Molecular Biology Growth Media

# Technical Information

NZCYM Growth Agar

## Product Code: G1018

NZCYM Growth Agar is used for lambda and filamentous phage.

Composition**		
Ingredients	Grams/Litre	
Casein enzymic hydrolysate	10.00	
Casein acid hydrolysate	1.00	
Yeast extract	5.00	
MgSO <sub>4</sub> . 7H <sub>2</sub> O	2.00	
Sodium chloride	5.00	
Agar	15.00	

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

## Methodology

Suspend 38 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. Mix well and dispense as desired.

## Principle and Interpretation:

NZCYM Growth Agar is used for lambda and filamentous phage. This medium was developed by Blattner and colleagues as a rich medium for the propagation of bacteriophages (1). Cells grow very fast in this medium as this medium provides all the amino acids, vitamins and other metabolites required for cell growth (2). Casein enzymic hydrolysate and casein acid hydrolysate provide nitrogen, amino acids, and carbon sources for the cells. Yeast extract functions as the source of vitamins and trace elements. Sodium chloride provides sodium ions for transport and osmotic balance and Magnesium sulfate is a source of magnesium ions required in a variety of enzymatic reactions, including DNA replication (3). This medium contains agar as solidifying agent. NZCYM Growth Agar allows the cells to grow more rapidly as they do not have to synthesize nucleotide precursors and other factors required for growth.

# **Quality control:**

#### Appearance of Powder :

Cream to yellow coloured, homogeneous, free flowing powder. Gelling: Firm, comparable with 1.5% Agar gel. Colour and Clarity : Light amber coloured, clear to slight opalescent gel forms in the tube. Cultural Response : Cultural characteristics observed after an incubation at 35-37°C for 18 - 48 hours. Organisms (ATCC) Growth

Escherichia coli ATCC 23724 Escherichia coli ATCC 25922 Escherichia coli MTCC 1652 Growth good-luxuriant good-luxuriant good-luxuriant

# Storage and Shelf Life

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



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## Reference :

(1)Blattner, F. R., B. G. Williams, A. E. Blechl, K. Denniston-Thompson, H. E.Faber, L. A. Furlong, D. J. Grunwald, D. O. Kiefer, D. D. Moore, J. W. Schumm,E.L. Sheldon, and O. Smithies. 1977. Charon phages: Safer derivatives ofbacteriophage for DNA cloning. Science 196:161. (2)Ausubel, F. M., R. Brent, R. E. Kingston, D. D. Moore, J. G. Seidman, J. A.Smith, and K. Struhl. 1994. Current protocols in molecular biology, vol. 1.Current Protocols, New York, NY.

(3)Sambrook J., E. F. Fritsch, and T. Maniatis. 1989. Molecular cloning: a laboratorymanual, 2nd ed. Cold Spring Harbor Laboratory, Cold Spring Harbor, NY.

### **Disclaimer**:

- User must ensure suitability of the product(s) in their application prior to use.
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