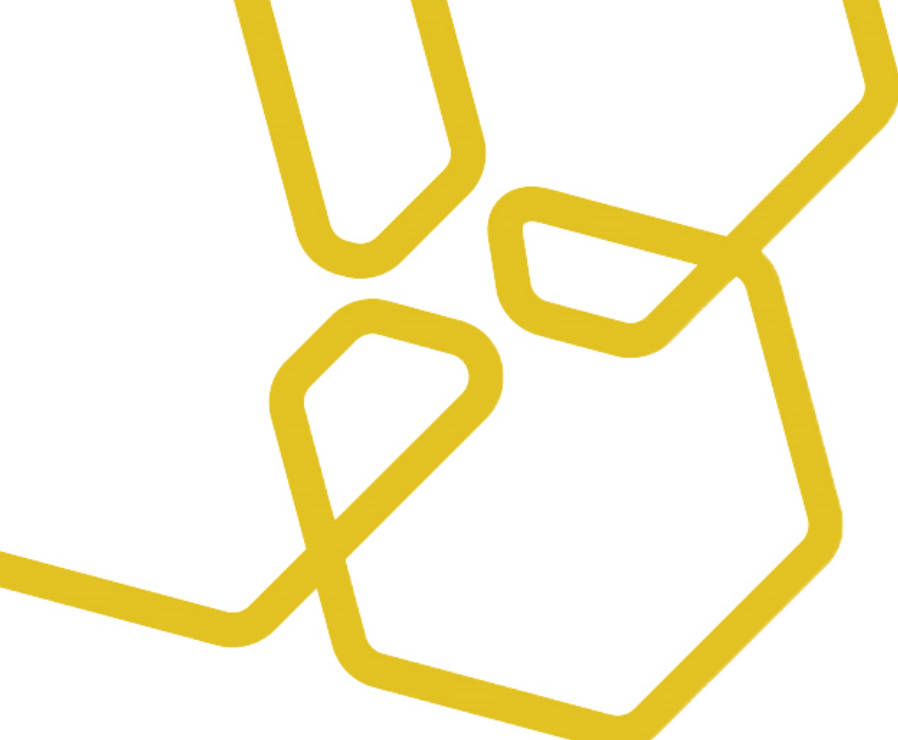


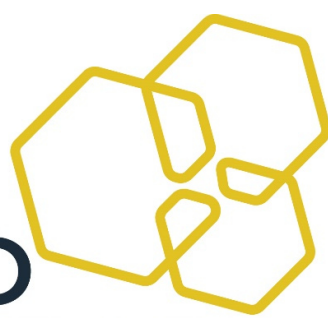


Extended User Instructions for Path-Chek Hygiene swabbing kits

Path-Chek Hygiene Pathogen detection is intended for the detection of important food pathogens on food contact surfaces in food handling and manufacturing environments after cleaning and sanitizing has been performed.

Cat. No. PCS-100

For 100 tests



**GOLD
STANDARD
DIAGNOSTICS**

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1 INTRODUCTION

The Food Safety Pyramid (**Figure 1.**) provides food manufacturers and processors at all levels with a simple structure to enable the protection against microbial contamination (which have the potential to cause food spoilage and food poisoning) for both the products manufactured by the company, and the customers who may later consume these products.

Figure 1.: Food Safety pyramid



Microorganisms are always present in food handling environments. These microorganisms can be characterised as belonging to 2 distinct groups: transient and resident. Transient microorganisms are usually introduced into the food environment through raw materials, water and employees. Normally the routine application of good sanitation practices are able to kill these organisms. However, if contamination levels are high or sanitation procedures are inadequate, transient microorganisms may be able to establish themselves, multiply and become resident. Organisms such as *Coliforms* and *Salmonella* spp. and *Listeria* spp. have a well-established history of becoming residents in food handling environments.

2 ENVIRONMENTAL MONITORING

Food processors should employ environmental sampling programs to monitor for general levels of hygiene (the efficacy of general cleaning and sanitation for the removal of transient microorganisms) or indicator testing may be achieved through a variety of methods including visual inspection, ATP monitoring or the detection of surface protein residues. In addition, pathogen specific environmental sampling should be undertaken to monitor for the presence of specific pathogens that may be present as transient or resident microorganisms. The detection of specific pathogens serves two important roles. Firstly, it highlights the presence of important food pathogens which may have been introduced into a food handling environment but may not have been eliminated by routine sanitation practices.

Therefore may be passed onto other food materials being processed. Secondly, it assists in determining sources of these important pathogens that may be resident.

3 ENVIRONMENTAL SAMPLING PROGRAMS

3.1 Introduction

All food handling companies and establishments should employ an environmental sampling program to monitor for food spoilage microorganisms and food poisoning pathogens. Such a program, if well designed will enable the detection of unacceptable microbial contamination in a timely manner. Over the last decade environmental monitoring has changed from essentially random sampling, employing imaginary grids over a production area and testing points within each grid, to current methods that are focused on risk assessment to determine the most appropriate methods for monitoring. Sampling programs should include the collection of samples during production on a regular basis from work surfaces in a randomised manner which will reflect the differing working conditions. In addition, samples should be taken from these sites after sanitising and from sites which may serve as harbors of resident organisms. Sampling should not only be conducted on food contact surfaces, but the evaluation of non-food contact surfaces such as conveyor belts, rollers, walls, drains and air is equally as important as there are many ways (aerosols and human intervention) in which microorganisms can migrate from non-food contact surfaces to food. The results of these samples should be tabulated as soon as available and in such a way that they can be compared with previous results in order to highlight trends.

3.2 Determining the Frequency of Monitoring

The development of an effective environmental monitoring program should reflect a balance between employing the available resources efficiently and monitoring at sufficient intervals so as to ensure that a meaningful picture of the levels and nature of bacterial contamination can be obtained. When establishing an environmental monitoring program, the frequency of monitoring different areas may be determined based on “Criticality Indexes” relevant to each specific processing area or environment, or by using a Zones of Risk System.

3.2.1 Critical Indexes

The development of a criticality program on which monitoring frequencies can be based should be focused on the targeting of the critical steps in the manufacturing process. Therefore, the final manufactured product should receive more monitoring than early manufacturing steps i.e. end product testing of product. The use of “criticality indexes” provides a means whereby the frequency of monitoring can be assigned to each designated critical area. The assessment of risk should be based on the potential impact any risk may have on the final quality or safety of the products being manufactured e.g. exposure to low temperatures would constitute a low risk whilst the presence of water or warmer temperatures would constitute a high risk. The scheme employed by each food handling or manufacturing environment will necessarily be unique to the processes and types of food being handled in that environment. An example of how such a scheme can be constructed is shown in Table 1.

In the development of such a scheme, all manufacturing areas should be evaluated against a series of guiding questions which may include:

Higher Weighting should be given to:

- Dirtier activities.
- Areas where dirty activities are performed in close relative proximity to clean areas.
- Areas which are often wet.
- Areas with open drains.
- Areas with high levels of staff activity.

Higher Monitoring Frequencies should be assigned to:

- Warm or ambient handling areas as opposed to cold rooms.
- Areas with sinks, drains or ongoing wetness as opposed to dry areas.
- Areas where unprocessed raw foods are handled.
- Product filling.
- Packaging

Table 1: Criticality Indexes and Monitoring Frequency

Criticality Index	Frequency of Monitoring
1	Daily or Each Batch
2	Weekly
3	Fortnightly
4	Monthly
5	Three Monthly
6	Six Monthly

Once the critical factors have been established a final Monitoring Schedule can be developed (**Table 2.**). Most food manufacturing processes involve one or more steps that effectively kill pathogenic bacteria, although manufacturing involving the production of fresh (salads etc.), frozen products (vegetables, meats, poultry, and fish) or some dairy products may not. In manufacturing processes designed to kill bacteria, the challenge is to prevent the processed food from becoming re-contaminated. In these situations, food handling surfaces and possibly equipment become contaminated by bacteria travelling through the food processing environment through a series of steps before finally coming into contact with the food that has been processed.

