

70186 EMB Agar (Eosin Methylene Blue Agar)

EMB Agar is a very versatile solid medium. Originally developed by Levine for the differentiation of *Escherichia coli* and *Aerobacter aerogenes*, it turned out to be effective for the rapid identification of *Candida albicans* and was found useful for the identification of coagulase-positive Staphylococci.

Composition:

Ingredients	Grams/Litre
Peptone	10.0
Lactose	10.0
Dipotassiummonohydrogenphosphate	2.0
Methylene Blue	0.065
Eosine Y	0.4
Agar	15.0
Final pH 7.1 +/- 0.2 at 25°C	

Store prepared media below 8°C and protected from direct light. Store dehydrated powder in a dry place, in tightly-sealed containers at 2-25°C.

Appearance: Faintly violet to pink, homogeneous, hygroscopic powder.

Gelling: Firm

Color and Clarity: Deep red-brown, clear to slightly turbid

Directions:

Suspend 37.5 g in 1 litre of distilled water. Heat to dissolve completely, sterilize by autoclaving at 121°C for 15 minutes. Cool to 60°C and shake the medium in order to oxidize the methylene blue and to suspend the precipitate.

Principle and Interpretation:

The presence of the colorants Eosine Y and Methylene Blue inhibits the growth of most of the common accompanying Gram-positive microorganisms. Levine described this classical method to identify *E.coli* from other coliforms as *Aeorobacter aerogenes* (2). Lactose is added as distinctive carbon and energy source. In combination with the added dyes it allows to distinguish between lactose-positive and lactose-negative organisms. Lactose positive cultures are generally dark violet (*Enterobacter, Klebsiella, E.coli*), while lactose negative organisms (*Salmonella, Shigella*) have only peptone as energy source are colourless.

Some gram-positive bacteria, such as fecal streptococci, staphylococci will may grow on this medium as inhibited small colonies. A number of nonpathogenic, lactose-nonfermenting gram-negative bacteria will grow on this medium and must be distinguished from the pathogenic strains by additional biochemical tests.



Cultural characteristics after 24-48 hours at 35°C.

Organisms (ATCC)	Growth	Appearance of Colony
Staphylococcus aureus (25923)	-/+	colorless
Escherichia coli (11775)	+++	Ø 2-3mm, dark violet cultures with black center and green metallic shine
Escherichia coli (25922)	+++	Ø 2-3mm, dark violet cultures with black center and green metallic shine
Enterobacter aerogenes (13048)	+++	large, pink with blue center
Klebsiella pneumoniae (33495)	+++	large, pink with blue center
Salmonella abony (NCTC 6017)	+++	large, colorless
Shigella flexneri (12022)	+++	large, colorless
Enterococcus faecalis (29212)	+	punctiform, dark violet cultures
Pseudomonas aeruginosa (27853)	+++	irregular, colorless
Candida albicans (10231)*	++	colorless, cotton like (other yeasts look classical)

^{*} key: incubated in 10% carbodioxide

References:

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- 4. APHA-AWWA-WPCF, 1995, Standard Methods for the Examination of Water and Wastewater, 19th ed. APHA Washington D.C.
- 5. Y. Henis et al., 1989, Microb. Ecol. 17, 171
- 6. Weld, Julia, 1953, Candida albicans: Rapid Identification in cultures made directly from human materials, Arch. Dermat. Syph. 67(5), 473-478
- 7. Windle Taylor, E. 1958, The Examination of Water and Water Supplies, Churchill Ltd. 7th ed. London
- 8. U.S. Pharmakopoeia, Microbial Limit Test, The United States Pharmacopeial Convention, Rockville, Md. (2002)

Precautions and Disclaimer

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