

**Final Report** 

# Introduction of a *Campylobacter* proficiency testing scheme for food laboratories FS101219

# **August 2022**

# UK Health Security Agency – Food and Environmental Proficiency testing Unit

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Special thanks to The Food Standards Agency for appointing FEPTU to deliver this EQA programme.

### **Abbreviations**

Abbreviations	Definition
CFU	Colony forming units
CPT	Campylobacter Proficiency Testing
EQA	External Quality Assessment
FEPTU	Food and Environmental Proficiency Testing Unit
FSA	Food Standards AgencySARS-2-Cov
UKAS	United Kingdom Accreditation Service
UKHSA	UK Health Security Agency
UN	United Nation

### **Executive Summary**

In 2018, the Food Standards Agency (FSA) commissioned UK Health Security Agency (UKHSA; previously known as Public Health England) Food and Environmental Proficiency Testing Unit (FEPTU) to provide a bespoke external quality assessment (EQA) scheme for the detection and enumeration of *Campylobacter* spp. from a simulated matrix of uncooked chicken.

The EQA was sent to a maximum of 20 UK laboratories selected by the FSA that provided an accredited test for *Campylobacter* spp. testing in foods.

In summary, 13 dispatches of this EQA scheme were sent throughout the period March 2019 – November 2021. Three samples were sent with each delivery, representing a total of 39 simulated samples; 26 for enumerating the levels of *Campylobacter* spp. (if detected) and 13 for detection of the organism. Each simulated sample contained varying levels of *Campylobacter* strains (if included), and other microbiological flora found in raw chicken was also included as part of the sample design. The testing methods which laboratories used to enumerate and detect *Campylobacter* spp. in foods was also captured.

Due to the SARS-2-CoV pandemic this EQA exercise was suspended from March 2020 until January 2021, with agreement from the FSA.

Results reported were assessed using a unique UKHSA scoring system and categorised as 2, 1 and 0.

This method of UKHSA scoring system enabled laboratories to be provided with ongoing performance assessment over a period of time.

Individualised reports were generated for each laboratory which included the results reported for each examination and the overall results submitted by all laboratories. After each distribution a confirmation e-mail was sent to the FSA with a list of activities and dates when the work was executed. A file containing the anonymised performance data of the laboratories and a blank summary scheme report was also provided.

Overall, laboratories returned an acceptable or questionable result for a minimum of 80% of samples examined for enumeration or detection of *Campylobacter* spp.. This shows that laboratories can undertake testing for *Campylobacter* spp. in foods to a using both enumeration and detection methods.

EQA provides laboratories with an independent external assessment of their performance. Regular participation in EQA schemes is an important part of laboratories quality procedures and helps to ensure that the results of their tests are accurate. Satisfactory performance with EQA can provide assurance to laboratories that they are compliant with testing standards, thereby meeting and maintaining accreditation requirements. It also provides an assurance step for their clients.

### 1. Introduction

In 2018, the Food Standards Agency (FSA) commissioned UK Health Security Agency's (UKHSA; previously known as Public Health England) Food and Environmental Proficiency Testing Unit (FEPTU) to provide a bespoke external quality assessment (EQA) scheme for the detection and enumeration of *Campylobacter* spp. from a simulated matrix representing uncooked chicken.

UKHSA exists to protect and improve the nation's health and wellbeing and reduce health inequalities. UKHSA is an executive agency, sponsored by the Department of Health and Social Care. FEPTU has over 32 years of experience in providing microbiology EQA schemes to food, water and environmental laboratories both in the UK and in over 70 countries internationally. The full comprehensive services offered by FEPTU can be found on this link: <u>Proficiency testing for food, water and</u> <u>environmental microbiology</u>.

This EQA scheme was available to food laboratories in the UK that provide an accredited test for *Campylobacter* spp. in foods. The selection of laboratories was done by the FSA.

The objective of this exercise was to provide UK testing laboratories with EQA samples that simulated a matrix of uncooked chicken for *Campylobacter* spp. testing. EQA is a useful tool to help laboratories identify process gaps and highlight areas for quality improvements, this will provide the FSA with a better understanding of the performance of laboratories in isolating and enumerating *Campylobacter* spp. in real samples and to make sure that they were all operating at the same standards of detection.

# 1.1 Set up of the exercise

FEPTU is a UKAS accredited (Schedule of Accreditation 0006) EQA provider complying with ISO/IEC 17043:2010 (Conformity assessment - General requirements for proficiency testing).

### **1.2 Confidentiality**

The procedures for the organisation of this EQA ensures that the identity of the laboratories and the association of their performance data is treated as confidential. The laboratories were provided with a unique laboratory identification, log-in and password details. After each distribution a confirmation e-mail was sent to the FSA with a list of activities and dates when the work was executed. A file containing the anonymised performance data of the laboratories and a blank summary scheme report was also provided.

### 1.3 Time frame

In January 2019 laboratories were contacted by the FSA by email to determine if they were interested in taking part in this funded EQA programme. 20 laboratories registered their interest for the first year (March 2019 – January 2020) of this tender which included six dispatches.

Renewal of tender participation for year 2 and 3 was done in December 2019 prior to the start of the next tender period the following year. Laboratories had the option to participate in selected distributions in year 2 and 3.

The 13 dispatches were sent on the following dates:

- 18 March, 7 May, 8 July, 9 September and 11 November 2019
- 13 January and 17 February 2020
- 25 January, 8 March, 10 May, 5 July, 6 September and 1 November 2021

Due to the SARS-2-CoV pandemic this EQA tender was suspended from March 2020 until January 2021 following discussion and agreement with the FSA. Additionally, the FSA also agreed to a temporary suspension of *Campylobacter* sampling (March-May 2020) by retailers for their own testing due to issues arising from Covid-19. All retailers re-commenced sampling by 1 June 2020.

### **1.4 Distribution**

13 dispatches were sent from March 2019 – November 2021. Each dispatch contained three samples, two for reporting a *Campylobacter* spp. enumeration result and one a detection result. Each dispatch and each sample had unique numbers allocated.

Samples were sent as freeze-dried material which, when reconstituted, represented raw chicken. Each simulated sample design contained varying levels of *Campylobacter* strains (if included). Microbiological flora normally found in raw chicken was also included as part of the sample design (see Annex 1).

The simulated freeze-dried samples were dispatched at ambient temperature in approved United Nation (UN) containers. Samples were normally received by the participating laboratory within 72 hours of dispatch, and laboratories were informed to store samples at 4°C until they were ready to process them.

Laboratories were given three weeks to report their results using a secure webbased reporting platform.

Three weeks after the distribution closed, the results were published, and reports were published within three weeks of the closing date.

### **1.5 Instructions to participants**

Paperwork was included with the samples dispatched in the UN boxes, which included:

- specimen details (Annex 2; includes web links to safety instructions and how to report results via a secure web portal)
- instructions on storing and processing of samples (Annex 3)

Laboratories were informed to examine the samples using the methods they currently follow. Details of the method, media, incubation conditions and enrichment broth were provided by the laboratories to FEPTU when the EQA results were reported.

For the second EQA year only two distributions were dispatched by UKHSA. UKHSA is a category one responder to national incidents and therefore UKHSA staff were diverted to support the pandemic.

Following the SARS CoV-2 pandemic (March 2020), whereby administration staff were mandated to work from home, limited paperwork was included in the United Nations (UN) box when the EQA service was re-instated January 2021. Instead, emails were sent a week prior to the dispatch date and included all the paperwork required to process the samples. Laboratories were informed that if they did not receive the samples by the end of the week to contact FEPTU. No formal confirmation of receipt of samples was requested, and instead the confirmation of the laboratory receiving the samples was assumed when results were returned.

# 2. Test Materials

### 2.1 Preparation

The selection of the strains was chosen by the FEPTU team: different species of *Campylobacter* at varying colony forming units (CFU) per mL were included in the repertoire, and background organisms were included to simulate uncooked chicken.

Strains of *Campylobacter* spp. and background organisms were taken from a bank of organisms held in the FEPTU laboratory as fully characterised isolates; these strains were fully characterised in-house using conventional methods and an analytical profile index system (if available). Strains of *Campylobacter* spp. were obtained from an UKHSA *Campylobacter* reference laboratory.

Samples were prepared as freeze-dried material at least five weeks before the dispatch date. This method of preparing samples has been extensively validated and proven to preserve organisms over long period of time and is a method used for other well established accredited schemes provided by UKHSA. The freeze-drying matrix used was inositol serum broth.

Samples in freeze dried format were stored at 4°C.

The simulated samples that contained strains of *Campylobacter* spp. had varied CFU values, from  $1.0x10^2 - 1.0x10^5$ .

### 3. Homogeneity and stability

After sample preparation, post-preservation testing was done to confirm if the sample content matched the expected sample design.