Listeria species for example can multiply rapidly to high numbers on wet areas such as floors and drains and then be transferred to conveyor belts and benches through human intervention or the use of high-pressure water hose that both can result in the production aerosols.

Any processed food that subsequently touches these surfaces may become re-contaminated. In the development of any monitoring program, post-processing environmental monitoring should always be considered as likely areas where pathogens may reappear and contaminate food post-processing. Generally, these post processing environments should be relatively free of bacteria when production commences. After periods of production, it should be expected that the level of bacterial contamination of these areas should increase. However, the presence of microorganisms normally present in the pre-processed foods should not be expected.

Table 2: Monitoring Schedule Based on the Determination of Criticality Factors

Criticality Index	Likelihood of Impact on Finished Product	Definition	Frequency of Monitoring
1	Highly Likely	Mixing and Filling Machines workplaces are sanitized daily	Daily or Each Batch
2	Likely	Packaging areas or areas in which final handling is performed	Weekly
3	Moderately Likely	Areas where process food is exposed to the environment	Fortnightly
4	Unlikely	Cold areas where little or no processing is performed	Monthly
5	Very Unlikely	Areas in which indirect exposure to prepared and packaged product is unlikely	Three Monthly
6	Highly Unlikely	Any are that is uncontrolled or where microbial contamination is very unlikely such as freezer	Six Monthly

3.3 Zones of Risk System Approach

A more simplistic approach that can be adopted is to employ a program based around zones that have different levels of risk. A typical three zone plan for example would divide a production facility into zones that cover low, medium and high risk areas within the production facility, with high risk zones being those in which there are direct contact between food products and surfaces (International Commission on Microbiological Specifications of Foods. 2002. Microorganisms in Foods. 7. Microbiological Testing in Food Safety Management. Blackwell Scientific. London.). These high risk areas would be those most stringently monitored (See Table 3.).

Table 3: Zones of Risk Analysis

Zone	Food Contact Surfaces
1	After dryer Pipes Conveyor belts Silos
2	Lids Covers External Surfaces of Silos
3	Floors Walls Pipes

4 WHAT TO MONITOR

Organisations involved in food handling should employ environmental monitoring as a means of:

1. Monitoring the general levels of hygiene within the environment in question. The monitoring of the general level of hygiene provides an overall impression of the level of cleanliness within the test environment – it measures the efficiency of the general cleaning and sanitation procedures in place and their ability to remove food residues and transient microorganisms. A variety of methods are available to achieve this task, including general physical inspections, ATP Monitoring Systems and the detection of the presence of food residues (generally protein).
2. Environmental microbiological monitoring for the presence of specific pathogens within the processing environment. The detection of specific pathogens serves two important roles:
 - a) It highlights the presence of important food pathogens which may have been introduced into the food handling environment generally through human contact or from raw ingredients, but which may not have been eliminated by routine cleaning and sanitation procedures.
 - b) Secondly, it highlights the sources of these pathogens that may be resident in the environments being tested.

Microbiological environmental monitoring should be used to indicate either unacceptable conditions or practices which in turn should aid in controlling pathogenic bacteria such as Salmonella and Listeria.

The presence of Coliforms may also be valuable as they will provide an indication of the general levels of microbiological cleanliness within test environments.

5 HOW TO COLLECT SWAB SAMPLES

At this point in time no detailed standard methods (AOAC, USDA FSIS, USFDA etc.) exist for the performance of microbiological environmental monitoring. ISO 18593:2004(E) Microbiology of food and animal feeding stuffs – Horizontal methods for sampling techniques from surfaces using contact plates and swabs does however provide a general platform for the critical steps that should be considered in the development of testing procedures. The key elements of this standard that should be taken into consideration include:

1. Moistened swabs should be employed for all sampling of surfaces. (Item 4.1.)
2. The solution used to moisten swabs should neutralise any detergents and sanitisers employed. (Item 6.1.)
3. Swab moisturizer solution must preserve the integrity of the sample i.e. bacterial numbers should remain constant until the sample collected onto the swab can be evaluated. (Item 6.1. and 7.)
4. Wherever possible the size of the area sampled should be greater than 100cm². (Item 8.2.)
5. The analysis of the samples for specific pathogens is achieved by transferring the swabs into an appropriate enrichment broth. (Item 8.2)
6. After enrichment transfer a sample to an appropriate agar plate medium for the target organism being sought. (Item 8.3.4.)
7. Report the target microorganism as present or absent. (Item 9.2.5.)
8. The contact plate method (including dipslides, RODAC plates and 3M Petrifilm™) shall not be used for the specific detection of pathogenic microorganisms. (Item 8.1.)

6 DETECTION OF TARGET MICROORGANISMS

As discussed above, the examination of environmental swabs for specific food pathogens is not described in any specific standard methods, however the general principle that the analysis of the samples for specific pathogens can be achieved by transferring swabs into an appropriate enrichment broth can be applied to any specific pathogen being sought.

In such instances, methods such as AOAC and USDA FSIS methods can be applied to any specific pathogen. In doing so however, it must be recognized that different methods employ different culture media and brands (and hence recipes) of different culture media will vary in terms of their selectivity, sensitivity and specificity. On this basis, although methods such as these may be applied, results may vary.

Therefore, it should be standard practice that prior to the adoption of any method it should be thoroughly evaluated to ensure that it is capable of recovering low numbers (<10 CFU) of the target organism whilst inhibiting high numbers (>103 CFU) of potentially competing organisms. Any system should also be evaluated to ensure that it is capable of recovering damaged organisms in low numbers.

The recovery of *Listeria monocytogenes* for example may be achieved using one of the following combinations employed in various standard methods (Table 4.).

Table 4.: Culture media suitable for the selective enrichment of Listeria spp.

Method	24 Hours	48 Hours
USDA FSIS	UVM Broth	Fraser Broth
Health Canada (HFLP-38,2022)	PALCAM Broth	UVM2 Broth
USFDA (BAM) AOAC/FDA	LEB	LEB

7 EVALUATION OF THE RESULTS

Following sampling and microbiological analysis a series of results will be available that provide an indication of the overall levels of hygiene in the processing environments evaluated. This body of information provides a valuable tool for maintaining and improving the quality and safety of products. In addition, the detection of specific pathogens such as *Salmonella* and *Listeria* is critical in ensuring food safety for the consumer.

It is not uncommon for food manufacturers only to react to unacceptable results when these pathogens appear when final food products are evaluated however, it is important (particularly in high risk products) that on-going environmental sampling practices are implemented and performed. The evaluation of samples and sampling plans and the test data generated over extended periods should lead to changes in test sample frequency and location, which in turn should lead to improvements in cleaning and sanitising practises.

8 PATH-CHEK HYGIENE PATHOGEN DETECTION

The Path-Chek Hygiene Pathogen Detection product range is a range of microbiological environmental monitoring products that combine all of the requirements of the ISO Standard, ISO 18593:2004(E) Microbiology of food and animal feeding stuffs – Horizontal methods for sampling techniques from surfaces using contact plates and swabs and the pathogen isolation, and detection methods employed in pathogen isolation and detection methods such as those described in AOAC, USDA FSIS, USFDA methods into a single convenient package.

