



Technical Data

Vanco T Supplement

FD028

An antibiotic supplement recommended for the selective isolation of *Neisseria gonorrhoeae* and *Neisseria meningitidis*.

Composition

Per vial sufficient for 500 ml medium

*Ingredients	Concentration
Vancomycin	1mg
Colistin sulphate	3.750mg
Amphotericin B	0.500mg
Trimethoprim lactate	1.500mg

Directions:

Rehydrate the contents of 1 vial aseptically with 10 ml sterile distilled water. Mix gently. Avoid frothing of the solution to dissolve. Aseptically add the contents of one vial to 490 ml sterile, molten, cooled (45-50°C) GC Agar Base [M434](#)/ GC HiVeg™ Agar Base [MV434](#) along with FO Growth Supplement [FD022](#) and GC Selective Supplement [FD021](#). Also add sterile rehydrated contents of 1 vial of Yeast Autolysate Supplement [FD027](#) or Vitamino Growth Supplement [FD025](#). Aseptically add the contents of one vial to 930 ml of sterile, molten, cooled (45-50°C) Martin Lewis Agar Base [M2085](#) alongwith one vial of of Vitamino Growth Supplement [FD025](#). 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

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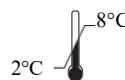
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**In vitro diagnostic
medical device**



CE Marking



Storage temperature



**Do not use if
package is damaged**

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