3M Petrifilm™

Interpretation Guide

The 3M™ Petrifilm™ Select *E. coli* Count Plate is a sample-ready-culture medium system which contains selective agents, nutrients, a cold-water-soluble gelling agent, and an indicator of glucuronidase activity, 5-bromo-4-chloro-3-indolyl-D-glucuronide (BCIG), which facilitates colony enumeration.
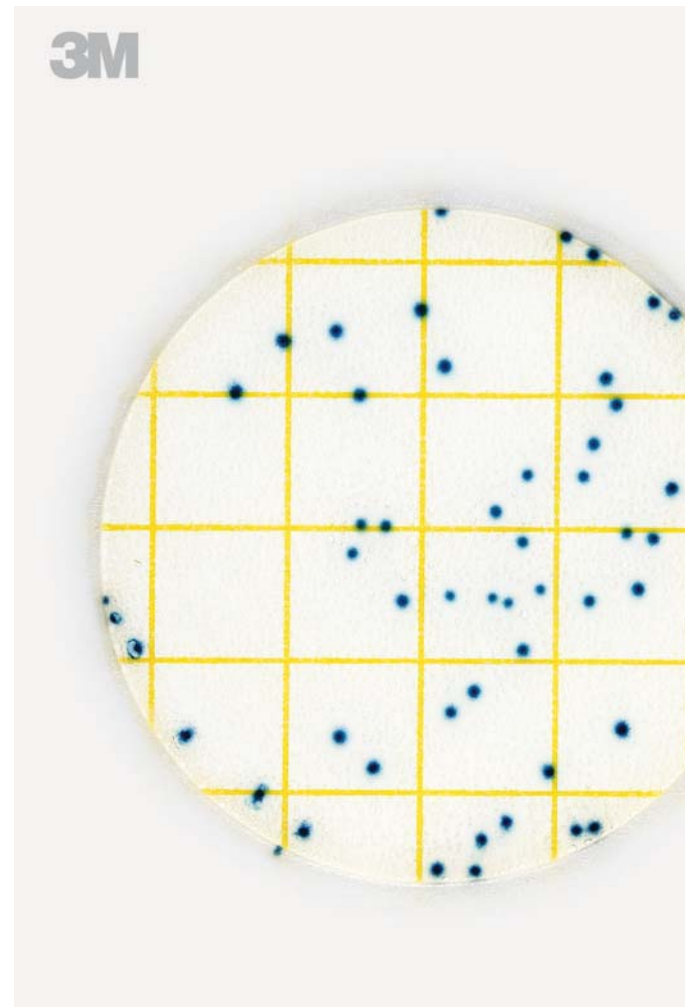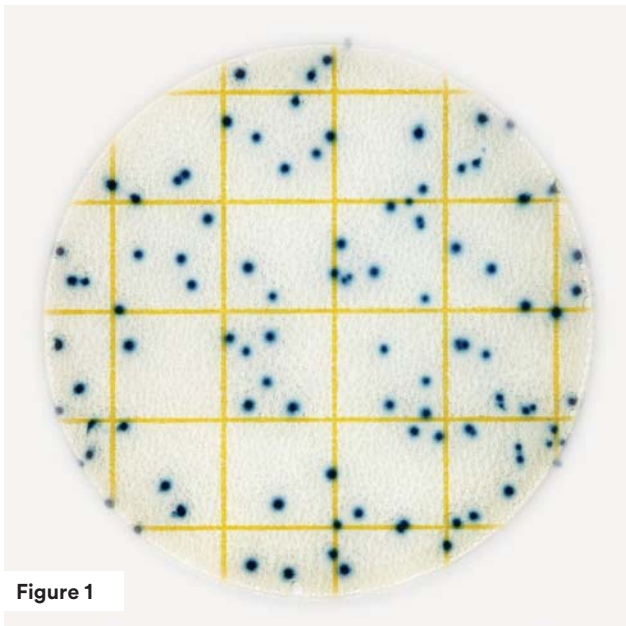

SEC
Select *E. coli* Count Plate

Figure 1

E. coli count = 97

About 97% of *E. coli* strains are thermotolerant and produce beta-glucuronidase, an enzyme that reacts with BCIG indicator in 3M™ Petrifilm™ Select *E. coli* Count Plates to produce dark green to blue-green colonies.