If these results were acceptable, then samples were processed further quantitatively to determine homogeneity for the *Campylobacter* spp. in the sample only. Replicate testing of the *Campylobacter* spp. from 10 randomised samples was tested. Homogeneity results were analysed using robust in-house statistical analyses and used to assess batch acceptance according to in-house procedures. If results were accepted, then they were included in a distribution and then tested for stability prior to an assigned dispatch date.

Stability testing was done on three samples two weeks prior to dispatch date. A further three samples were tested a week after the dispatch date on samples that had been through the postal system, and results were analysed using in-house statistical analyses and would confirm if transit conditions had any impacted on the stability of the *Campylobacter* spp. in the sample.

Samples were tested in the FEPTU laboratory according to international methods to replicate the methodology that laboratories would use on real samples:

- ISO 10272-1:2017 Microbiology of the food chain Horizontal method for detection and enumeration of *Campylobacter* spp. – Part 1 Detection method.
- ISO 10272-2:2017 Microbiology of the food chain Horizontal method for detection and enumeration of *Campylobacter* spp. – Part 2: Colony-count technique.

This is in accordance with food laboratories being accredited to ISO/IEC 17025:2010 (General requirements for the competence of testing and calibration laboratories).

### 4. Assigned values, standard uncertainties and scoring

Results for enumerations and samples were analysed according to ISO 13528:2015 (Statistical methods for use in proficiency testing by interlaboratory comparison).

### 4.1 Assigned values

Assigned values for enumerations (quantitative) were determined using robust statistics; the participants' median value was designated as the assigned value.

Assigned values for detection (qualitative) results was determined by confirmation of microbiological contents within the sample.

Performance assessment forms an integral component for a laboratory to confirm their performance with EQA samples. The data can show a trend in performance either with a specific parameter or as an overall examination in a scheme. Laboratories will receive a report comparing their performance to other participating laboratories. Laboratories can then determine whether their own performance on this PT scheme was acceptable and take appropriate action. UKHSA scoring system is designed to allow for this analysis of performance overtime.

# 4.2 Scoring and evaluation criteria

Scores were applied to reported results as an easy management tool.

Results reported were assessed using a unique UKHSA scoring system and is categorised as 2, 1 and 0. Scores of 2 and 0 are used for qualitative results and translate as 2 being satisfactory and 0 as unsatisfactory. The z-score formula is not used for allocating z-scores for qualitative tests. Participants who report a correct result are allocated a z-score of 0.

### 4.3. Enumeration results (quantitative)

Percentiles are used to identify results outside the expected range by ranking all participants' counts from lowest to highest and calculating the 5<sup>th</sup>, 10<sup>th</sup>, 90<sup>th</sup> and 95<sup>th</sup> percentiles. The 0.5 log<sub>10</sub> rule, which is based on microbiological criteria and has been adapted from a publication by Basil Jarvis (Sampling for Microbiological Analysis in 'The Microbiological Safety and Quality of Food' Volume II, 1999, edited by Lund,

Baird-Parker and Gould), is then applied before scores are allocated so counts within 0.5 log<sub>10</sub> units of the consensus median are re-classified as satisfactory and allocated the maximum score. The value of the maximum score is scheme dependent. As a general rule, questionable and unsatisfactory results should be investigated.

For quantitative schemes a score of 2 is acceptable and results are within the expected range. 1 is used for quantitative results just outside the expected range - these results are questionable as they are considered partially incorrect. 0 is given and considered unsatisfactory when results reported are completely outside the expected ranges (Table 1).

The statistics used to calculate the UKHSA scores for enumeration results is shown in the Table 1

Result type	Calculation	UKHSA scores for enumeration
Expected range	Median ± 0.5 log <sub>10</sub> units or counts within 11 <sup>th</sup> to 89 <sup>th</sup> percentiles	2 (acceptable)
Outlying results (1)	Median ± >0.5 log <sub>10</sub> units and in 6 <sup>th</sup> to 10 <sup>th</sup> or 90 <sup>th</sup> to 95 <sup>th</sup> percentiles	1 (questionable)
Outlying results (2)	Median ± >0.5 log₁₀ units and in 0 <sup>th</sup> to 5 <sup>th</sup> or 96 <sup>th</sup> to 100 <sup>th</sup> percentiles	0 (unsatisfactory)

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UKHSA scores are devised in such a way that they can be combined over a period of time; the cumulative results are summarised in the individualised distribution reports for the last six distributions. Those laboratories with cumulative scores that are less than 80% of the maximum possible score are likely to have some underlying problems with their examinations which should be investigated for a root cause. Participants with cumulative scores of between 80% and 99% are also encouraged to assess their lower scores. In the individual scheme reports z-scores are also calculated and shown, however they have not been used to assess performance of the laboratories as part of this report.

Z-scores provide a statistical means of standardising data points on a single scale so they can be compared. Each z-score corresponds to a point in a normal distribution, describing how far each data point deviates from the consensus median. Table 2 shows the formula used. this information is presented in individual scheme reports and is not used to assess performance of the laboratories as part of this report.

Table 2 Shows the z-scores formulation

 $Z = \underline{(x_{j} - X_{pt})} \sigma_{pt}$ 

Z scores formulation	Details	
Xi	participants' result (expressed as a log <sub>10</sub> value)	
X <sub>pt</sub>	assigned value (participants' consensus median expressed as a log10 value)	
$\sigma_{ ho t}$	the fixed standard deviation for the examination (calculated by FEPTU)	

The  $\sigma_{pt}$  -value expresses the acceptable difference between the individual participant's result and the participants' consensus median. The  $\sigma_{pt}$  -values have been calculated by FEPTU staff using data from previous distributions and the value of  $\sigma_{pt}$  = 0.35 is use.

Therefore, z-scores used in EQA are interpreted as follows:

- z = -1.99 to +1.99 satisfactory
- z = -2 to -2.99 or +2 to +2.99 **questionable**

unsatisfactory

• z = <-3.00 or >+3.00

# 4.4 Detection results (qualitative)

Scores awarded for detection results is shown in Table 3

### Table 3 Shows the criteria for score awarded for qualitative results

Result	UKHSA score	Z-score
Fully correct result	2	0
False positive or false negative result	0	4

# 5. Results and Discussion

### 5.1 General observations

Not all laboratories consistently took all the distributions that were available, therefore the final numbers reporting a result varied by sample and by distribution.

One laboratory did not undertake testing using a detection method and only returned results for enumeration examinations.

Only one laboratory obtained an overall performance of 100% for all the distributions and samples examined.

Non-return of results have been excluded from the all performance calculations as this is not a measurement of a laboratory's testing capabilities.

Testing of three samples for stability after dispatch confirmed that the *Campylobacter* spp. in the samples were stable during the distribution period.

The overall performance by sample, the expected range and the levels of *Campylobacter* spp. (if present) is shown in Annex 4. Performance assessments were designed to identify laboratories with on-going problems with their examinations and were undertaken after every distribution. Scores were allocated to results reported for every sample to help assess participants' performance.

A laboratory's actual identification number was changed to maintain anonymity in Graphs 2, 4 and 5.

### 5.1.1. Statistical evaluation

The organiser advises that for a robust statistical evaluation at least 20 reported results are required for a parameter. Where statistical calculation is based on 10 - 19 results, the results should be interpreted with caution as they may be overly influenced by outlying values. For this bespoke scheme the number of laboratories participating in each sample distribution was usually less than 19.

### 5.2. Performance – enumeration examination (quantitative)

The laboratories were required to report the number of CFU obtained per mL. The expected range was then calculated using the participants' median  $\pm$  0.5 log<sub>10</sub> units, or counts within 11<sup>th</sup> to 89<sup>th</sup> percentiles. A score of 2 was given to laboratories returning results within this expected range (Table 1). The sample content and the overall performance by sample obtained in the expected range is shown in Annex 4.

For enumeration results, a maximum of 19 laboratories (varied for each sample) reported a CFU count for the 26 enumeration samples that were sent out. Three enumeration samples (CPT009, CPT011 and CPT023) did not contain any isolates of *Campylobacter*.

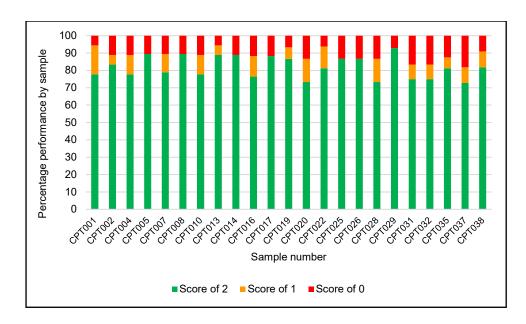
The enumeration performance was analysed in two ways; one by sample (5.2.1) and the other by laboratory (5.2.2).

