

Lethen Broth Base Modified without Tween (NCM0129)

Intended Use

Lethen Broth Base Modified without Tween is used with Polysorbate 80 for the recovery of microorganisms from cosmetics in a laboratory setting. Lethen Broth Base Modified without Tween is not intended for use in the diagnosis of disease or other conditions in humans.

Description

In 1948, Weber and Black described the value of a highly nutritional solid medium containing neutralizing agents for quaternary ammonium compounds in sanitizers. The addition of Lecithin and Polysorbate 80 to Tryptone Glucose Extract (TGE) Agar resulted in a medium that effectively neutralizes quaternary ammonium compounds in testing of germicidal activity. Total neutralization of disinfectants is critical. Disinfectant residues can result in a false negative (no-growth) test.

Lethen Broth Base Modified without Tween is based on the formula described in FDA Bacteriological Analytical Manual, and a modification of Lethen Broth Base. Lethen Broth Base Modified is recommended by the FDA for use in the microbiological testing of cosmetics.

Typical Formulation

| | |
|-----------------------------------|----------|
| Lethen Broth Base | 20.7 g/L |
| Enzymatic Digest of Casein | 5.0 g/L |
| Enzymatic Digest of Animal Tissue | 10.0 g/L |
| Yeast Extract | 2.0 g/L |
| Sodium Bisulfite | 0.1 g/L |

Final pH: 7.2 ± 0.2 at 25°C

Formula may be adjusted and/or supplemented as required to meet performance specifications.

Supplement / Liter (7992)

Tween 80 (Polysorbate 80) 5g

Precaution

Refer to SDS

Preparation

1. Dissolve 37.8 g of the medium and 5 g of Tween 80 (7992) in one liter of purified water.
2. Heat with frequent agitation to completely dissolve the medium.
3. Autoclave at 121°C for 15 minutes.

Quality Control Specifications

Dehydrated Appearance: Powder is homogeneous, free flowing, and light beige.

Prepared Appearance: Prepared medium is orange to amber and clear to lightly hazy.

Expected Cultural Response: Cultural response in Lethen Broth Base Modified (supplemented with Tween) incubated aerobically at 35 ± 2°C and examined for growth after 18 - 24 hours.

| Microorganism | Approx. Inoculum (CFU) | Expected Results |
|---|------------------------|------------------|
| <i>Enterococcus faecalis</i> ATCC® 29212 | 10 - 300 | Growth |
| <i>Escherichia coli</i> ATCC® 25922 | 10 - 300 | Growth |
| <i>Pseudomonas aeruginosa</i> ATCC® 27853 | 10 - 300 | Growth |
| <i>Staphylococcus aureus</i> ATCC® 25923 | 10 - 300 | Growth |

The organisms listed are the minimum that should be used for quality control testing.

Effective Date: 5/20/2019

Revision: 0

Test Procedure

1. Prepare and dilute samples in Lethen Broth Base Modified (with Tween) in accordance with established guidelines. Proceed with procedure using Lethen Agar Base Modified, Product No. 7495.
2. Using the spread plate technique, inoculate in duplicate 0.1 mL of the diluted samples onto Lethen Agar Base Modified, Potato Dextrose Agar containing Chlorotetracycline, Baird Parker Agar, Anaerobic Agar, and a second set of Lethen Agar Base Modified plates.
3. Incubate one set of Lethen Agar Base Modified (with Tween) at 30°C for 48 hours and the other set at 35°C under anaerobic conditions for 2 - 4 days. Incubate the Potato Dextrose Agar plates at 30°C for 7 days and the Baird Parker Agar plates, if inoculated, at 35°C for 48 hours.
4. Incubate diluted samples from Step 1 at 35°C for 7 days. Subculture enriched samples onto Lethen Agar Base Modified only if there is no growth on the primary Lethen Agar Base, Modified.

Results

Refer to appropriate references for results

Expiration

Refer to expiration date stamped on the container. The dehydrated medium should be discarded if not free flowing, or if appearance has changed from the original color. Expiry applies to medium in its intact container.

Limitation of the Procedure

Due to nutritional variation, some strains may grow poorly or fail to grow on this medium.

Storage

Store dehydrated culture media at 2-30°C away from direct sunlight. Once opened and recapped, place container in a low humidity environment at the same storage temperature. Protect from moisture and light by keeping container tightly closed.

References

1. Weber, G. R., and L. A. Black. 1948. Relative efficiency of quaternary inhibitors. Soap and Sanit. Chem. 24:134-139.
2. www.fda.gov/Food/ScienceResearch/LaboratoryMethods/BacteriologicalAnalyticalManualBAM/default.htm
3. Quisno, R., I. W. Gibby, and M. J. Foter. 1946. A neutralizing medium for evaluating the germicidal potency of the quaternary ammonium salts. Am. J. Pharm. 118:320-323.
4. Erlandson, A. L., Jr., and C. A. Lawrence. 1953. Inactivating medium for hexachlorophene (G-11) types of compounds and some substituted phenolic disinfectants. Science. 118:274-276.
5. Brummer, B. 1976. Influence of possible disinfectant transfer on *Staphylococcus aureus* plate counts after contact sampling. Appl. Environ. Microbiol. 32:80-84.
6. Favero (chm.). 1967. Microbiological sampling of surfaces-a state of the art report. Biological Contamination Control Committee, American Association for Contamination Control.