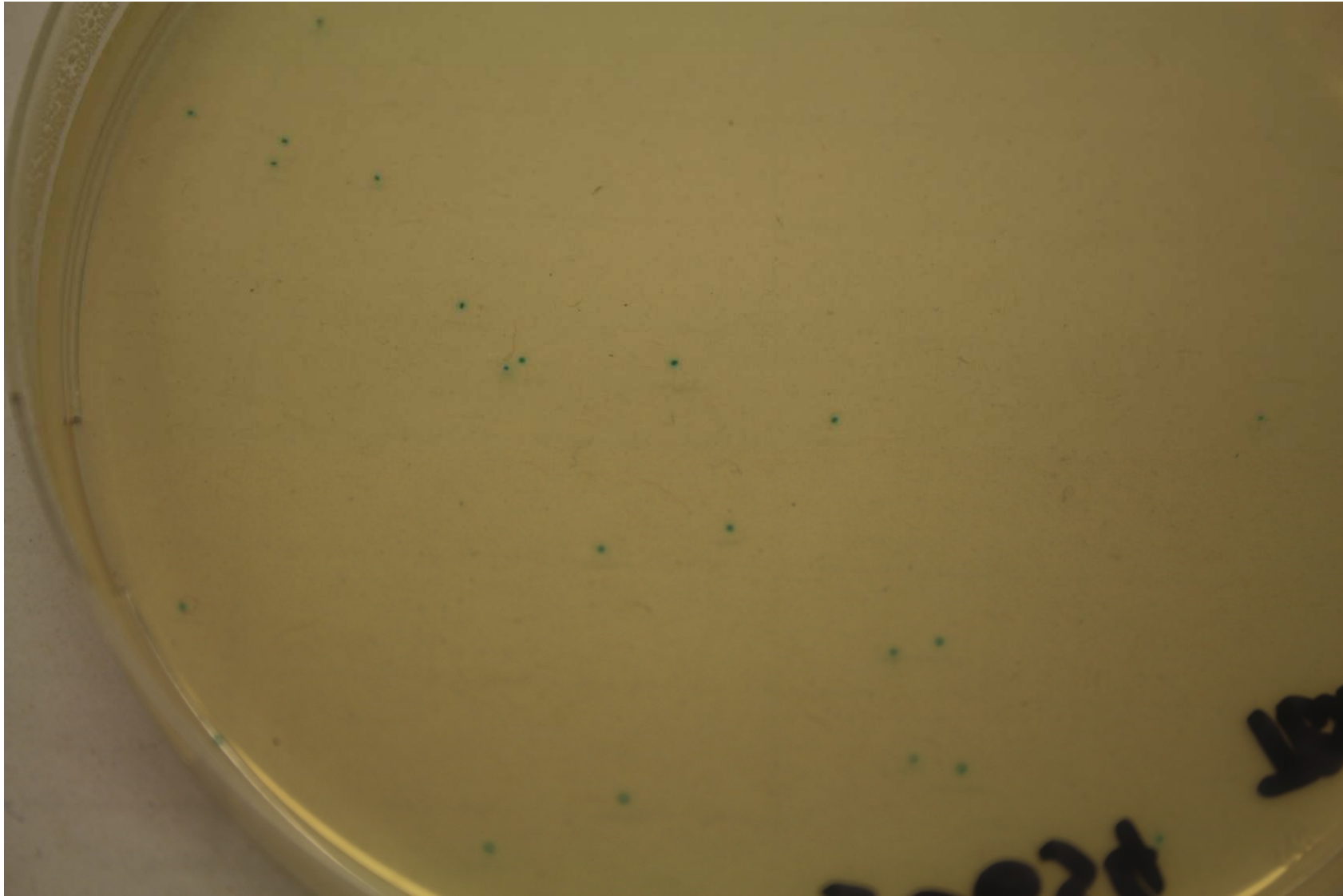
Standard Scheme

| STANDARD SCHEME food microbiology examinations | | | | |
|---|----------------|---------------|---------------------|---|
| Sample schedule for 2019 | | | | |
| Distribution number | Sample numbers | Dispatch date | Date results due by | Examinations and enumerations required Up to three sets of results can be reported |
| 312 | S0661 S0662 | 07/01/2019 | 15/02/2019 | <i>Campylobacter</i> spp. detection <i>Campylobacter</i> spp. enumeration <i>Escherichia coli</i> O157 detection <i>Salmonella</i> spp. detection Aerobic colony count Enterobacteriaceae enumeration |
| 314 | S0665 S0666 | 18/02/2019 | 29/03/2019 | Presumptive <i>Bacillus cereus</i> enumeration Coagulase-positive staphylococci enumeration <i>Listeria</i> spp. (including <i>L. mono</i>) enumeration <i>Listeria monocytogenes</i> enumeration Aerobic colony count Coliform enumeration |
| 316 | S0669 S0670 | 08/04/2019 | 17/05/2019 | <i>Cronobacter</i> spp. detection <i>Listeria</i> spp. (including <i>L. mono</i>) detection <i>Listeria monocytogenes</i> detection <i>Clostridium perfringens</i> enumeration Aerobic colony count <i>Escherichia coli</i> enumeration |
| 318 | S0673 S0674 | 03/06/2019 | 12/07/2019 | <i>Campylobacter</i> spp. detection <i>Campylobacter</i> spp. enumeration <i>Escherichia coli</i> O157 detection <i>Salmonella</i> spp. detection Aerobic colony count Enterobacteriaceae enumeration <i>Yersinia enterocolitica</i> |
| 320 | S0677 S0678 | 29/07/2019 | 06/09/2019 | Presumptive <i>Bacillus cereus</i> enumeration Coagulase-positive staphylococci enumeration <i>Listeria</i> spp. (including <i>L. mono</i>) enumeration <i>Listeria monocytogenes</i> enumeration Aerobic colony count Coliform enumeration |
| 322 | S0681 S0682 | 30/09/2019 | 08/11/2019 | <i>Listeria</i> spp. (including <i>L. mono</i>) detection <i>Listeria monocytogenes</i> detection <i>Salmonella</i> spp. detection <i>Clostridium perfringens</i> enumeration Sulfite-reducing anaerobic bacteria Aerobic colony count <i>Escherichia coli</i> enumeration |

Note for this scheme molecular methods can be used for pathogen examination or as a confirmation test



What is growing on this TBX plate?



Standard Scheme: Sample S0634

| Distribution number | Sample numbers | Dispatch date | Date results due by | Examinations and enumerations required |
|---------------------|----------------|---------------|---------------------|--|
| | S0634 | | | <i>L. monocytogenes</i> detection <i>Salmonella</i> detection <i>Clostridium perfringens</i> enumeration Aerobic colony count <i>E. coli</i> enumeration |

Distribution 298 – October 2017

- Sample S0634
- Contents:
 - *Listeria seeligeri* (8.3×10^2)
 - *Listeria monocytogenes* (90)
 - *Salmonella* Crewe (37)
 - *Clostridium sporogenes* (5.4×10^3)
 - *Norcadia farcinica* (1.3×10^3)
 - *Staphylococcus capitis* (2.4×10^3)
- All levels are presented as colony forming units (cfu) per ml reconstituted sample

Escherichia coli

- 39/154 (25%) reported a count for *E. coli* when the sample did not have this organism – **false positive**
- Counts reported were from 1 – 1500 cfu per gram

- Methods used:
 - 24/39 (61%) had used TBX plate
 - 3/39 (8%) had used a chromogenic agar
 - 12/39 (31%) has stated 'other' of which seven were TEMPO users

- Microbiology of food and animal feeding stuffs -- Horizontal method for the enumeration of beta-glucuronidase-positive *Escherichia coli* -- Part 2: Colony-count technique at 44 degrees C using 5-bromo-4-chloro-3-indolyl beta-D-glucuronide
- 3.1 – bacteria which at 44°C form typical blue colony on tryptone-bile-glucuronide medium (TBX)

- Microbiology of food and animal feeding stuffs -- Horizontal method for the enumeration of beta-glucuronidase-positive *Escherichia coli* -- Part 2: Colony-count technique at 44 degrees C using 5-bromo-4-chloro-3-indolyl beta-D-glucuronide
- 3.1 – bacteria which at 44°C form **typical** blue colony on tryptone-bile-glucuronide medium (TBX)