8.1 Features

The following is a list of the key features of the Path-Chek Hygiene Pathogen Systems (Table 5.):

Table 5.: Key Features of Path-Chek Hygiene Pathogen Systems

PRODUCT FEATURE	BENEFITS TO USERS
Pre-Moistened Swabs	- Meets the requirements of ISO 18593:2004(E) Microbiology of food and animal feeding stuffs – Horizontal methods for sampling techniques from surfaces using contact plates and swabs. - This improves the recovery of organisms from both wet and dry surfaces, increasing the sensitivity of the test.
Swab Moisturiser Neutralises Detergents and Sanitisers	The wetting agent neutralises the effects of residual detergents and sanitisers remaining on surfaces after cleaning, this maintains integrity of the sample if such residues are present.
Swab Moisturizer Preserves the Integrity of the Sample	The wetting agent ensures that the samples introduced into the various Pathogen Detection Broths are representative of the sample taken. This is especially important if there are delays in transferring the sample swabs to the testing laboratory.
Swab sticks have special “break point”	The shaft of the swabs used have a special “break point” to simplify the transfer of samples into the Pathogen Detection Broths.
A Range of Individual Growth Media	Three different individual Pathogen Detection Broths are currently available: - Coliforms – generally used as an indicator of overall hygiene conditions. - Salmonella – an important cause of food poisoning. - Listeria – a very important foodborne organism that causes a range of diseases with potentially high mortality rates.

PRODUCT FEATURE	BENEFITS TO USERS
High Sensitivity and Specificity	- Sensitivity – able to detect <1 organism per 10cm² of surface tested. - Specificity – high levels of specificity minimize problems with false positive tests.
Detection Media and Confirmation Methods Compliant with International Standards	The Specific Growth Media, the Methods of Use and the Recommended Confirmation Methods all comply with recognised Food Standards such as USDA/ FSIS, US FDA etc. Meets laboratory accreditation requirements.

8.2 Kit Description

Path-Chek Hygiene Pathogen Systems are a range of screening tests intended for use in food handling and manufacturing environments and on food contact surfaces for the detection of *Coliforms*, *Salmonella ssp.* *Listeria* species. Path-Chek Hygiene Pathogen Systems should be considered as a fundamental component of Good Manufacturing Practice and an integral component of any HACCP (Hazard Analysis Critical Control Points) plan, providing a highly sensitive and specific indication of the presence of the foodborne pathogen being tested for.

8.3 Kit Contents

Cat. no. PC010 (100 reactions)

- 100x **Coliform** Detection Broth (3 mL)
- Instructions for Use

Cat. no. PC020 (100 reactions)

- 100x **Salmonella** Detection Broth (3 mL)
- Instructions for Use

Cat. no. PC080 (100 reactions)

- 100x **Listeria** Detection Broth (3 mL)
- Instructions for Use

Cat. no. PCS-100 (100 reactions)

- 100x **Pre-moistened swabs** (100 piece)
- Instructions for Use

8.4 Additional Materials

- Incubator set to appropriate temperature
- Rack to hold tubes during incubation
- Template 10 x 10 cm to assist with sampling

Note: The Path-Chek Hygiene Detection Broths (3 mL) and Pre-moistened Path-Chek Hygiene swabs, (CODE: PCS-100 per 100 swabs) are purchased separately and combined to create the appropriate Path-Chek system.

8.5 Storage and Shelf Life

The Path-Chek Hygiene Detection Broths should be stored at 2 – 8 °C when not in use. The pre-moistened sample swabs should be stored at 4 – 25 °C. Both components should not be used after the expiry date printed on the carton label.

8.6 Procedure

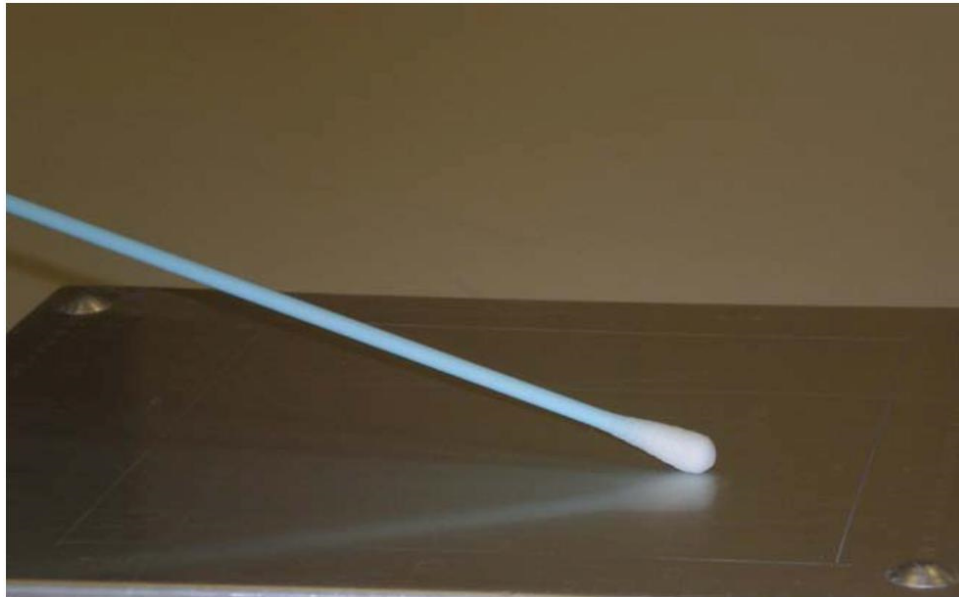
STEP 1

Carefully remove the cap from the pre-moistened Path-Chek Hygiene swab.



STEP 2

Thoroughly swab a standard sample area (10 x 10 cm), rotating the swab as the sample is being collected. If sample areas are irregular develop a standard sampling procedure which is documented and used consistently.



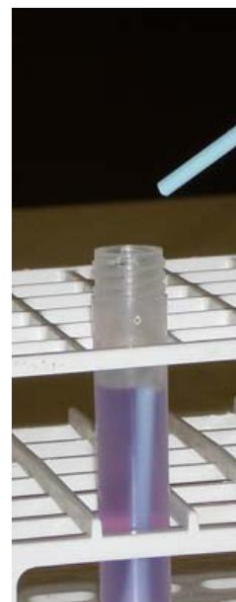
STEP 3

After swabbing the test area, aseptically remove the cap from the Path-Chek Hygiene Detection Broth and carefully place the swab into the tube. If the swab cannot be transferred immediately into the Path-Chek Hygiene Detection Broth, return it to its holding tube and store in a cool place. Label the Path-Chek Hygiene Detection Broth or the holding tube for the swab.



NOTES:

1. Swabs should be placed into the Path-Chek Hygiene Detection Broth at an angle of 45° with the tip of the swab against the side of the tube. Press down on the shaft of the swab. The shaft of the swab will break at breakpoint of the swab, 45 mm from the swab tip.



2. If the swab cannot be transferred immediately into the Path-Chek Hygiene Detection Broth, the swab should be returned to its holding tube and store in a cool place. Swabs may be held at a maximum temperature of 20 °C for up to 24 hours.

STEP 4

Place inoculated tubes into a suitable rack and incubate at 35-37 °C, for 18-24 hours for Coliforms and Salmonella, and at 28-30 °C, for 24-48 hours, for Listeria.

Note: If Path-Chek Hygiene Listeria is incubated at 35-37 °C there will, in certain circumstances, be an increased risk of false positives.

