



S Selective Supplement

FD068

An antibacterial supplement recommended for selective isolation of Salmonella species.

Composition

Per vial sufficient for 500 ml medium

| *Ingredients | Concentration |
|------------------------|---------------|
| Sodium sulphaacetamide | 500mg |
| Sodium mandelate | 125mg |

Directions:

Rehydrate the contents of one vial aseptically with 5.0 ml sterile distilled water. Mix well and aseptically add to 500 ml sterile, molten, cooled (45-50°C) Brilliant Green Agar Base, Modified M016/ Brilliant Green Agar W1.2% Agar M016A/ Brilliant Green HiVeg[™] Agar Base, Modified MV016/ Brilliant Green HiCynth[™] Agar Base, Modified MCD016/ Brilliant Green Agar Base w/ 1.2% Agar, Granulated GM016A/ Brilliant Green HiVeg[™] Agar Base, w/1.2% Agar MV016A/ Brilliant Green Agar Base w/Phosphates M971/ M971S/ Brilliant Green Agar Base w/Phosphates, Granulated GM971/ Brilliant Green HiVeg[™] Agar Base w/Phosphates M971. Brilliant Green Agar w/ Phosphates M9711. Mix well and pour into sterile petri plates.

Type of specimen

Clinical samples - faeces; Food samples

Specimen Collection and Handling

For clinical samples follow appropriate techniques for handling specimens as per established guidelines (1,2). For food samples follow appropriate techniques for handling specimens as per established guidelines (3). After use, contaminated materials must be sterilized by autoclaving before discarding.

Warning & Precautions

In Vitro diagnostic use. 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.

3. Salfinger Y., and Tortorello M.L., 2015, Compendium of Methods for the Microbiological Examination of Foods, 5th Ed., American Public Health Association, Washington, D.C.

* Not For Medicinal Use

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EC REP

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



IVD



-8°C 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.

In vitro diagnostic

medical device

HiMedia Laboratories Pvt. Ltd. Corporate ice Plot No. C-40, Road No.21Y, MIDC, Wagle Industrial Area, Thane (W) - 400604, India. Customer care No. 022-614 1919 Email techhelp himedialabs.com Website www.himedialabs.com