### 5.2.1 Analysis of enumeration examinations by sample type

For 26 of the samples in this scheme, an enumeration examination was required. However, only 23 samples contained a *Campylobacter* species, while three samples did not contain any *Campylobacter* isolates. The performance of each sample using the assessment criteria given in Table 1 shown in Figure 1.

The overall performance for each sample was calculated by combining the scores awarded (as defined in Table 1) for each sample examined, and then calculating the percentage out of the maximum score available for the number of samples analysed. It needs to be noted that the total number of data sets analysed for each sample varied as not all laboratories (between 11 - 19) returned results for every sample.

Figure 1: Overall performance by samples containing Campylobacter spp.. based on assessment criteria defined in Table 1.



The percentage of laboratories achieving expected results (score of 2) for each sample was between 73-93%, while the percentage of laboratories achieving scores of 1 ranged between 0-17%, and percentage of laboratories achieving scores of 0 ranged from 6-18%.

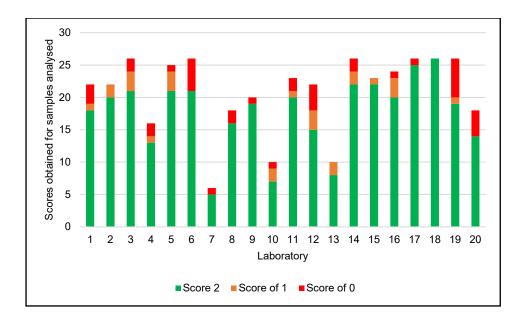
For the three samples (CPT009, 011 and 023) that did not contain isolates of *Campylobacter*, the overall performance was 100%.

### 5.2.2 Analysis of enumeration examinations by laboratory

A total of 20 laboratories were registered for this scheme. However, the number of laboratories that analysed each sample varied.

Figure 2 shows the scores obtained by each laboratory for the samples they analysed (maximum of 26). Criteria for scores awarded is defined in Table 1.

# Figure 2 Scores awarded for each laboratory for samples analysed that required an enumeration examination



Only one laboratory reported all their counts within the expected range across all samples examined (laboratory 18).

As explained, not all laboratories returned results for all enumeration samples: for example, laboratories 4, 7, 10 and 13 did not analyse all enumeration samples sent to them so total scores for individual laboratories were calculated form the maximum samples the laboratory did analyse. All laboratories reported some counts outside the expected range, either in the questionable range (score of 1 given), or in the unsatisfactory range (score of 0 given) however the root cause of this was not within the Organiser's remit to determine.

Laboratory 19 contacted FEPTU early in the programme (before CPT4 dispatch) for some advice as their results were consistently outside the expected range. Following FEPTU advice, their performance significantly improved with most enumeration results being reported in the expected range from CPT4 onwards.

Information on the method used by laboratories was provided when they reported their EQA results online. Numerous options on method parameters were available for laboratories to select from (Annex 2). For the enumeration examinations all laboratories used ISO 10272-2:2017 Microbiology of the food chain — Horizontal method for detection and enumeration of *Campylobacter* spp. - Part 2: Colony-count technique. The media used was modified charcoal cefoperazone deoxycholate agar (mCCDA/CCDA) with an incubation at 41.5 °C for 48 hours.

# 6. Performance – Detection results (qualitative)

The sample content and the overall performance by sample is shown in Annex 4. From the 13 detection samples sent to participating laboratories, Only one sample (CPT030) did not contain a *Campylobacter* species.

For detection results, a maximum of 19 laboratories (varied laboratory participation for each sample) reported a result.

The detection performance was analysed in two ways; one by sample (5.3.1) and the other by laboratory (5.3.2).

# 6.1 Analysis of detection examinations by sample

For 13 of the 39 samples in this scheme a detection examination was required. The performance of each sample using the assessment criteria given in Table 3 is shown in Figure 3.

Figure 3 shows the scores obtained by laboratory for the samples analysed (maximum of 26).

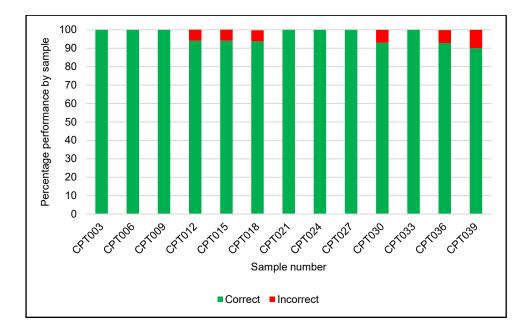


Figure 3 Performance by sample for a detection method.

The number of laboratories that reported results for each detection sample varied between 9-18. For the 13 samples sent to laboratories for detection examination, the percentage of laboratories obtaining the correct result (score 2) was between 90 –

100% and the percentage of laboratories obtaining the incorrect result (score 0) was between 0 – 10% as shown in Figure 3. Only 12 out of 13 detection samples contained isolates of *Campylobacter* spp.. The average percentage of laboratories achieving the correct result for these samples was 97%. For the one sample (CPT030) that did not contain *Campylobacter* spp. 13/14 (93%) laboratories reported the correct result. The overall average percentage of laboratories achieving the correct score for all 13 samples was 97%

Two laboratories (14 and 15) reported a false negative result for two samples: Laboratory 14 for samples CPT036 and CPT039 and Laboratory 15 for samples CPT015 and CPT018. Two laboratories (10 and 12) reported a false negative result for one sample.

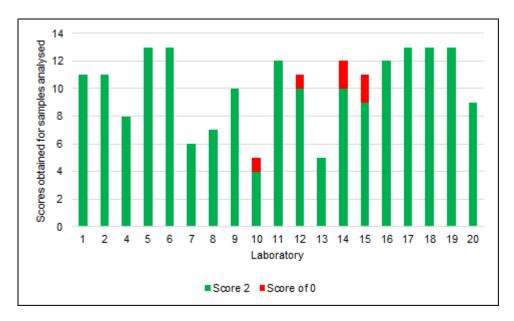
Laboratory 12 also reported a false positive result for the one sample (CPT030) that did not contain a *Campylobacter* spp..

### 6.2 Analysis of detection examinations by laboratory

A total of 19 laboratories were registered for this scheme. However, the number of laboratories that analysed each sample varied.

Figure 4 shows the scores obtained by laboratory for the samples analysed (maximum of 13). Criteria for scores awarded was defined in Table 3.

Figure 4 Scores awarded for each laboratory for samples analysed that required an detection examination.



The overall laboratory performance across all 13 samples is 97% with fifteen laboratories obtaining a score of 2 for all samples they examined.

Four laboratories reported an incorrect detection result, two laboratories on one occasion and two laboratories on two occasions as shown in above Figure 4

As explained, not all laboratories returned results for all detection examinations: for example, laboratories 4, 7, 8, 10 and 13 did not analyse all enumeration samples sent to them. Total scores for individual laboratories were calculated form the maximum samples the laboratory did analyse. Laboratories 10, 12, 14 and 15 reported results outside the expected range in the unsatisfactory range (score of 0 given), however the root cause of this was not within the Organiser's remit to determine.

Laboratory 3 did not examine any samples using a detection method (only did enumeration tests) so is not shown in Figure 4.

Annex 2 lists the numerous options for method parameters available for laboratories to select from. For detection examination all laboratories used ISO 10272-2:2017 Microbiology of the food chain — Horizontal method for detection and enumeration of *Campylobacter* spp. – Part 1: Detection method without any variation to the method. A mixture of media was used by the laboratories but all had used mCCDA/CCDA, and six laboratories also used an additional media. Generally, the enrichment broths used were either Preston or Bolton.

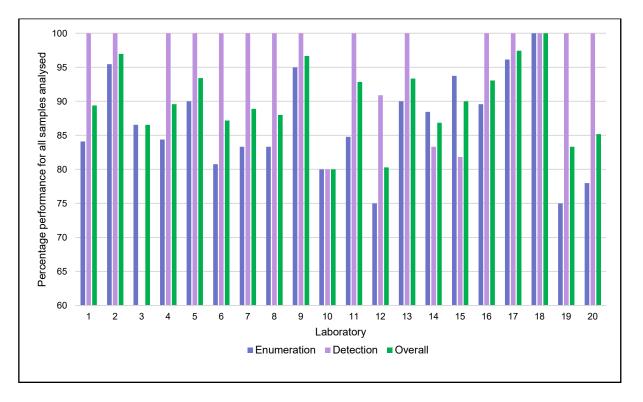
# 6.3 Combining enumeration and detection examinations to find overall performance

The percentage performance of laboratories for all samples and examinations (where reported) is shown in

An overall performance across all samples examined was calculated. This was done by totalling all the scores awarded for both enumeration and detection examination for samples examined (including scores in the questionable range) and a percentage calculated from the total maximum score available.