STEP 5

Observe for colour changes and record the results. A positive result may be interpreted as early as 18 hours however, results must not be considered as negative until the Path-Chek Hygiene Detection Broth has been incubated for up to 24 hours for the Coliform and Salmonella spp. systems and 48 hours for the Listeria spp. system.

Listeria

(+)

(-)

Coliforms

(+)

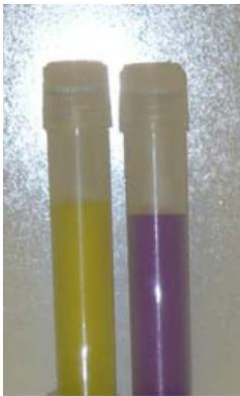
(-)

Salmonella

(+)

(-)




STEP 6
Interpretation:

Table 6.: Interpretation Criteria for Path-Chek Hygiene Detection Broths.

System	Positive	Negative
<i>Coliform</i>	Yellow	Purple
<i>Salmonella</i>	Black	Purple/ Yellow
<i>Listeria</i>	Black	Straw Color

STEP 7
OPTIONAL CONFIRMATION PROCEDURES

Presumptive positive tests may be confirmed by sub-culturing a drop of the growth medium onto an appropriate selective agar plate medium for the organism being tested.
 The use of the following media will provide compliance with standard testing methods such as BAM and USDA/FSIS etc

Table 7.: Culture media for Confirmation of Positive Detection Broths.

System	Media
<i>Coliform/ E.coli</i>	mENDO VRBA
<i>Salmonella</i>	XLD Bismuth Sulphite
<i>Listeria</i>	Oxford Palcam ALOA

After incubation at 35 - 37°C for 24 – 48 hours, plates should be examined for colonies resembling the targets being sought.

Any suspect colonies should be further identified using more definitive tests such as microscopy and biochemical tests such as the Microgen® GNA ID (MID-64) and GNB ID (MID-65) and Listeria ID (MID-67).

9 PERFORMANCE CHARACTERISTICS

9.1 Pre-Moistened Swabs

Preservative Efficiency of Neutralizing Buffer

Eight species of commonly encountered environmental bacteria and food pathogens bacteria were grown overnight on Tryptone Soya Agar plates and suspended in 10 mL Ringers solution to an approximate turbidity of Browns Opacity Standard No 1. 0. 1 mL of these dilutions were transferred into the 100 mL neutraliser. Six sponges were inoculated with 5ml neutraliser for each type of bacteria. The sponges were incubated at 22 °C for the duration of the test.

RESULTS

Organism	0 hrs	24 hrs	48 hrs	72 hrs	168 hrs
<i>S. enteritidis</i> (NCIMB 50073)	100%	100%	70%	70%	70%
<i>B. cereus</i> (ATCC 11778)	100%	70%	70%	50%	30%
<i>S. typhimurium</i> (ATCC 14028)	100%	100%	100%	70%	50%
<i>S. aureus</i> (NCTC 6571)	100%	100%	50%	25%	0%
<i>E. coli</i> (NCIMB 11943)	100%	100%	100%	70%	70%
<i>L. innocua</i> (NCTC 11288)	100%	100%	100%	100%	100%
<i>L. monocytogenes</i> (NCTC 11994)	100%	100%	100%	70%	70%

Organism	0 hrs	24 hrs	48 hrs	72 hrs	168 hrs
<i>L. monocytogenes</i> (ATCC 7645)	100%	100%	70%	70%	70%

DISCUSSION

All of the organisms in this test maintained constant numbers up to 24 hours when stored at 22 °C. Beyond 24 hours a gradual decrease in the numbers of organisms recovered occurred with all species tested, however *S. aureus* was the only organism to suffer significant reductions in numbers.

The results show that the swab neutralising buffer is able to preserve the viability of the bacteria used in the test whilst also preventing over growth.

9.2 Path-Chek Hygiene Coliform Detection Broth

SENSITIVITY

Organism	Dilution	CFU / Swab	24 Hours
<i>E. coli</i> MBCC 69	10 ⁴	TNC	Positive
	10 ³	230	Positive
	10 ²	30	Positive
	10 ¹	2	Positive
	10 ⁰	0	Negative
<i>E. coli</i> MBCC 70	10 ⁴	TNC	Positive
	10 ³	290	Positive
	10 ²	50	Positive
	10 ¹	3	Positive
	10 ⁰	0	Negative

Organism	Dilution	CFU / Swab	24 Hours
<i>E. coli</i> MBCC 71	10^4	TNC	Positive
	10^3	356	Positive
	10^2	35	Positive
	10^1	5	Positive
	10^0	0	Negative
<i>E. coli</i> MBCC 72	10^4	TNC	Positive
	10^3	430	Positive
	10^2	20	Positive
	10^1	1	Positive
	10^0	0	Negative
<i>E. coli</i> MBCC 73	10^4	TNC	Positive
	10^3	190	Positive
	10^2	10	Positive
	10^1	0	Positive
	10^0	0	Positive
<i>K. oxytoca</i> MBCC 46	10^4	500	Positive
	10^3	60	Positive
	10^2	6	Positive
	10^1	<1	Positive
<i>K. pneumoniae</i> MBCC 47	10^4	TNC	Positive
	10^3	150	Positive
	10^2	15	Positive
	10^1	1	Positive
<i>K. pneumoniae</i> MBCC 161	10^4	TNC	Positive
	10^3	100	Positive
	10^2	12	Positive (weak)
	10^1	1	Purple

CFU = Colony Forming Unit

TNC = Too Numerous to Count

Pre-moistened Path-Chek swabs were inoculated with a range of coliform species at dilutions down to zero CFU. The swabs were transferred into Path-Chek Hygiene Coliform Detection Broth and the broths incubated at 35 – 37 °C for 24 hours and then observed for any colour changes.

RESULTS

Based on these studies, the Path-Chek Hygiene Coliform Detection Broth is capable of detecting levels as low as 1 coliform organism per swab.

SPECIFICITY

A total of 31 isolates of *E. coli* were inoculated onto pre-moistened swabs at a level of $10 < \text{CFU}$. Inoculated swabs were then transferred into Path-Chek Hygiene Coliform Detection Broth and the broths incubated at $35 - 37^\circ\text{C}$ for 24 hours and then observed for any colour changes.

Target Bacteria	Number	Positive	Negative	Comment
<i>E. coli</i>	31	30	1	-
<i>Klebsiella sp.</i>	9	9	0	-
<i>Enterobacter sp.</i>	9	7	2	2x non lactose fermenting strains
<i>Citrobacter sp.</i>	13	8	5	5x non lactose fermenting strains
Total Coliform	62	54	1	-

The Path-Chek Hygiene Coliform Detection Broth detected 30/31 (97%) of the *E. coli* examined after 24 hours incubation at initial levels of $< 10 \text{ CFU}$ per swab. The one isolate that was not detected was of clinical origins and may have developed unique resistance patterns. In addition, a range of other “Coliform” organisms were examined, some of which were non lactose fermenting and failed to be detected by the indicator system employed in the Path-Chek Hygiene Coliform Detection Broth.

In a second study, 49 non-*Coliform* species comprising both Gram negative and Gram positive organisms were inoculated onto pre-moistened swabs at a level of 10^2 CFU . Inoculated swabs were then transferred into Path-Chek Hygiene Coliform Detection Broth and the broths incubated at $35 - 37^\circ\text{C}$ for 24 hours and then observed for any colour changes.