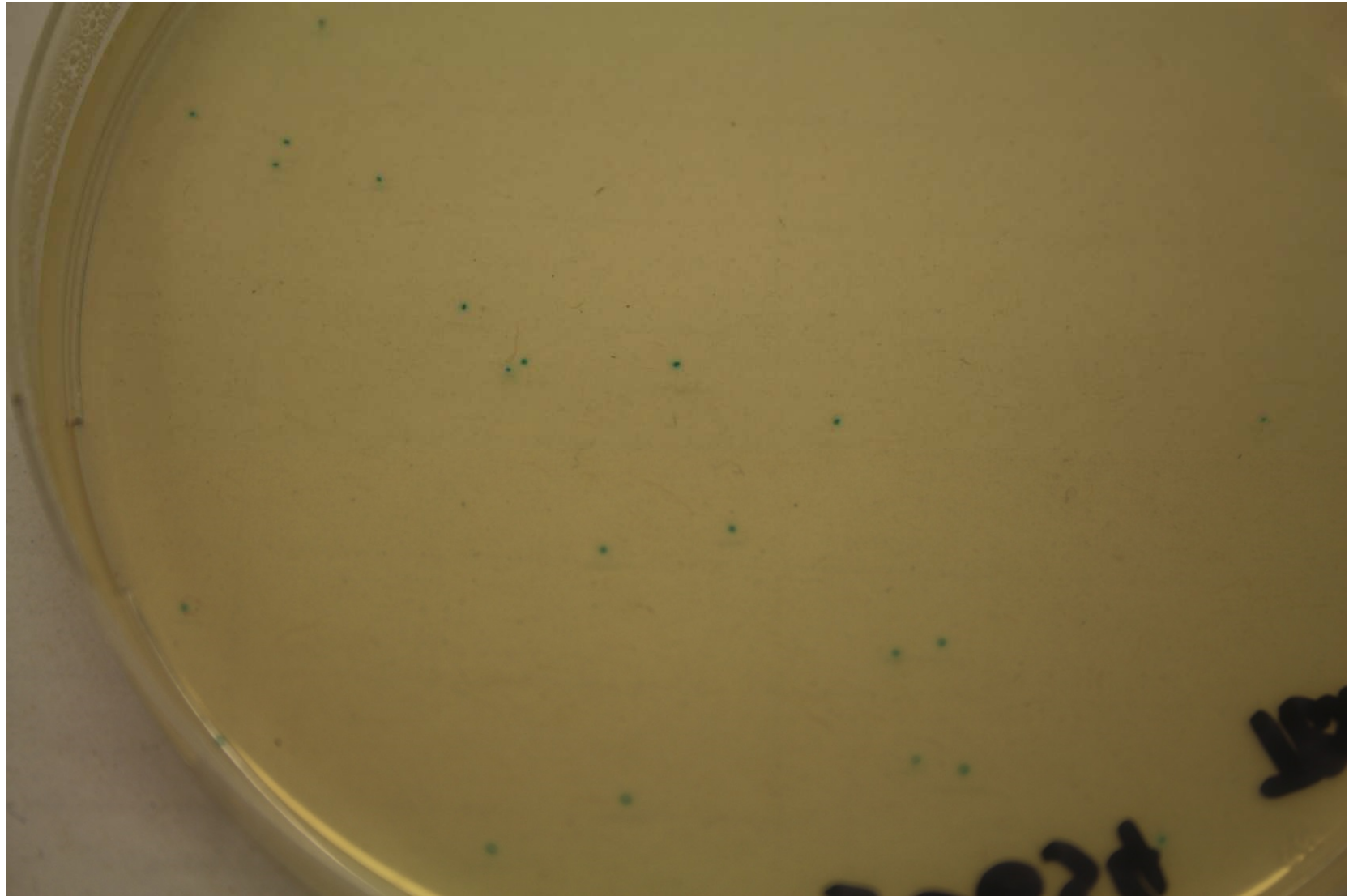


ISO 16649-2: 2001

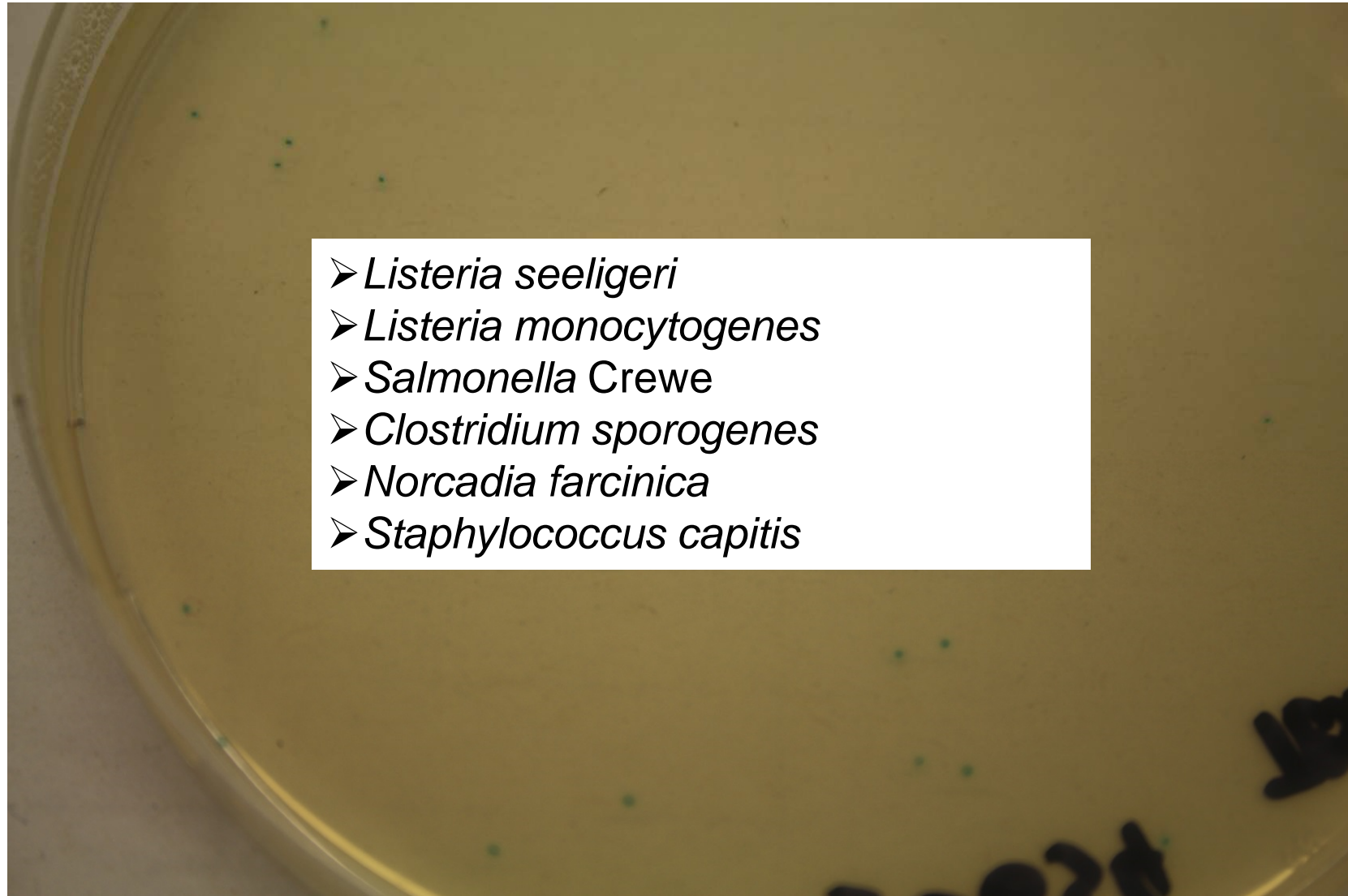
- Microbiology of food and animal feeding stuffs -- Horizontal method for the enumeration of beta-glucuronidase-positive *Escherichia coli* -- Part 2: Colony-count technique at 44 degrees C using 5-bromo-4-chloro-3-indolyl beta-D-glucuronide
- 3.1 – bacteria which at 44°C form **typical** blue colony on tryptone-bile-glucuronide medium (TBX)
- Only description found when various information sheets from different suppliers were analysed was: Blue/green



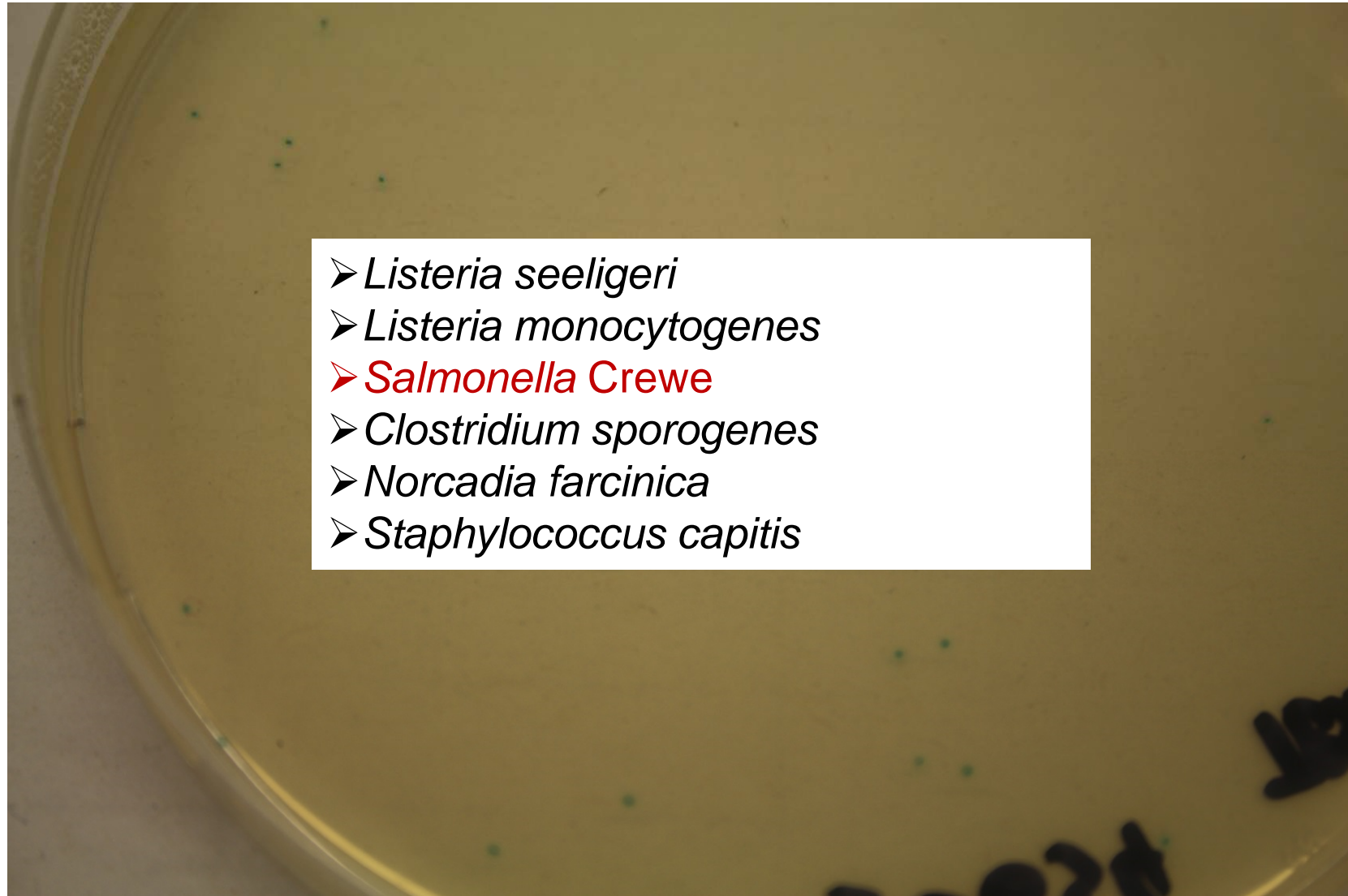
What is growing on this TBX plate?