The laboratory's score for each sample and overall performance for all samples examined is shown in Annex 5.

# Figure 5 Overall percentage performance by laboratory for all the samples analysed



The number of laboratories reporting results for each sample varied between 9 to 19.

The overall percentage performance of laboratories where a result was reported and a score of either 2 (acceptable) or 1 (questionable) was awarded ranged from 80 – 100%. On average, laboratories returned an acceptable or questionable result for 90% of samples examined.

Some of the obvious findings for some laboratories determined are:

Only one laboratory (18) reported all their counts within the expected range, obtaining an overall performance of 100% for all the distributions and samples examined.

Laboratory 5 reported a count outside the expected range for four samples out of the 25 enumeration samples analysed. On three occasions these were in the questionable range (score of 1 given) and once in the unsatisfactory range (score of 0 given). Their overall performance for the 38 samples analysed (including any questionable results scoring 1/2) was 71/76 (93%).

Laboratory 19 contacted FEPTU early in the programme (before CPT4 dispatch) for some advice as their results were consistently outside the expected range. Following FEPTU advice, their performance significantly improved with most enumeration results being reported in the expected range from CPT4 onwards.

Laboratory 20 only joined this scheme from CPT 3 onwards and reported CFU counts outside the expected range (score of 0 or 1) on four occasions. Their overall performance for the 18 samples they analysed was 46/54 (85%).

# 7. Conclusion

Food testing laboratories play a vital role in protecting the public's health, by helping to ensure public health monitoring is accurately done.

FSA commissioned FEPTU and UKHSA to provide an EQA for 20 laboratories selected by the FSA to ensure the results of their testing for *Campylobacter* in chicken was accurate. Distributions sent to the laboratories contained samples that could be used for either enumeration or detection examinations.

There were no issues encountered with the preparation of the simulated specimens/samples. Homogeneity, stability and viability were consistent throughout all the stages of production and distribution.

The *Campylobacter* spp. strains chosen to simulate food samples varied and contained the common species(*jejuni* as well as *lari* and *coli*). High levels of background organisms were included to simulate the contents of a raw chicken. There was no evidence, through quality control checks done in the FEPTU laboratory, that the background flora included in the samples competed with the *Campylobacter* spp..

There was variation in the enumerations results reported, even though all laboratories claimed the same method was used. However, it is widely known that variations in results can be obtained with *Campylobacter* spp. testing, as the media used has inherent performance variability.

One laboratory at the start of the EQA programme consistently reported high counts for *Campylobacter* spp., (CPT1, CPT2 and CPT3) and FEPTU's advice was sought on the methodology. Since this intervention, the laboratory improved their performance and most counts reported since CPT4 were within the expected range.

This shows the educational benefits that can be gained by addressing failures in an EQA. It is an important consideration for laboratories who are looking for continuous improvement.

Laboratories not returning a result or not examining samples were excluded from the overall performance calculations because it was not a measurement of their testing capabilities: one laboratory did not take part in this EQA from distribution CPT4 onwards, two laboratories did not take part from CPT7 onwards and four laboratories did not register for all the distributions available.

Further analysis was done to determine if the counts reported outside the expected range was attributed to the levels of *Campylobacter* spp. in the sample. *Campylobacter* spp. are a group of fastidious organisms, and it is known that media used in enumeration methods can vary from batch to batch as well as variability existing between different manufacturers. The conclusion drawn was that there was no substantive evidence that *Campylobacter* spp. presence in samples was a root cause for reporting results outside the expected range.

Overall, the performance of laboratories participating in the 2019 - 2021 EQA was greater than 80% . There were no issues identified for the detection of *Campylobacter* spp.. However, it must be noted that these samples contained varying levels of this organism to simulate the natural levels found in raw chicken, therefore for samples with high levels of a *Campylobacter* spp. the detection method/s may not have been challenged sufficiently to identify issues if they did exist.

This scheme has been useful to demonstrate the capabilities of laboratories for detecting and enumerating samples for *Campylobacter* spp.. Only one laboratory obtained a total performance of 100% for all samples examined suggesting that there is room for improvement for all the other laboratories.

EQA provides laboratories with an independent external assessment of their performance. Regular participation in EQA schemes is an important part of laboratories quality procedures and helps to ensure that the results of their tests are accurate. Satisfactory performance with EQA can provide assurance to laboratories

that they are compliant with testing standards, thereby meeting and maintaining accreditation requirements. It also provides an assurance step for their clients.

Regular EQA participation will allow laboratories to keep a check with any changes to operating conditions such as media, staffing levels. Addressing gaps identified through an EQA will help ensure that public health incidents are detected early and managed effectively.

### 8. References

None

### 9. Annexes/Appendices

### **Annex 1: Sample information**

A total of 13 distributions were dispatched each containing three samples.

Total of 39 samples were sent, 26 for enumeration testing and 13 for detection examination.

All levels are presented as colony forming units (CFU) per ml reconstituted sample unless indicated otherwise.

Wild strains of organisms were used unless indicated otherwise.

Sample number	CPT001	CPT002	СРТ003
Examination	<i>Campylobacter</i> spp. enumeration	<i>Campylobacter</i> spp. enumeration	<i>Campylobacter</i> spp. detection
Contents:	Campylobacter jejuni (1.1x10 <sup>2</sup> ) Enterococcus faecalis (3.5x10 <sup>3</sup> ) Escherichia coli (7.9x10 <sup>3</sup> ) Pseudomonas aeruginosa (5.0x10 <sup>2</sup> )	Campylobacter coli (8.3x10 <sup>(</sup> ) Cryptococcus albidus (1.6x10 <sup>3</sup> ) Pantoea agglomerans (3.4x10 <sup>4</sup> ) Pseudomonas Iuteola (1.2x10 <sup>5</sup> )	Campylobacter jejuni (2.3x10 <sup>2</sup> ) Escherichia coli O157 (4.4x10 <sup>2</sup> ) Salmonella Essen (35 per disc) Escherichia coli (1.1x10 <sup>4</sup> ) Klebsiella oxytoca (1.7x10 <sup>4</sup> ) Lactococcus lactis (8.7x10 <sup>4</sup> )

### CPT2: 7 May 2019

Sample number	CPT004	СРТ005	CPT006
Examination	<i>Campylobacter</i> spp. enumeration	<i>Campylobacter</i> spp. enumeration	<i>Campylobacter</i> spp. detection
Contents:	Campylobacter coli (5.4x10²) Enterococcus gallinarum (3.2x10⁴) Hafnia alvei (5.7x10³) Micrococcus sp. (1.5.x10⁵)	Campylobacter jejuni (1.4x10 <sup>4</sup> ) Aerococcus viridans (2.3x10 <sup>3</sup> ) (NCTC 8251) Citrobacter braakii (3.2x10 <sup>3</sup> )	Campylobacter coli (5.6x10 <sup>3</sup> ) Enterococcus faecium (1.4x10 <sup>5</sup> ) Pantoea agglomerans (9.1x10⁴)

### CPT3: 8 July 2019

Sample number	CPT007	CPT008	СРТ009
Examination	<i>Campylobacter</i> spp. enumeration	<i>Campylobacter</i> spp. enumeration	Campylobacter spp. detection
Contents:	Campylobacter jejuni (6.8x10 <sup>3</sup> ) Enterococcus faecalis (3.9x10 <sup>5</sup> ) (NCTC 5957) Escherichia coli (5.3x10 <sup>5</sup> ) Pseudomonas putida (4.9x10 <sup>2</sup> )	Campylobacter coli (2.4x10 <sup>4</sup> ) Lactobacillus paracasei (1.3x10 <sup>5</sup> ) Pseudomonas aeruginosa (7.2x10 <sup>4</sup> )	Campylobacter jejuni (8.4x10 <sup>3</sup> ) Klebsiella pneumonia (1.5x10 <sup>5</sup> ) Providencia rettgeri (6.8x10 <sup>5</sup> ) (NCTC 7475) Staphylococcus aureus (2.7x10 <sup>5</sup> )

### CPT4: 9 September 2019

Sample number	CPT010	CPT011	CPT012
Examination	<i>Campylobacter</i> spp. enumeration	<i>Campylobacter</i> spp. enumeration	<i>Campylobacter</i> spp. detection
Contents:	Campylobacter jejuni (2.6x10 <sup>3</sup> ) Micrococcus luteus (1.3x10 <sup>5</sup> ) (NCTC 4819) Serratia liquefaciens (2.0x10 <sup>4</sup> ) Staphylococcus epidermidis (5.0x10 <sup>4</sup> )	Penicillium chrysogenum (2.8x10 <sup>3</sup> ) Enterobacter cloacae (2.1x10 <sup>4</sup> ) Escherichia coli (5.2x10 <sup>4</sup> ) Enterococcus faecalis (6.8x10 <sup>3</sup> ) Lactobacillus paracasei (2.2x10 <sup>4</sup> )	Campylobacter coli (2.2x10 <sup>3</sup> ) Leuconostoc mesenteroides (1.8x10 <sup>4</sup> ) Pantoea species (1.0x10 <sup>5</sup> ) Pseudomonas aeruginosa (6.1x10 <sup>4</sup> )

