Technical Information

TN Agar

Product Code: DM 1950

Application: - TN Agar is used for isolation and cultivation of *Vibrios* from food samples.

Composition**

Ingredients	Gms / Litre
Casein enzymic hydrolysate	10.000
Sodium chloride	10.000
Agar	15.000
Final pH (at 25°C)	7.2±0.2

**Formula adjusted, standardized to suit performance parameters

Principle & Interpretation

Members of the genus *Vibrio* are defined as gram-negative, asporogenous rods that are straight or have a single rigid curve. Three species of *Vibrio,* namely *Vibrio cholerae, Vibrio parahaemolyticus* and *Vibrio mimicus* are well-documented as human pathogens.
V. cholerae, the type species of the genus *Vibrio* is the causative agent of cholera outbreaks and epidemics.
V. cholerae can be differentiated from other *Vibrio* species except *V. mimicus ,* because of its obligate requirement for sodium ions (Na+).TN Agar is formulated according to APHA [(1)] for cultivation of *Vibrio* species from foods.
Casein enzymic hydrolysate provides nitrogenous, carbonaceous compounds, sulphur, vitamin B complex and other essential growth nutrients. Sodium chloride improves the selectivity of the medium.
Weigh 25 grams of food sample such as seafood or vegetable and blend or cut into small pieces into two jars. To one jar add 225 ml Alkaline Peptone Water (DM1618) and to another jar add 225 ml Gelatin Phosphate Salt Broth. Incubate at 35 ± 2°C for 6 to 8 hours. Transfer a loopful from surface growth of each broth culture to two plated media, i.e. TCBS Agar (DM1189) and Gelatin Phosphate Salt Agar (DM1921), and incubate at 35 ± 2°C for 18-24 hours. Subculture 3-4 colonies from each plating medium to TN Agar. Growth from TN Agar is further confirmed by inoculating Kligler Iron Agar slants (DM1078) and stabbing the butt. After incubation, *V. cholerae* cultures will have an alkaline (red) slant and an acid (yellow) butt, no gas, and no blackening (H_2S production) in the butt. Presumptive test for suspected strains of *V. cholerae* from TN Agar is carried out by using string test [(2).]

Methodology

Suspend 35 grams of powder media in 1000 ml distilled water. Shake well & heat to dissolve the medium completely. Dispense in tubes. Sterilize by autoclaving at 15 lbs pressure (121°C) for 15 minutes. Allow the tubed medium to solidify in an inclined position (long slants).

Quality Control

Physical Appearance

Cream to yellow homogeneous free flowing powder

Gelling

Firm, comparable with 1.5% Agar gel.

Colour and Clarity of prepared medium

Light yellow coloured clear to slightly opalescent gel forms in tubes as long slants.

Reaction

Reaction of 3.5% w/v aqueous solution at 25°C. pH : 7.2±0.2

pH range

7.00-7.40

Cultural Response/Characteristics

DM 1950: Cultural characteristics observed after an incubation at 35-37°C for 18-24 hours.

Organism	Inoculum(CFU)	Growth
Vibrio cholerae ATCC 15748	50-100	good-luxuriant
Vibrio parahaemolyticus ATCC 17802	50-100	good-luxuriant

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: 2-8^0 in sealable plastic bags for 2-5 days.

Further Reading

1. Downes F. P. and Ito K., (Eds.), 2001, Compendium of Methods for the Microbiological Examination of Foods, 4th Ed., APHA, Washington, D.C.
2. Smith H. L. Jr., 1970, Bull. World Health Organization, 42:817.

Disclaimer :

- User must ensure suitability of the product(s) in their application prior to use.
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