

# **Technical Data**

M1979

## SS Agar w/sucrose

## **Intended Use:**

It is used for the selective isolation and differentiation of *Salmonella* and *Shigella* species from clinical and non-clinical samples.

## Composition\*\*

| Ingredients             | Gms / Litre |
|-------------------------|-------------|
| HM extract $\ominus$    | 3.000       |
| Tryptone \$             | 4.000       |
| Peptone                 | 4.000       |
| Sodium citrate          | 5.000       |
| Sodium thiosulphate     | 2.000       |
| Ferric ammonium citrate | 1.000       |
| Lactose                 | 10.000      |
| Saccharose(Sucrose)     | 10.000      |
| Bile salt               | 5.000       |
| Neutral red             | 0.020       |
| Agar                    | 15.000      |
| Final pH ( at 25°C)     | $7.4\pm0.2$ |

<sup>\*\*</sup>Formula adjusted, standardized to suit performance parameters

#### **Directions**

Suspend 59.03 grams in 1000 ml purified / distilled water. Heat to boiling with frequent agitation to dissolve the medium completely. Cool to 45-50°C. Mix and pour into sterile Petri plates. DO NOT AUTOCLAVE OR OVERHEAT. Overheating may destroy the selectivity of the medium.

## **Principle And Interpretation**

Salmonella and Shigella are gram-negative, facultatively anaerobic, non-sporulating rods in the family Enterobacteriaceae. The media is recommended as differential and selective medium for the isolation of Salmonella and Shigella species from pathological specimens (5), suspected foodstuffs (2, 7, 9, 10) and for microbial limit test (8). SS

Agar is a moderately selective medium in which gram-positive bacteria are inhibited by bile salts.

HM extract, tryptone and peptone provides nitrogen, vitamins, minerals and amino acids essential for growth. Lactose and sucrose are the fermentable carbohydrates providing carbon and energy. Bile salts selectively inhibit gram-positive and coliform organisms. Sodium thiosulphate is reduced by certain species of enteric organisms to sulphite and H<sub>2</sub>S gas. This reductive enzymatic process is attributed to thiosulphate reductase. Production of H<sub>2</sub>S gas is detected as an insoluble black precipitate of ferrous sulphide, formed upon reaction of H<sub>2</sub>S with ferric ions or ferric citrate, indicated by black centered colonies. On fermentation of lactose by few lactose-fermenting normal intestinal flora, acid is produced which is indicated by change of colour from yellow to red by the pH indicator neutral red. Thus these organisms grow as red-pigmented colonies. Lactose non-fermenting organisms grow as translucent colourless colonies with or without black centers. *Salmonella* species exhibit colourless colonies with black centers resulting from H<sub>2</sub>S production. *Shigella* species form colourless colonies, which do not produce H<sub>2</sub>S.Agar acts as a solidifying agent.

## Type of specimen

Clinical samples - Stool; Food and dairy samples.

## **Specimen Collection and Handling**

For clinical samples follow appropriate techniques for handling specimens as per established guidelines (3,4). For food and dairy samples, follow appropriate techniques for sample collection and processing as per guidelines (1,7,9). After use, contaminated materials must be sterilized by autoclaving before discarding.

 $<sup>\</sup>ominus$  - Equivalent to Meat extract

<sup>\$ -</sup> Equivalent to Pancreatic digest of casein

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#### **Limitations:**

1. This medium is general purpose medium and may not support the growth of fastidious organisms.

2. Further biochemical and serological tests must be carried out for further identification.

#### **Performance and Evaluation**

Performance of the medium is expected when used as per the direction on the label within the expiry period when stored at recommended temperature.

## **Quality Control**

#### **Appearance**

Light yellow to pink homogeneous free flowing powder

#### Gelling

Firm, comparable with 1.5% Agar gel

#### **Colour and Clarity of Prepared Medium**

Reddish orange coloured clear to slightly opalescent gel forms in Petri plates

#### Reaction

Reaction of 5.93% w/v aqueous solution at 25°C. pH: 7.4±0.2

#### pН

7.20-7.60

## **Cultural Response**

Cultural characteristics observed after an incubation at 35-37°C for 18-24 hours.

| Organism                                   | Inoculum<br>(CFU) | Growth         | Recovery | Colour of colony             |
|--|-------------------|----------------|----------|------------------------------|
| Escherichia coli ATCC<br>25922 (00013*)    | 50-100            | luxuriant      | >=70%    | pink with bile precipitate   |
| Salmonella Typhimurium ATCC 14028 (00031*) | 50-100            | good-luxuriant | >=50%    | colourless with black centre |
| Shigella sonnei ATCC 2593                  | <i>l</i> 50-100   | luxuriant      | 50-70%   | colourless                   |

Key: \*Corresponding WDCM numbers.

## Storage and Shelf Life

Store between 10-30°C in a tightly closed container and the prepared medium at 20-30°C. Use before expiry date on the label. On opening, product should be properly stored dry, after tightly capping the bottle in order to prevent lump formation due to the hygroscopic nature of the product. Improper storage of the product may lead to lump formation. Store in dry ventilated area protected from extremes of temperature and sources of ignition Seal the container tightly after use. Product performance is best if used within stated expiry period.

## **Disposal**

User must ensure safe disposal by autoclaving and/or incineration of used or unusable preparations of this product. Follow established laboratory procedures in disposing of infectious materials and material that comes into contact with clinical sample must be decontaminated and disposed of in accordance with current laboratory techniques (3,4).

## Reference

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- 4. Jorgensen, J.H., Pfaller, M.A., Carroll, K.C., Funke, G., Landry, M.L., Richter, S.S and Warnock., D.W. (2015) Manual of Clinical Microbiology, 11th Edition. Vol. 1.
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- 7. Salfinger Y., and Tortorello M.L., 2015, Compendium of Methods for the Microbiological Examination of Foods, 5th Ed., American Public Health Association, Washington, D.C.
- 8. The United States Pharmacopoeia, 2006, USP29/NF24, The United States Pharmacopoeial Convention. Rockville, MD.
- 9. Wehr H. M. and Frank J. H., 2004, Standard Methods for the Microbiological Examination of Dairy Products, 17th Ed., APHA Inc., Washington, D.C.

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In vitro diagnostic medical



CE Marking



Storage temperature



Do not use if package is damaged



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