NO 5 Selective Supplement

FD096

An antibiotic supplement for the presumptive identification of *Aeromonas hydrophila*.

Composition

Per vial sufficient for 250/1000 ml medium

*Ingredients	Concentration
Novobiocin	5.00mg

Directions:

Rehydrate the contents of 1 vial aseptically with 2 ml of sterile distilled water and aseptically add to 1000 ml of sterile, molten, cooled (45-50 C) RS Medium Base M576 /RS Medium HiVeg Base ~~MV57 or~~ 250 ml of sterile Modified EC Broth Base M1285/ Modified EC Broth Base, Granulated GM1285/ Semisolid Rappaport Vassiliadis Medium Base, Modified M1282/ Soyabean Bile Broth Base M1286 / ~~Soy~~abean HiVeg Broth Base MV1286. Mix well and pour into sterile Petri plates.

Type of specimen

Clinical samples - Stool, urine, etc; Food samples; Water 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).

For water samples follow appropriate techniques for handling specimens as per established guidelines (4).

After use, contaminated materials must be sterili ed 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
3. of Clinical Microbiology,11th Edition. Vol. 1.
4. Salfinger Y., and Tortorello M.L. Fifth (Ed.), 2015, Compendium of Methods for the Microbiological Examination of Foods, American Public Health Association, Washington, D.C.
5. Baird R.B., Eaton A.D., and Rice E.W., (Eds.), 2015, Standard Methods for the Examination of Water and Wastewater 23rd ed., APHA, Washington, D.C.

*** Not For Medicinal Use**

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EC REP

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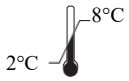
IVD ***In vitro*** **diagnostic medical device**

CE Marking



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.