

# **Technical Data**

## **Yeast Autolysate Supplement**

**FD027** 

An enrichment supplement recommended for the cultivation of Neisseria species.

### Composition

Per vial sufficient for 500 ml medium

Ingredients	Concentration
Yeast autolysate	5g
Dextrose	0.500g
Sodium bicarbonate	0.075g

### **Directions:**

Rehydrate the contents of 1 vial aseptically with 10 ml sterile distilled water. Mix well & aseptically add to sterile, molten, cooled (45-50°C) 490 ml of GC Agar Base M434 / GC HiVeg<sup>TM</sup> Agar Base MV434 containing FO Growth Supplement FD022 along with GC Selective supplement FD021 or In Chocolate Agar Base M103 / Chocolate HiVeg<sup>TM</sup> Agar Base MV103 along with 5% v/v defibrinated blood (after addition of blood mix and heat at 80°C for 10mins) or In 320 ml of NYC Agar Base M1348 alongwith 100ml sedimented horse blood cells, 60ml citrated Horse plasma and NYC supplement FD150. Mix well and pour into sterile petri plates.

### Type of specimen

Clinical samples - Stool, urine, respiratory exudates, etc.

### **Specimen Collection and Handling**

For clinical samples follow appropriate techniques for handling specimens as per established guidelines (1,2). After use, contaminated materials must be sterilized by autoclaving before discarding.

### **Warning & Precautions**

In Vitro diagnostic use only. For professional 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.

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

Store at 2 - 8°C. Use before expiry date on the label.

### **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 (1,2).

#### Reference

- 1. Isenberg (Ed.),2004, Clinical Microbiology Procedures Handbook, Vol.3, American Society for Microbiology, Washington. D.C.
- 2. 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.
- \* Not For Medicinal Use

Revision :02/2022

HiMedia Laboratories Technical Data





HiMedia Laboratories Pvt. Limited, Plot No.C-40, Road No.21Y, MIDC, Wagle Industrial Area, Thane (W) -400604, MS, India

CEpartner4U, Esdoornlaan 13, 3951DB Maarn, NL www.cepartner4u.eu



In vitro diagnostic medical device





Storage temperature

Do not use if package is damaged

### Disclaimer :

User must ensure suitability of the product(s) in their application prior to use. Products conform solely to the information contained in this and other related HiMedia™ publications. The information contained in this publication is based on our research and development work and is to the best of our knowledge true and accurate. HiMedia™ Laboratories Pvt Ltd reserves the right to make changes to specifications and information related to the products at any time. Products are not intended for human or animal or therapeutic use but for laboratory, diagnostic, research or further manufacturing use only, unless otherwise specified. Statements contained herein should not be considered as a warranty of any kind, expressed or implied, and no liability is accepted for infringement of any patents.