

Technical Information

EMB Agar Base

Product Code: DM 1301

Application: - EMB Agar Base is a basal medium to which different carbohydrates and other test substances may be added for differentiation and study of various enteric bacteria.

Composition**

Ingredients	Gms / Litre
Peptic digest of animal tissue	10.000
Dipotassium phosphate	2.000
Eosin - Y	0.400
Methylene blue	0.065
Agar	15.000
Final pH (at 25°C)	7.3±0.2

**Formula adjusted, standardized to suit performance parameters

Principle & Interpretation

Levine^(1, 2) formulated Levine EMB Agar used for the differentiation of *Escherichia coli* and *Enterobacteriaceae* and for the detection, enumeration and differentiation of members of the coliform group and also for the rapid identification of *Candida albicans* by American Public Health Association⁽³⁻⁵⁾. Some gram-positive bacteria such as faecal Streptococci, yeasts grow on this medium and form pinpoint colonies. EMB Agar Base is a modification of EMB Agar, Levine without lactose. This facilitates the use of the medium as a basal agar to which desired carbohydrates are be added to differentiate between various enteric bacteria.

Eosin-Y and methylene blue make the medium slightly selective and inhibit certain gram-positive bacteria. These dyes also differentiate between lactose fermenters and nonfermenters. The ratio of eosin-methylene blue is adjusted to approximately 6:1. Coliforms produce purplish black colonies due to uptake of methylene blue-eosin dye complex, when the pH drops. Non-fermenters probably raise the pH of surrounding medium by oxidative de-amination of protein, which solubilizes the methylene blue-eosin complex resulting in formation of colourless colonies⁽⁶⁾.

Peptic digest of animal tissue serves as source of carbon, nitrogen, and other essential growth nutrients. Eosin-Y and methylene blue serve as differential indicators. Phosphate buffers the medium.

The test sample can be directly streaked on the medium plates. Inoculated plates should be incubated, protected from light. However standard procedures should be followed to obtain isolated colonies. A non-selective medium should be inoculated in conjunction with EMB Agar. Confirmatory tests should be further carried out for identification of isolated colonies.

Methodology

Suspend 27.5 grams of powder media in 1000 ml distilled water. Shake well & heat to dissolve the medium completely. Add desired carbohydrate in desired concentration before sterilization. Sterilize by autoclaving at 15 lbs pressure (121°C) for 15 minutes. AVOID OVERHEATING. Cool to 50°C and shake the medium in order to oxidize the methylene blue (i.e. restore its blue colour) and to suspend the precipitate, which is an essential part of the medium. Precaution: Store the medium away from light to avoid photooxidation.

Quality Control

Physical Appearance

Light pink to purple homogeneous free flowing powder

Gelling

Firm, comparable with 1.5% Agar gel

Colour and Clarity of prepared medium

Reddish purple coloured, opalescent gel with greenish cast and finely dispersed precipitate forms in Petri plates

Reaction

Reaction of 2.75% w/v aqueous solution at 25°C. pH : 7.3±0.2

pH range: 7.1-9.5

Cultural Response/ characteristics

DM 1301: Cultural characteristics observed with added carbohydrate after an incubation at 35-37°C for 18-24 hours (Fungal cultures incubated at 25-30°C for 24-48 hours).

Organism	Inoculum (CFU)	Growth	Recovery	Colour of colony
<i>Escherichia coli</i> ATCC 8739				
<i>Candida albicans</i> ATCC 10231	50-100	luxuriant (incubated in 10% CO ₂)	≥50%	colourless
<i>Escherichia coli</i> ATCC 25922	50-100	luxuriant	≥50%	blue-black with metallic sheen
<i>Enterobacter aerogenes</i> ATCC 13048	50-100	good	40-50%	pink-red
<i>Enterococcus faecalis</i> ATCC 29212	50-100	non-poor	≤10%	colourless
<i>Pseudomonas aeruginosa</i> ATCC 27853	50-100	luxuriant	≥50%	colourless
<i>Salmonella Typhimurium</i> ATCC 14028	50-100	luxuriant	≥50%	colourless
<i>Saccharomyces cerevisiae</i> ATCC 9763	50-100	none-poor	≤10%	cream
<i>Staphylococcus aureus</i> ATCC 25923	50-100	none-poor	≤10%	colourless

***Pseudomonas aeruginosa* ATCC 9027**

***Staphylococcus aureus* ATCC 6538**

***Escherichia coli* ATCC 25922**

***Escherichia coli* NCTC 9002**

***Pseudomonas aeruginosa* ATCC 27853**

***Enterobacter aerogenes* ATCC 13048**

***Proteus mirabilis* ATCC 25933**

***Salmonella Enteritidis* ATCC 13076**

***Shigella boydii* ATCC 12030**

***Staphylococcus aureus* ATCC 25923**

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.



Dehydrated Culture Media
Bases / Media Supplements

Further Reading

1. Levine M., 1918, J. Infect. Dis., 23:43.
2. Levine M., 1921, Bull. 62, Iowa State College Engr. Exp. Station.
3. Greenberg A. E., Trussell R. R. and Clesceri L. S. (Eds.), 1998, Standard Met for the Examination of Water and Wastewater, 20th ed., APHA, Washington, D.C.
4. Marshall R. (Ed.), 1992, Standard Methods for the Examination of Dairy,, Products, 16th ed., APHA Inc., New York.
5. Downes F. P and Ito K. (Ed.), 2001, Compendium of Methods for the Microbiological Examination of Foods, 4th ed., APHA, Washington, D.C.
6. Howard B. J., 1994, Clinical and Pathogenic Microbiology, 2nd Ed., Mosby Year Book, Inc

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