What is growing on this TBX plate?



What is growing on this TBX plate?



What is growing on this TBX plate?

- *Listeria seeligeri*
- *Listeria monocytogenes*
- ***Salmonella* Crewe**
- *Clostridium sporogenes*
- *Norcadia farcinica*
- *Staphylococcus capitis*

Literature search shows that 30% of *Salmonella* spp. possess the enzyme β -glucuronidase



beAWARE

Interesting? Yes

- Can give you a better understanding of how suitable your processes are
- Can provide a useful insight into your laboratory's ability to accurately produce test results
- Gain more in depth information on the limitations of your media/methods used
- Exposure to 'unusual' organisms especially if they are not frequently encountered
- Create learning opportunities for your staff



Nuisance? Yes

- You have to investigate failures – this means you may have action/s (inconvenient)
- Might have to provide an explanation to your accreditation body or service users
- Additional cost implications for investigations
- Repeat testing of the PT sample to identify a root cause
- Might have to do a written report
- Might have to update documentation/s and then re-train staff
- You might not be awarded a score even if you got the right result



Distribution 284 – September 2016

- Samples S0605 and S0606 were exactly the same
- Contents:
 - *Staphylococcus aureus* (5.2×10^3) (wild strain)
 - *Listeria ivanovii* (7.2×10^2) (wild strain)
 - *Listeria monocytogenes* (3.8×10^2) (wild strain)
 - *Salmonella* Derby 1,4,[5],12:f,g:[1,2] (34) (wild strain)
 - *Enterobacter cloacae* (9.0×10^3) (wild strain)
- **Therefore the expectation is that participants should obtain comparable enumeration results for the specific examinations**
- All levels are presented as colony forming units (cfu) per mL reconstituted sample

Comparable: accuracy and precision

Accurate
Precise



Not Accurate
Precise



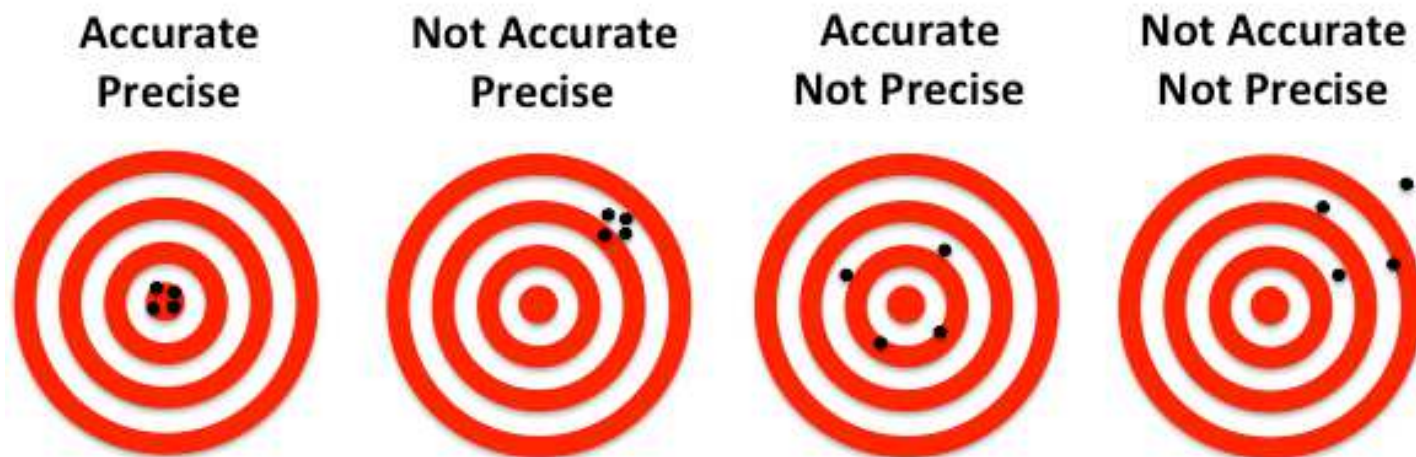
Accurate
Not Precise



Not Accurate
Not Precise

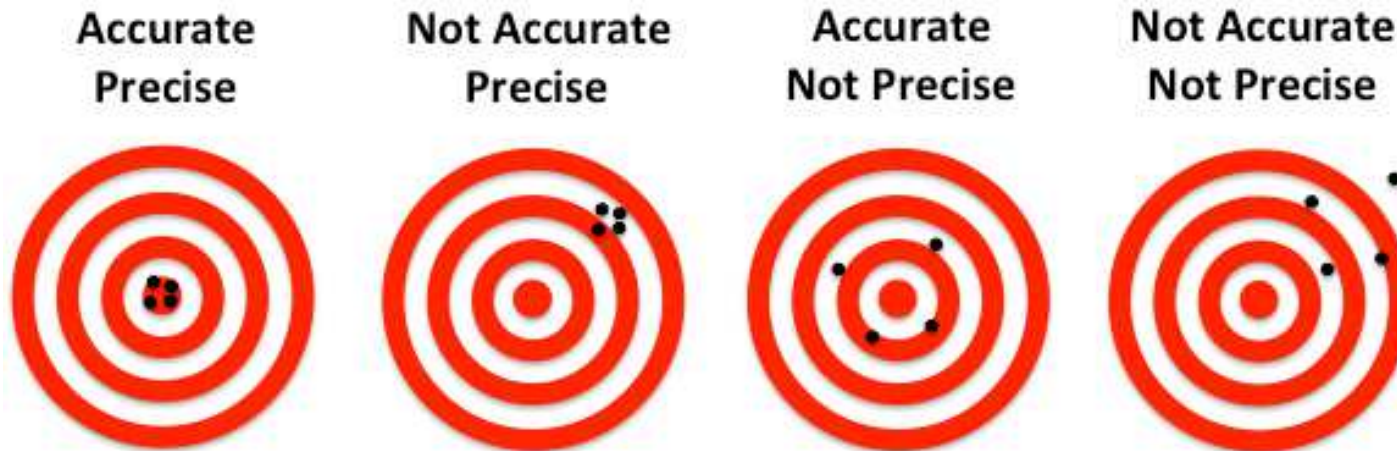


Comparable: accuracy and precision



Accuracy could be defined as how close your results are to the participants' median

Comparable: accuracy and precision



Accuracy could be defined as how close your results are to the participants' median

Precision could be defined as being in the expected range
– but even this can be anywhere and wide for PT data

The expected precision of culture-based microbial methods is typically derived mathematically based on the assumption that bacteria are distributed randomly in a well-mixed sample and follow a Poisson distribution

Expected range

| Examination | Expected range (cfu g ⁻¹) | |
|---|---|---|
| | S0605 | S0606 |
| Coagulase-positive staphylococci | 1.5x10 ³ – 1.5x10 ⁴ | 1.5x10 ³ – 1.5x10 ⁴ |
| <i>Listeria</i> spp. (including <i>L. monocytogenes</i>) | 3.5x10 ² – 3.5x10 ³ | 3.5x10 ² – 3.5x10 ³ |
| <i>L. monocytogenes</i> | 1.3x10 ² – 1.3x10 ³ | 1.2x10 ² – 1.2x10 ³ |
| Aerobic colony count | 1.2x10 ⁴ – 1.2x10 ⁵ | 1.1x10 ⁴ – 1.1x10 ⁵ |
| Coliform | 3.7x10 ³ – 4.4x10 ⁴ | 2.4x10 ³ – 4.1x10 ⁴ |

Expected range: participants' median $\pm 0.5 \log_{10}$ units or counts within the 11th to 89th percentile

