

Technical Information

Dextrose Tryptone MiVeg Agar

Product Code : VM1092

Application:- Dextrose Tryptone MiVeg Agar is used for the detection and enumeration of mesophilic and thermophilic aerobic microorganisms in foods.

Composition

Ingredients	Gms / Litre
MiVeg hydrolysate	10.0
Dextrose	5.00
Bromo cresol purple	0.04
Agar	15.00
Final pH (at 25°C)	6.7±0.2

** Formula adjusted, standardized to suit performance parameters.

Principle & Interpretation

Dextrose Tryptone MiVeg Agar is prepared by using MiVeg hydrolysate instead of Casein enzymic hydrolysate thereby making the media free from BSE/TSE risks. This medium is the modification of the medium evolved by Williams (1) for cultivation and enumeration of the thermophilic bacteria. Use of this medium for routine culture purpose is recommended by Cameron (2) and the Association of Official Agricultural Chemists (3). It is used for the examination of canned food, sugar and starch for thermophilic bacteria such as *Bacillus stearothermophilus* ('flat sour' spoilage bacteria) (4) and also for plate count of mesophilic or thermophilic aerobes in sweetening agents used in frozen desserts (5) and for counts of aerobic microorganisms in liquid sugar. This medium contains MiVeg hydrolysate which supplies nutrients to the organisms. Dextrose is the energy source. Bromo cresol purple in the medium serves as an pH indicator. Acid producing organisms produce yellow coloured colony. The plates should be incubated at 55°C for 48 hours in a humid incubator. It is useful for enumeration of mesophilic organisms, thermophiles in cereals and cereal products, dehydrated fruits vegetables and spices (6)

Methodology

Suspend 30 grams of powder media in 1000 ml distilled water. Mix thoroughly. Heat to boiling to dissolve the medium completely. Sterilize by autoclaving at 15 lbs pressure (121°C) for 15 minutes.

Quality Control

Physical Appearance

Yellow coloured, may have slightly greenish tinge, homogeneous, free flowing powder.

Gelling

Firm, comparable with 1.5% Agar gel

Colour and Clarity of prepared medium

Purple coloured, slightly opalescent gel forms in petri plates.

Reaction

Reaction of 3.0% w/v aqueous solution pH: 6.7 ±0.2 at 25°C

pH range

6.5-6.9

Cultural Response/Characteristics

Cultural characteristics observed after an incubation at 55°C for 48 hours.



Dehydrated Culture Media
Bases / Media Supplements

Organisms (ATCC)	Inoculum (CFU)	Growth	Recovery	Colour of colony on the Agar Media
<i>Bacillus coagulans</i> (8038)	10^2-10^3	Good-luxuriant	>70%	yellow
<i>Bacillus stearothermophilus</i> (7953)	10^2-10^3	Good-luxuriant	>70%	yellow
<i>Bacillus brevis</i> (8246)	10^2-10^3	Good-luxuriant*	>70%	yellow

Key : * = with or without dextrose fermentation

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. Williams O.B., 1936, Food Res., 1:217.
2. Cameron E.J., 1936, J. Assoc. Official Agr. Chem., 19:433.
3. Association of Official Agriculture Chemists, 1945, Official and Tentative Methods of Analysis, Washington.
4. Tanner F.W., 1944., The Microbiology of Foods, 2nd ed., Garrard Press, Champaign, P.762 and 1127.
5. Standard Methods for the Examination of Dairy Products. 17th Edition, 2004 Edited by H. Michael Wehr and Joseph H. Frank.
6. Downes FP and Ito K (Eds.), 2001, Compendium of Methods For The Microbiological Examination of Foods, 4th ed., APHA, Washington, D.C.

Disclaimer :

- User must ensure suitability of the product(s) in their application prior to use.
- The product conform solely to the technical information provided in this booklet and to the best of knowledge research and development work carried at CDH is true and accurate
- **Central Drug House Pvt. Ltd.** reserves the right to make changes to specifications and information related to the products at any time.
- Products are not intended for human or animal diagnostic or therapeutic use but for laboratory, research or further manufacturing of diagnostic reagents extra.
- Statements contained herein should not be considered as a warranty of any kind, expressed or implied, and no liability is accepted for infringement of any patents. Do not use the products if it fails to meet specifications for identity and performance parameters.