3M Petrifilm Select *E. coli* Count Plates will not detect *E. coli* O157 as most strains are atypical. They are glucuronidase negative, and will not produce a blue-green colony.
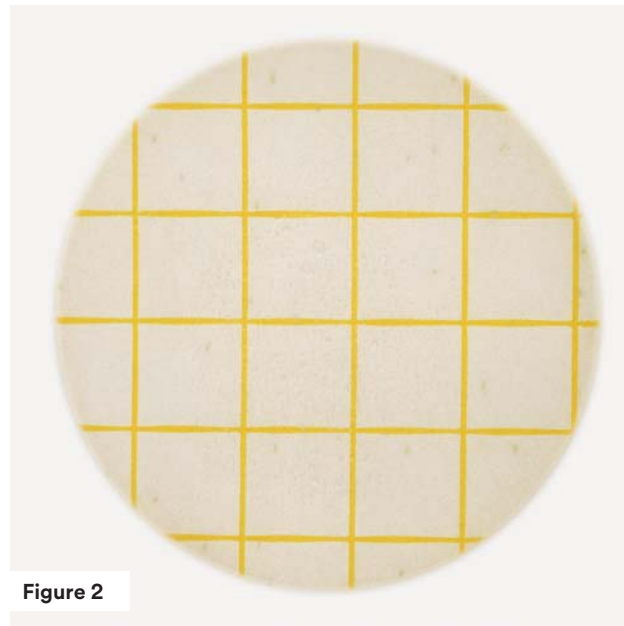

Figure 2

E. coli count = 0

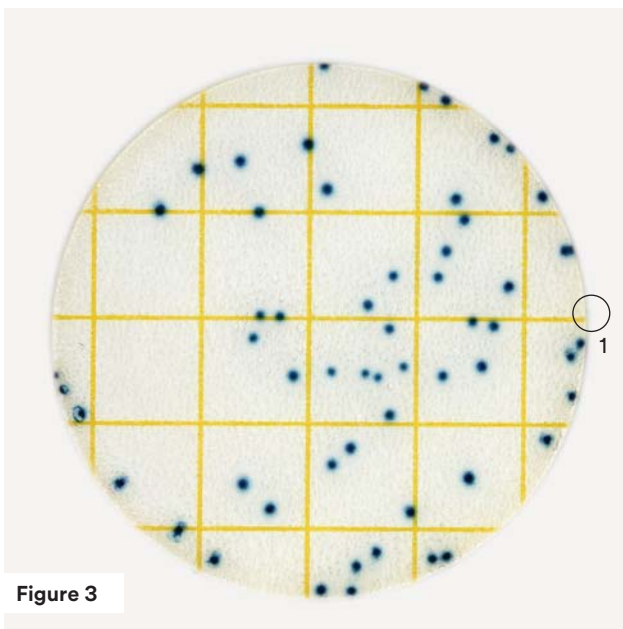Colonies other than *E. coli* are difficult to see because they are colorless to a light grey-beige.