Expected range

| Examination | Expected range (cfu g ⁻¹) | |
|---|---|---|
| | S0605 | S0606 |
| Coagulase-positive staphylococci | 1.5x10 ³ – 1.5x10 ⁴ | 1.5x10 ³ – 1.5x10 ⁴ |
| <i>Listeria</i> spp. (including <i>L. monocytogenes</i>) | 3.5x10 ² – 3.5x10 ³ | 3.5x10 ² – 3.5x10 ³ |
| <i>L. monocytogenes</i> | 1.3x10 ² – 1.3x10 ³ | 1.2x10 ² – 1.2x10 ³ |
| Aerobic colony count | 1.2x10 ⁴ – 1.2x10 ⁵ | 1.1x10 ⁴ – 1.1x10 ⁵ |
| Coliform | 3.7x10 ³ – 4.4x10 ⁴ | 2.4x10 ³ – 4.1x10 ⁴ |

Reason for variations:

- Number examining is different
- Actual counts reported varies



Can impact the participants' median

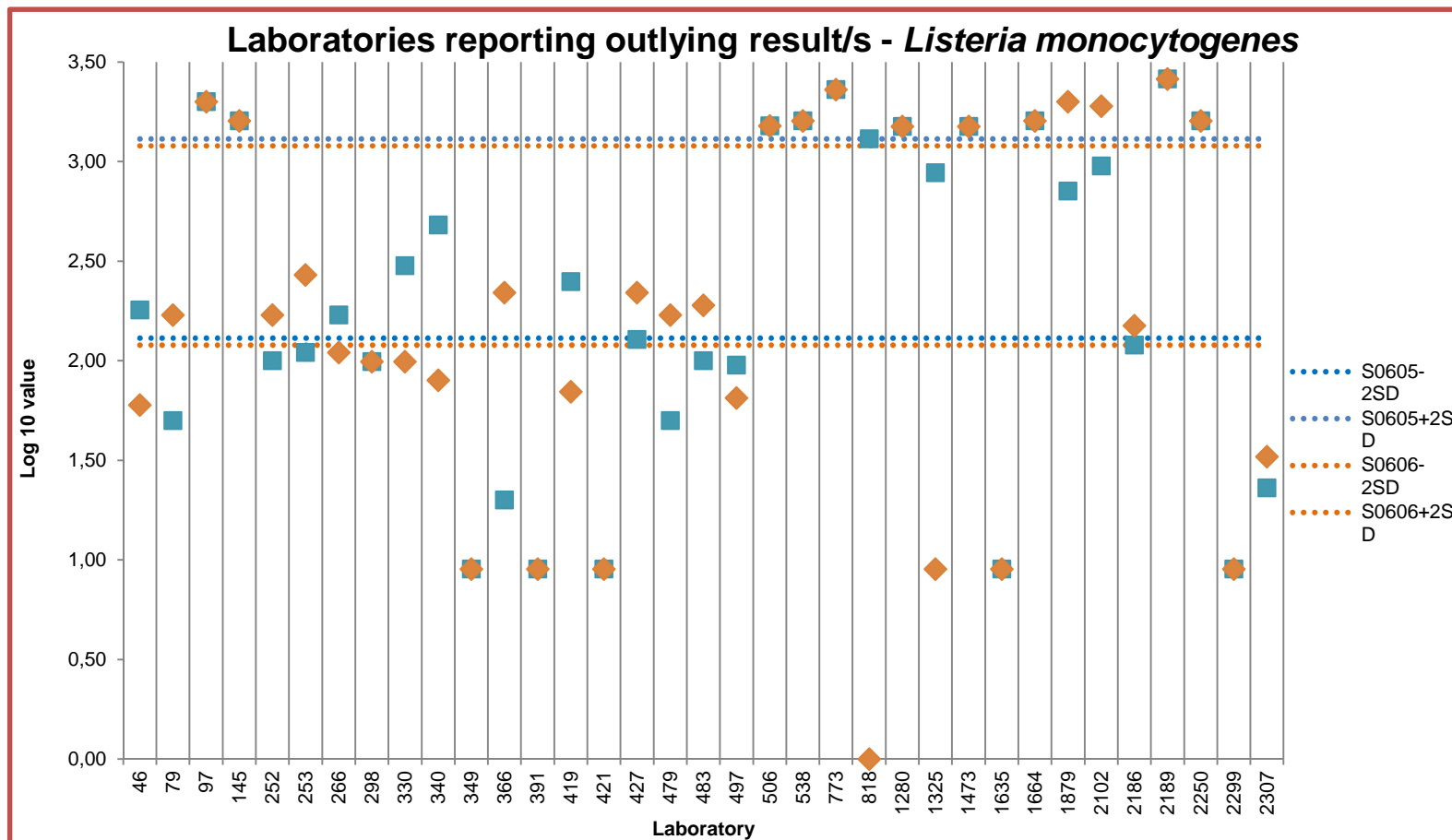
Results reported outside the range or a difference of >3SD

| Examination (data sets analysed) | Number of results | |
|--|--|-------|
| | S0605 | S0606 |
| Coagulase-positive staphylococci (143) | One | Four |
| | Two for both | |
| | Five reported a result with >3SD difference | |
| <i>Listeria</i> spp. (including <i>L. monocytogenes</i>) (97) | Two | Two |
| | Six for both | |
| | Six reported a result with >3SD difference (three in the expected range) | |
| Aerobic colony count (153) | Seven | Six |
| | Four for both | |
| | Ten reported a result with >3SD difference (three in the expected range) | |

Results reported outside the range or a difference of >3SD

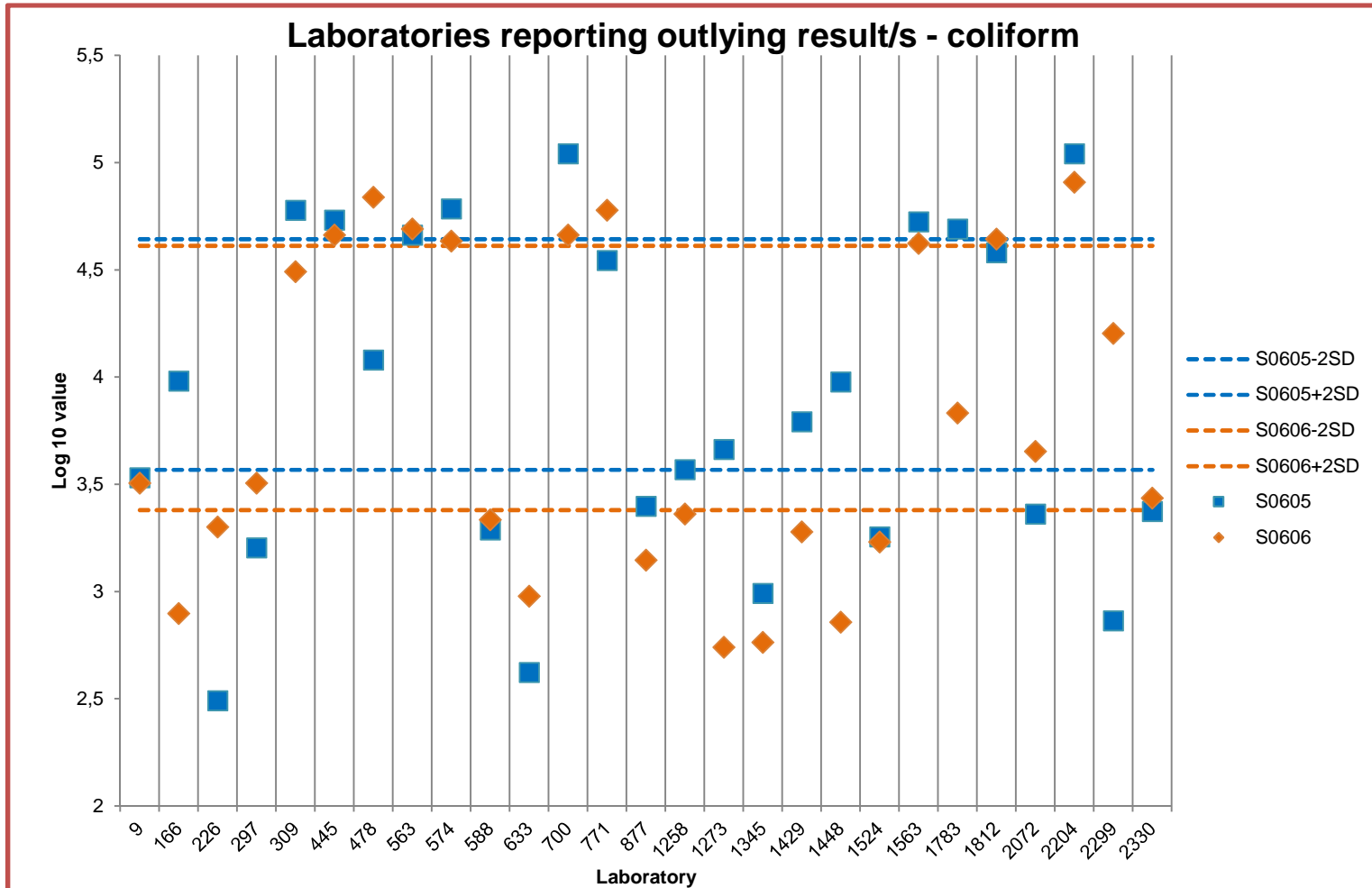
| Examination (data sets analysed) | Number of results | |
|----------------------------------|---|-------|
| | S0605 | S0606 |
| <i>L. monocytogenes</i> (114) | Nine | Nine |
| | 17 for both | |
| | Four reported a result with >3SD difference (one in the expected range) | |
| Coliform (112) | Seven | Eight |
| | 12 for both | |
| | Nine reported a result with >3SD difference (two in the expected range) | |

Listeria monocytogenes



- Nine laboratories reported a result outside the range for S0605 and nine for S0606.
- 17 laboratories reported a result outside the expected range for both samples.