Non-Target Bacteria	Number	Positive	Negative
<i>Salmonella spp.</i>	5	0	5
<i>Listeria spp.</i>	4	0	4
<i>Enterococcus spp.</i>	4	0	4
<i>Staphylococcus spp.</i>	5	0	5
<i>Streptococcus spp.</i>	3	0	3
<i>Bacillus spp.</i>	6	0	6
<i>Pseudomonas spp.</i>	3	0	3
<i>Burkholderia spp.</i>	8	0	8
<i>Morganella spp.</i>	3	0	3
<i>Serratia spp.</i>	2	0	2
<i>Providencia spp.</i>	2	0	2
<i>Acinetobacter spp.</i>	1	0	1
<i>Aeromonas spp.</i>	1	0	1
<i>Proteus spp.</i>	1	0	1
<i>Micrococcus spp.</i>	1	0	1
Total Non-Coliform	49	0	49

9.3 Competitor Analysis

The Path-Chek Coliform and the Medical Wire® Coliform Environmental swab system were compared to determine both their efficiency in the recovery of Coliform organisms and their specificity.

Individual swabs of each system were challenged with <10 CFU of a total 28 different species of Coliform, or 100 CFU of a range of non-Coliform species. Both tests were read after 24 hours incubation.

Sensitivity Comparison

Organism	Total	Path-Chek Hygiene Coliform	Medical Wire® Coliform
<i>E. Coli</i>	14	14	14
<i>K. pneumoniae</i>	6	6	6
<i>K. oxytoca</i>	1	1	1
<i>C. youngii</i>	1	1	1
<i>C. freundii</i>	2	2	2
<i>E. cloacae</i>	4	4	4
Total	28	28	28

Both the Path-Chek Hygiene Coliform and the Medical Wire® Coliform were able to detect 28/28 (100%) of the isolates tested after 24 hours incubation.

Specificity Comparison

Organism	Path-Chek Hygiene Coliform	Medical Wire® Coliform
<i>L. monocytogenes</i>	Negative	Positive
<i>L. innocua</i>	Negative	Negative
<i>L. grayi</i>	Negative	Weak Positive
<i>Micrococcus sp.</i>	Negative	Negative
<i>B. cereus</i>	Negative	Negative
<i>B. licheniformis</i>	Negative	Negative
<i>E. avium</i>	Negative	Negative
<i>E. faecalis</i>	Negative	Positive
<i>E. faecium</i>	Negative	Positive
<i>E. gallinarum</i>	Negative	Negative
<i>Bacillus sp.</i>	Negative	Negative
<i>Staphylococcus sp.</i>	Negative	Positive
<i>M. morgani</i>	Negative	Positive
<i>S. maltophilia</i>	Negative	Negative
<i>A. baumannii</i>	Negative	Negative
<i>S. seftenberg</i>	Negative	Positive
<i>P. mirabilis</i>	Negative	Positive
<i>P. stuartii</i>	Negative	Positive

Organism	Path-Chek Hygiene Coliform	Medical Wire® Coliform
<i>P. aeruginosa</i>	Negative	Negative
<i>S. typhimurium</i>	Negative	Positive
<i>S. marcesens</i>	Negative	Positive
Total	21	10

A total of 21 species of both Gram negative and Gram positive organisms were examined. The Path-Chek Coliform inhibited the growth of all species tested i.e. 100% specificity.

The Medical Wire® Coliform failed to inhibit the growth of 11 species which were also able to produce positive reactions similar to Coliforms. “False positive” results were caused by a range of both Gram negative and Gram positive organisms. On the basis of this trial, the specificity of the Pat-Chek Coliform was 94% and the Medical Wire® Coliform was 48%.

9.4 Path-Chek Hygiene Salmonella Detection Broth

Sensitivity

Pre-moistened Path-Chek swabs were inoculated with a range of Salmonella species at dilutions down to zero CFU. The swabs were transferred into Path-Chek Hygiene Salmonella Detection Broth and the broths incubated at 35 – 37 °C for 24 hours and then observed for any color changes.

RESULTS

Organism	Dilution	CFU / Swab	Results
<i>Salmonella typhimurium</i> MBCC 215	10 ⁴	TNC	Black
	10 ³	TNC	Black
	10 ²	120	Black
	10 ¹	10	Black
	10 ⁰	0	Black
<i>Salmonella bispeberg</i> MBCC 274	10 ⁴	TNC	Black
	10 ³	250	Black
	10 ²	20	Black
	10 ¹	2	Black
	10 ⁰	0	Purple

Organism	Dilution	CFU / Swab	Results
<i>Salmonella java</i> MBCC 275	10 ⁴	TNC	Black
	10 ³	180	Black
	10 ²	20	Black
	10 ¹	1	Black
	10 ⁰	0	Purple
<i>Salmonella virginia</i> MBCC 277	10 ⁴	TNC	Black
	10 ³	420	Black
	10 ²	30	Black
	10 ¹	2	Black
	10 ⁰	0	Purple
<i>Salmonella saint-paul</i> MBCC 278	10 ⁴	500	Black
	10 ³	40	Black
	10 ²	10	Black
	10 ¹	1	Black
	10 ⁰	0	Purple
<i>Salmonella derby</i> MBCC 281	10 ⁴	TNC	Black
	10 ³	270	Black
	10 ²	30	Black
	10 ¹	1	Black
	10 ⁰	0	Purple
<i>Salmonella senftenberg</i> MBCC 282	10 ⁴	TNC	Black
	10 ³	290	Black
	10 ²	50	Black
	10 ¹	3	Black
	10 ⁰	0	Purple
<i>Salmonella derby</i> MBCC 281	10 ⁴	TNC	Black
	10 ³	270	Black
	10 ²	30	Black
	10 ¹	1	Black
	10 ⁰	0	Purple
<i>Salmonella senftenberg</i> MBCC 282	10 ⁴	TNC	Black
	10 ³	290	Black
	10 ²	50	Black
	10 ¹	3	Black
	10 ⁰	0	Purple

Organism	Dilution	CFU / Swab	Results
<i>Salmonella senftenberg</i> MBCC 282	10 ⁴	TNC	Black
	10 ³	250	Black
	10 ²	30	Black
	10 ¹	2	Black
	10 ⁰	0	Purple
<i>Salmonella rostock</i> MBCC 283	10 ⁴	TNC	Black
	10 ³	500	Black
	10 ²	70	Black
	10 ¹	10	Black
	10 ⁰	1	Black

TNC = Too Numerous to Count

This study demonstrated that the Path-Chek Hygiene Salmonella Detection Broth is capable of detecting levels as low as 1 *Salmonella* spp. organism per swab.

Specificity

A total of 97 different serotypes of *Salmonella* spp. were inoculated onto pre-moistened swabs at a level of 10 < CFU. Inoculated swabs were then transferred into Path-Chek Hygiene Salmonella Detection Broth and the broths incubated at 35 – 37 °C for 24 hours and then observed for any colour changes.

Target Bacteria	Number	Positive	Negative	Comment
<i>S. enteritidis</i>	5	5	0	-
<i>S. typhimurium</i>	8	8	0	-
<i>S. dublin</i>	7	7	0	-
<i>Salmonella</i> spp.	77	76	1	1 <i>Salmonella</i> spp. H ₂ S negative
Total Salmonella	97	96	1	-

The Path-Chek Hygiene Salmonella Detection Broth detected 99% of the *Salmonella* spp. examined after 24 hours of incubation at initial levels of < 10 CFU per swab. The one isolate that was not detected was found to be H₂S negative.

In a second study, 88 *non-Salmonella* spp., comprising both Gram negative and Gram-positive organisms were inoculated onto pre-moistened swabs at a level of 102 CFU.

