

Dehydrated Culture Media Bases / Media Supplements

# **Technical Information**

# AC Agar

# Product Code: DM 1337

Application: AC Agar is recommended for cultivation of wide variety of microorganisms particularly for sterility testing.

Composition**			
Ingredients	Gms / Litre		
Proteose peptone	20.000		
Beef extract	3.000		
Yeast extract	3.000		
Malt extract	3.000		
Dextrose	5.000		
Ascorbic acid	0.200		
Agar	1.000		
Final pH (at 25°C)	7.2±0.2		
**Formula adjusted, standardized to suit perf	ormance parameters		

### **Principle & Interpretation**

AC Medium support an early and luxuriant growth of aerobic, anaerobic and microaerophilic microorganisms. Many pathogenic and saprophytic aerobes can also be isolated using AC Medium <sup>(1)</sup>. This medium can also be used for sterility testing of solutions and biological products not containing mercurial preservatives. AC Agar does not exhibit the toxicity as shown by some media containing sodium thioglycollate for some organisms <sup>(2,3)</sup> Earlier studies performed have reported the usefulness of this medium for the cultivation of a wide variety of organisms <sup>(4, 5)</sup>. Kolb Schneither <sup>(6)</sup> used AC Agar to test the viability of *Bacillus anthracis* after exposure to methyl bromide to test the efficiency of methyl bromide as a germicidal and sporicidal agent. In addition to being a source of vitamins and cofactors. proteose peptone, beef extract, yeast extract and malt extract also serve as the carbon and nitrogen sources. Dextrose provider the fermentable carbohydrate source of energy. Ascorbic acid helps to improve the clarity of the medium.

# Methodology

Suspend 35.2 grams of powder media in 1000 ml of distilled water. Shake well and heat to dissolve the medium completely. Distribute in tubes or bottles to give the desired depth and sterilize by autoclaving at 15 lbs pressure (121°C) for 15 minutes. If the medium is not used on same day, it is advisable to drive off dissolved gases by boiling or steaming in the autoclave and cool without agitation before use.

### **Quality Control**

 Physical Appearance

 Cream to yellow homogeneous free flowing powder

 Gelling

 Semisolid, comparable with 0.1 % Agar gel.

 Colour and Clarity of prepared medium

 Medium amber coloured clear to slightly opalescent solution

 Reaction

 Reaction of 3.52% w/v aqueous solution at 25°C. pH : 7.2±0.2

 pH Range:- 7.00-7.40

 Cultural Response/Characteristics

DM 1357: Cultural characteristics observed after an incubation at 35-37°C for 18-24 hours (Clostridium species incubated anaerobically).





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Organism	lnoculum (CFU)	Growth
Clostridium perfringens ATCC 12919	50-100	luxuriant
Escherichia coli ATCC25922	50-100	Luxuriant
Neisseria meningitides ATCC 13090	50-100	Luxuriant
Staphylococcus aureus ATCC 25923	50-100	Luxuriant
Streptococcus mitis ATCC9811	50-100	Luxuriant
Streptococcus pneumoniae ATCC 6303	50-100	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° in sealable plastic bags for 2-5 days.

### **Further Reading**

1. MacFaddin J. F., 1985, Media for Isolation-Cultivation-Identification-Maintenance of Medical Bacteria, Vol. I. Williams & Wilkins, Baltimore, Md.

- 2. Christensen, 1944, Paper read at New York Meeting, American Public Health Association.
- 3. Malin and Finn, 1951, J. Bacteriol., 62:349.
- 4. Reed and Orr, 1943, J. Bacteriol., 45:309.
- 5. Schneiter, Dunn and Caminita, 1945, Public Health Rep., 60:789.
- 6. Kolb and Schneiter, 1950, J. Bacteriol., 59:401.

#### **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 a at **CDH** is true and accurate
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