Figure 3

E. coli count = 56

Do not count colonies on the foam dam because they are removed from the selective influence of the medium. See Circle 1.
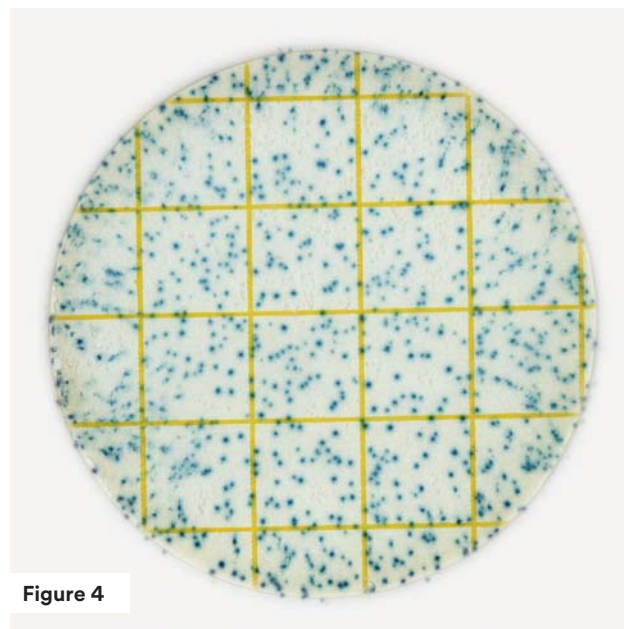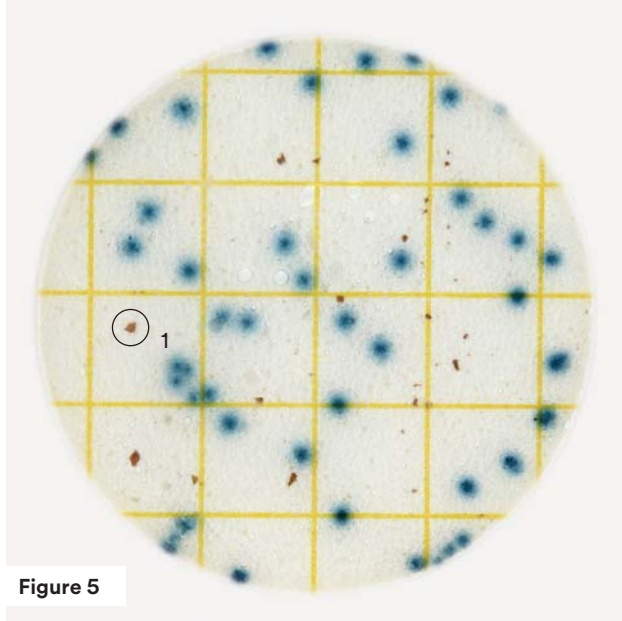

Figure 4

Estimated *E. coli* count = 740

Estimates can be made on plates containing greater than 150 colonies by counting the number of colonies in two or more representative squares and determining the average number per square. Multiply the average number by 20 to determine the estimated count per plate. The circular growth area is approximately 20 cm².

For a more accurate count, further dilution of the sample may be necessary.

Interference from food products

3M Petrifilm Select *E. coli* Count Plates have been evaluated using samples from many, but not all foods. Foods tested include certain fresh and frozen meats, vegetables and seafood; frozen prepared meals; and fresh, fermented and dry dairy foods. In a limited number of cases, such as liver, the food may interfere with enumeration.
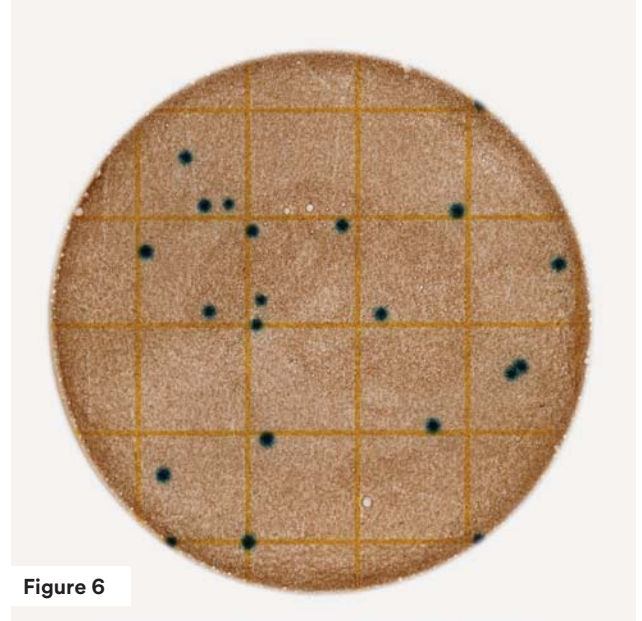
To reduce such food interference, dilute the sample further.

