

Reference: 01-068

Scharlau Microbiology - Technical data sheet

Product:

**EOSIN METHYLENE BLUE AGAR (EMB AGAR)** 

### Also known as

EMB Agar

## **Specification**

Selective differential medium for the isolation and enumeration of coliforms according to ISO 21150 standard and USP.

## Formula \* in q/L

10,000
10,000
2,000
0,400
0,065
15,000

Final pH 7.1 ±0,2 at 25 °C

#### **Directions**

Add 37,5 g to 1 L of distilled water. Bring to the boil and distribute in suitable containers. Sterilize in the autoclave at 121°C for 15 minutes.

### **Description**

A very versatile medium originally developed for the differentiation of *E.coli* and *Enterobacter aerogenes*. It has also proved very effective in the rapid identification of *Candida albicans* and demostrates a high correlation with the coagulase test for staphylococci.

It has been repeatedly recommended for the detection, enumeration and differentiation of members of the coliform group of bacteria.

## **Technique**

The Weld method for the identification of Candida albicans uses this medium with chlortetracycline (100 mg/L) in a 10% CO2 envferment. The method's efficacy has been tested with a variety of samples, such as sputum, oral secretions, faeces, nails and vaginal secretions, all of which provide definitive results within 24-48 hours. Staphylococci are also easily identified, particularly coagulase-positive strains. These have a very characteristic appearance: small colourless colonies with a central red nucleus. The medium's prevailing application is in the differentiation of E. coli and E. aerogenes.

The medium should be sterilized once distributed into tubes containing 20 mL of product each, and then refrigerated. Melt in a boiling water bath before use and stir until it acquires a dark purple colour. Pour a tube into each sterile plate and allow it to solidify. It is advisable to dry the medium's surface before use, leaving the plate open but inverted.

For each doubtful lactose broth tube, inoculate one plate by streaking, and incubate for 24 hours à 35±2°C.

- Escherichia coli and Citrobacter form flat colonies of 2-3 mm in diameter and are dark violet in colour with a black centre which produces a distinctive green metallic sheen when light is reflected on it.
- Enterobacter and Klebsiella form convex colonies which are twice as big as the very smooth E. coli, have no metallic sheen and are pink in colour with a dark blue centre. Non-lactose fermenting organisms produce colourless colonies.
- Candida albicans colonies incubated in a CO2 atmosphere have a very peculiar cotton-like appearance which distinguishes them from other Candida species that produce classical yeast like colonies.

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<sup>\*</sup> Adjusted and /or supplemented as required to meet performance criteria



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# **Quality control**

Incubation temperature: 30 - 35 °C Incubation time: 24 - 48 h

**Inoculum:** Streak isolation or ≥ 10<sup>3</sup> CFU (specificity) according to ISO 11133:2014/Amd 1:2018. Spiral plate Methods or Loop spreading.

## Microorganism

Salmonella abony NCTC® 6017
Escherichia coli ATCC® 11775
Escherichia coli ATCC® 25922
Escherichia coli ATCC® 8739
Salmonella typhimurium ATCC® 14028
Pseudomonas aeruginosa ATCC® 27853
Candida albicans ATCC® 10231

### Growth

Good

Good to very good Good to very good Good to very good Good to very good Good to very good Good to very good

#### Remarks

Colorless colonies w/o green metalic shine Dark violet colonies w. green metalic shine Dark violet colonies w. green metalic shine Dark violet colonies w. green metalic shine Colorless ccolonies w/o green metalic shine Colorless ccolonies w/o green metalic shine Cotton-like colonies in CO2



Escherichia coli ATCC 8739



Escherichia coli ATCC 8739 "Detail"



Salmonella typhimurium ATCC 14028

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### References

- ·CLESCERI, L.S., A.E. GREENBERG & A.D. EATON. (1998) Standard Methods for the Examination of Water and Wastewater. 20<sup>th</sup> edition. APHA-AWWA-WEF. Washington D. C.
- ·HOLT-HARRIS, J. E. y TEAGUE O.A. (1916) A New Culture Medium for the Isolation of *Bacillus typhosus* from Stools J. Infect. Dis. 18:596-600.
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- ·LEVINE, M (1918) Diferentation of *E. coli* and *A. aerogenes* on simplified Eosin-ethylene Blue Agar. J. Infect. Dis. 23:43 -47.
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- ·WELD, J. (1953) *Candida albicans*: Rapid Identification in Cultures made directly from Human materials Arch. Dermat. Syph. 67(5):473-478.
- ·WINDLE TAYLOR, E. (1958)The Examination of Water and Water Supplies. Churchill Ltd. 7th ed. Londres.
- ·US-FDA (1998) Bacteriological Analytical Manual 8<sup>th</sup> ed. Revision A. AOAC International. Gaithersburg. Md. USA: L-EMB Agar (On line BAM Media M80)
- ·USP Convention (2019) Dietary Supplement Compendium. Vol. 1 USP-NF Dietary Supplementary Monographs. <2022> Microbiological Procedures for Absence of Specified Microorganisms. Nutritional and Dietary Supplements.
- ·USP 29 NF 24 (2006) 2<sup>nd</sup> Suppl. <61> Microbial Tests. USP Con. Inc. Rockville, MD, USA
- ·USP 43 NF 38 (2019) 1st Suppl. <61> Microbial Tests. USP Con. Inc. Rockville, MD, USA

# Storage

For laboratory use only. Keep tightly closed, away from bright light, in a cool dry place (+4 °C to 30 °C).