Coliforms



- Seven laboratories reported a result outside the range for S0605 and eight for S0606.
- 12 laboratories reported a result outside the expected range for both samples.

Overall performance



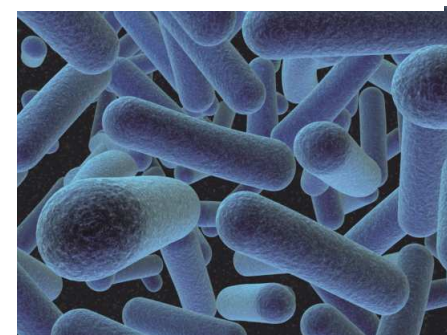
Total number of laboratories that reported a result outside the expected range or a difference of $>3SD$ for the number of data sets analysed is shown below

| Examination | |
|---|--------------|
| Coagulase-positive staphylococci | 5/143 (4%) |
| <i>Listeria</i> spp. (including <i>L. monocytogenes</i>) | 13/97 (13%) |
| <i>L. monocytogenes</i> | 36/114 (32%) |
| Aerobic colony count | 20/153 (13%) |
| Coliform | 29/112 (26%) |

Challenging organism or failure to follow ISO method?

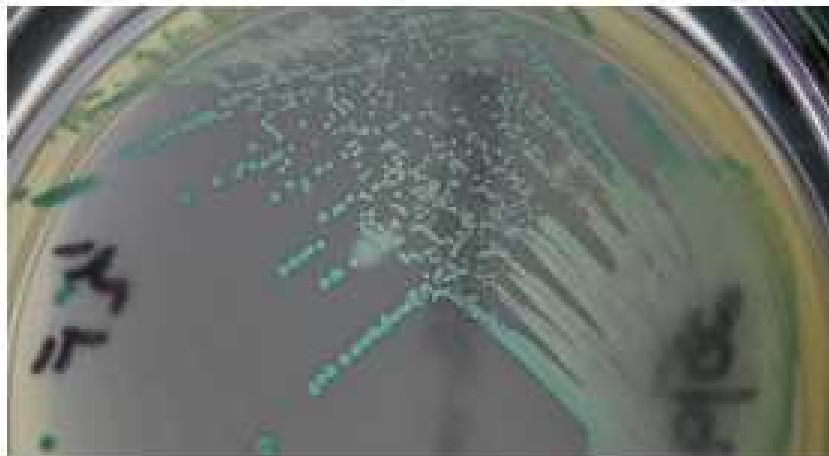
- Food distribution 274, sample S0586
 - Contents: *Listeria monocytogenes* (3.0×10^2) (wild strain)
 - *Listeria welshimeri* (9.3×10^2) (wild strain)
 - *Clostridium paraputrificum* (3.1×10^2) (wild strain)
- Examination was for detection of *Listeria monocytogenes* in 25g

| <i>L.monocytogenes</i> | |
|---|----------|
| Total participants reporting for <i>L.monocytogenes</i> | 113 |
| Participants reporting correctly | 82 (73%) |

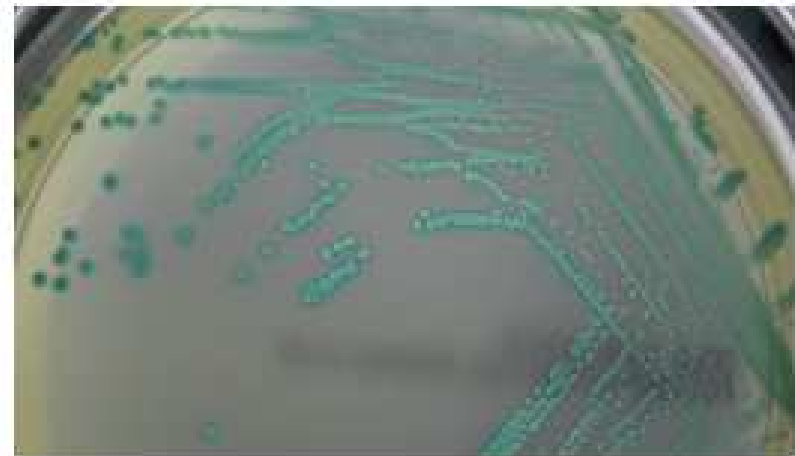


Findings - S0586

- 31/113 (27%) failed to detect the *L. monocytogenes* in this sample
- In the FEPTU laboratory the *L. monocytogenes* was isolated from a subculture of the half Fraser broth
- No *L. monocytogenes* was isolated following sub-culture of the full Fraser broth



Half Fraser sub on ALOA



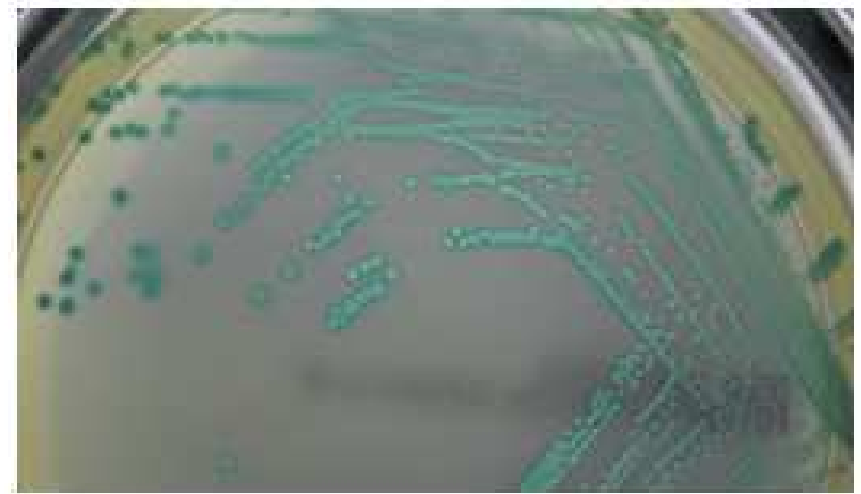
Full Fraser sub on ALOA

Findings - S0586

- 31/113 (27%) failed to detect the *L. monocytogenes* in this sample
- In the FEPTU laboratory the *L. monocytogenes* was isolated from a subculture of the half Fraser broth
- No *L. monocytogenes* was isolated following sub-culture of the full Fraser broth



Half Fraser sub on ALOA



Full Fraser sub on ALOA

ISO 11290-1 - method

- ISO 11290-1 Microbiology of food and animal feeding stuffs - Horizontal method for the detection and enumeration of *Listeria monocytogenes* - Part 1:
- Detection method recommends that 'broths are sub-cultured after each enrichment stage onto Ottaviani and Agosti agar and a second selective media'

| <i>L.monocytogenes</i> Method | <i>L.monocytogenes</i> Media | <i>L.monocytogenes</i> Enrichment | No. Participants detected | No. Participants not detected |
|----------------------------------|--|--------------------------------------|---------------------------------|-------------------------------------|
| ISO 11290 - 1 | Ottaviani and Agosti agar (ALOA) | Fraser broth (half followed by full) | 9 | 3 |
| ISO 11290 - 1 | Oxford Listeria selective agar; Ottaviani and Agosti agar (ALOA) | Fraser broth (half followed by full) | 8 | 5 |
| ISO 11290 - 1 | PALCAM Listeria selective agar; Ottaviani and Agosti agar (ALOA) | Fraser broth (half followed by full) | 6 | 4 |

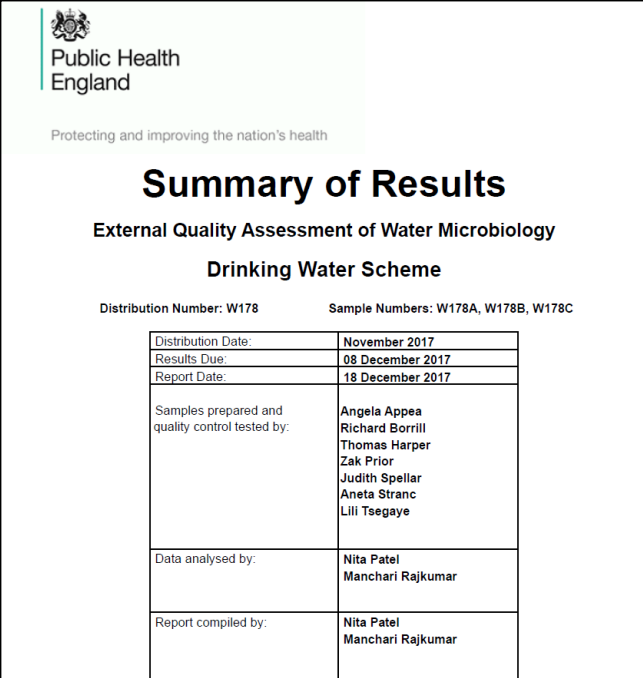
| <i>Listeria</i> spp. (including <i>L.mono</i>) | |
|--|----------|
| Total participants reporting for <i>Listeria</i> spp. (including <i>L.mono</i>) | 98 |
| Participants reporting correctly | 97 (99%) |

Points to note

- It is important to follow the ISO method as instructed – these methods have been extensively validated to increase the chance of isolating the target organism
- *Listeria* spp. other than *monocytogenes* in food samples can compete with *L. monocytogenes* if present in low numbers – therefore sub culturing of the half Fraser broth is extremely important
- If you deviate from an ISO method you must understand the impact on results – potentially at risk of not isolating the target organism
- New ISO 11290:2017 – detection method
 - incubation of the first enrichment has been changed from a minimum of 22 h to a minimum of 24 h
 - incubation of the second enrichment has been reduced to 24 h if only *L. monocytogenes* is sought

Personalised report will contain

- performance data including on-going performance over a year
- actions to take
- educational information on difficult and challenging samples
- PHE and z-scores
- contents and levels



The image shows the cover page of a 'Summary of Results' report from Public Health England. The report is titled 'External Quality Assessment of Water Microbiology' and 'Drinking Water Scheme'. It includes a table with key dates and personnel involved in the analysis and reporting.