Figure 5

Figure 6

E. coli count = 42

E. coli colonies can easily be distinguished from food particles which are often irregularly shaped and variable in size and color. Circle 1 shows nut particles.
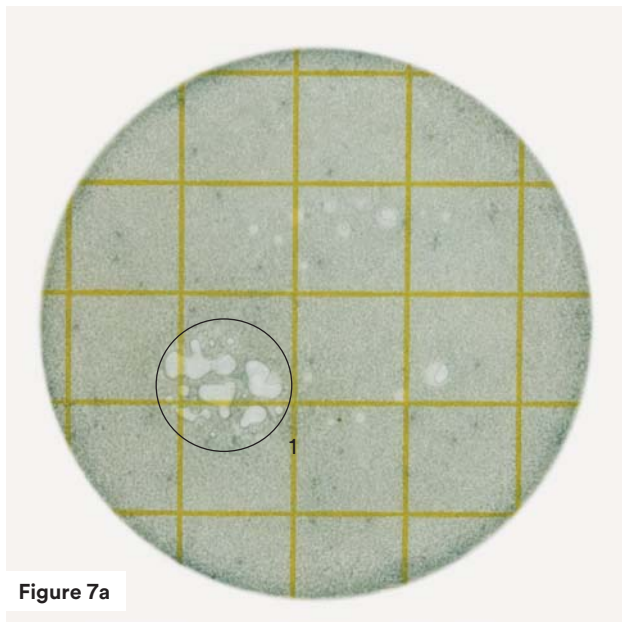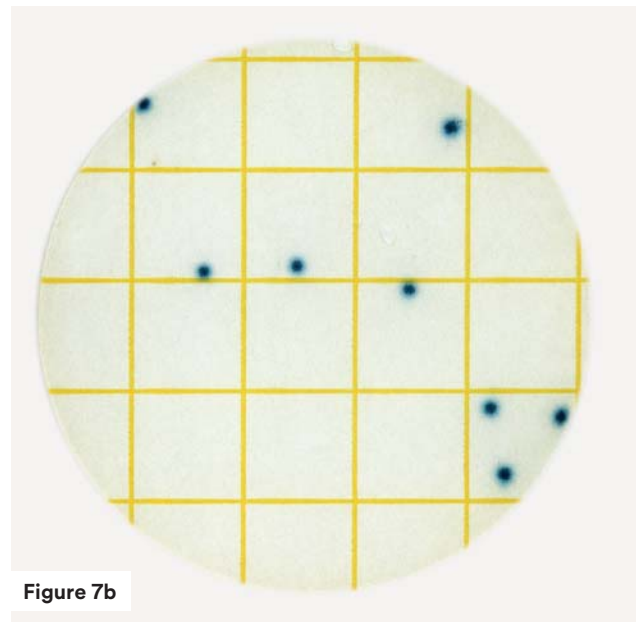
E. coli count = 21

Some dark foods may produce a colored background that makes *E. coli* colonies less distinguishable. Further dilutions will lighten the background color making the *E. coli* colonies easier to count.

Figure 6 shows cocoa powder diluted 1:50.

Figure 7a

Figure 7b

Raw liver contains ß-glucuronidase which produces a blue-green background color on the growth area on the plates, making the *E. coli* colonies less distinguishable. Further dilution will lighten the background color making the *E. coli* colonies easier to count and will help to distinguish food interference from TNTC plates that have confluent colonies (see Figure 13). Artifact bubbles may result from improper inoculation of the plate or from trapped air from the sample. See Circle 1.