Inoculated swabs were then transferred into Path-Chek Hygiene Salmonella Detection Broth and the broths incubated at 35 – 37 °C for 24 hours and then observed for any colour changes.

Non-Target Bacteria	Number	Positive	Negative
<i>E. coli</i>	15	0	15
<i>Klebsiella spp.</i>	5	0	5
<i>Enterobacter spp.</i>	6	0	6
<i>Citrobacter spp.</i>	14	11	3
<i>Listeria spp.</i>	9	0	9
<i>Enterococcus spp.</i>	4	0	4
<i>Staphylococcus spp.</i>	5	0	5
<i>Streptococcus spp.</i>	3	0	3
<i>Bacillus spp.</i>	6	0	6
<i>Pseudomonas spp.</i>	3	0	3
<i>Burkholderia spp.</i>	6	0	6
<i>Morganii spp.</i>	3	0	3
<i>Maltophilia spp.</i>	2	0	2
<i>Serratia spp.</i>	2	0	2
<i>Providencia spp.</i>	2	0	2
<i>Acinetobacter spp.</i>	1	0	1
<i>Aeromonas spp.</i>	1	0	1
<i>Proteus spp.</i>	1	0	1
<i>Micrococcus spp.</i>	1	0	1
Total Non-Salmonella	88	11	77

The Path-Chek Hygiene Salmonella Detection Broth demonstrated a high degree of selectivity for the Gram-positive isolates tested and most of the Gram negative isolates. Some isolates of *Citrobacter spp.* were not inhibited by the selective agents incorporated into the Path-Chek Salmonella Detection Broth, resulting in false positive results.

The occurrence of false positive results due to detection of *Citrobacter spp.* should still be considered as a significant result. The detection of significant levels of *Citrobacter spp.* from surfaces is an important indication of fecal contamination and/ or poor cleaning and sanitising and should therefore be investigated.

Recovery from Surfaces

A study was performed using the method of G. Moore and C. Griffith (A Comparison of Surface Sampling Methods for Detecting Coliforms on Food Contact Surfaces. Food Microbiology. 2002: 19, - 73), to determine the efficiency of the swabbing process combined with the Path-Chek Salmonella Detection Broth when used to sample both wet and dry surfaces.

Organism	Wet Surface		Dry Surface	
	Results	CFU	Results	CFU
<i>S. tranora MBCC 171</i>	Positive	306	Positive	520
	Positive	40	Positive	52
	Positive	10	Positive	4
	Negative	2	Negative	0

The combination of the pre-moistened swabs and the Path-Chek Salmonella Detection Broth was successful in detecting < 10 CFU recovered from both wet and dry sample surface areas of 100 cm².

10 COMPETITOR ANALYSIS

No competitor products are available for comparison purposes.

10.1 Path-Chek Hygiene Listeria Detection Broth

Sensitivity

Pre-moistened Path-Chek swabs were inoculated with a range of Listeria species at dilutions down to zero CFU. The swabs were transferred into Path-Chek Hygiene Listeria Detection Broth and the broths incubated at 35 – 37 °C for up to 48 hours and then observed for any colour changes.

Note: If Path-Chek Hygiene Listeria is incubated at 35-37 °C, there will, in certain circumstances, be an increased risk of false positives, compared to incubation at 28-30 °C.

RESULTS

Organism	Dilution	CFU / Swab	24 Hours	48 Hours
<i>L. monocytogenes</i> MBCC 97	10 ⁴	TNC	Positive	Positive
	10 ³	800	Positive	Positive
	10 ²	110	Positive	Positive
	10 ¹	10	Negative	Positive
	10 ⁰	0	Negative	Negative
<i>L. monocytogenes</i> MBCC 98	10 ⁴	TNC	Positive	Positive
	10 ³	250	Positive	Positive
	10 ²	20	Positive	Positive
	10 ¹	2	Negative	Positive
	10 ⁰	0	Negative	Positive
<i>L. monocytogenes</i> MBCC 99	10 ⁴	TNC	Positive	Positive
	10 ³	560	Positive	Positive
	10 ²	60	Positive	Positive
	10 ¹	2	Negative	Positive
	10 ⁰	0	Negative	Positive
<i>L. innocua</i> MBCC 93	10 ⁴	TNC	Positive	Positive
	10 ³	700	Positive	Positive

Organism	Dilution	CFU / Swab	24 Hours	48 Hours
	10 ²	45	Positive	Positive
	10 ¹	5	Positive	Positive
	10 ⁰	0	Positive	Negative
<i>L. innocua</i> MBCC 94	10 ⁴	TNC	Positive	Positive
	10 ³	440	Positive	Positive
	10 ²	20	Positive	Positive
	10 ¹	3	Positive	Positive
	10 ⁰	0	Negative	Negative
<i>L. innocua</i> MBCC 95	10 ⁴	TNC	Positive	Positive
	10 ³	450	Positive	Positive
	10 ²	30	Positive	Positive
	10 ¹	2	Positive	Positive
	10 ⁰	0	Negative	Negative
<i>L. seeligeri</i> MBCC 110	10 ⁴	TNC	Negative	Positive
	10 ³	400	Negative	Positive
	10 ²	60	Negative	Positive
	10 ¹	0	Negative	Negative
	10 ⁰	0	Negative	Negative
<i>L. ivanovii</i> MBCC 111	10 ⁴	TNC	Negative	Positive
	10 ³	560	Negative	Positive
	10 ²	40	Negative	Positive
	10 ¹	3	Negative	Positive
	10 ⁰	0	Negative	Negative
<i>L. welshmeri</i> MBCC 114	10 ⁴	TNC	Positive	Positive
	10 ³	880	Positive	Positive
	10 ²	100	Positive	Positive
	10 ¹	10	Positive	Positive
	10 ⁰	1	Positive	Positive

This study demonstrated that the Path-Chek Hygiene Listeria Detection Broth is capable of detecting levels as low as 1 Listeria spp. organism per swab.

Specificity

A total of 97 different species of Listeria were inoculated onto pre-moistened swabs at a level of 10 < CFU. Inoculated swabs were then transferred into Path-Chek Hygiene Listeria Detection Broth and the broths incubated at 35 – 37 °C for up to 48 hours and then observed for any colour changes.

Target Bacteria	Number	Positive	Negative
<i>L. monocytogenes</i>	42	42	0
<i>L. innocua</i>	24	24	0
<i>L. seeligeri</i>	12	12	0
<i>L. ivanovii</i>	8	8	0
<i>L. welshimeri</i>	11	11	0
Total Listeria	97	97	0

This investigation demonstrated that 100% of the *Listeria spp.* examined could be detected after 48 hours of incubation at initial levels < 10 CFU per swab.

In a second study, 76 non *Listeria spp.* comprising both Gram negative and Gram positive organisms were inoculated onto pre-moistened swabs at a level of > 5 x 10² CFU. Inoculated swabs were then transferred into Path-Chek Hygiene Listeria Detection Broth and the broths incubated at 35 - 37°C for 48 hours and then observed for any colour changes.

