

## TECHNICAL DATA SHEET

# MANNITOL SALT AGAR

### ENUMERATION OF PATHOGENIC STAPHYLOCOCCI

## 1 INTENDED USE

Mannitol Salt agar is used for the selective isolation, detection and enumeration of pathogenic staphylococci in filterable water as in swimming pools, potable water or spas. It is also used for the detection of *Staphylococcus aureus* according to the Pharmacopeia and in cosmetic products.

The typical composition responds to the formulation defined in the European Pharmacopeia, and in the Directive NF T90-421 for the water control and in the Directive NF EN ISO 22718 in cosmetics.

## 2 HISTORY

The experiments of Koch showed that staphylococci could tolerate hypersalted media at 7.5%. Chapman confirmed these initial results and observed that staphylococci which coagulated rabbit plasma formed yellow colonies on this medium, while most other bacteria were inhibited.

## 3 PRINCIPLES

The high sodium chloride concentration inhibits the growth of most bacteria other than staphylococci.

Mannitol fermentation, shown by the color change of the pH indicator (phenol red) to yellow orients the diagnosis.

The demonstration of pathogenic staphylococci is generally confirmed by a coagulase test.

## 4 TYPICAL COMPOSITION

The composition can be adjusted to obtain optimal performance.

For 1 liter of media:

- Tryptone .....	5,0 g
- Peptic digest of meat.....	5,0 g
- Meat extract.....	1,0 g
- Mannitol.....	10,0 g
- Sodium chloride .....	75,0 g
- Phenol red.....	25,0 mg
- Bacteriological agar.....	15,0 g

pH of the ready-to-use media at 25 °C: 7,4 ± 0,2.

## 5 PREPARATION

- Suspend 111,0 g of dehydrated medium (BK030) in 1 liter of distilled or deionized water.
- Slowly bring to boiling, stirring with constant agitation until complete dissolution.
- Dispense in tubes or flasks.
- Sterilize in an autoclave at 121°C for 15 minutes.

✓ **Reconstitution:**  
111,0 g/L

✓ **Sterilization:**  
15 min at 121 °C

- Cool and maintain at 44-47°C.
- Pour into sterile Petri dishes (Ø 55 mm or 90 mm according to the application) and let solidify on a cold surface.

## 6 INSTRUCTIONS FOR USE

### Detection of *Staphylococcus aureus* (Pharmacopeia, NF EN ISO 22718)

- Dry in an incubator the 90 mm plates with the covers partially removed.
- Transfer 0.1 mL of the product to analyze to the surface of the plates.
- Spread the inoculum on the surface of the medium with a sterile triangle.
- Incubate at 30-35 °C for 18 to 24 hours.

✓ **Inoculation:**  
0,1 mL on surface

✓ **Incubation:**  
18-24 h at 30-35 °C

### Water testing by membrane filtration

- Aseptically filter the appropriate volume of water through a nitrocellulose membrane.
- Place the membrane, filtered side up, on the surface of the agar prepared as described (or use ready-to-use plates BM148), insuring a complete contact.
- Incubate at (36 ± 2) °C for (44 ± 4) hours.

✓ **Inoculation:**  
Membrane filtration

✓ **Incubation:**  
44 h at 36 °C

## 7 RESULTS

Count the characteristic colonies.

Pathogenic staphylococci form luxuriant pigmented colonies, surrounded by a yellow ring due to the fermentation of mannitol.

Non-pathogenic staphylococci generally give rise to small red colonies which do not change the color of the medium.

Several strains of *Staphylococcus epidermidis* can ferment mannitol.

After 48 hours of incubation, several strains of enterococci, *Bacillus*, *Micrococcus* and *Serratia* may grow.

See ANNEX 1: PHOTO SUPPORT.

