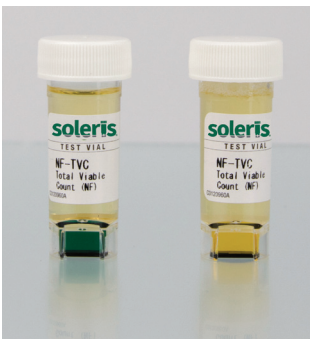




# soleris

## Non-Fermenting Total Viable Count

Product Number: NF-TVC



*Pictured:* Soleris® vial uninoculated (left) and inoculated vial (right).

### Introduction

The Total Viable Count (NF-TVC) Vial (9 mL) is suitable for sterility testing in a variety of matrices. The vial has broad inclusivity and an assay time of 24 hours or less for most applications. As organisms grow in the broth medium, the carbon dioxide (CO<sub>2</sub>) produced diffuses through a membrane layer into a soft agar plug containing a dye indicator. The color change in the dye is read by the Soleris® instrument. The membrane layer also serves as a barrier, eliminating product interference with the reading frame.

In an AOAC Research Institute Performance Tested Method SM License #071203 study, the Soleris NF-TVC vial was found to be an effective procedure for semi-quantitative determination of total aerobic count in the following sample types: raw chicken, deli ham, lettuce, almonds, black pepper, cheesecake, ice cream mix, nonfat dry milk, and cocoa powder. Test duration is 24 hours, with the exception of cocoa powder, which is 27 hours.

### Materials Required

NF-TVC, Non-fermenting Total Viable Count (NF-TVC) vial

### Dependent on Sample Tested

1. Butterfield's Phosphate Buffer (BPB-99)
2. Sterile 1 N to 5 N sodium hydroxide (NaOH) and/or hydrochloric acid (HCl)
3. pH meter or pH paper
4. For USP Testing: Tryptic Soy Broth (BLX-TSB90)
  - a. If required, use a designated neutralization broth, such as D/E Neutralizer, TAT Broth, Modified Lethen Broth, etc.

### Vial Specifications

1. Vial pH is  $7.3 \pm 0.2$
2. Vial sample capacity up to 1.0 mL

### Vial Preparation

1. Remove NF-TVC vials from the refrigerator and allow to equilibrate to room temperature.



### Sample Preparation

1. Add sample directly or, if using dilute-to-specification, complete the dilution required.
  - a. For USP testing, perform 1:10 dilution by adding 10 g of sample in 90 mL of Tryptic Soy Broth or designated neutralization broth.
    - i. Check pH and adjust, if necessary, to  $7.0 \pm 1.0$ .
  - b. For all other testing, perform 1:10 dilution by adding 11 g of sample in 99 mL of Butterfield's Phosphate Buffer.
    - i. Check pH and adjust, if necessary, to  $7.0 \pm 1.0$ .
2. If necessary, use Butterfield's Phosphate Buffer to create the dilutions to the appropriate specification.

### Sterility Testing

Preincubation for sterility testing

1. Preincubate the sample for 48–72 hours at 32–37°C as optimized for the customer.
2. Shake the container to thoroughly mix any organisms present in the product.
3. After the preincubation, test the sample by adding up to 1.0 mL of the sample to the vial.

### Inoculation of Vial

1. Inoculate the vial with no more than 1.0 mL and no less than 0.10 mL of the sample to be tested. If using dilute-to-specification method, add the volume of the appropriate dilution required.
2. Cap the vial and gently invert 3 times to mix sample. Keep cap tight.
3. Insert the vial into the Soteris instrument set at 32–35°C or as indicated by trainer. The incubation temperature and test duration can be optimized within the listed ranges for different product types. It is not recommended to adjust parameters without consulting NEOGEN® Technical Services.

### Algorithm Utilized:

Test	Threshold	Skip	Shuteye	Test Duration	Temperature
NF-TVC	10	1	30*	24 hours*	32°C–35°C

\*If shuteye detections are observed at 2.8 hours, the parameters may need to be adjusted based on product matrix. Please consult NEOGEN Technical Services for assistance.

**Caution:** Products containing CO<sub>2</sub>-releasing compounds (e.g., ascorbic acid, calcium carbonate, or calcium ascorbate) need to be carefully validated, as reactions with the vial chemistry may occur causing false positive results.

### Disclaimers:

Information provided is based on validation procedures that NEOGEN performed in NEOGEN laboratories. Deviation from procedures is possible, but should be discussed with NEOGEN Technical Services.

Appearance of the vials should be inspected prior to use.

Certain product matrices may require new parameters. For more information, contact NEOGEN Technical Services.

If shuteye detections are observed, the threshold may need to be adjusted based on the product matrix. Certain product matrices may require parameter adjustments, including increased test duration. For more information contact NEOGEN Technical Services.