### CPT5: 11 November 2019

Sample number	CPT013	CPT014	CPT015
Examination	<i>Campylobacter</i> spp. enumeration	<i>Campylobacter</i> spp. enumeration	<i>Campylobacter</i> spp. detection
Contents:	Campylobacter jejuni (4.5x10 <sup>4</sup> ) Citrobacter braakii (1.6x10 <sup>4</sup> ) Aerococcus viridans (3.8x10 <sup>3</sup> ) (NCTC 8251)	Campylobacter coli (2.5x10⁵) Pseudomonas aeruginosa (3.6x10⁵) Staphylococcus sciuri (9.3x10⁵)	Campylobacter lari (4.0x10 <sup>3</sup> ) (NCTC 11845) Cryptococcus albidus (2.0x10 <sup>4</sup> ) Enterobacter amnigenus (5.3x10 <sup>5</sup> )

### CPT6: 13 January 2020

Sample number	CPT016	CPT017	CPT018
Examination	<i>Campylobacter</i> spp. enumeration	<i>Campylobacter</i> spp. enumeration	<i>Campylobacter</i> spp. detection
Contents:	Campylobacter jejuni (1.4x10 <sup>4</sup> ) Escherichia coli (5.2x10 <sup>5</sup> ) Klebsiella oxytoca (3.7x10 <sup>3</sup> ) Lactobacillus brevis (2.7x10 <sup>5</sup> )	Campylobacter coli (3.0x10 <sup>4</sup> ) Enterobacter aerogenes (3.4x10 <sup>5</sup> ) Enterococcus faecalis (3.4x10 <sup>5</sup> ) Escherichia coli (8.0x10 <sup>4</sup> ) Pseudomonas fluorescens (3.5x10 <sup>5</sup> )	Campylobacter coli (4.8x10⁵) Pseudomonas fluorescens (3.6x10⁵) Staphylococcus sciuri (9.3x10⁵)

### CPT7: 17 February 2022

Sample number	CPT019	CPT020	CPT021
Examination	<i>Campylobacter</i> spp. enumeration	<i>Campylobacter</i> spp. enumeration	<i>Campylobacter</i> spp. detection
Contents:	Campylobacter jejuni (6.9x10 <sup>3</sup> ) Escherichia coli (1.0x10 <sup>5</sup> ) Pseudomonas aeruginosa (2.4x10 <sup>5</sup> ) Staphylococcus saprophyticus (2.4x10 <sup>5</sup> )	Campylobacter coli (1.5x10 <sup>5</sup> ) Enterobacter cloacae (1.2x10 <sup>5</sup> ) (NCTC 10005) Kocuria kristinae (7.1x10 <sup>4</sup> ) (NCTC 11038) Pseudomonas fluorescens (1.1x10 <sup>6</sup> ) (NCTC 3756)	Campylobacter lari (4.0x10 <sup>3</sup> ) (NCTC 11845) Cryptococcus albidus (2.0x10 <sup>4</sup> ) Enterobacter amnigenus (5.3x10 <sup>5</sup> )

### CPT8: 25 January 2021

Sample number	CPT022	CPT023	CPT024
Examination	<i>Campylobacter</i> spp. enumeration	<i>Campylobacter</i> spp. enumeration	Campylobacter spp. detection
Contents:	Campylobacter jejuni (1.4 x10 <sup>4</sup> ) Citrobacter braakii (1.5x10 <sup>3</sup> ) Pseudomonas aeruginosa (3.6x10 <sup>4</sup> ) Staphylococcus epidermidis (7.1x10 <sup>3</sup> )	Candida tropicalis (1.3x10 <sup>5</sup> ) Escherichia. coli (9.5x10 <sup>3</sup> ) Enterobacter aerogenes (5.5x10 <sup>3</sup> ) Enterococcus faecalis (3.4x10 <sup>4</sup> ) Lactobacillus brevis (2.2x10 <sup>4</sup> )	Campylobacter lari (7.3x10³) (NCTC 11845) Bacillus cereus (3.4x10⁴) Escherichia coli (6.1x10⁴)

### CPT9: 8 March 2021

Sample number	CPT025	CPT026	CPT027
Examination	<i>Campylobacter</i> spp. enumeration	<i>Campylobacter</i> spp. enumeration	<i>Campylobacter</i> spp. detection
Contents:	Campylobacter coli (1.3x10 <sup>4</sup> ) Candida tropicalis (4.8x10 <sup>4</sup> ) Citrobacter braakii (1.6x10 <sup>5</sup> ) Escherichia coli (6.1x10 <sup>5</sup> )	Campylobacter jejuni (4.8x10 <sup>2</sup> ) Enterococcus faecalis (1.9x10 <sup>5</sup> ) Lactococcus lactis (2.4x10 <sup>5</sup> ) Staphylococcus epidermidis (1.6x10 <sup>5</sup> )	Campylobacter coli (2.1x10 <sup>3</sup> ) Klebsiella oxytoca (3.8x10 <sup>5</sup> ) Lactobacillus brevis (1.5x10 <sup>4</sup> ) Pseudomonas aeruginosa (1.4x10 <sup>6</sup> )

### CPT10: 10 May 2021

Sample number	CPT028	СРТ029	СРТ030
Examination	<i>Campylobacter</i> spp. enumeration	<i>Campylobacter</i> spp. enumeration	<i>Campylobacter</i> spp. detection
Contents:	Campylobacter coli (1.9x10 <sup>3</sup> ) Klebsiella oxytoca (4.9x10 <sup>4</sup> ) Lactobacillus brevis (1.5x10 <sup>4</sup> )	Campylobacter jejuni (3.4x10 <sup>3</sup> ) Escherichia coli (3.4x10 Lactobacillus paracasei (2.8x10 <sup>5</sup> ) Pseudomonas aeruginosa (2.1x10 <sup>5</sup> )	Aspergillus fumigatus (4.3x10 <sup>3</sup> ) Citrobacter braakii (1.3x10 <sup>4</sup> ) Escherichia coli (2.8x10 <sup>4</sup> ) Lactococcus lactis (7.6x10 <sup>5</sup> ) Enterococcus faecalis (3.4x10 <sup>4</sup> )

### CPT11: 5 July 2021

Sample number	CPT031	CPT032	СРТ033
Examination	<i>Campylobacter</i> spp. enumeration	<i>Campylobacter</i> spp. enumeration	<i>Campylobacter</i> spp. detection
Contents:	Campylobacter jejuni (4.9x10⁵) Escherichia coli (4.5x10³) Pediococcus pentosaceus (6.9x10⁴)	(1.9x10⁴́)	Campylobacter jejuni (1.1x10 <sup>3</sup> ) Candida tropicalis (1.4x10 <sup>4</sup> ) Enterococcus faecium (2.4x10 <sup>3</sup> )

### CPT12: 6 September 2021

Sample number	СРТ034	СРТ035	СРТ036
Examination	<i>Campylobacter</i> spp. enumeration	<i>Campylobacter</i> spp. enumeration	<i>Campylobacter</i> spp. detection
Contents:	Escherichia coli (1.6x10 <sup>3</sup> ) Enterobacter cloacae (1.8x10 <sup>3</sup> ) Enterococcus durans (8.0x10 <sup>2</sup> ) Pseudomonas fluorescens (1.5x10 <sup>4</sup> )	Campylobacter coli (3.0x10 <sup>3</sup> ) Enterococcus faecalis (4.0x10 <sup>2</sup> ) Serratia liquefaciens (8.1x10 <sup>3</sup> ) Aspergillus fumigatus (1.0x10 <sup>3</sup> )	Campylobacter jejuni (1.0x10 <sup>2</sup> ) Escherichia coli (2.9x10 <sup>4</sup> ) Klebsiella oxytoca (2.9x10 <sup>4</sup> )

### CPT13: 1 November 2021

Sample number	CPT037	CPT038	СРТ039
Examination	<i>Campylobacter</i> spp. enumeration	<i>Campylobacter</i> spp. enumeration	<i>Campylobacter</i> spp. detection
Contents:	Campylobacter coli (7.3x10 <sup>3</sup> ) Bacillus circulans (2.2x10 <sup>4</sup> ) Escherichia coli (1.9x10 <sup>4</sup> ) Staphylococcus capitis (4.5x10 <sup>3</sup> )	Campylobacter jejuni (1.1x10 <sup>3</sup> ) Candida tropicalis (1.4x10 <sup>4</sup> ) Enterococcus faecium (2.4x10 <sup>3</sup> )	Campylobacter jejuni (4.9x10⁵) Escherichia coli (4.5x10³) Pediococcus damnosus (6.9x10⁴)

