

Lauryl Sulfate Chromogenic Broth

Cat. 1465

Enrichment medium for the simultaneous detection of total Coliforms and E. coli in water, foods and dairy products by the fluorogenic procedure.

Practical information

| Applications | Categories |
|-----------------------|------------------|
| Selective enumeration | Coliforms |
| Selective enumeration | Escherichia coli |
| Selective enrichment | Coliforms |
| Selective enrichment | Escherichia coli |
| Detection | Coliforms |
| Detection | Escherichia coli |

Industry: Water / Food / Dairy products



Principles and uses

Lauryl Sulfate Chromogenic Broth allows the detection of total Coliform and E. coli count at the same time due to the Chromogenic-Fluorogenic Mix.

The combination of chromogenic compounds within Lauryl Sulfate Broth provide a double indicator system. This medium contains a phosphate buffer to ensure the high growth of the total number of Coliforms. Lauryl sulfate inhibits gram-positive bacteria. Coliforms and E. coli contain β -galactosidase which cleaves the chromogenic substrate. The enzyme which cleaves MUG is highly specific to E. coli, making the simultaneous detection of total Coliforms and E. coli possible. IPTG stimulates the synthesis and increases the activity of β -galactosidase.

The color change from amber to blue-greenish due to the reaction of the chromogenic substrate indicates the presence of coliforms. Blue fluorescence under UV light allows the rapid detection of E. coli due to the MUG.

Tryptophane promotes the indol reaction after adding Kovac's reagent (Cat. 5205). This reactive detects the microorganism capable of cleaving the tryptophane. When E. coli is present in the medium, indol is liberated and reacts with 4-dimethylaminobenzaldehyde to form a dark red dye.

Formula in g/L

| | | | |
|-----------------------------|------|-----------------------|-----|
| Chromogenic-fluorogenic mix | 0,23 | Dipotassium phosphate | 2,7 |
| Monopotassium phosphate | 2 | Sodium chloride | 5 |
| Sodium lauryl sulfate | 0,1 | Sorbitol | 1 |
| Tryptophan | 1 | Tryptose | 5 |

Preparation

Suspend 17 grams of the medium in one liter of distilled water. Mix well and dissolve by heating with frequent agitation. Boil for one minute until complete dissolution. Dispense into appropriate containers and sterilize in the autoclave at 121 °C for 15 minutes.

Instructions for use

Inoculate and incubate at 35±2 °C during 18-24 hours. Check the tubes under UV light (366 nm). Light blue fluorescence indicates the presence of E. coli.

Quality control

| Solubility | Appearance | Color of the dehydrated medium | Color of the prepared medium | Final pH (25°C) |
|------------|-------------|--------------------------------|------------------------------|-----------------|
| w/o rests | Fine powder | Beige | Clear amber | 6,8±0,2 |

Microbiological test

Incubation conditions: (35±2 °C / 18-24 h).

| Microrganisms | Specification | Characteristic reaction |
|-----------------------------------|---------------|---|
| Shigella flexneri ATCC 12022 | Good growth | Medium color without changes, Fluorescence (-) |
| Enterobacter aerogenes ATCC 13048 | Good growth | Blue-greenish medium, Fluorescence (-), Indol (-) |
| Klebsiella pneumoniae ATCC 13883 | Good growth | Blue-greenish medium, Fluorescence (-), Indol (-) |
| Salmonella typhimurium ATCC 14028 | Good growth | Medium color without changes, Fluorescence (-) |
| Escherichia coli ATCC 25922 | Good growth | Blue-greenish medium, Fluorescence (+), Indol (+) |
| Citrobacter freundii ATCC 8090 | Good growth | Blue-greenish medium, Fluorescence (-) |
| Escherichia coli ATCC 8739 | Good growth | Blue-greenish medium, Fluorescence (+), Indol (+) |

Storage

Temp. Min.:2 °C

Temp. Max.:8 °C

Bibliography

MANAFI, M., KNEIFEL, F., a. BASCON, S.: Fluorogenic and chromogenic substrates used in bacterial diagnosis. Microbiol. Rev. 55; 335-348 (1991). OSSMER, R.: Simultaneous Detection of Total Coliforms and E. coli-Fluorocult LMX-Broth. - 15th international Symposium/FOOD MICRO 1993. The International Committee on Food Microbiology and Hygiene, Bingen/Rhine (1993).