

Bases / Media Supplements

# **Technical Information**

## Semisolid Nutrient Agar

### Product Code: DM 2191

Application: - Semisolid Nutrient Agar is recommended for detection of *Salmonella* species on the basis of motility and hydrogen sulphide (H<sub>2</sub>S) production.

Composition**							
Ingredients	Gms / Litre						
Peptic digest of animal tissue	5.000						
Beef extract	3.000						
Agar	4.000						
Final pH (at 25°C)	7.0±0.2						
**Formula adjusted, standardized to suit performance parameters							

### Principle & Interpretation

Nutrient Agar is a basic culture medium used in water and food studies. It is generally used for maintenance and to check the purity of subcultures <sup>(1)</sup>. Bacterial motility is an important feature in making a final species identification. Tubes containing semisolid media are most often used to determine motility. Motile organisms form a diffuse zone of growth flaring out from the line of inoculation. Non-motile organisms grow along the stabline. Certain bacterial species have ability to liberate sulphur from sulphur-containing amino acids or other compounds in the form of  $H_2S$  (hydrogen sulphide). Lead acetate paper strips are used as the  $H_2S$  indicators <sup>(2)</sup>. Semisolid Nutrient Agar couples these two tests in a single medium. It is also recommended by ISO Committee <sup>(3)</sup> for the detection of *Salmonella* species.

Peptic digest of animal tissue and beef extract provide essential growth nutrients. The motile cultures grow away from stabline while nonmotile grow along the stabline. After inoculation, with the test organism, insert a lead acetate paper strip (031043) between the plug and inner wall of tube. Lead acetate strip incorporation helps to detect H<sub>2</sub>S production.

### Methodology

Suspend 12 grams of powder media in 1000 ml distilled water. Shake well & heat to the medium completely. Dispense in tubes. Sterilize by autoclaving at 15 lbs pressure (121°C) for 15 minutes. Allow the tubes to cool in an upright position.

Note: Due to the presence of calcium carbonate, the prepared medium forms opalescent solution with white precipitate.

### **Quality Control**

#### Physical Appearance

Cream to yellow homogeneous free flowing powder.

#### Gelling

Semisolid, comparable with 0.4% Agar gel.

#### Colour and Clarity of prepared medium

Light yellow coloured clear gel forms in tubes as butts.

#### Reaction 🕠

Reaction of 1.2% w/v aqueous solutions at 25°C. pH : 7.0±0.2

#### pH Range 6.80-7.20

#### Cultural Response/ characteristics

**DM 2191:** Culture characteristic observed after an incubation at 35-37<sup>0</sup>C for 18-24 hours.

Organism	Inoculu m CFU)	Growth	Motility	H <sub>2</sub> S (with lead acetate strip)
Escherichia coli ATCC 25922	50-100	Luxuriant	positive, growth away from stabline causing turbidity	negative reaction, no blackening
Salmonella Typhi ATCC 6539	50-100	Luxuriant	positive, growth away from stabline causing turbidity	Positive reaction, no blackening of the lower portion of the strip





Dehydrated Culture Media Bases / Media Supplements

Salmonella Enteritidis ATCC 13076

50-100

Luxuriant

positive, growth away from stabline Positive reaction, no causing turbidity blackening of the lower portion of the strip

## Storage and Shelf Life

Dried Media: Store below 30°C in tightly closed container and use before expiry date as mentioned on the label. Prepared Media: 2-8° in sealable plastic bags for 2-5 days.

### Further Reading

1. Lapage S. P., Shelton J. E. and Mitchell T. G., 1970, Methods in Microbiology, Norris J. R. and Ribbons D. W., (Eds.), Vol. 3 A, Academic Press, London.

2. Koneman E. W., Allen S. D., Janda W. M., Schreckenberger P. C., Winn W. C. Jr., 1992, Colour Atlas and Textbook of Diagnostic Microbiology, 4th Ed., J. B. Lippinccott Company.

3. International Organization for Standardization (ISO), 1993, Draft ISO/DIS 6579.

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