Public Health England
Protecting and improving the nation's health

Summary of Results
External Quality Assessment of Water Microbiology
Drinking Water Scheme

Distribution Number: W178 Sample Numbers: W178A, W178B, W178C

| | |
|---|---|
| Distribution Date: | November 2017 |
| Results Due: | 08 December 2017 |
| Report Date: | 18 December 2017 |
| Samples prepared and quality control tested by: | Angela Appea Richard Borrill Thomas Harper Zak Prior Judith Spellar Aneta Stranc Lili Tsegaye |
| Data analysed by: | Nita Patel Manchari Rajkumar |
| Report compiled by: | Nita Patel Manchari Rajkumar |

Swab Sample: ES0023

- **Sample type:** Swab sample from the inside of a milk bottle containing some milk residue. Late evening on 30 July 2018 the local health authority received complaints from 15 tourists who had violent vomiting, nausea and stomach cramps. All were staying in a local hotel that had served raw artisan milk cheese made by a small local dairy. Food samples and swab were taken from the dairy.
- **Request:** Examine samples following your routine protocol for pathogens based on the outbreak scenario provided
- **Contents:** Staphylococcus aureus 1.4×10^4 (wild strain), Aerococcus viridans 1.7×10^3 (wild strain), Enterococcus faecalis 7.0×10^2 (wild strain)

Expected Results:

| Examination | Expected Result | Your Result | Score for performance assessment | Z-score |
|----------------------------------|--|-------------|----------------------------------|---------|
| <i>Listeria monocytogenes</i> | Not Detected | | | |
| <i>Salmonella</i> spp. | Not Detected | | | |
| Coagulase-positive staphylococci | 9.9×10^2 - 7.8×10^4 cfu per swab | | | |
| <i>Campylobacter</i> spp. | Not Detected | | | |

Swab Sample: ES0023

Comments on Performance:

| <i>Listeria monocytogenes</i> | |
|---|--|
| Total participants reporting for <i>Listeria monocytogenes</i> | 25 |
| Participants reporting correctly | 24 (96%) |
| <i>Salmonella</i> spp. | |
| Total participants reporting for <i>Salmonella</i> spp. | 27 |
| Participants reporting correctly | 26 (96%) |
| Coagulase-positive staphylococci | |
| Total participants reporting for Coagulase-positive staphylococci | 28 |
| Assigned value (participants' median) | 8.8x10 ³ cfu per swab (3.94 log ₁₀) |
| Uncertainty of assigned value ($U(X_{p,t}) = \log_{10}$ cfu per swab) | 0.11 |
| No. of outlying counts | 6 (6 low / 0 high) |
| Participants' mean | 6.4x10 ³ cfu per swab (3.81 log ₁₀) |
| *Standard deviation of participants' results | 0.47 log ₁₀ cfu per swab |
| FEPTU QC median | 1.4x10 ⁴ cfu per swab (4.14 log ₁₀) |
| <i>Campylobacter</i> spp. | |
| Total participants reporting for <i>Campylobacter</i> spp. | 6 |
| Participants reporting correctly | 6 (100%) |
| Total sent samples | 38 |
| Non-returns | 5 |
| Not examined | 1 |

Swab Sample: ES0023

Comments for distribution ES12

Sample ES0023

28 laboratories analysed the sample from this distribution. The pathogen in this sample was a coagulase positive staphylococci.

The table below shows the additional examinations carried out by the laboratories and the reported results.

| Additional examinations | Number of laboratories examining | Reported results |
|--------------------------------|----------------------------------|---|
| <i>Bacillus cereus</i> | 6 | 1100 (1) 0 (1) <100 (2) <200 (1) Not detected (1) |
| <i>Clostridium perfringens</i> | 6 | 0 (2) <10 (2) <100 (2) |
| <i>Escherichia coli</i> O157 | 3 | Not examined (1) Not detected (2) |

Swab Sample: ES0023

It is important for participant to be aware that outbreaks relating to artisan cheese made from raw milk can occur with this organism.

[https://www.journalofdairyscience.org/article/S0022-0302\(18\)30078-X/abstract](https://www.journalofdairyscience.org/article/S0022-0302(18)30078-X/abstract)

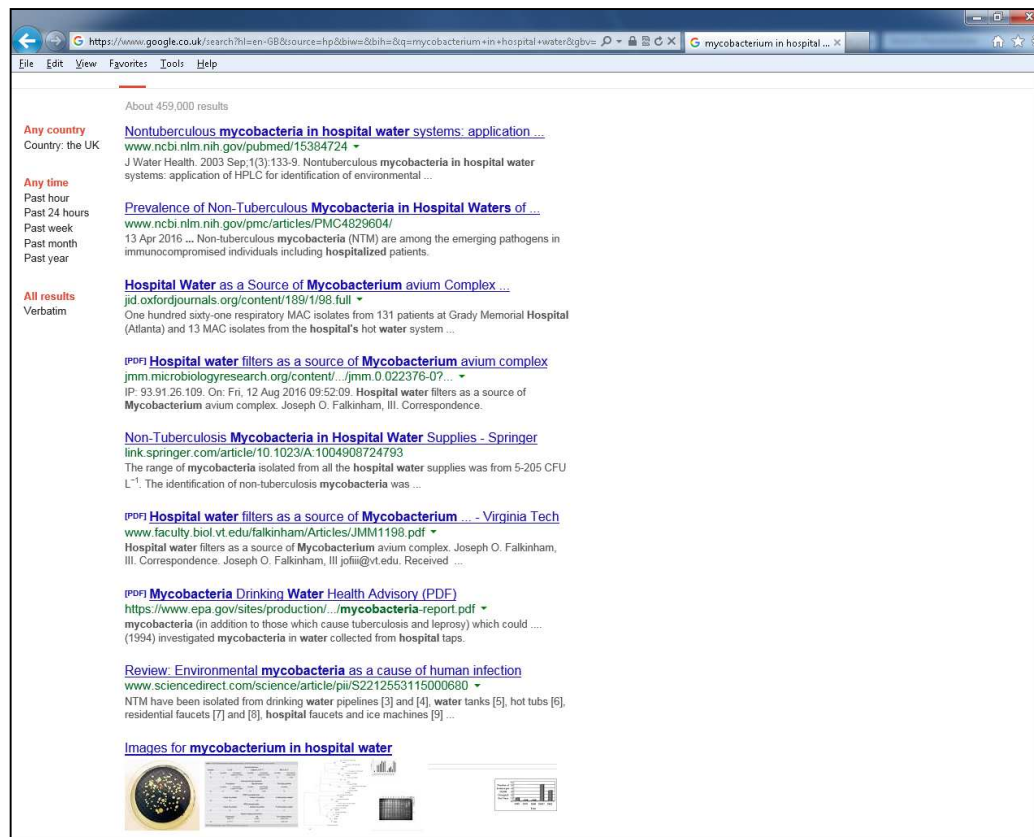
The word 'artisan' or 'artisanal' implies that a cheese is produced primarily by hand, in small batches, with particular attention paid to the tradition of the cheesemaker's art and thus using as little mechanisation as possible in production of the cheese. Artisan, or artisanal, cheese may be made from all types of milk including raw milk.

Read this article regarding a study undertaken about '*Staphylococcus aureus* Entrance into the Dairy Chain: Tracking *S. aureus* from Dairy Cow to Cheese'

<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC5061776/>

New and emerging organism – *Mycobacterium* spp.

