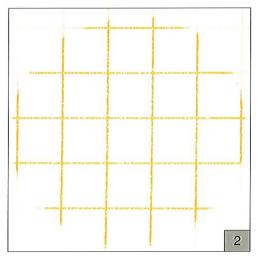
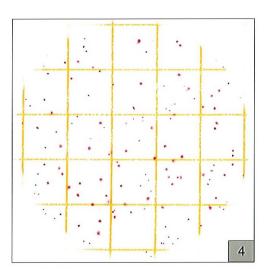
$3M^{\mathsf{TM}}$ Petrifilm Aerobic Count Plate



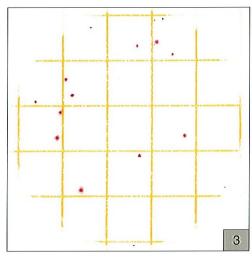
Count = 0

It is easy to interpret the Petrifilm AC plate. Figure 2 shows a Petrifilm AC plate without colonies.



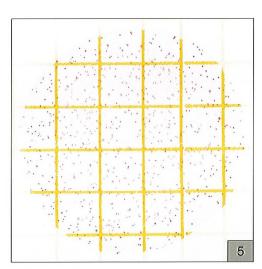
Count = 143

The preferable counting range on a Petrifilm AC plate is 25–250 colonies. See figure 4.



Count = 16

Figure 3 shows a Petrifilm AC plate with a few bacterial colonies.



Estimated Count = 560

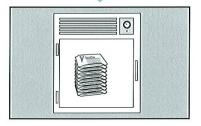
When colonies number more than 250, as in figure 5, estimate the count. Determine the average number of colonies in one square (1 cm²) and multiply it by 20 to obtain the total count per plate. The inoculated area on a Petrifilm AC plate is approximately 20 cm².

3M Petrifilm[™] Aerobic Count Plates

Reminders for Use

For detailed CAUTIONS, DISCLAIMER OF WARRANTIES / LIMITED REMEDY, LIMITATION OF 3M LIABILITY, STORAGE AND DISPOSAL information, and INSTRUCTIONS FOR USE see Product's package insert.

Storage



Store unopened packages at ≤8°C (≤46°F). Use before expiration date on package. In areas of high humidity where condensate may be an issue, it is best to allow packages to reach room temperature before opening.



To seal opened package, fold end over and tape shut.

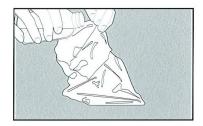


3 Keep resealed packages at ≤25°C (≤77°F) and ≤50%RH. Do not refrigerate opened packages. Use Petrifilm plates within one month after opening.

Sample Preparation



Prepare a 1:10 or greater dilution of food sample. Weigh or pipette food product into an appropriate container such as a stomacher bag, dilution bottle, Whirl-Pak® bag, or other sterile container.



Add appropriate quantity of one of the following sterile diluents: Butterfield's phosphate buffer (IDF phosphate buffer, 0.0425 g/L of KH₂PO₄ adjusted to pH 7.2), 0.1% peptone water, peptone salt diluent (ISO method 6887), buffered peptone water (ISO method 6579), saline solution (0.85 - 0.90%), bisulfate-free letheen broth, or distilled water.

Do not use buffers containing citrate, bisulfite, or thiosulfate; they can inhibit growth.

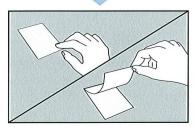


Blend or homogenize sample per current procedure.

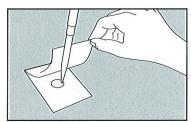
Adjust pH of the diluted sample between 6.6 and 7.2:

- for acid products, use 1 N NaOH,
- for alkaline products, use 1NHCI.

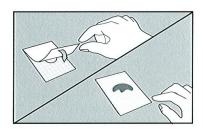
Inoculation



Place Petrifilm plate on **level** surface. Lift top film.

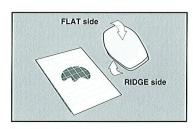


With pipette perpendicular to Petrifilm plate, place 1 mL of sample onto center of bottom film.

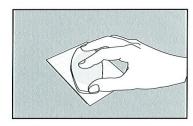


Palease top film; allow it to drop.
Do not roll top film down.

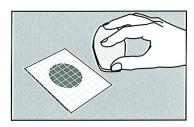
Continued - over



10 With **ridge** side down, place spreader on top film over inoculum.

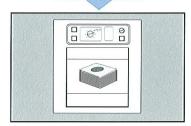


11 Gently apply pressure on spreader to distribute inoculum over circular area. Do not twist or slide the spreader.



12 Lift spreader. Wait at least one minute for gel to form.

Incubation



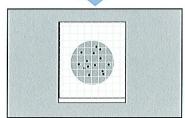
13 Incubate plates with clear side up in stacks of no more than 20. It may be necessary to humidify incubator to minimize moisture loss.

Incubation time and temperature varies by method. Most common approved methods:

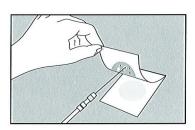
- AOAC® Official Method 986.33 & 989.10 (milk and dairy products) Incubate 48h ± 3h at 32°C ± 1°C
- AOAC° Official Method 990.12 Incubate 48h ± 3h at 35°C ± 1°C
- AFNOR Validated Method 3M 01/1-09/89

Incubate 72h ± 3h at 30°C ± 1°C

Interpretation



14 Petrifilm plates can be counted on a standard colony counter or other magnified light source. Refer to the *Interpretation Guide* section when reading results.



15 Colonies may be isolated for further identification. Lift top film and pick the colony from the gel.

Additional Comments

- Questions? U.S., call 1-800-328-6553.
- To order Petrifilm Plates in the U.S., call 1-800-328-1671.
- 3M Microbiology offers a full line of products to accomplish a variety of your microbial testing needs. For more information, visit us at www.3M.com/microbiology

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3M Microbiology 3M Center, Bldg. 275-5W-05 St. Paul, MN 55144-1000 USA 1-800-328-6553 microbiology@mmm.com

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