Non-Target Bacteria	Number	Positive	Negative	Comment
<i>E. coli</i>	4	0	4	-
<i>Klebsiella spp.</i>	10	4	6	-
<i>Enterobacter spp.</i>	5	0	5	-
<i>Citrobacter spp.</i>	3	0	3	-
<i>Salmonella spp.</i>	9	0	9	-
<i>Enterococcus spp.</i>	6	4	2	6/6 Negative when challenged with <5 x 10 ²
<i>Staphylococcus spp.</i>	5	0	5	-
<i>Streptococcus spp.</i>	3	0	3	-
<i>Bacillus spp.</i>	5	1	4	6/6 Negative when challenged with <5 x 10 ²
<i>Pseudomonas spp.</i>	4	0	4	-
<i>Burkholderia spp.</i>	7	0	7	-
<i>Lactobacillus spp.</i>	3	0	3	-
<i>Carnobacterium spp.</i>	2	0	2	-
<i>Cornebacterium spp.</i>	1	0	1	-
<i>Kurthia spp.</i>	1	0	1	-
<i>Acinetobacter spp.</i>	1	0	1	-
<i>Achromobacter spp.</i>	1	0	1	-
<i>Proteus spp.</i>	2	0	2	-
<i>Rhodococcus equi</i>	1	0	1	-
<i>Micrococcus spp.</i>	1	0	1	-
<i>Morganella spp.</i>	1	0	1	-
<i>Stenotrophomonas spp.</i>	1	0	1	-
Total Non-Listerias	76	9	76	-

Path-Chek Hygiene Listeria Detection Broth demonstrated a high degree of selectivity for the all of the Gram positive isolates tested with the exception of 2/6 *Enterococcus spp.* and 1/4 *Bacillus spp.* which produced positive reactions at levels $> 5 \times 10^2$ but failed to grow at lower levels ($< 1 \times 10^2$) after 48 hours of incubation.

With the exception of *Klebsiella spp.*, all other Gram-negative isolates either failed to grow or produced negative results after 48 hours of incubation. The Path-Chek Listeria Detection Broth, resulting in false positive results.

The occurrence of false positive results due to detection of *Enterococcus spp.* should still be considered as a significant result. The detection of significant levels of *Enterococcus spp.* from surfaces is an important indication of fecal contamination and/ or poor cleaning and sanitising and should therefore be investigated.

Recovery from Surfaces

A study was performed using the method of G. Moore and C. Griffith (A Comparison of surface Sampling Methods for Detecting Coliforms on Food Contact Surfaces. Food Microbiology. 2002: 19, - 73), to determine the efficiency of the swabbing process combined with the Path-Chek Salmonella Detection Broth when used to sample both wet and dry surfaces.

Organism	Wet Surface		Dry Surface	
	Results	CFU	Results	CFU
<i>L. monocytogenes</i> <i>MBCC 305</i>	Positive	300	Positive	100
	Positive	30	Positive	10
	Positive	6	Negative	1
	Positive (weak)	2	Negative	0

The combination of the pre-moistened swabs and the Path-Chek Listeria Detection Broth was successful in detecting < 10 CFU recovered from both wet surfaces and 10 CFU from dry sample surface areas of 100 cm².

Competitor Analysis

The Path-Chek Listeria and the Medical Wire® Listeria Environmental swab system were compared to determine both their efficiency in the recovery of *Listeria spp.* and their specificity.

Individual swabs of each system were challenged with < 10 CFU of a total 62 different species of *Listeria* species, or 100 CFU of a range of non-*Listeria* species. Both tests were read after 24 hours and 48 hours incubation.

Sensitivity Comparison

Organism	Total	Path-Chek Hygiene Listeria		Medical Wire® Listeria	
		24 Hours	48 Hours	24 Hours	48 Hours
<i>L. monocytogenes</i>	30	29	30	20	30
<i>L. innocua</i>	18	18	18	16	18
<i>L. ivanovii</i>	4	3	4	0	4
<i>L. seeligeri</i>	4	4	4	0	4
<i>L. welshimeri</i>	6	6	6	0	6
Total	62	60	62	36	62

The Path-Chek Hygiene Listeria detected 60 (97%) of the Listeria isolates tested after 24 hours incubation and 62/62 (100%) of the isolates after 48 hours incubation. The Medical Wire® Listeria was able to detect 36/62 (58%) of the isolates tested after 24 hours of incubation and 62/62 (100%) after 48 hours incubation. The Medical Wire® was only able to detect *L. monocytogenes* and *L. innocua* species after 24 hours of incubation.

Specificity Comparison

Organism	Path-Chek Listeria		Medical Wire® Listeria	
	24 Hours	48 Hours	24 Hours	48 Hours
<i>C. diversus</i>	Negative	Negative	Negative	Negative
<i>S. enteritidis</i>	Negative	Negative	Negative	Negative
<i>S. enteritidis</i>	Negative	Negative	Negative	Negative
<i>S. hadar</i>	Negative	Negative	Negative	Negative
<i>S. indiana</i>	Negative	Negative	Negative	Negative
<i>S. infantis</i>	Negative	Negative	Negative	Negative
<i>S. typhimurium</i>	Negative	Negative	Negative	Negative
<i>P. mirabilis</i>	Negative	Negative	Negative	Negative
<i>P. stuartii</i>	Negative	Negative	Negative	Negative
<i>P. aeruginosa</i>	Negative	Negative	Negative	Negative
<i>E. aerogenes</i>	Negative	Negative	Negative	Negative
<i>E. coli</i>	Negative	Negative	Negative	Negative
<i>E. coli</i>	Negative	Negative	Negative	Negative
<i>K. oxytoca</i>	Negative	Negative	Negative	Negative
<i>K. oxytoca</i>	Negative	Negative	Negative	Negative
<i>K. pneumoniae</i>	Negative	Negative	Negative	Negative
<i>K. pneumoniae</i>	Negative	Negative	Negative	Negative
<i>K. pneumoniae</i>	Negative	Negative	Negative	Negative
<i>K. pneumoniae</i>	Positive	Positive	Positive	Positive
<i>K. pneumoniae</i>	Negative	Positive	Negative	Positive
<i>K. pneumoniae</i>	Positive	Positive	Negative	Negative
<i>K. pneumoniae</i>	Negative	Negative	Negative	Negative
<i>K. pneumoniae</i>	Positive	Positive	Negative	Negative
<i>E. cloacae</i>	Negative	Negative	Negative	Negative
<i>E. cloacae</i>	Negative	Negative	Negative	Negative
<i>E. cloacae</i>	Negative	Negative	Negative	Negative
<i>E. cloacae</i>	Negative	Negative	Negative	Negative
<i>Carnobacterium divergens</i>	Negative	Negative	Positive	Positive
<i>Carnobacterium piscicola</i>	Negative	Negative	Negative	Negative
<i>Lactobacillus casei</i>	Negative	Negative	Negative	Negative
<i>Lactobacillus lactis</i>	Negative	Negative	Negative	Negative
<i>Lactobacillus</i>	Negative	Negative	Negative	Negative