- If you did a search on the internet, you would find a long list of articles on *Mycobacterium* in hospital waters going back over 20 years



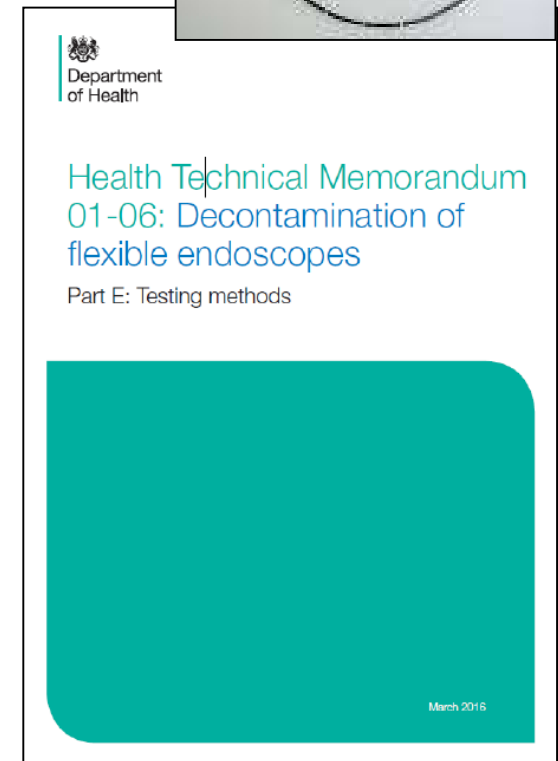
Rinse-water

Environmental non-pathogenic mycobacteria present a particular problem when they occur in the final rinse-water of some instruments used for diagnosis

Mycobacteria that occur in water, for example *Mycobacterium kansasii* and *Mycobacterium chelonae*, are opportunistic pathogens

For equipment rinsed after the disinfection stage, there should be no recovery of mycobacteria from 100 mL of final rinse-water

This group of organism is now included annually in the Endoscope Rinse Water Scheme

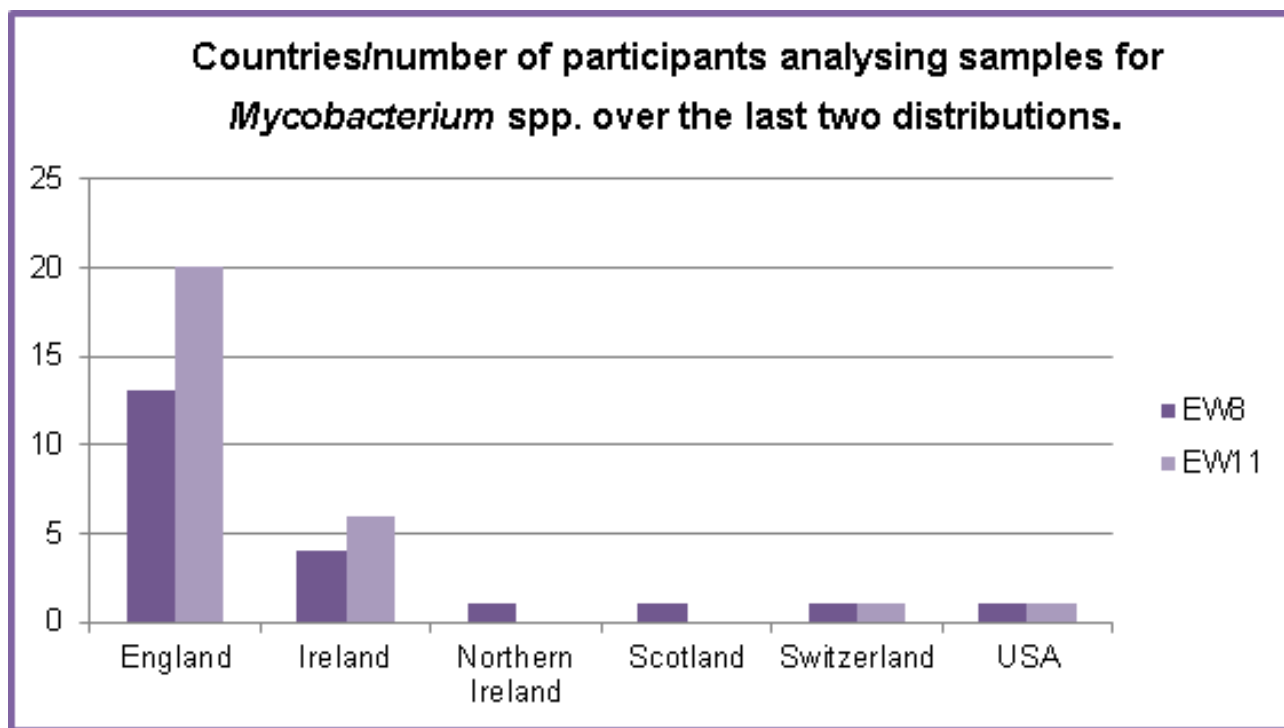


Mycobacterium spp.

- Main objectives of the scheme:
- To produce LENTICULE® discs at sufficient levels of *Mycobacterium* spp. to be isolated using membrane filtration methods
- To determine testing compliance with HTM 01-06 (CFPP 01-06):
 - media used
 - volume of water used for examination
 - incubation temperature and period
- To provide suitable samples that challenge a laboratory's ability to isolate and identify *Mycobacterium* spp. from waters
- To give a laboratory confidence in the method they use
- To confirm performance of culture media used such as Middlebrook 7H10
- To determine variability of the methods/media used
-

Mycobacterium spp.

- Number of participants examining for *Mycobacterium* in the Endoscope Rinse Water scheme over the last two distributions



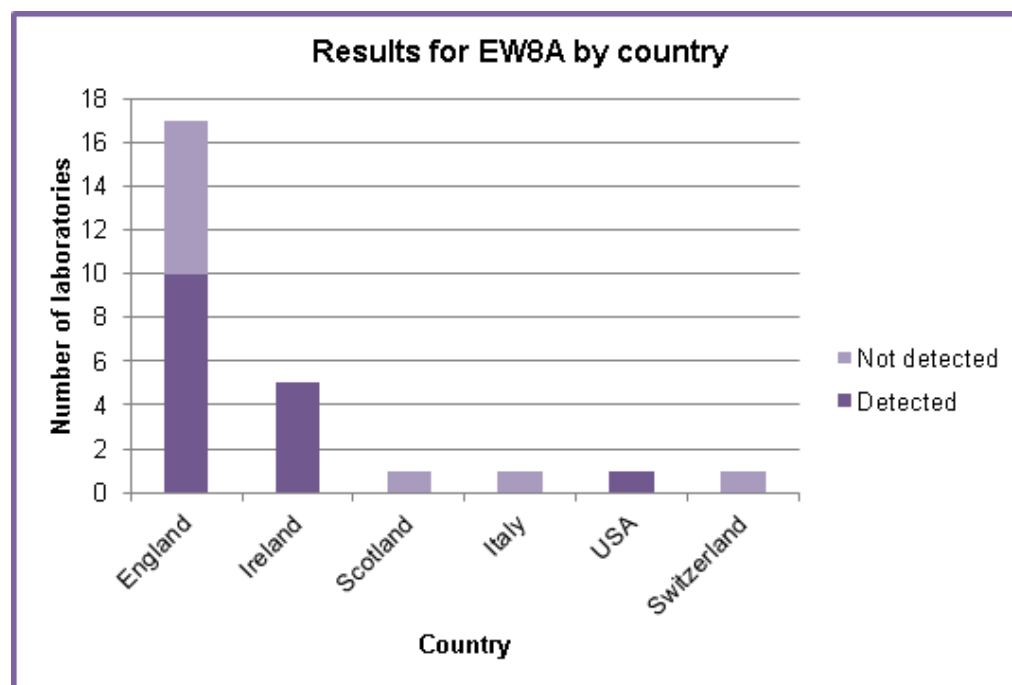
- EW8 – August 14
- EW11 – August 15
- EW14 – closes October 16

EW8A

26 / 60 (43%) laboratories examined the samples for *Mycobacterium* spp.

This sample contained *Mycobacterium chelonae* (1.1×10^2 cfu per 100mL) and *Pseudomonas aeruginosa*

16 / 26 (62%) laboratories correctly reported that *Mycobacterium* spp. was present in 100mL of this sample



26 / 26 (100%) laboratories correctly reported that there was no *Mycobacterium* spp. present in sample EW8B

EW8A – test conditions

| Results for EW8A | Guideline followed | Culture medium used | Filter size | Incubation temperature | Incubation period |
|------------------|--------------------|-----------------------|----------------|------------------------|-------------------|
| Not isolated | HTM2030 | Middlebrooks | 0.2 | 35 | 16 |
| Negative | ISO 15883-4 | 7H11 Middlebrook | 0.45 | 30 | 28 |
| Not detected | HTM2030 and CFPP | 7H10 Middlebrook | 0.45 | 35 | 28 |
| Negative | ISO 15883-4 | | 0.45 | | |
| Not isolated | | MP bottles bioMerieux | Not applicable | | |
| Not isolated | CFPP 01-06 | 7H10 Middlebrook | 0.45 | 37 | 28 |
| Negative | Local guideline | LJ and MGIT (BD) | | 37 | 45 |
| Negative | | VCAT agar | 0.45 | 30 | 7 |
| Not detected | HTM2030 | Middlebrooks | 0.45 | 30 | 22 |
| Not isolated | CFPP 01-06 | 7H10 Middlebrook | 0.45 | 30 | 28 |

- Findings:

- Non validated methods and media was used
- Length of incubation was incorrect – should be up to 28 days (no growth)
- Incorrect filter size used; mycobacteria rods are 1 -10um in length and 0.2 - 0.8um in width (varies depending on the species)
- Temperature for incubation in CFPP is incorrect it should be 30°C not 35°C

EW11A

28 / 71 (39%) laboratories examined the samples for *Mycobacterium* spp.