### Annex 2: Example document of sample details

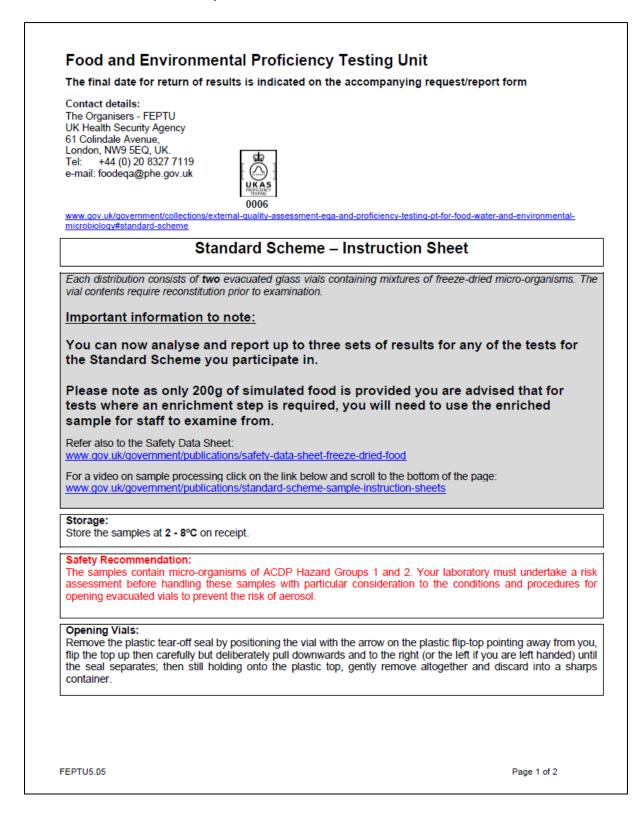
This was provided either in the UN box or sent electronically via email

	b No> Dispatch date: 01 November 2021 Final date for return of results: 22 November 2021				
Contact details: The Organisers - FEPTU JK Health Security Agency 51 Colindale Avenue, ondon, NW9 5EQ, UK. Fel: +44 (0) 20 8327 7119 -mail: foodeqa@phe.gov.uk UKAS 0006					
(	Campylobacter spp. Scheme - Request/Report Form				
Distribution No.: CPT13	Sample numbers: CPT037, CPT038 and CPT039				
Download the sample instruction sheet. A video on sample processing is located further down the webpage:					
	www.gov.uk/government/publications/safety-data-sheet-freeze-dried-food				
Download the safety data sheet:	www.gov.uk/government/publications/safety-data-sheet-freeze-dried-food				
Download the safety data sheet:	www.gov.uk/government/publications/safety-data-sheet-treeze-dned-tood				

	Not examined		ichment media used / nditions of incubation	Media used		Method used	Analyst	CPT037	CPT038	CPT03
			41.5°C/48h	Chromogenic agar - please state		ISO 10272-2:2017	Nominated result:			
Campylobacter spp. (*cfu g <sup>-1</sup> )			Other- please specify	Modified charcoal cefoperazone deoxycholate agar (mCCDA/CCDA)		Other – please specify	Result 2:			
( sid g )				Preston agar						
				Other- please specify			Result 3:			
			Bolton Broth	Chromogenic agar - please state		ISO 10272-1:2017	Nominated result:			
Campylobacter spp.			Preston broth	Modified charcoal cefoperazone deoxycholate agar (mCCDA/CCDA)		PCR	Result 2:			
oampyiobacier spp.	1		Other – please specify	Preston agar	•	VIDAS	·			
				Other- please specify		Other – please specify	Result 3:			
olony forming unit Microbiolog	ist's commen	ts:								

### **Annex 3: Sample instruction sheet**

Available from the weblink provided in document in Annex 2



#### Reconstitution:

- The samples are reconstituted in 200mL nutrient broth (pre-warmed to 30 ± 2°C) as follows:
  Aseptically remove the rubber stopper from the vial of the freeze-dried sample.
  Transfer approximately 1mL of the 200mL warmed nutrient broth to the vial and allow to stand for 2 5 minutes. •
  - Transfer the vial contents back to the remaining nutrient broth. Rinse the vial a further 3 4 times with the **200mL** nutrient broth sample. .
- . Shake the reconstituted samples before examination.

#### Examination:

- Each reconstituted sample is equivalent to a 200g food sample.
- Undertake the examinations listed on the request/report form between 30 - 45 minutes after reconstitution. •
- Examine in accordance with routine procedures.

#### Return of Results:

- Return your results on-line using the web-based reporting system.
- The distribution will be closed on the deadline date for return of results.
- Results cannot be accepted after the distribution has closed. •

### Annex 4: Overall performance by sample

Includes the expected range and the levels of Campylobacter spp. (if present)

CFU per g = colony forming units per gram

### N/A = Not applicable

Expected range has been calculated as: Median  $\pm$  0.5 log10 units or counts within 11th to 89<sup>th</sup> percentiles (Table 1)

Distribution number Date	Sample number	<i>Campylobacter</i> spp. (levels shown are CFU per g)	Examination	Expected range (CFU per g)	Overall percentage performance for a fully correct score of 2 N (%)
CPT1	CPT001	<i>Campylobacter jejuni</i> (1.1x10²)	Enumeration	25 - 4.8x10²	14/18 (78)
18 March	CPT002	<i>Campylobacter coli</i> (8.3x10 <sup>3</sup> )	Enumeration	8.5x10² - 8.5x10³	15/18 (83)
2019	CPT003	<i>Campylobacter jejuni</i> (2.3x10²)	Detection	N/A	17/17 (100)
CPT2	CPT004	<i>Campylobacter coli</i> (5.4x10²)	Enumeration	82 - 8.9x10²	14/18 (78)
	CPT005	Campylobacter jejuni (1.4x10⁴)	Enumeration	1.0x10³ - 1.8x10⁴	17/19 (89)
7 May 2019	CPT006	<i>Campylobacter coli</i> (5.6x10³)	Detection	N/A	18/18 (100)
СРТ3	CPT007	<i>Campylobacter jejuni</i> (6.8x10³)	Enumeration	1.2x10³ - 1.2x10⁴	15/19 (79)
8 July 2019	CPT008	Campylobacter coli (2.4x10⁴)	Enumeration	7.1x10³ - 7.1x10⁴	17/19 (89)
	CPT009	<i>Campylobacter jejuni</i> 8.4x10 <sup>3</sup>	Detection	N/A	18/18 (100)
CPT4	CPT010	<i>Campylobacter jejuni</i> (2.6x10³)	Enumeration	2.7x10² - 5.2x10³	14/18 (78)
9 September	CPT011	No Campylobacter spp.	Enumeration	N/A	18/18 (100)
2019	CPT012	<i>Campylobacter coli</i> (2.2x10 <sup>3</sup> )	Detection	N/A	16/17 (94)
CPT5	CPT013	Campylobacter jejuni (4.5x10⁴)	Enumeration	4.4x10³ - 4.5x10⁴	16/18 (89)

Distribution number Date	Sample number	<i>Campylobacter</i> spp. (levels shown are CFU per g)	Examination	Expected range (CFU per g)	Overall percentage performance for a fully correct score of 2 N (%)
11	CPT014	Campylobacter coli (2.5x10⁵)	Enumeration	3.1x10⁴ - 3.1x10⁵	16/18 (89)
November 2019	CPT015	<i>Campylobacter lari</i> (4.0x10³)	Detection	N/A	16/17 (94)
CPT6	CPT016	<i>Campylobacter jejuni</i> (1.4x10⁴)	Enumeration	1.9x10³ - 2.3x10⁴	13/17 (76)
13 January	CPT017	<i>Campylobacter coli</i> (3.0x10 <sup>4</sup> )	Enumeration	3.5x10³ - 3.5x10⁴	15/17 (88)
2020	CPT018	Campylobacter coli (4.8x10⁵)	Detection	N/A	15/16 (94)
CPT7	CPT019	<i>Campylobacter jejuni</i> (6.9x10³)	Enumeration	1.5x10² - 1.3x10⁴	13/15 (87)
17 February	CPT020	<i>Campylobacter coli</i> (1.5x10⁵)	Enumeration	4.3x10³ - 1.0x10⁵	11/15 (73)
2020	CPT021	<i>Campylobacter lari</i> (4.0x10 <sup>3</sup> )	Detection	N/A	13/13 (100)
CPT8	CPT022	<i>Campylobacter jejuni</i> (1.4x10 <sup>4</sup> )	Enumeration	5.4x10² - 6.0x10³	13/16 (81)
25 January	CPT023	No Campylobacter spp.	Enumeration	N/A	16/16 (100)
2021	CPT024	<i>Campylobacter lari</i> (7.3x10 <sup>3</sup> )	Detection	N/A	15/15 (100)
CPT9	CPT025	<i>Campylobacter coli</i> (1.3x10 <sup>4</sup> )	Enumeration	2.9x10³ - 2.9x10⁴	13/15 (87)
8 March	CPT026	<i>Campylobacter jejuni</i> (4.8x10 <sup>2</sup> )	Enumeration	20 - 7.6x10²	13/15 (87)
2021	CPT027	<i>Campylobacter coli</i> (2.1x10 <sup>3</sup> )	Detection	N/A	15/15 (100)
CPT10	CPT028	<i>Campylobacter coli</i> (1.9x103)	Enumeration	1.1x10² - 2.6x10³	11/15 (73)
10 May	CPT029	<i>Campylobacter jejuni</i> (3.4x10 <sup>3</sup> )	Enumeration	2.1x10 <sup>2</sup> - 3.0x10 <sup>3</sup>	13/14 (93)
2021	CPT030	No Campylobacter spp.	Detection	N/A	14/15 (93)

