Technical Information

Luria Agar

Product Code : G1557

Recommended for routine cultivation and estimation of not particularly fastidious microorganisms.

Composition**

Ingredients	Grams/Litre
Tryptone	10.000
Yeast extract	5.000
Sodium chloride	5.000
Agar	15.000
Final pH (at 25°C)	7.0±0.2

** Formula adjusted, standardized to suit performance parameters

Methodology

Suspend 35.0 grams in 1000 ml purified/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

Luria Agar is prepared as described by Lennox (3) for cultivation and maintenance of recombinant strains of Escherichia coli. The media is generally used for molecular and genetic studies, because of its nutritive capacity and simple composition, which can be easily altered as per specific requirements. The medium is nutritionally rich for the growth of pure cultures of recombinant strains. Strains which are generally derived from Escherichia coli K12 are deficient in Vitamin B synthesis and are further modified by specific mutation to create auxotrophic strains that are unable to grow on nutritionally deficient media. Tryptone provides peptides and peptones while Vitamin B complex is provided by yeast extract. Sodium chloride provides sodium ions for the membrane transport and maintains osmotic equilibrium of the medium.

Quality control

Appearance of Powder :
Cream to yellow homogeneous free flowing powder
Gelling
Firm, comparable with 1.5% Agar gel
Colour and Clarity of the prepared medium:
Yellow to amber coloured, clear to slightly opalescent gel forms in Petri plates
Reaction:
Reaction of 3.5% w/v aqueous solution at 25°C. pH: 7.0±0.2
pH:
6.80-7.20
Cultural Response:
Cultural characteristics observed after an incubation at 35-37°C for 18-24 hours.
Specimen:
Isolated Microorganisms

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

Limitations:
1.This medium is general purpose medium and does not support the growth of fastidious organisms

Storage and Shelf Life

Store between 10-30°C in a tightly closed container and the prepared medium at 20-30°C. Use before expiry date 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. 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 ofinfectious materials and material that comes into contact with sample must be decontaminated and disposed of in accordance with current laboratory techniques (1,2).

Warning and Precautions

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 specimens. Safety guidelines may be referred in individual safety data sheets.

References

1.Isenberg, H.D. Clinical Microbiology Procedures Handbook 2nd Edition.
2.Jorgensen, J.H., Pfaller, M.A., Carroll, K.C., Funke, G., Landry, M.L., Richter, S.S andWarnock., D.W. (2015) Manual of Clinical Microbiology, 11th Edition. Vol. 1.
3.Lennox E.S., 1955, Transduction of Linked Genetic Characters of the host bybacteriophage P1., Virology, 1:190.

Disclaimer :

- User must ensure suitability of the product(s) in their application prior to use.
- The product conforms solely to the technical information provided in this booklet and to the best of knowledge research and development work carried at **CDH** is true and accurate.
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