

Technical Information

Tryptone Soya Agar Plate with 1% glycerol

Product Code: PM 1290AGT

Application: Used for cultivation of a wide variety of microorganisms

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Composition				
Ingredients	Gms / Litre			
Pancreatic digest of casein	15.000			
Papaic digest of soyabean meal	5.000			
Sodium chloride	5.000			
Agar	15.000			
Glycerol	10.000			
**Formula adjusted, standardized to suit performance	parameters			

Principle & Interpretation

Soyabean Casein Digest Agar is a widely used medium, which supports the growth of wide variety of organisms even that of fastidious ones such as Neisseria, Listeria, and Brucella etc. The medium with addition of blood provides perfectly defined haemolysis zones, while preventing the lysis of erythrocytes due to its sodium chloride content. It has been frequently used in the health industry to produce antigens toxins etc. Its simple and inhibitor-free composition makes it suitable

for the detection of antimicrobial agents in the food and other products. Tryptone Soya Agar is recommended by various pharmacopoeias as sterility testing medium (1, 2). Tryptone Soya Agar conforms as per USP (1) and is used in microbial limit test and antimicrobial preservative effective test. Gunn et al (3) used this medium for the growth of fastidious organisms and study of haemolytic reaction after addition of 5% v/v blood. The combination of Pancreatic digest of casein and papaic digest of soyabean meal makes this media nutritious by providing amino acids and long chain peptides for the growth of microorganisms. Sodium chloride maintains the osmotic balance. Soyabean Casein Digest Agar does not contains X and V growth factors. It can be conveniently used in determining the requirements of these growth factors by isolates of Haemophilus by the addition of X-factor (DD020) V-factor (DD021), and X+V factor discs (DD022) factor to inoculated TSA plates (4).

Warning and Precautions

Read the label before opening the container. Wear protective gloves/protective clothing/eye protection/face protection. Follow good microbiological lab practices while handling specimens and culture. Standard precautions as per established guidelines should be followed while handling clinical specimens. Safety guidelines may be referred in individual safety data sheets.

Limitations

- Individual organisms differ in their growth requirement and may show variable growth patterns on the medium.
- Each lot of the medium has been tested for the organisms specified on the COA. It is recommended to users to validate the medium for any specific microorganism other than mentioned in the COA based on the user's unique requirement.
- It is recommended to store the plates at 24-30°C to avoid minimum condensation .



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 .

Methodology

Either streak, inoculate or surface spread the test inoculum (50-100 CFU) aseptically on the plate

Quality Control

Appearance

Sterile Soyabean Casein Digest Agar in 90mm plates

Light yellow coloured medium

Quantity of Medium

30ml of medium in 90mm plates

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7.10- 7.50

Dose of irradiation

10.00- 25.00

Cultural Response

Recovery rate is considered 100% for bacteria growth on Blood Agar and fungus growth on Sabouraud Dextrose Agar.

Growth promoting properties

Growth of microorganism comparable to that previously obtained with previously tested and approved lot of medium occurs at the specified temperature for not more than the shortest period of time specified inoculating <=100 cfu (at 30-35°C for 18 hours).

Sterility Test

Passes release criteria.

Cultural Response

PM1290AGT: Growth Promotion was carried out in accordance with the harmonized method and growth was observed afteran incubation at 30-35°C for 18-24 hours.

Oragnism	Inoculum	Growth (CFU)	Observed Lot value (CFU)	Recovery	Incubation temperature	Incubation period	
Growth promoting							
Bacillus subtilis	50-100	luxuriant	35-100	>=70 %	30 -35 °C	18 -24 hrs	
ATCC 6633							
Staphylococcus aureus	50-100	luxuriant	35-100	>=70 %	30 -35 ℃	18 -24 hrs	
ATCC 25923							
Staphylococcus aureus	50-100	luxuriant	35-100	>=70 %	30 -35 °C	18 -24 hrs	
ATCC 6538							
Escherichia coli	50-100	luxuriant	35-100	>=70 %	30 -35 °C	18 -24 hrs	
ATCC25922							
Escherichia coli	50-100	luxuriant	35-100	>=70 %	30 -35 °C	18 -24 hrs	
ATCC 8739							
Escherichia coli	50-100	luxuriant	35-100	>=70 %	30 -35 °C	18 -24 hrs	
NCTC 9002							
Pseudomonas	50-100	luxuriant	35-100	>=70 %	30 -35 °C	18 -24 hrs	
Aeruginosa ATCC 27853							
Pseudomonas	50-100	luxuriant	35-100	>=70 %	30 -35 °C	18 -24 hrs	
Aeruginosa ATCC 9027							
Salmonella Abony	50-100	luxuriant	35-100	>=70 %	30 -35 ℃	18 -24 hrs	
NCTC6017							
Micrococcus luteus	50-100	luxuriant	35-100	>=70 %	30 -35 °C	18 -24 hrs	
ATCC9341							
Streptococcus	50-100	luxuriant	35-100	>=70 %	30 -35 °C	18 -24 hrs	
Pneumoniae ATCC 6305							



50-100	luxuriant	35-100	>=70 %	30 -35 °C	18 -24 hrs
50-100	luxuriant	35-100	>=70 %	30 -35 °C	<=5 d
50-100	luxuriant	35-100	>=70 %	30 -35 °C	<=5 d
50-100	good-luxuriant	25-70	50-70 %	30 -35 °C	<=5 d
50-100	luxuriant	35-100	>=70 %	20 -25 °C	<=5 d
	50-100 50-100 50-100	50-100 luxuriant 50-100 luxuriant 50-100 good-luxuriant	50-100 luxuriant 35-100 50-100 luxuriant 35-100 50-100 good-luxuriant 25-70	50-100 luxuriant 35-100 >=70 % 50-100 luxuriant 35-100 >=70 % 50-100 good-luxuriant 25-70 50-70 %	50-100 luxuriant 35-100 >=70 % 30 -35 °C 50-100 luxuriant 35-100 >=70 % 30 -35 °C 50-100 good-luxuriant 25-70 50-70 % 30 -35 °C

Storage and Shelf Life

Store at 15-25°C. Use before expiry date on the label.

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 (2,4).

Further Reading

- The United States Pharmacopoeia / National Formulary, 2008, USP 31, The United States Pharmacopoeial Convention Inc., Rockville,
- Indian Pharmacopoeia, 2007, Govt. of India, Ministry of Health and Family Welfare, New Delhi, India.
- Gunn B. A., Ohashi D K., Gaydos C. A., Holt E. S., 1977, J. Clin. Microbiol., 5(6): 650.
- Forbes B. A., Sahm A. S. and Weissfeld D. F., 1998, Bailey and Scotts Diagnostic Microbiology, 10th Ed., Mosby Inc.St. Louis, Mo

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.
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- Do not use the products if it fails to meet specifications for identity and performens parameters.