This sample contained *Mycobacterium chelonae* (54 cfu per 100mL) and *Staphylococcus capitis*

23 / 28 (82%) laboratories correctly reported that *Mycobacterium* spp. was present in 100mL of this sample



22 / 24 (92%) laboratories correctly reported that there was no *Mycobacterium* spp. present in sample EW11B. Four laboratories reported that they were unable to report a result due to overgrowth of fungi on the culture plates

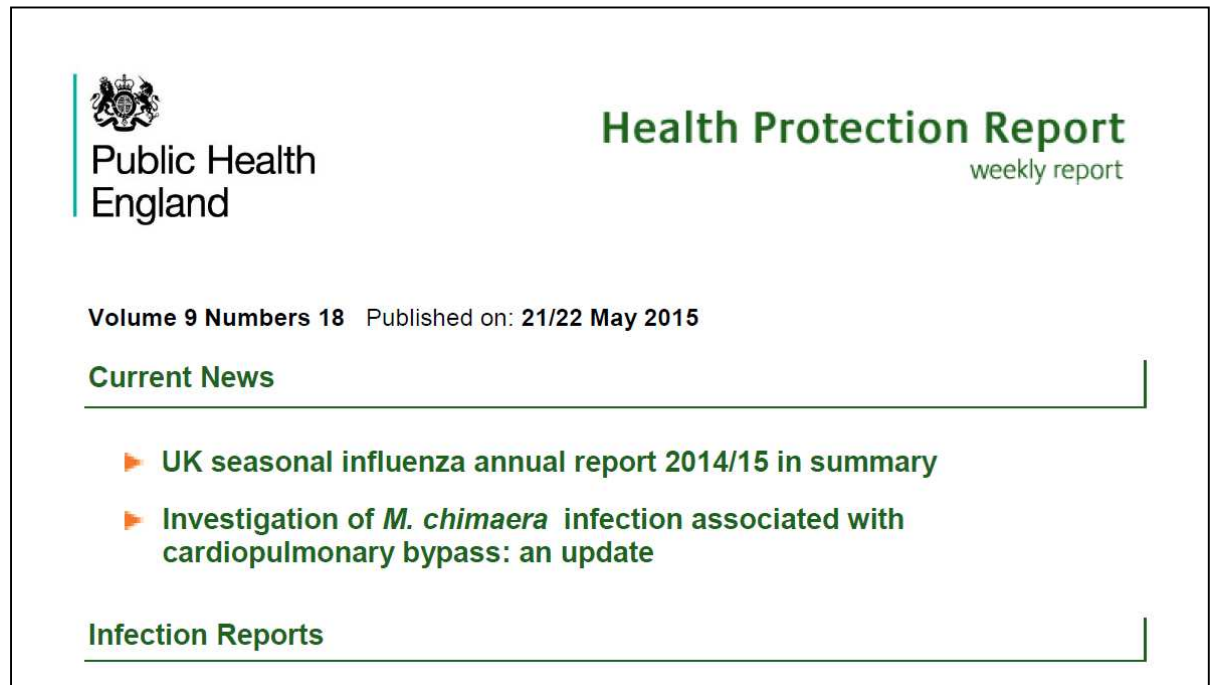
EW8 and EW11 methods used


| Guideline or standard used | Number of laboratories using the guideline / standard | |
|---|---|------|
| | EW8A | EW11 |
| CFPP 01-06* | 6 | 13 |
| HTM 2030 | 2 | |
| HTM 2030 / CFPP 01-06 | 1 | |
| HTM 2030 / CFPP 01-06 / ISO 15883-1:2006 | 2 | 2 |
| HTM 2030 and ISO 15883 | 1 | |
| In-house or local guideline | 2 | 1 |
| ISO 15883-1:2006 | 2 | |
| ISO 15883-4:2008 | 3 | 8 |
| ISO 15883-1:2006 and ISO 15883-4:2008 | 1 | |
| ISO 15883-1:2006 and ISO 15668:2008 | | 1 |
| HTM 04-01 | | 1 |
| Public Health England Healthcare document | 3 | 1 |
| Not specified | 3 | 1 |

- ISO 15883-4:2008 Washer-disinfectors - Part 4: Requirements and tests for washer-disinfectors employing chemical disinfection for thermolabile endoscopes
- *Replaced by Health Technical Memorandum 01-06: Decontamination of flexible endoscopes Part E: Testing methods

Mycobacterium chimaera ? A new problem

- As reported on 30 April, cases of invasive *Mycobacterium chimaera* infection have been reported in patients who have undergone cardiac surgery in Switzerland and the Netherlands. A Swiss investigation has been published attributing these infections to aerosol generated by contaminated heater cooler units (HCUs) used during cardiopulmonary bypass
- A case of similar infection has also been reported in Germany




Public Health
England

Health Protection Report
weekly report

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Current News

- ▶ UK seasonal influenza annual report 2014/15 in summary
- ▶ Investigation of *M. chimaera* infection associated with cardiopulmonary bypass: an update

Infection Reports



Available online at www.sciencedirect.com

Journal of Hospital Infection

journal homepage: www.elsevier.com/locate/jhin



Emerging *Mycobacteria* spp. in Cooling Towers

To the Editor: The importance of nontuberculous mycobacteria (NTM) in various clinical situations recently has increased. *Mycobacterium avium* complex cause a high percentage in persons with acquired immunodeficiency syndrome. So considered emerging pathogens.

Nosocomial pseudo-outbreak of *Mycobacterium gordonae* associated with a hospital's water supply contamination: a case series of 135 patients

Marija Zlojtro, Mateja Jankovic, Miroslav Samarzija, Ljiljana Zmak, Vera Katalinic Jankovic, Mihaela Obrovac, Igor Zlojtro and Marko Jakopovic

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Detection limit of *Mycobacterium chimaera* in water samples for monitoring medical device safety: insights from a pilot experimental series

ANN IST SUPER SANITA 2010 | VOL. 46, No. 3: 254-258
DOI: 10.4415/ANN_10_03_05

Non-tuberculous mycobacteria and microbial populations in drinking water distribution systems

Rossella Briancesco, Maurizio Semproni, Simonetta Della Libera, Simona Sdanganelli and Lucia Bonadonna

Istituto Superiore di Sanità, Istituto Superiore di Sanità, Rome, Italy

Summary. Data on the occurrence of non-tuberculous mycobacteria (NTM), in parallel with those used for bacterial indicators and amoebae, are presented with the aim to collect information on the spread of NTM in drinking water distribution systems in Italy. Samples were collected from taps in hospitals and households in Central and Southern Italy. The concentration values obtained for more traditional microbial parameters complied with the mandatory requirements for drinking water. Conversely, moderate-to-high microbial loads (up to 300 CFU/L) were observed for the NTM. Positive samples were obtained from 62% of the investigated water samples. Analogous results were observed for amoebae showing a higher percentage of positive samples (76%). In terms of public health, the presence of mycobacteria in water distribution systems may represent a potential risk especially for vulnerable people such as children, the elderly or immunocompromised individuals.

Prevalence of Non-Tuberculous *Mycobacteria* in Hospital Waters of Major Cities of Khuzestan Province, Iran

Azar Dokht Khosravi^{1,2}, Abdolrazagh Hashemi Shahraki^{1,3}, Mohammad Hashemzadeh^{2,4*}, Rasa Shekari¹

¹ Health Research Center, Ahvaz, Ahvaz, Iran, ² Jundishapur University of Medical Sciences, Ahvaz, Iran, ³ Jundishapur University of Medical Sciences, Ahvaz, Iran, ⁴ Jundishapur University of Medical Sciences, Ahvaz, Iran

AN OUTBREAK OF BACTEREMIAS ASSOCIATED WITH *MYCOBACTERIUM MUCOGENICUM* IN A HOSPITAL WATER SUPPLY

Susan Kline, MD, MPH; Sarah Cameron, MT; Andrew Streifel, MPH; Mitchell A. Yakrus, MS, MPH; Frank Kairis, MT; Keith Peacock, BA; John Besser, MS; Robert C. Cooksey, PhD

What can be learnt from challenging samples

- Exposure to **new organisms** of public health concern – raising awareness of their existence and allowing you to assess suitability of your current method/s or validating new ones
- Raising awareness of **atypical organisms** that exist in the environment and a greater understanding of the impact on laboratory testing and results
- Helps you to understand the **limitations of methods/media** used
- Helps you to understand the **limitations of confirmation tests**
- Allows you to understand **gaps in your procedures** – especially if an approved method is not followed
- Helps your laboratory understand how accurate your test results are



Why participate in PHE Schemes?

- PHE PT samples are designed to challenge your testing procedures therefore will include challenging organisms – so beware
- We extensively test the samples using ISO methods – so your results should align with our results including confirmatory test results
- Process PT samples the same as other routine samples. Otherwise nothing will be learnt about your quality system
- We are not here to trick you but to:
 - raise awareness of the limitation/s of your procedure or method
 - provide an insight into staffs' knowledge and experience
 - endorse the requirement to carry out confirmatory tests
 - encourage the use of approved methods
 - give you an opportunity to examine samples containing organisms less frequently encountered that are of public health concern

We are (possibly) the only ones that

- Use wild strains
- Investigate failures with PT samples properly – a supplementary report and support provided
- Give you access to technical and scientific experts
- Provide unique schemes that supports EU legislation
- Use a unique PHE scoring system to calculate performance as well as including z-scores in reports
- Robust statistic (a lot of participant of each scheme around the world)
- Free of charge repeats



Public Health England Proficiency Tests



Gracias – Thank you