

# **ANTIBIOTIC BROTH N.3**

Liquid medium used for the determination potency of antibiotics by the microbial assay technique following the USP specifications.

TYPICAL FORMULA (g/L)				
Peptone	5.0			
Yeast Extract	1.5			
Beef Extract	1.5			
Sodium Chloride	3.5			
Dextrose	1.0			
Dipotassium Phosphate	3.68			
Monopotassium Phosphate	1.32			
Final pH 7.0 ± 0.2				

## DESCRIPTION

**ANTIBIOTIC BROTH N.3** is a liquid medium for the determination potency of antibiotics by the microbial assay technique following the USP (United Staes Pharmacopeia) specifications.

### PRINCIPLE

## Cylinder Plate Assay

This method is used in the assay of commercial preparations of antibiotics and in the quantitative determination of antibiotics in body fluids, animal feeds and other materials. It is based on the diffusion of an antibiotic solution from a cylinder placed on the surface of an inoculated agar medium. The diameter of a zone of inhibition after incubation depends, in part, on the concentration or activity of the antibiotic.

## **Turbidimetric Assay**

This method is based on the inhibition of growth of a microbial culture in a fluid medium containing a uniform solution of an antibiotic. Turbidimetric determinations have the advantage of requiring a short incubation period, providing test results after 3 or 4 hours. However, the presence of solvents or other inhibitory materials may influence turbidimetric assays more markedly than cylinder plate assays. Use of this method is appropriate only when test samples are clear.

### PREPARATION

Suspend 17.5 g of powder in 1 litre of distilled or deionized water. Heat with frequent agitation and boil for 1 minute to completely dissolve the powder. Dispense into culture tubes. Autoclave at 121°C for 15 minutes.

#### TECHNIQUE

Prepare the inoculum for assay by washing growth from a fresh 24-48 hour agar slant using sterile purified water and further dilute the culture to obtain the desired organism concentration.

### **Cylinder Plate Assay**

Use 20  $\times$  100 mm glass or plastic Petri dishes with sufficient depth so that cylinders used in the assay will not be pushed into the medium by the cover. Use stainless steel or porcelain assay cylinders having the following dimensions ( $\pm$  0.1 mm): 8 mm outside diameter, 6 mm inside diameter and 10 mm long.

To assure accurate assays, work on a level surface to obtain uniformly thick base and seed layers in the Petri dish. Allow the base layer to solidify and then overlay the seed layer containing a proper concentration of the test organism. The amount of medium in the layers varies for different antibiotics, with most assays specifying a 21 mL base layer and a 4 mL seed layer. In any case, dishes with flat bottoms are required to assure complete coverage of the bottom of the dish when small amounts of base medium are used. Tilt the plate to obtain even coverage of the base layer and allow it to solidify in a level position. Plates should be used the same day as prepared.

## Turbidimetric Assay

Use glass or plastic test tubes (i.e., 16 × 125 mm or 18 × 150 mm) that are relatively uniform in length, diameter and thickness. Prepare working dilutions of the antibiotic reference standards in specific concentrations. To a 1 mL quantity of each solution in a suitable tube, add 9 mL of inoculated broth, as required. Prepare similar solutions of the assay materials containing approximately the same amounts of antibiotic activity and place in tubes. Incubate the tubes for 3-4 hours at the required temperature, generally in a water bath. At the end of the incubation period, stop growth by adding 0.5 mL of 1:3 formalin. Determine the amount of growth by measuring light transmittance with a suitable spectrophotometer. Determine the concentration of the antibiotic by comparing the growth obtained with that given by reference standard solutions.

### INTERPRETATION OF RESULTS

Refer to appropriate procedures for results.

### STORAGE

The powder is very hygroscopic: store the powder at 10-30°C, in a dry environment, in its original container tightly closed until the expiry date indicated on the label or until signs of deterioration or contamination are evident.

### Store prepared media at 2-8°C. WARNING and PRECAUTIONS

The product is not classified as hazardous by current legislation and does not contain harmful substances in concentrations of  $\geq$ 1%. The product must be used only by properly trained operators.

## DISPOSAL of WASTE

Disposal of waste must be carried out according to national and local regulations in force.

## REFERENCES

- 1. Grove and Randall. (1995). Assay methods of antibiotics. Medial Encyclopedia, Inc. New York, N.Y.
- 2. United States Pharmacopeial Convention, Inc. (2001). The United States pharmacopeia 25/The national formulary 20-2002. United States Convention, Inc., Rockville, Md.
- 3. Horwitz (ed.). (2000). Official methods of analysis of AOAC International, 17th ed., vol. 1.AOAC International, Gaithersburg, Md.
- 4. Foster and Woodruff. (1943). J. Bacteriol. 46:187.



# Liofilchem s.r.l.

Via Scozia, Zona industriale - 64026 Roseto degli Abruzzi (Te) Italy - Tel. +39.0858930745 - Fax +39.0858930330 Web site: <u>http://www.liofilchem.net</u> - E-mail: liofilchem@liofilchem.net



## NAME

# ANTIBIOTIC BROTH N.3

# PRESENTATION

Dehydrated culture medium

# STORAGE

10-30°C

#### PACKAGING

Code	Content	Packaging		
610316	500 g	500 gr of powder in plastic bottle		

## pH OF THE MEDIUM

 $7.0\pm0.2$ 

#### USE

**ANTIBIOTIC BROTH N.3** is a liquid medium for the determination potency of antibiotics by the microbial assay technique following the USP (United Staes Pharmacopeia) specifications.

### TECHNIQUE

Refer to technical sheet of the product.

# APPEARANCE of the MEDIUM

Dehydrated medium Appearance: free-flowing, homogeneous. Colour: light beige <u>Prepared medium</u> Appearance: clear to slightly hazy Colour: light to medium amber

### SHELFLIFE

4 years

## QUALITY CONTROL

1. Control of general characteristics, label and print

- Sterility control 7 days at 25 ± 1°C, in aerobiosis 7 days at 36 ± 1°C, in aerobiosis
- Microbiological control Inoculum for productivity: 10-100 CFU/ml Inoculum for specificity: ≤ 10<sup>4</sup> CFU/ml

Incubation conditions: at 25 ± 2°C for the Saccharomyces cerevisiae and 35 ± 2°C for the remaining organisms for 7 days.

Microorganisms		Inoculum CFU	Recovery
Bacillus subtilis	ATCC 6633	≤ 10 <sup>3</sup>	Good
Escherichia coli	ATCC 10536	≤ 10 <sup>3</sup>	Good
Micrococcus luteus	ATCC 9341	≤ 10 <sup>3</sup>	Good
Saccharomyces cerevisiae	ATCC 9763	≤ 10 <sup>3</sup>	Good
Staphylococcus aureus	ATCC 6538P	≤ 10 <sup>3</sup>	Good

## TABLE OF SYMBOLS

LOT Batch code	Temperature limitation	Manufacturer	$\sum$	Contains sufficient for <n> tests</n>
<b>REF</b> Catalogue number	Keep away from heat	Use by	[]i	Caution, consult accompanying documents