Variability in *E. coli* colony appearance

Glucuronidase-positive *E. coli* colonies may vary in size, color intensity and shape, depending on the strain itself, the food and the influence of external factors such as testing protocols. The blue-green *E. coli* colonies may have gas bubbles associated with them.
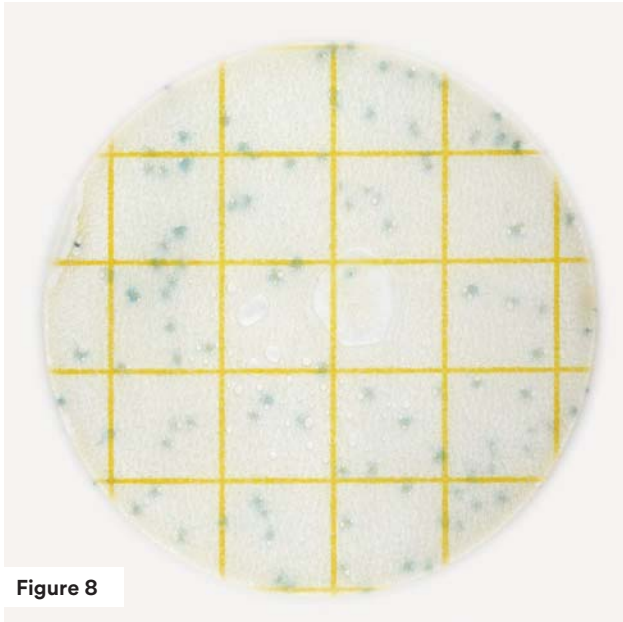

Figure 8

Figure 9

E. coli count = 92

Pale green colonies may result either from *E. coli* that are weak producers of glucuronidase or from an interaction with food such as those containing high acid or high sugar.

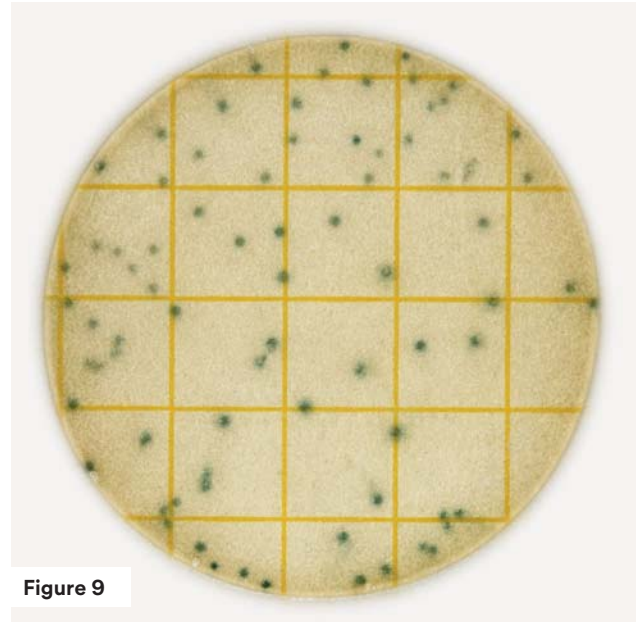Figure 8 shows a highly acidic fermented dairy product.

E. coli count = 75

Greenish-brown color variability may appear with some foods.
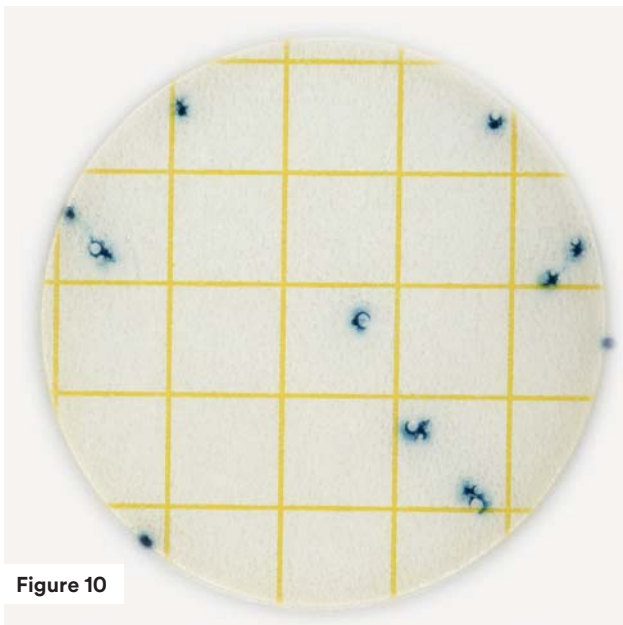
Figure 9 shows a kidney sample.

