

Technical Information

Acetamide Agar, Modified (Twin Pack)

Product Code: DM 2867

Application: - Acetamide Agar, Modified is used for confirmation of Pseudomonas aeruginosa in water samples.

Composition**

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Ingredients	Gms / Litre		
Part A	-		
Acetamide	3.000		
Part B	-		
Sodium chloride	5.000		
Yeast extract	0.500		
Monopotassium phosphate	1.000		
Phenol red	0.030		
Dextrose	0.200		
Agar	15.000		
Final pH (at 25°C)	6.3±0.2		
**Formula adjusted, standardized to suit performance parameters			

Principle & Interpretation

Acetamide Agar, Modified is formulated as per recommendation of Standard Methods for Examination of Water and Wastewater (1). Gilardi and others showed that a wide variety of non-fermenting organisms were capable of utilizing acetamide by using basal mineral media (2, 3). However very few organisms growing in the medium metabolize acetamide by the process of deamination (acrylamidase activity) (4, 5). This unique ability is useful in identification of various non-fermenting gram- negative organisms (6, 7, 8). This ability is shown by Pseudomonas aeruginosa, Pseudomonas aciovorans Group III (Achromobacter xylosoxidans) and Alcaligenes odorans (9). Acetamide deamination leads to the liberation of ammonia, which thereby increases the pH of the medium, leading to a subsequent colour change of the phenol red indicator from yellow orange to purplish red. Some strains require upto seven days to exhibit a positive reaction as they deaminate acrylamide slowly. However, only about 40% of apyocyanogenic strains of *Pseudomonas aeruginosa* exhibit a positive reaction. It is therefore, not advisable to rely on this test as the only criterion for identification.

The medium contains inorganic salts and acetamide a sole as a source of carbon and nitrogen. Sodium chloride maintains the osmotic equilibrium. Phenol red is the pH indicator.

Methodology

Suspend 21.73 grams of dehydrated culture media part B in 1000 ml distilled water. Add 3.0 grams of Part A. Shake well and heat to boiling to dissolve the medium completely. Dispense in tubes or as desired. Sterilize by autoclaving at 15 lbs pressure (121°C) for 15 minutes. Cool the tubes in a slanted position.





Quality Control

Appearance

Part A: Colourless deliquescent crystals Part B: Light yellow to brick red homogeneous free flowing powder

Gelling

Firm, comparable with 1.5% Agar gel

Colour and Clarity

Orange coloured clear to slightly opalescent gel forms in tubes as slants.

Reaction

Reaction of the medium (Mixture of 0.3% w/v Part A and 2.17% Part B) at 25°C. pH : 6.30±0.2

pH Range

6.10-6.50

Cultural Response

DM2867: Cultural characteristics observed after an incubation at 35-37°C for 4-7 days.

Organism	Inoculum (CFU)	Growth	Deamination
Cultural Response Stenotrophomonas maltophila ATCC 13637	50-100	good-luxuriant	negative reaction ,no purplish red colour within 7 days
Pseudomonas aeruginosa ATCC 27853	50-100	good-luxuriant	positive reaction ,no purplish red colour within 7 days

Storage and Shelf Life

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

Further Reading

- 1. Eaton A. D., Clesceri L. S., and Greenberg A. W., (Eds.), 1995, Standard Methods for the Examination of Water and Waste water, 21st Ed., APHA, Washington, D.C.
- 2. Gilardi, 1974, Antonie Van Leeuwenhoek, J. Microbiology Serol., 39:229.
- 3. Stainier, Palleroni and Doudoroff, 1966, J. Gen Microbiol., 43:159.
- 4. Pickett M. J. and Pedersen M.M., 1970, Can. J. Microbiol., 16:351.
- 5. Pickett M. J. and Pedersen M.M., 1970, Can. J. Microbiol., 16:401.
- 6. Hedberg, 1969, Appl. Microbiol., 17: 481
- 7. Smith and Dayton, 1972, Appl. Microbiol., 24: 143
- 8. Buhlmann, Vischer and Bruhin, 1961, J. Bacteriol., 82:787
- 9. Oberhofer and Rowen, 1974, Appl. Microbiol., 28:720.

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