Organism	Path-Chek Listeria		Medical Wire® Listeria	
	24 Hours	48 Hours	24 Hours	48 Hours
<i>plantarum</i>				
<i>Bacillus mycoides</i>	Negative	Negative	Negative	Negative
<i>Kurthia zopfii</i>	Negative	Negative	Negative	Negative
<i>Micrococcus spp.</i>	Negative	Negative	Negative	Negative
<i>Rhodococcus equi</i>	Negative	Negative	Negative	Negative
<i>Enterococcus durans</i>	Negative	Negative	Negative	Positive
<i>Pseudomonas fluorescens</i>	Negative	Negative	Negative	Negative
<i>Corynebacterium renale</i>	Negative	Negative	Negative	Negative
<i>Bacillus cereus</i>	Negative	Negative	Negative	Negative
<i>Bacillus licheniformis</i>	Negative	Negative	Negative	Negative
<i>Enterococcus avium</i>	Negative	Negative	Positive	Positive
<i>Enterococcus faecalis</i>	Negative	Negative	Positive	Positive
<i>Enterococcus faecium</i>	Negative	Negative	Positive	Positive
<i>Enterococcus gallinarum</i>	Negative	Negative	Negative	Positive
<i>Streptococcus spp.</i>	Negative	Negative	Negative	Negative
<i>Streptococcus spp.</i>	Negative	Negative	Negative	Negative
<i>Streptococcus spp.</i>	Negative	Negative	Negative	Negative
<i>Bacillus spp.</i>	Negative	Negative	Negative	Negative
<i>Bacillus spp.</i>	Negative	Negative	Negative	Negative
<i>Bacillus spp.</i>	Negative	Negative	Negative	Negative
<i>Streptococcus spp.</i>	Negative	Negative	Negative	Negative
<i>S. saprophyticus</i>	Negative	Negative	Negative	Negative
<i>S. aureus</i>	Negative	Negative	Negative	Negative
<i>S. aureus</i>	Negative	Negative	Negative	Negative
<i>S. hyicus ss hyicus</i>	Negative	Negative	Negative	Negative
<i>B. stabilis</i>	Negative	Negative	Negative	Negative
<i>B. cenocepacia</i>	Negative	Negative	Negative	Negative
<i>B. multivorans</i>	Negative	Negative	Negative	Negative
<i>B. cenocepacia</i>	Negative	Negative	Negative	Negative

Organism	Path-Chek Listeria		Medical Wire® Listeria	
	24 Hours	48 Hours	24 Hours	48 Hours
<i>A. xylosoxidans</i>	Negative	Negative	Negative	Negative
<i>B. cepacia</i>	Negative	Negative	Negative	Negative
<i>R. mannitolilytica</i>	Negative	Negative	Negative	Negative
<i>B. cenocepacia</i>	Negative	Negative	Negative	Negative
<i>P. aeruginosa</i>	Negative	Negative	Negative	Negative
<i>P. putida</i>	Negative	Negative	Negative	Negative
<i>B. vietnamiensis</i>	Negative	Negative	Negative	Negative
Total	58	58	54	54

A total of 62 species of both Gram negative and Gram positive organisms were examined. The Path-Chek Listeria failed to inhibit the growth of 4 species which were also able to produce positive reactions similar to *Listeria spp.* These “false positive” results were caused by *K. pneumoniae* (3) and *Enterococcus spp.* (1).

The Medical Wire® Listeria failed to inhibit the growth of 8 species which were also able to produce positive reactions similar to *Listeria spp.* “False positive” results were caused by *K. pneumoniae* (1), *Enterococcus spp.* (6) and *C. divergens* (1).

On the basis of this trial, the specificity of the Pat-Chek Listeria was 94% and the Medical Wire® Listeria was 87%.

11 POSITIVE RESULTS – WHAT DO THEY MEAN?

With any of the Path-Chek Hygiene Pathogen systems, a positive result for a specific pathogen can mean one of two things:

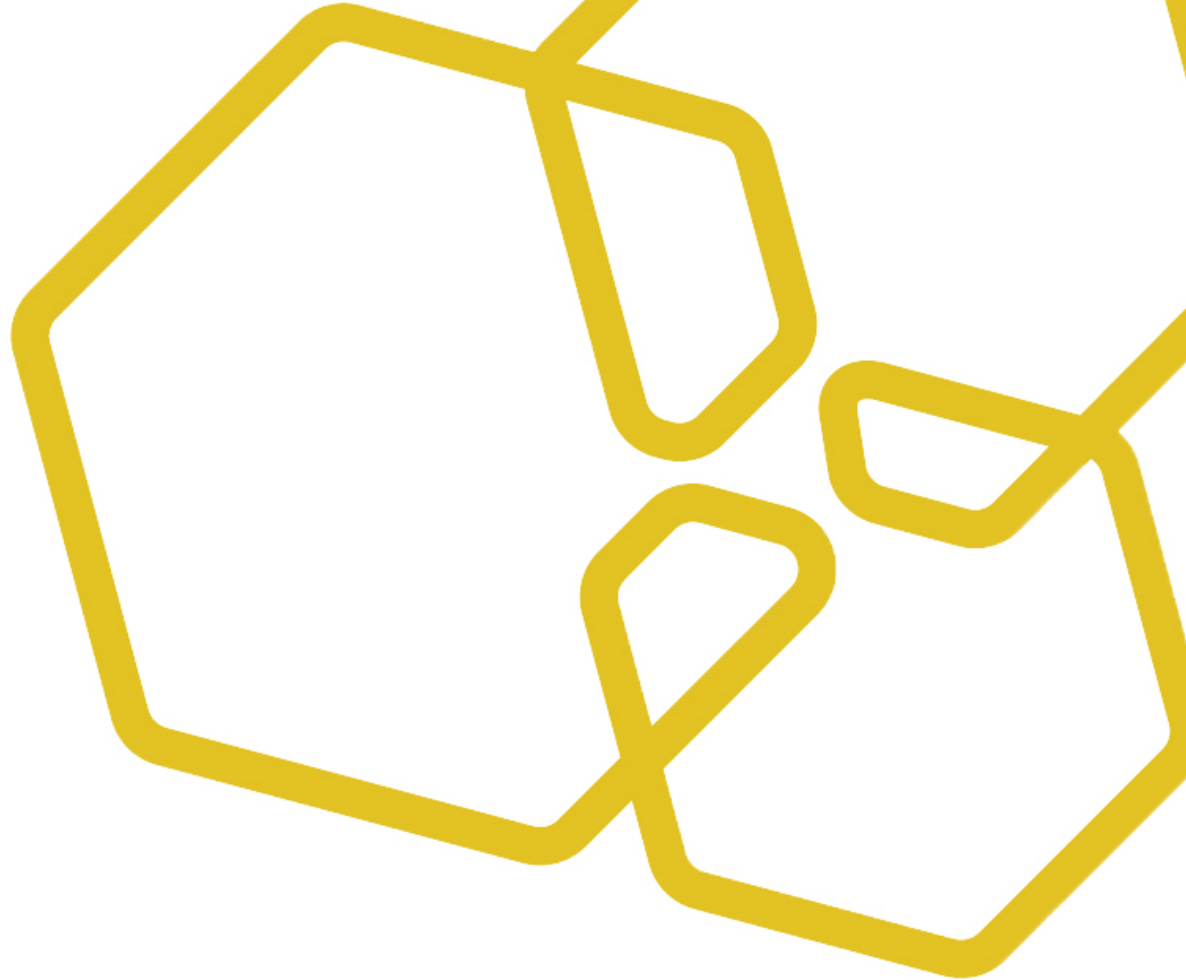
1. The presence of a specific pathogen such as Salmonella or Listeria at levels as low as **1 – 10 CFU** per sampled area which should be at least 100 cm², OR
2. The presence in high numbers, usually > 102 of potentially cross-reacting organisms

Table 8.: Most Probable Cause of “False Positive Tests”.

Path-Chek Detection System	Most Probable Cause of “False Positive Tests”
Coliform	<i>Nil</i>
Salmonella	<i>Citrobacter spp.</i>
Listeria	<i>K. pneumoniae, Enterococcus spp.</i>

In each of these cases, the potential causes of “False Positive” tests are organisms of fecal origin.

As such, ALL Positive Tests should be confirmed by subculture and identification of suspicious colonies. If a specific target pathogen cannot be isolated, the results should still be considered as significant as they indicate the presence of significant levels of organisms of fecal origin on the test surfaces. These organisms should not be present if adequate cleaning and sanitising procedures are in place and being performed correctly.



TECHNICAL SUPPORT SERVICE

For technical assistance and more information please contact Gold Standard Diagnostics Budapest's Customer Service or your local distributor.

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