Figure 10

Figure 11

E. coli count = 10

E. coli colonies may have gas bubbles associated with them depending on the *E. coli* strain and the food. Count all colonies with or without gas.
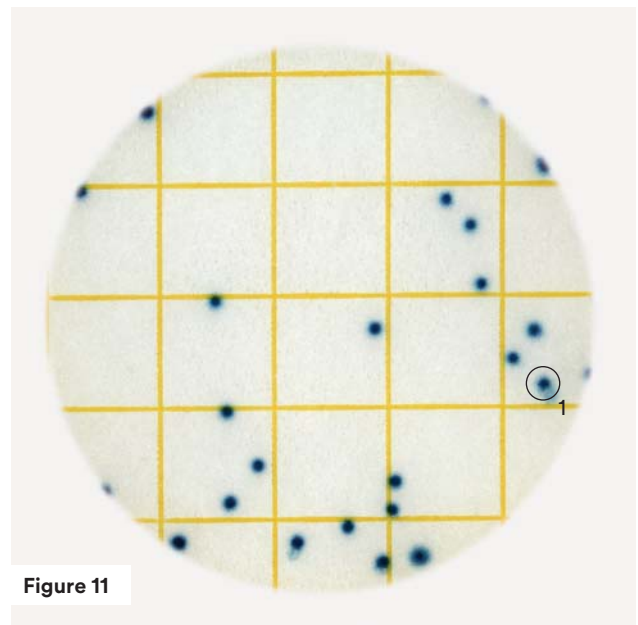
E. coli count = 21

Smeared colonies may appear. See Circle 1.

To minimize the production of smeared colonies spread immediately after inoculation, pressing gently on the center of the spreader.
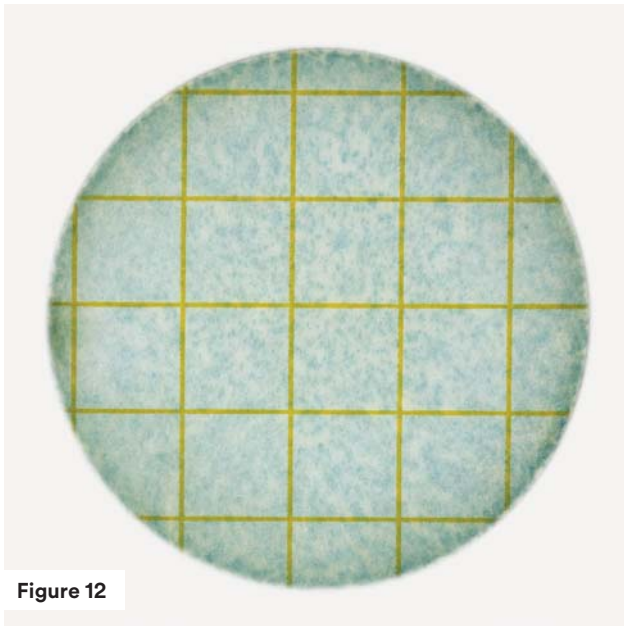
Figure 12

E. coli count = Too numerous to count (TNTC)

When present in large numbers, *E. coli* may appear as small, indistinct colonies.

