

Technical Information

Calcium M Pretein Agar

Product Code: DM 2309

Application: - Calcium Caseinate Agar is used for the detection and enumeration of proteolytic microorganisms in foodstuffs and other materials.

Composition**

Ingredients	Gms / Litre					
Peptic digest of animal tissue	4.000					
Meat extract	2.000					
Casein enzymic hydrolysate	2.000					
Calcium caseinate	3.500					
Calcium chloride.2H ₂ O	0.200					
Tri-potassium citrate, H₂0	0.350					
Disodium hydrogen phosphate	0.105					
Potassium dihydrogen phosphate	0.035					
Sodium chloride	5.000					
Agar	13.000					
Final pH (at 25°C)	7.0±0.2					
**Formula adjusted, standardized to suit performance parameters						

Principle & Interpretation

Protein hydrolysis by microorganisms in foods may produce a variety of odour and flavour defects. On the other hand, microbial proteolytic activity may be desirable in certain foods such as in the ripening of cheese, where it contributes to the development of flavour, body and texture. In some foods the level of proteolytic microorganisms may be of value to predict refrigerated storage life and to assess processing methods (1, 2). Calcium Caseinate Agar is a modification of the original formulation of Frazier and Rupp (3) and is used for the detection and enumeration of proteolytic microorganisms in foodstuffs and other materials. Casein enzymic hydrolysate supplies nitrogenous, carbonaceous nutrients along with vitamins and amino acids. Phosphates are added to buffer the medium. Sodium chloride helps to maintain the osmotic equilibrium. Casein in the medium is degraded by the proteolytic organisms. This results in formation of clear zones around the proteolytic colonies, in the otherwise opaque medium.

The test sample can be directly surface inoculated or the inoculation can be carried out by the pour plate technique. After an incubation for 24-48 hours, proteolytic organisms, if present will form clear zones on the medium. For better visualization of the zones, the plates can be flooded with 5-10% acetic acid.

Methodology

Suspend 30.19 grams of dehydrated media in 1000 ml distilled water. Mix thoroughly & heat gently while frequently shaking until the suspension boils. Boil for 10 minutes. Sterilize by autoclaving at 15 lbs pressure (121°C) for 15 minutes. Mix thoroughly & while pouring into sterile Petri plates to suspend the precipitate. If desired, to increase turbidity, add 5-10 grams of skim milk powder before heating.

Quality Control

Appearance

Cream to yellow homogeneous free flowing powder

Firm, comparable with 1.3% Agar gel.





Colour and Clarity

Whitish coloured, turbid gel forms in Petri plates

Reaction

Reaction of 3.02 % w/v aqueous solution at 25°C. pH: 7.0±0.2

Ph Range

6.80-7.20

Cultural Response

DM 2309: Cultural characteristics observed after an incubation at 35-37°C for 2 4 -48 hours.

Organism	Inoculum	Growth	Recovery	Proteolytic	
	(CFU)			activity	
Bacillus cereus ATCC 14579	50-100	good-luxuriant	>=70%	positive, clear zone surrounding colonies	
Escherichia coli ATCC 25922	50-100	good-luxuriant	>=70%	negative, no clear zone surrounding colonies	
Proteus vulgaris ATCC 13315		good-luxuriant	>=70%	negative, no clear zone surrounding colonies	
Pseudomonas aeruginosa ATCC 27853	50-100	good-luxuriant	>=70%	positive, clear zone surrounding colonies	

Storage and Shelf Life

Dried Media: Store below 30°C in tightly container and prepared medium at 2-8°C. Use before expiry period on the label. **Prepared Media**: 2-8° in sealable plastic bags for 2-5 days.

Further Reading

- 1. Chai T., Chen C., Rosen A. and Levin R. E., 1968, Appl. Microbiol., 16: 1738.
- 2. Martely F. G., Jayashankar S. R. and Lawrence, R. C., 1970, J. Appl. Bacteriol., 33: 363.
- 3. Frazier W. C. and Rupp P., 1928, J. Bacteriol., 16, 57-63

Disclaimer:

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