Distribution number Date	Sample number	<i>Campylobacter</i> spp. (levels shown are CFU per g)	Examination	Expected range (CFU per g)	Overall percentage performance for a fully correct score of 2 N (%)
CPT11	CPT031	<i>Campylobacter jejuni</i> (9.4x10 <sup>4</sup> )	Enumeration	7.7x10² - 4.8x10⁴	9/16 (75)
5 July 2021	CPT032	<i>Campylobacter coli</i> (2.1x10 <sup>3</sup> )	Enumeration	1.6x10² - 2.1x10³	9/16 (75)
	CPT033	<i>Campylobacter jejuni</i> (1.1x10 <sup>3</sup> )	Detection	N/A	9/9 (100)
CPT12	CPT034	No Campylobacter spp.	Enumeration	N/A	16/16 (100)
6 Sontombor	CPT035	<i>Campylobacter coli</i> (3.0x10 <sup>3</sup> )	Enumeration	1.3x10² - 2.5x10³	13/16 (81)
September 2021	CPT036	<i>Campylobacter jejuni</i> (1.0x10 <sup>2</sup> )	Detection	N/A	14/15 (93)
CPT13	CPT037	<i>Campylobacter coli</i> (8.6x10²)	Enumeration	1.3x10² - 2.5x10³	8/11 (73)
1 November	CPT038	<i>Campylobacter jejuni</i> (2.5x10 <sup>4</sup> )	Enumeration	1.5x10³ - 2.1x10⁴	9/11 (82)
2021	CPT039	<i>Campylobacter jejuni</i> (4.9x10 <sup>5</sup> )	Detection	N/A	9/10 (90)

### **Annex 5 - Performance Assessment Sheets for each laboratory**

Performance assessments are designed to identify laboratories with on-going problems with their examinations and are undertaken after every distribution. Scores are allocated to results reported for every sample to help assess participants' performance. See Tables 1 and 2.

Non return of results have been excluded from the overall performance calculations.

### NE = Not examined NR = No return of results

N/A = Did not take distribution

A summary of the performance of each laboratory is shown below:

#### Laboratory 1:

CPT	CPT0	CPT																		
001	002	003	004	005	006	007	008	009	010	011	012	013	014	015	016	017	018	019	020	021
2	2	2	1	2	2	2	2	2	2	2	2	2	0	2	2	0	2	2	2	2

CPT0 22	CPT0 23	CPT0 24	CPT0 25	CPT0 26	CPT0 27	CPT0 28	CPT0 29	CPT0 30	CPT0 31	CPT0 32	CPT0 33	CPT0 34	CPT0 35	CPT0 36	CPT0 37	CPT0 38	CPT0 39	Tot al	O ut of	Overall percent age
0	2	2	2	2	2	2	2	2				2	2	2				59	66	89

Laboratory 2:

CPT	CPT0	CPT																		
001	002	003	004	005	006	007	008	009	010	011	012	013	014	015	016	017	018	019	020	021
2	2	2	2	2	2	2	2	2	1	2	2	2	2	2	2	2	2	2	2	2

CPT0 22	CPT0 23	CPT0 24	CPT0 25	CPT0 26	CPT0 27	CPT0 28	CPT0 29	CPT0 30	CPT0 31	CPT0 32	CPT0 33	CPT0 34	CPT0 35	CPT0 36	CPT0 37	CPT0 38	CPT0 39	Tot al	O ut of	Overall percent age
2	2	2	2	2	2	1	2	2				2	2	2				64	66	97

Laboratory 3:

CPT	CPT0	CPT																		
001	002	003	004	005	006	007	008	009	010	011	012	013	014	015	016	017	018	019	020	021
2	2		2	2		1	2		0	2		2	2		2	2		1	2	

CPT0 22	CPT0 23	CPT0 24	CPT0 25	CPT0 26	CPT0 27	CPT0 28	CPT0 29	CPT0 30	CPT0 31	CPT0 32	CPT0 33	CPT0 34	CPT0 35	CPT0 36	CPT0 37	CPT0 38	CPT0 39	Tot al	O ut of	Overall percent age
2	2		2	2		2	2		0	2		2	2		2	1		45	52	87

Laboratory 4:

CPT	CPT0	CPT																		
001	002	003	004	005	006	007	008	009	010	011	012	013	014	015	016	017	018	019	020	021
2	2	2	2	2	2	2	2	2				1	2	2	2	2	2	2	0	2

CPT0 22	CPT0 23	CPT0 24	CPT0 25	CPT0 26	CPT0 27	CPT0 28	CPT0 29	CPT0 30	CPT0 31	CPT0 32	CPT0 33	CPT0 34	CPT0 35	CPT0 36	CPT0 37	CPT0 38	CPT0 39	Tot al	O ut of	Overall percent age
						2	2	2				2	0	2				43	48	90

CPT	CPT	CPT	CPT	CPT	CPT	CPT	CPT0	CPT		CPT	CPT	CPT	CPT	CP						
001	002	003	004	005	006	007	008	009	010	011	012	013	014	015	016	017	018	019	020	02
1	1	2		0	2	2	2	2	2	2	2	2	2	2	2	2	2	2	1	2
												-								•
CPT0	CPT0	CPT0	CPT0	CPT0	CPT0	CPT0	CPT0	CPT0	CPT0	CPT0	CPT0	CPT0	CPT0	CPT0	CPT0	CPT0	CPT	D Tot	0	Over
22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	al	ut	perce
																			of	age
2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	71	76	93
abora	atory 6																•			
		ODT	ODT	ODT	ODT	ODT	CPT0	CPT	CPT	CPT	CPT	CF								
СРТ	CPT	CPT	CPT	CPT	CPT	CPT	CPIU	OFI		OFI										
-	CPT 002	003	004	005	006	007	008	009	010	011	012	013	014	015	016	017	018	019	020	
CPT 001 2		-														017 2	018 2	019 2		02
001	002	003	004	005	006	007	008	009	010	011	012	013	014	015	016	-			020	02

CPT0 22	CPT0 23	CPT0 24	CPT0 25	CPT0 26	CPT0 27	CPT0 28	CPT0 29	CPT0 30	CPT0 31	CPT0 32	CPT0 33	CPT0 34	CPT0 35	CPT0 36	CPT0 37	CPT0 38	CPT0 39	Tot al	O ut of	Overall percent age
2	2	2	2	2	2	2	2	2	0	0	2	2	2	2	2	2	2	68	78	87

Laboratory 7:

CPT	CPT0	CPT																		
001	002	003	004	005	006	007	008	009	010	011	012	013	014	015	016	017	018	019	020	021
2	2	2	2	2	2	2	0	2												

CPT0 22	CPT0 23	CPT0 24	CPT0 25	CPT0 26	CPT0 27	CPT0 28	CPT0 29	CPT0 30	CPT0 31	CPT0 32	CPT0 33	CPT0 34	CPT0 35	CPT0 36	CPT0 37	CPT0 38	CPT0 39	Tot al	O ut of	Overall percent age
																		16	18	89

Laboratory 8:

CPT	CPT0	CPT																		
001	002	003	004	005	006	007	008	009	010	011	012	013	014	015	016	017	018	019	020	021
2	2	2	2	2	2	2	2	2	0	2	2									

CPT0 22	CPT0 23	CPT0 24	CPT0 25	CPT0 26	CPT0 27	CPT0 28	CPT0 29	CPT0 30	CPT0 31	CPT0 32	CPT0 33	CPT0 34	CPT0 35	CPT0 36	CPT0 37	CPT0 38	CPT0 39	Tot al	O ut of	Overall percent age
2	2	2				2	2	2	2	0		2	2		0	2	2	44	50	88