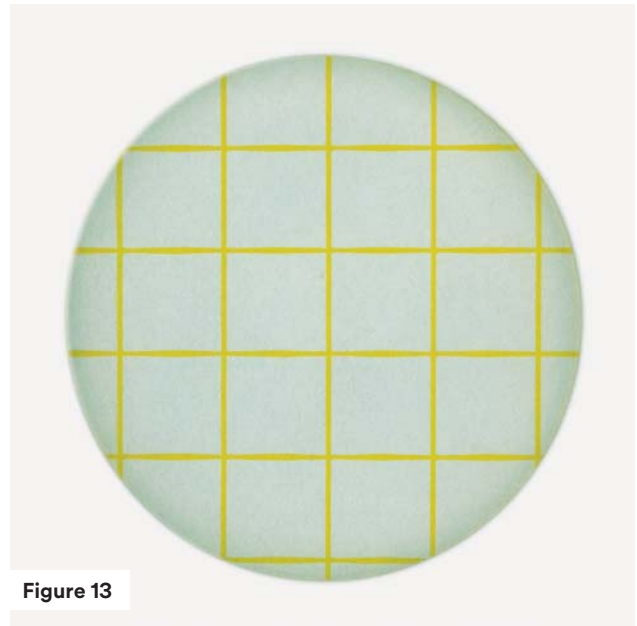*For a more accurate count, further dilution of the sample may be necessary.*
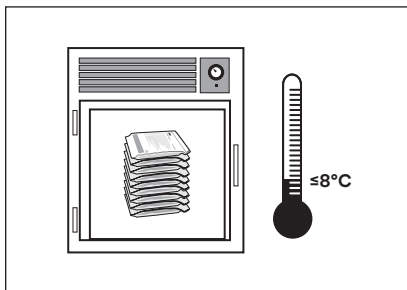

Figure 13

E. coli count = TNTC

High concentrations of *E. coli* will cause the entire growth area to turn blue-green.
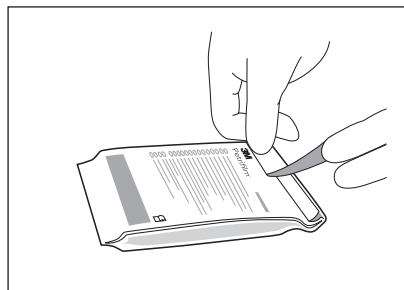
For a more accurate count, further dilution of the sample may be necessary.

Reminders for Use

Storage

Inoculation

1 Store unopened packages at ≤8°C (≤46°F). Use before expiration date on package. Just prior to use, allow unopened pouches to come to room temperature before opening.
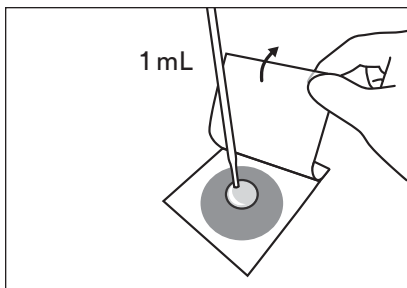

2 To seal opened package, fold the end of the pouch over and apply adhesive tape. Store resealed pouches in a cool dry place for no longer than four weeks. **To prevent exposure to moisture, do not refrigerate opened pouches.**
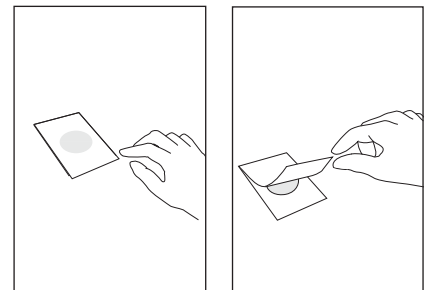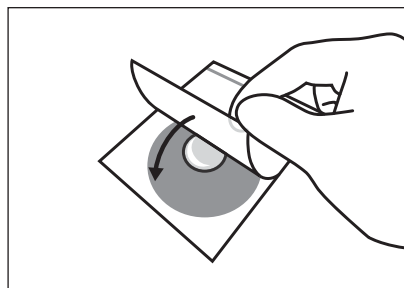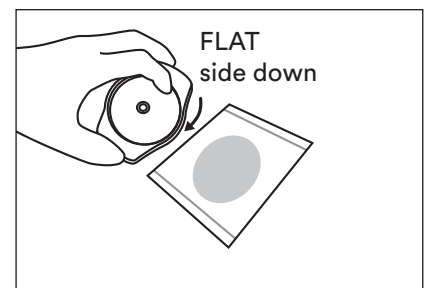

