

# **Technical Information**

### **BYE Agar**

### Product Code: DM 1470

**Application:** BYE Agar is a simplified medium developed for the cultivation of *Mycoplasma*, or Pleuropneumonia like organisms and L-forms of bacteria.

## Composition\*\*

Ingredients	Gms / Litre	
Proteose peptone	10.000	
Calf brain, infusion from	200.000	
Beef heart, infusion from	250.000	
Dextrose	2.000	
Sodium chloride	5.000	
Disodium phosphate	2.500	
Yeast extract	2.000	
Agar	13.000	
Final pH ( at 25°C)	7.9±0.2	
**Formula adjusted, standardized to suit performance pa	rameters	

### **Principle & Interpretation**

Mycoplasmas the smallest free-living microorganisms were designated pleuropneumonia like organism (PPLO), because of similarities to Mycoplasma mycoides (subsp. mycoides), the causative agent of bovine pleuropneumonia <sup>(1, 2)</sup>. BYE media are simple media developed to study cultivation distribution, habitat and possible pathogenesis of Mycoplasma. Pleuropneumonia like organisms and L-forms of bacteria by Barile, Yaguchi and Eveland <sup>(3)</sup>. These media can be used for isolation of L-forms of Corynebacterium, Neisseria, and Streptococcus PPLOs from urethritis, penile ulcerations and cervical specimens and are also used for detecting PPLO contamination of tissue culture cell-lines and membrane filter work <sup>(4, 5)</sup>.

BYE Agar contains brain and heart infusion along with yeast extract, which provide carbon, nitrogen, vitamins and other growth factors required for the metabolism of Mycoplasma - Pleuropneumonia like organisms. Inoculations are made in duplicates. One set is incubated aerobically while the other anaerobically for 48 hours or more. Usually growth occurs within 3-5 days; however, negative results are reported after 10 days. Anaerobic conditions are most important for the first 3 days while secondary transfers can be incubated aerobically.

## Methodology

Suspend 52 grams of powder media in 850 ml distilled water. Shake well & heat the medium completely. Sterilize by autoclaving at 15 lbs pressure (12 1°C) for 15 minutes. Cool to 50°C and aseptically add 150 ml of sterile human or animal blood or serum. Mix gently and pour into sterile Petri plates.





## **Quality Control**

#### Physical Appearance

Cream to yellow homogeneous free flowing powder

#### Gelling

Firm, comparable with 1.3% Agar gel.

#### Colour and Clarity of prepared medium

Yellow coloured, clear to slightly opalescent gel forms in Petri plates

#### Reaction

Reaction of 5.2% w/v aqueous solution at 25°C. pH: 7.9±0.2

pH range 7.70-8.10

#### Cultural Response/Characteristics

DM1470: Cultural characteristics observed with added serum under humidified anaerobic conditions, after an incubation at 35-37°C for 5-10 days.

Organism	Growth
Mycoplasma bovis ATCC 25523	good-luxuriant
Mycoplasma gallinarium ATCC 19708	good-luxuriant
Mycoplasma pneumonia ATCC 15531	good-luxuriant
Streptococcus pneumonia ATCC 6303	good-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. Murray P.R., Baron E. J., Pfaller M.A., Tenover F.C., Yolken R.H.(Eds.), 1995, Manual of Clinical Microbiology, 6th Ed., ASM Press.
- 2. Collee J.G, Fraser A.G., Marmion B.P., Simmons. A (Eds.), 1996, Mackie and McCartney Practical Medical Microbiology, 14th Ed, Churchill Livingstone.
- 3. Barile, Yaguchi, Eveland, 1958, Am. J. Clin. Path. 30:171.
- 4. Barile, 1962, National Cancer Institute Monograph, No.7: 5.5. Barile, 1962, J. Bacteriol., 83:430.

### Disclaimer:

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