Laboratory 9:

CPT	CPT0	CPT																		
001	002	003	004	005	006	007	008	009	010	011	012	013	014	015	016	017	018	019	020	021
2	2	2	2	2	2	2	2	2	2	2	2	2	2	2				2	2	2

CPT0 22	CPT0 23	CPT0 24	CPT0 25	CPT0 26	CPT0 27	CPT0 28	CPT0 29	CPT0 30	CPT0 31	CPT0 32	CPT0 33	CPT0 34	CPT0 35	CPT0 36	CPT0 37	CPT0 38	CPT0 39	Tot al	O ut of	Overall percent age
2	2	2	2	2	2	0	2	2	2	2	2							58	60	97

Laboratory	1	0:
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CF	CPT	CPT	CPT	CPT	CPT	CPT	CPT0	CPT												
00	002	003	004	005	006	007	008	009	010	011	012	013	014	015	016	017	018	019	020	021
			1	2	2	2	0	2	2	2	0	2	2	2	1	2	2			

CPT0 22	CPT0 23	CPT0 24	CPT0 25	CPT0 26	CPT0 27	CPT0 28	CPT0 29	CPT0 30	CPT0 31	CPT0 32	CPT0 33	CPT0 34	CPT0 35	CPT0 36	CPT0 37	CPT0 38	CPT0 39	Tot al	O ut of	Overall percent age
																		24	30	80

Laboratory 11:

CPT 001	CPT 002	CPT 003	CPT 004	CPT 005	CPT 006	CPT 007	CPT0 008	CPT 009	CPT 010	CPT 011	CPT 012	CPT 013	CPT 014	CPT 015	CPT 016	CPT 017	CPT 018	CPT 019	CPT 020	CPT 021
2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2

CPT0 22	CPT0 23	CPT0 24	CPT0 25	CPT0 26	CPT0 27	CPT0 28	CPT0 29	CPT0 30	CPT0 31	CPT0 32	CPT0 33	CPT0 34	CPT0 35	CPT0 36	CPT0 37	CPT0 38	CPT0 39	Tot al	O ut of	Overall percent age
2	2	2	2	2	2	1		2				2	2	2	0	0	2	65	70	93

Laboratory 12:

CPT	CPT0	CPT																		
001	002	003	004	005	006	007	008	009	010	011	012	013	014	015	016	017	018	019	020	021
1	2	2	0	2	2	2	2	2	2	2	2	2	2	2	0	0	2			

CPT0 22	CPT0 23	CPT0 24	CPT0 25	CPT0 26	CPT0 27	CPT0 28	CPT0 29	CPT0 30	CPT0 31	CPT0 32	CPT0 33	CPT0 34	CPT0 35	CPT0 36	CPT0 37	CPT0 38	CPT0 39	Tot al	O ut of	Overall percent age
2	2	2	2	0	2	2	2	0	1	1	2	2	2	2				53	66	80

Laboratory 13:

		•																		
CPT 001	CPT 002	CPT 003	CPT 004	CPT 005	CPT 006	CPT 007	CPT0 008	CPT 009	CPT 010	CPT 011	CPT 012	CPT 013	CPT 014	CPT 015	CPT 016	CPT 017	CPT 018	CPT 019	CPT 020	CPT 021
2	2	2	2	2	2				1	2	2	2	2	2	1	2	2			

CPT0 22	CPT0 23	CPT0 24	CPT0 25	CPT0 26	CPT0 27	CPT0 28	CPT0 29	CPT0 30	CPT0 31	CPT0 32	CPT0 33	CPT0 34	CPT0 35	CPT0 36	CPT0 37	CPT0 38	CPT0 39	Tot al	O ut of	Overall percent age
																		28	30	93

Laboratory 14:

CPT	CPT0	CPT																		
001	002	003	004	005	006	007	008	009	010	011	012	013	014	015	016	017	018	019	020	021
1	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	0	2	

CPT0 22	CPT0 23	CPT0 24	CPT0 25	CPT0 26	CPT0 27	CPT0 28	CPT0 29	CPT0 30	CPT0 31	CPT0 32	CPT0 33	CPT0 34	CPT0 35	CPT0 36	CPT0 37	CPT0 38	CPT0 39	Tot al	O ut of	Overall percent age
2	2	2	0	2	2	2	2	2	2	2	2	2	1	0	2	2	0	66	76	87

Laboratory	/ 1	5:
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CPT 001	CPT 002	CPT 003	CPT 004	CPT 005	CPT 006	CPT 007	CPT0 008	CPT 009	CPT 010	CPT 011	CPT 012	CPT 013	CPT 014	CPT 015	CPT 016	CPT 017	CPT 018	CPT 019	CPT 020	CPT 021
2	2	2	2	2	2	2	2	2	2	2	2	2	2	0	2	2	0	2	1	2
		1	1	1	1	1	1	1	T	1	T									
CPT0	CPT0	CPT0	CPT0	CPT0	CPT0	CPT0	CPT0	CPT0	CPT0	CPT0	Tot	0	Overall							
22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	al	ut	percent
																			of	ade

																of	age
2	2	2	2	2	2		2	2	2	0	2	2	2	2	63	70	90

### Laboratory 16:

CPT	CPT0	CPT																		
001	002	003	004	005	006	007	008	009	010	011	012	013	014	015	016	017	018	019	020	021
2	0	2	2	2	2	1	2	2	2	2	2	2	2	2	2	2	2	2	2	

CPT0 22	CPT0 23	CPT0 24	CPT0 25	CPT0 26	CPT0 27	CPT0 28	CPT0 29	CPT0 30	CPT0 31	CPT0 32	CPT0 33	CPT0 34	CPT0 35	CPT0 36	CPT0 37	CPT0 38	CPT0 39	Tot al	O ut of	Overall percent age
1	2	2	2	2	2				2	2	2	2	2	2	1	2	2	67	72	93

Laboratory 17:

CPT	CPT0	CPT																		
001	002	003	004	005	006	007	008	009	010	011	012	013	014	015	016	017	018	019	020	021
2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2

CPT0 22	CPT0 23	CPT0 24	CPT0 25	CPT0 26	CPT0 27	CPT0 28	CPT0 29	CPT0 30	CPT0 31	CPT0 32	CPT0 33	CPT0 34	CPT0 35	CPT0 36	CPT0 37	CPT0 38	CPT0 39	Tot al	O ut of	Overall percent age
2	2	2	2	2	2	2	0	2	2	2	2	2	2	2	2	2	2	76	78	97

Laboratory 18

CPT	CPT0	CPT																		
001	002	003	004	005	006	007	008	009	010	011	012	013	014	015	016	017	018	019	020	021
2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2

(	CPT0 22	CPT0 23	CPT0 24	CPT0 25	CPT0 26	CPT0 27	CPT0 28	CPT0 29	CPT0 30	CPT0 31	CPT0 32	CPT0 33	CPT0 34	CPT0 35	CPT0 36	CPT0 37	CPT0 38	CPT0 39	Tot al	O ut of	Overall percent age
	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	78	78	100

Laboratory 19:

CPT	CPT0	CPT																		
001	002	003	004	005	006	007	008	009	010	011	012	013	014	015	016	017	018	019	020	021
0	0	2	0	0	2	0	2	2	2	2	2	2	0	2	2	2	2	2	2	2

CPT0 22	CPT0 23	CPT0 24	CPT0 25	CPT0 26	CPT0 27	CPT0 28	CPT0 29	CPT0 30	CPT0 31	CPT0 32	CPT0 33	CPT0 34	CPT0 35	CPT0 36	CPT0 37	CPT0 38	CPT0 39	Tot al	O ut of	Overall percent age
1	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	2	65	78	83

Labora	Laboratory 20:																			
CPT 001	CPT 002	CPT 003	CPT 004	CPT 005	CPT 006	CPT 007	CPT0 008	CPT 009	CPT 010	CPT 011	CPT 012	CPT 013	CPT 014	CPT 015	CPT 016	CPT 017	CPT 018	CPT 019	CPT 020	CPT 021
001	002	003	004	005	006	007	000	009	010	011	012	013	014	015	010	017	010	019	020	021
						2	2	2	2	2	2	2	2	2	0	2	2	2	2	2
			[	r		T	Т		1	T	T	1	1		T	-1	-1			
CPT0	CPT0	CPT0	CPT0	CPT0	CPT0	CPT0	CPT0	CPT0	CPT0	CPT0	CPT0	CPT0		CPT0	CPTC		-		0	Overall
22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	al	ut of	percent age
2	2	2	0	0	2	0	2	2				2	2	2				46	54	85

End of report