

# **Technical Data**

# НіСтоте<sup>тм</sup> М-ТЕС Agar

# M1571

# **Intended Use**

Recommended by the U.S. Environmental Protection Agency (USEPA) for differentiation and enumeration of thermotolerant *Escherichia coli* in water by the membrane filtration technique. It can also be used to isolate *E.coli* from clinical samples.

# **Composition\*\***

Ingredients	Gms / Litre	
Proteose peptone	5.000	
Yeast extract	3.000	
Lactose	10.000	
Sodium chloride	7.500	
Dipotassium hydrogen phosphate	3.300	
Potassium dihydrogen phosphate	1.000	
Sodium lauryl sulphate	0.200	
Sodium deoxycholate	0.100	
Chromogen	0.500	
Agar	15.000	
Final pH ( at 25°C)	7.3±0.2	
**Formula adjusted standardized to suit performance parameters		

\*\*Formula adjusted, standardized to suit performance parameters

## Directions

Suspend 45.6 grams in 1000 ml distilled water. Heat to boiling to dissolve the medium completely. Sterilize by autoclaving at 15 lbs pressure (121°C) for 15 minutes. Cool to 45-50°C. Mix well and pour into sterile Petri plates.

# **Principle And Interpretation**

HiCrome<sup>TM</sup> M-TEC Agar is a chromogenic media used for detection and enumeration of thermo-tolerant *E.coli* (TEC)in water by membrane filtration (5), developed by Dufour (2). The modified medium contains the chromogen, 5-bromo-6-chloro-3-indolyl- $\beta$ -D-glucuronide that is cleaved by enzyme  $\beta$ -D-glucuronidase to yield glucuronic acid, produced by *E.coli* strains. This imparts a purple-magenta colour to the colonies of *E. coli* only. *E.coli* can be isolated from clinical samples. Proteose peptone and yeast extract provides carbon, nitrogen substances, long chain amino acids, vitamins and essential nutrients. Lactose is the fermentable carbohydrate. Sodium chloride maintains osmotic equilibrium. Potassium dihydrogen phosphate and dipotassium hydrogen phosphate provide strong buffering system to control the pH in the presence of fermentative action. Sodium lauryl sulphate and sodium deoxycholate make the medium more selective by inhibiting gram positive bacteria. Membrane filter through which water sample has been passed is aseptically placed on the medium. The plates are then incubated at 44.5 ± 0.2°C for 22 - 24 hours. Following incubation *E.coli* will form purple to magenta coloured colonies on the membrane filters.

## **Type of specimen**

Clinical samples urine, Water samples.

# **Specimen Collection and Handling:**

For clinical samples follow appropriate techniques for handling specimens as per established guidelines (3,4). For water samples, follow appropriate techniques for sample collection, processing as per guidelines and local standards (1). After use, contaminated materials must be sterilized by autoclaving before discarding.

## Warning and Precautions :

In Vitro diagnostic Use only. Read the label before opening the container. Wear protective gloves/protective clothing/eye protection/ face protection. Follow good microbiological lab practices while handling specimens and culture. Standard precautions as per established guidelines should be followed while handling clinical specimens. Safety guidelines may be referred in individual safety data sheets.

## **Limitations :**

1. Due to variable nutritional requirements, some strains show poor growth on this medium.

#### **Performance and Evaluation**

Performance of the medium is expected when used as per the direction on the label within the expiry period when stored at recommended temperature.

## **Quality Control**

#### Appearance

Cream to yellow homogeneous free flowing powder

#### Gelling

Firm, comparable with 1.5% Agar gel

#### Colour and Clarity of prepared medium

Light amber coloured, clear to slightly opalescent gel forms in Petri plates

#### Reaction

Reaction of 4.56% w/v aqueous solution at 25°C. pH : 7.3±0.2

#### pН

#### 7.10-7.50

#### **Cultural Response**

M1571: Cultural characteristics observed after an incubation at 44.5±0.2°C for 22-24 hours.

Organism	Inoculum (CFU)	Growth	Colour of Colony
Escherichia coli ATCC 25922 (00013*) Enterococcus faecalis ATCC 29212 (00087*)	50-100 $C >= 10^3$	good to luxuriant inhibited	purple/ magenta
Proteus mirabilis ATCC 25933	50-100	good	colourless-light brown
Klebsiella pneumoniae ATCC 13883 (00097*)	50-100	good	tan-light purple

Key: \*Corresponding WDCM numbers.

#### **Storage and Shelf Life**

Store dehydrated powder and prepared medium at 2-8°C in tightly closed container. Use before expiry period on the label.On opening, product should be properly stored dry, after tightly capping the bottle in order to prevent lump formation due to the hygroscopic nature of the product. Improper storage of the product may lead to lump formation. Store in dry ventilated area protected from extremes of temperature and sources of ignition Seal the container tightly after use. Use before expiry date on the label. Product performance is best if used within stated expiry period.

## Disposal

User must ensure safe disposal by autoclaving and/or incineration of used or unusable preparations of this product. Follow established laboratory procedures in disposing of infectious materials and material that comes into contact with clinical sample must be decontaminated and disposed of in accordance with current laboratory techniques (3,4).

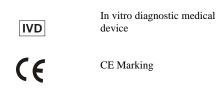
## Reference

- 1. Baird R.B., Eaton A.D., and Rice E.W., (Eds.), 2015, Standard Methods for the Examination of Water and Wastewater, 23rd ed., APHA, Washington, D.C
- 2. Dufour, Strickland and Cabelli, 1981, Appl. Environ. Microbiol. 41: 1152.

<sup>3</sup>·Isenberg, H.D. Clinical Microbiology Procedures Handbook. 2<sup>nd</sup> Edition.

- 4. Jorgensen, J.H., Pfaller, M.A., Carroll, K.C., Funke, G., Landry, M.L., Richter, S.S and Warnock., D.W. (2015) Manual of Clinical Microbiology, 11th Edition. Vol. 1.
- 5.U.S. Environmental Protection Agency, 2002, Method 1603; Publication EPA-821-R-02-023.

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2°C - 8°C



Do not use if package is damaged

Storage temperature



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