## 8 QUALITY CONTROL

**Dehydrated media:** cream to pinkish colored powder, free-flowing and homogeneous.

**Prepared media:** red agar.

Typical culture response after 18-24 hours of incubation at 30-35 °C

Microorganisms		Growth Productivity Ratio P <sub>R</sub>	Characteristics
<i>Staphylococcus aureus</i>	WDCM 00032	P <sub>R</sub> ≥ 50 %	Yellow colonies
<i>Staphylococcus aureus</i>	WDCM 00034	P <sub>R</sub> ≥ 50 %	Yellow colonies
<i>Staphylococcus epidermidis</i>	WDCM 00036	Good, score 2	Pink colonies
<i>Escherichia coli</i>	WDCM 00013	Inhibited, score 0	-

Typical culture response after 44 hours of incubation at 36 °C

Microorganisms		Growth R <sub>3</sub> acc. to NF T90-461	Characteristics
<i>Staphylococcus aureus</i>	WDCM 00035	66 % ≤ R <sub>3</sub> ≤ 150 %	Yellow colonies
<i>Enterococcus faecalis</i>	WDCM 00176	Inhibited	-

## 9 STORAGE / SHELF LIFE

**Dehydrated media:** 2-30 °C.

**Pre-poured media:** 2-8 °C.

The expiration dates are indicated on the labels.

**Prepared media in vials (\*):** 180 days at 2-8 °C.

**Prepared media in plates (\*):** 30 days at 2-8 °C.

(\*) Benchmark value, determined in standard conditions of preparation, following manufacturer's instructions.

## 10 PACKAGING

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### Dehydrated media:

500 g bottle ..... BK030HA

### Pre-poured media in Petri plates (Ø 55 mm):

20 plates ..... BM14808

## 11 BIBLIOGRAPHY

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Chapman, G.H. 1945. The significance of sodium chloride in studies of staphylococci. J. Bacteriol., 50: 201.

Chapman, G.H. 1948. An improved Stone medium for the isolation and testing of food poisoning *staphylococci*. Food Research, 13: 100-105.

NF T 90-421. Août 2006. Essais des eaux. Examens bactériologiques des eaux de piscines.

NF EN ISO 22718. Septembre 2009. Cosmétiques. Microbiologie. Détection de *Staphylococcus aureus*.

Pharmacopée européenne. Chapitre 2.6.13. Contrôle microbiologique des produits non stériles : Recherche de microorganismes spécifiés.

## 12 ADDITIONAL INFORMATION

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The information provided on the labels take precedence over the formulations or instructions described in this document and are susceptible to modification at any time, without warning.

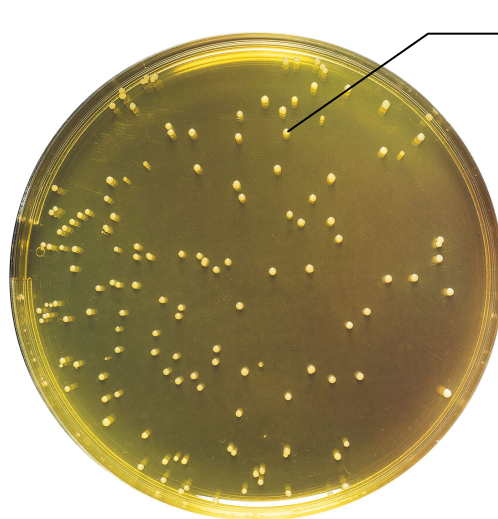
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## Mannitol Salt Agar

Detection and enumeration of pathogenic staphylococci

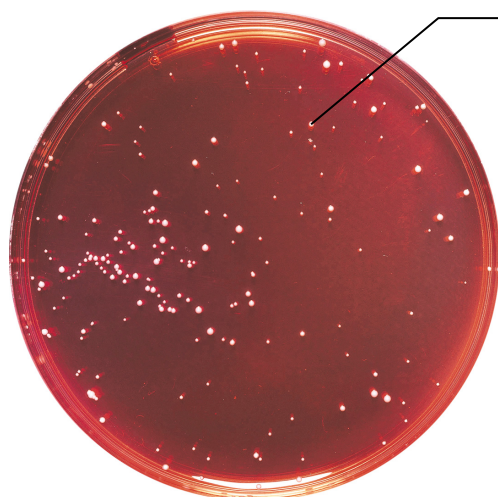
### Results:

Growth obtained after 24 hours of incubation at 37 °C (surface inoculation).



***Staphylococcus aureus***

Characteristic colony:  
yellow colony on yellow media, or  
surrounded by a yellow halo.



***Staphylococcus epidermidis***

Non characteristic colony:  
white colony on red agar.