



## Salt Agar, Modified

M1767

Salt Agar, Modified is used for isolation and differentiation of the enterococcal group D Streptococci from non-enterococcal group D streptococci based on salt tolerance.

### Composition\*\*

Ingredients	Gms / Litre
Peptic digest of animal tissue	10.000
Heart infusion	10.000
Glucose	1.000
Sodium chloride	65.000
Bromocresol purple	0.016
Agar	15.000
Final pH ( at 25°C)	7.2±0.2

\*\*Formula adjusted, standardized to suit performance parameters

### Directions

Suspend 101.01 grams in 1000 ml distilled water. Heat to boiling to dissolve the medium completely. Sterilize by autoclaving at 15 lbs pressure (121°C) for 15 minutes. Cool to 45-50°C. Mix well and pour into sterile Petri plates.

### Principle And Interpretation

Salt Agar, Modified is used for differentiating enterococcal group D streptococci from non-enterococcal group D streptococci. Medium containing 6.5% sodium chloride is used to differentiate Enterococci by determining salt tolerance of bile esculin positive and catalase negative cocci (2). High salt content of this medium acts as a differential and selective agent by interfering with membrane permeability and osmotic equilibrium (1). Enterococcal group D *Streptococcus* species ( *Enterococcus faecalis* , *Enterococcus faecium* , *Enterococcus durans* and *Enterococcus avium* ) can be easily differentiated from the non-enterococcal species like *Streptococcus bovis* , *Streptococcus equines* , by the 6.5% sodium chloride tolerance test.

Heart infusion and peptic digest of animal tissue provide essential nitrogenous nutrients while glucose is the carbohydrate source in the medium. Bromocresol purple is the pH indicator which turns yellow from purple at acidic pH (2). Sodium chloride serves as differential and selective agent. Growth is indicated by turbidity and sometimes changes in colour of the indicator. A change in colour from purple to yellow also may occur due to utilization of glucose and thereby acid production. Serological group D streptococci or bile esculin positive isolate may be easily identified as an *Enterococcus* species.

### Quality Control

#### Appearance

Cream to greenish yellow homogeneous free flowing powder

#### Gelling

Firm, comparable with 1.5% Agar gel

#### Colour and Clarity

Purple coloured clear to slightly opalescent solution

#### Reaction

Reaction of 10.1% w/v aqueous solution at 25°C. pH : 7.2±0.2

#### pH

7.00-7.40

#### Cultural Response

M1767: Cultural characteristics observed after an incubation at 35-37°C for 24-48 hours.

Organism	Inoculum (CFU)	Growth	Recovery
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#### Cultural Response

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<i>Streptococcus bovis</i> ATCC 9809	$\geq 10^3$	inhibited	0%
<i>Enterococcus faecalis</i> ATCC 29212	50-100	good	$\geq 70\%$

### Storage and Shelf Life

Store below 30°C in tightly closed container and the prepared medium at 2-8°C. Use before expiry date on the label.

### Reference

1. MacFaddin J. F., 1985, Media for Isolation-Cultivation-Identification- Maintenance of Medical Bacteria, Vol. 1, Williams Wilkins, Baltimore, Md.
2. Murray P. R., Baron J. H., Pfaller M. A., Jorgensen J. H. and Tenover F. C., (Ed.), 2003, Manual of Clinical Microbiology, 8th Ed., American Society for Microbiology, Washington, D.C.

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