3 Place 3M Petrifilm Select *E. coli* Count Plate on flat, level surface. Lift top film.

1 mL

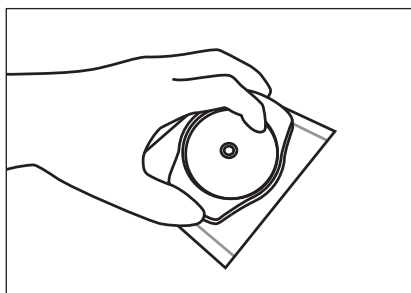**4** With pipette perpendicular to the plate, place 1 mL of sample onto center of bottom film.

5 Carefully roll top film down to avoid trapping air bubbles. Do not let top film drop.
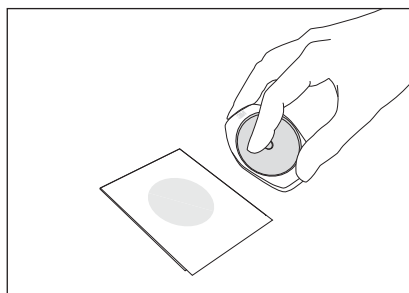

FLAT side down

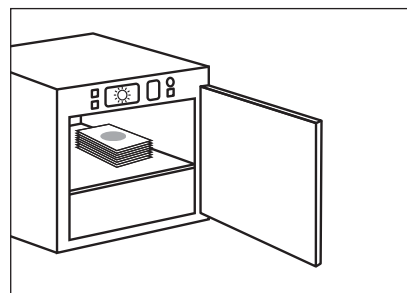**6** With flat side down, place 3M™ Petrifilm™ Spreader on top film over inoculum.

Interpretation

7 Gently apply pressure on spreader to distribute inoculm over circular area. Do not twist or slide the spreader.
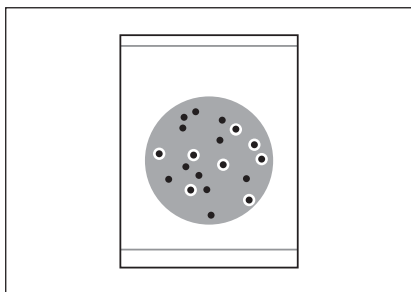

8 Lift spreader. Wait at least one minute to permit the gel to form.

9 Incubate plates with clear side up in stacks of up to 20. It may be necessary to humidify the incubator to minimize moisture loss. **Please refer to the product instructions for third party validated methods.**

Interpretation

10 3M Petrifilm Select *E. coli* Count Plates can be counted with the 3M™ Petrifilm™ Plate Reader, a standard colony counter or other illuminated magnifier. Colonies may be isolated for further identification. Lift top film and pick the colony from the gel.

Use Appropiate Sterile Diluents

Butterfield's phosphate buffered dilution water, 0.1% peptone water, peptone salt diluent, buffered peptone water, quarter-strength Ringer's solution, dipotassium hydrogen phosphate, saline solution (0.85-0.90%), bisulfite-free letheen broth or distilled water.

For optimal growth and recovery of the microorganisms, adjust the pH of the sample suspension to 6.5-7.5.

Do not use diluents containing citrate, bisulfite or thiosulfate with 3M Petrifilm Select *E. coli* Count Plates; they can inhibit growth.

If citrate buffer is indicated in the standard procedure, substitute with one of the buffers listed above, warmed to 40-45°C (104-113°F).

3M Food Safety offers a full line of products to accomplish a variety of your microbial testing needs. For more product information, visit us at **3M.com/foodsafety/Petrifilm** or call 1-800-328-6553.

![3M logo]

3M Food Safety
3M Center, Building 275-5W-05
St. Paul, MN 55144-1000 USA

1-800-328-6553
3M.com/foodsafety

3M Canada
Post Office Box 5757
London, Ontario N6A 4T1
Canada

1-800-364-3577