

# **Technical Data**

# **HiEncap**<sup>TM</sup> Luria Agar Base, Miller's Modification

EC1726D

HiEncap<sup>TM</sup> Luria Agar Base, Miller's modification is recommended for the cultivation and maintenance of recombinant strains of *Escherichia coli* with or without addition of glucose.

# Composition\*\*

Ingredients	<b>Gms / Litre</b>
Casein enzymic hydrolysate	10.000
Yeast extract	5.000
Sodium chloride	0.500
Agar	15.000
Final pH ( at 25°C)	7.0±0.2

<sup>\*\*</sup>Formula adjusted, standardized to suit performance parameters

#### **Directions**

Each capsule contains 15.25 gms of medium. Suspend 1 capsule in 500 ml (2 capsules in 1000 ml) distilled or purified 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. If desired add filter sterilized 20% v/v glucose solution i.e 5 ml in 500 ml or 10 ml in 1000 ml media. Mix well and pour into sterile Petri plates.

# **Principle And Interpretation**

This medium is based on original formula described by Miller for the growth and maintenance of *E.coli* strains used in molecular microbiology (1). Luria Agar Base, Miller is a nutritionally rich medium recommended for growth of pure cultures of recombinant strains. *E.coli* is grown in late log phase in LB medium. Some plasmid vectors may replicate to high copy numbers without selective amplification. Some vectors do not replicate so freely, and need to be selectively amplified. Chloramphenicol can be added to inhibit host synthesis and as a result prevent replication of the bacterial chromosome. (2)

Luria Agar Base, Miller's modification contains one tenth and one twentieth the sodium chloride level of the Lennox and Miller formulations of LB Agar respectively (1,2,3). This helps the user to select the optimal salt concentration for a specific strain. The medium may be aseptically supplemented with glucose, if desired.

Casein enzymic hydrolysate provides peptides and peptones while Vitamin B complex is provided by yeast extract. Sodium chloride provides sodium ions for membrane transport and also maintains the osmotic equilibrium of the medium. Agar acts as a solidifying agent.

#### **Quality Control**

#### **Appearance**

Gelatin capsule containing cream to yellow coloured granulated media

#### Gelling

Firm, comparable with 1.5% Agar gel

#### Colour and Clarity of prepared medium

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

#### Quantity

Each capsule contains 15.25 grams of medium sufficient for 500 ml media

#### Reaction

Reaction of 3.05% w/v aqueous solution at 25°C. pH: 7.0±0.2

#### рH

6.80-7.20

#### **Cultural Response**

Cultural characteristics observed after an incubation of 18-24 hours at 35-37°C with added 5 ml of 20% dextrose solution to 500 ml of EC1726D

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Cultural Response					
	Organism	Growth	Inoculum (CFU)	Recovery	
Cultural Response					
	Escherichia coli ATCC	luxuriant	50-100	>=70%	
	23724				
	Escherichia coli ATCC	luxuriant	50-100	>=70%	
	25922				
	Escherichia coli DH5 alpha	luxuriant	50-100	>=70%	
	MTCC 1652				

### **Storage and Shelf Life**

Store below 30°C in tightly closed container and the prepared medium at 2 - 8°C. Use before expiry date on the label.

#### Reference

- 1. Miller ,J.H. 1772. Experiments in molecular genetics. Cold spring Harbor Laboratory, Cold spring Harbor, New York.
- 2. Sambrook, J., E.F. Fritsch and T. Maniatis. 1989.Molecular cloning: A laboratory manual, 2nd ed., Cold Spring Harbor Laboartory, Cold Spring Harbor, New York.
- 3. Lennox E.S. 1955, Transduction of Linked Genetic Characters of the host by bacteriophage P1., Virology, 1:190.

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#### Disclaimer:

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