

TM 935 – FDA BROTH (ATCC BACTERIOSTASIS BROTH)

INTENDED USE

For routine antimicrobial testing of antiseptics and disinfectants.

PRODUCT SUMMARY AND EXPLANATION

ATCC Bacteriostasis Broth (FDA Broth) is useful for subcultures in phenol coefficient and dilution tests of bacteriostatic, germicidal, sporicidal activity and also as a base for the preparation of ATCC Bacteriostasis Agar.

COMPOSITION

Ingredients	Gms / Ltr
Peptic digest of animal tissue	10.000
Beef extract	5.000
Sodium chloride	5.000

PRINCIPLE

The medium consists of Peptic digest of animal tissue and beef extract that are sources of carbon, nitrogen, vitamins and minerals. Sodium chloride provides essential ions.

INSTRUCTION FOR USE

- Dissolve 20.0 grams in 1000 ml purified / distilled water.
- Heat if necessary to dissolve the medium completely.
- Sterilize by autoclaving at 15 psi pressure (121°C) for 15 minutes.
- Mix well and dispense as desired.

QUALITY CONTROL SPECIFICATIONS

- Appearance of Powder** : Cream to yellow homogeneous free flowing powder.
Appearance of prepared medium : Amber coloured clear solution in tubes.
pH (at 25°C) : 6.8 ± 0.2

INTERPRETATION

Cultural characteristics observed after incubation.

Microorganism	ATCC	Inoculum (CFU/ml)	Growth	Incubation Temperature	Incubation Period
<i>Escherichia coli</i>	25922	50-100	Good- luxuriant	35-37°C	18-24 Hours
<i>Pseudomonas aeruginosa</i>	27853	50-100	Good- luxuriant	35-37°C	18-24 Hours



<i>Staphylococcus aureus</i>	6538	50-100	Good- luxuriant	35-37°C	18-24 Hours
<i>Salmonella Typhi</i>	6539	50-100	Good- luxuriant	35-37°C	18-24 Hours

PACKAGING:

In pack size of 500 gm bottles.

STORAGE

Dehydrated powder, hygroscopic in nature, store in a dry place, in tightly-sealed containers between 25-30°C and protect from direct sunlight. Under optimal conditions, the medium has a shelf life of 4 years. When the container is opened for the first time, note the time and date on the label space provided on the container. After the desired amount of medium has been taken out replace the cap tightly to protect from hydration.













Product Deterioration: Do not use if they show evidence of microbial contamination, discoloration, drying or any other signs of deterioration.

DISPOSAL

After use, prepared plates, specimen/sample containers and other contaminated materials must be sterilized before discarding.

REFERENCES

- Williams (Ed.), 1995, Official methods of Analysis of AOAC, 16th ed. AOAC, Washington D.C.
- Tech. Manual of AATCC, 1985, Vol. 61, AATCC, Research Triangle Park,N.C.

 GMP Good Manufacturing Practices Certified	 IVD For In Vitro Diagnostic Use	 QTY. Quantity	 LOT/ B. NO. Lot / Batch Number	 REF Catalogue Number	 Manufacturer
 Temperature Unit	 EC REP Authorized Representative <small>MediMet GmbH Bockstrasse 10 48163 Muenster, Germany</small>	 European Conformity	 QR Code	 Consults Instructions for Use	 Best Before

NOTE: Please consult the Material Safety Data Sheet for information regarding hazards and safe handling Practices.

*For Lab Use Only
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