

# Neutralizing Fluid E.P.

Liquid medium for neutralizing antimicrobials, according to the European Pharmacopoeia

TYPICAL FORMULA	(g/l)
Peptone	1.0
Histidine Hydrochloride	1.0
Lecithin	3.0
Potassium Dihydrogen Phosphate	3.6
Disodium Hydrogen Phosphate Dihydrate	7.2
Sodium Chloride	4.3
Final pH 7.0 ± 0.2 at 25°C	

#### **DESCRIPTION**

Neutralizing Fluid E.P. is a liquid medium used for neutralizing the activity of antimicrobial agents.

The medium is formulated according to the European Pharmacopoeia specification for the microbiological examination of non-sterile products.

#### PRINCIPLE

Peptone supplies amino acids, nitrogen, carbon, minerals, vitamins and other nutrients which support the growth of microorganism. Histidine inactivates aldehydes. Lecithin neutralizes quaternary ammonium compounds. Phosphates serve as buffering agents. Sodium chloride maintains the osmotic balance of the medium.

Supplementation with polysorbate (Tween) 80 is effective against phenolic compounds and mercurial derivates.

#### **PREPARATION**

Suspend 20.1 g of powder in 1 liter of deionized or distilled water containing 30 g of polysorbate 80 (ref. 80031). Bring to boil and shake until completely dissolved. Pour into suitable containers. Sterilize at 121°C for 15 minutes.

#### **TECHNIQUE**

Neutralizing Fluid E.P. may be incorporated into diluents or media, such as Buffered Peptone Water EP, USP (ref. 400040), preferably before sterilization.

### INTERPRETATION OF RESULTS

Refer to technical sheet of the medium being used.

# STORAGE AND TRANSPORT CONDITIONS

The powder is very hygroscopic, store the powder at 10-30°C, in a dry environment, in its original container tightly closed and use it before the expiry date on the label or until sings of deterioration or contamination are evident. Store the prepared medium at 2-8°C away from light.

## WARNING AND PRECAUTIONS

The product does not contain hazardous substances in concentrations exceeding the limits set by current legislation and therefore is not classified as dangerous. it is nevertheless recommended to consult the safety data sheet for its correct use. The product is designed for professional use only and must be used by properly trained operators.

# DISPOSAL OF WASTE

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

## REFERENCES

 European Pharmacopeia (2007) Directorate for the Quality of Medicine of the Council of Europe. 2.6.13. Microbiological examination of non-sterile products: test for specific micro-organisms. Council of Europe Strasbourg.



# **PRODUCT SPECIFICATIONS**

## NAME

Neutralizing Fluid E.P.

## **PRESENTATION**

Dehydrated medium

## STORAGE

10-30°C

## **PACKAGING**

Ref.	Content	Packaging
610330	500 g	500 g of powder in plastic bottle
620330	100 g	100 g of powder in plastic bottle

# pH OF THE MEDIUM

7.0 ± 0.2

#### USE

Neutralizing Fluid E.P. is a liquid medium formulated according to the European Pharmacopoeia specification for the microbiological examination of non-sterile products

#### **TECHNIQUE**

Refer to technical sheet of the product

## APPEARANCE OF THE MEDIUM

Powder medium

Appearance: free-flowing, homogeneous

Colour: beige

Ready-to-use medium
Appearance: opalescent
Colour: light amber

## SHELFLIFE

4 years

# **QUALITY CONTROL**

Control of general characteristics, label and print

2. Microbiological control

Cultural response observed when subcultured on Tryptic Soy Agar after an incubation at 30°C for 3 hours in Neutralizing Fluid E.P. Inoculum for productivity: 50-100 CFU

Incubation Conditions: 18-24 h at 32.5 ± 2.5°C, in aerobiosis

Microorganism		Growth
Staphylococcus aureus	ATCC® 6538	Good
Escherichia coli	ATCC® 8739	Good
Pseudomonas aeruginosa	ATCC® 9027	Good
Bacillus subtilis	ATCC® 6633	Good
Salmonella typhimurium	ATCC® 14028	Good
Candida albicans	ATCC® 10231	Good
Aspergillus niger	ATCC® 16404	Good

#### **TABLE OF SYMBOLS** Fragile, handle with Batch LOT Do not reuse Manufacturer Use by code care Temperature Catalogue Contains sufficient Caution, consult REF number limitation for <n> tests instructions for use

