



INSTRUCTIONS FOR USE

VCNT ANTIMICROBIC SUPPLEMENT

Freeze-dried selective supplement

1 - INTENDED USE

In vitro diagnostic. Mixture of antimicrobials which is incorporated into culture media to permit the selective isolation of Neisseria gonorrhoeae from clinical specimens.

2 - COMPOSITION, TYPICAL FORMULA*

VIAL CONTENTS FOR 500 ML OF MEDIUM

1.50 mg Vancomycin Colistin 3.75 mg 6250 UI Nystatin Trimethoprim 2.5 mg

3 - PRINCIPLE OF THE METHOD AND EXPLANATION OF THE PROCEDURE

In 1964 Thayer and Martin¹ formulated a selective medium for the cultivation of Neisseria gonorrhoeae and Neisseria meningitidis, incorporating haemoglobin, yeast supplement B, polymyxin B and ristocetin into GC Agar. Thayer and Martin improved in 19662 the formulation substituting the original antibiotics with vancomycin, colistin and nystatin (VCN). In 1970 Martin and Lester3 modified the new Thayer-Martin Medium by increasing agar and glucose content and by incorporating an additional antibiotic, trimethoprim lactate; this improved medium is called Modified Thayer-Martin (MTM) medium.

VCNT Antimicrobic Supplement is a freeze-dried mixture of antimicrobials to be used as a supplement of GC medium Base for the selective isolation of Neisseria gonorrhoeae.

Vancomycin inhibits Gram-positive contaminants, colistin inhibits Gram-negative bacteria, including Pseudomonas species and almost all saprophytic Neisseria spp, nystatin is an anti-fungal agent and trimethoprim inhibits swarming of Proteus.

4- DIRECTIONS

Aseptically reconstitute the contents of one vial with 5 mL of sterile purified water and mix gently to dissolve. Use the reconstituted supplement for preparation of Modified Thayer-Martin (MTM) Medium as described below.

Suspend 19 g of GC Medium Base (REF 401520) in 500 mL of cold purified water; bring to boil stirring constantly and sterilize by autoclaving at 121°C for 15 minutes. Cool to 47-50°C and aseptically add 5% of defibrinated sheep blood and heat in a water bath at 80°C for 15 minutes. Cool to 47-50°C and add:

- the contents of one vial of Biovitex reconstituted with 5 mL of Restoring Fluid (ref. n° 4240009).
- the contents of one vial of VCNT Antimicrobic Supplement reconstituted as described above.

Instead of sheep blood, GC medium Base can be supplemented with sterile bovine haemoglobin solution: 5 g of bovine haemoglobin in 250 mL of water sterilized by autoclaving + 250 mL of autoclaved GC Medium Base at double concentration. Mix well and distribute into sterile plates.

Final pH of complete medium: 7.2 ± 0.2.

5 - PHYSICAL CHARACTERISTICS

Appearance of lyophilised product short, dense, yellow pastille Appearance of reconstituted product yellow, turbid solution

- MATERIALS PROVIDED - PACKAGING				
Product	Туре	REF	Packaging	
VCNT Antimicrobic Supplement	Supplement for culture media	4240008	10 vials, each for 500 mL of complete medium.	
			Secondary packaging: cardboard box	

7 - MATERIALS REQUIRED BUT NOT PROVIDED

GC Medium Base (REF 401520) or other suitable medium, bovine haemoglobin or defibrinated sheep blood, enrichment supplements Biovitex (ref 4240009) autoclave, water bath, incubator and other laboratory equipment. Flasks, sterile plates and tubes, loops and sterile swabs for microbiology, materials for the generation of a controlled incubation atmosphere with CO2 or CO2 incubator with humidifier, accessory culture media and reagents for the identification of colonies.

8 - SPECIMENS

Plates of Modified Thayer-Martin (MTM) medium can be directly inoculated with specimens from non-sterile human sites contaminated by mixed flora of bacteria and/or fungi (e.g., urogenital tract, upper respiratory tract, pus and exudates). 46 This medium is not useful for the isolation of Neisseria spp. from supposedly sterile sites.7

Good laboratory practices for collection, transport and storage of the clinical specimens should be applied; consult appropriate references for further information because Neisseria spp. are very sensitive to collection and storage procedures.

9 - TEST PROCEDURE

Allow plates to come to room temperature. The agar surface should be smooth and moist, but without excessive water.

Process the specimen as soon as possible after it is received in the laboratory to avoid loss of gonococci viability and overgrowth of contaminants.

Roll the swab over one quadrant of the surface then streak the specimen over the other quadrants of the plate to obtain well isolated colonies, ensuring that sections 1 and 4 do not overlap.

Alternatively, since swabs for gonococcal culture may contain only small numbers of organisms, roll the swabs directly on the medium in a large "Z" pattern to sufficiently transfer the specimen; cross-streak the "Z" pattern with a sterile loop.



^{*} The formula may be adjusted and/or supplemented to meet the required performances criteria.





Incubate at 35-36.5°C in a moist atmosphere supplemented with 3-7% CO₂; cultures should be examined daily for growth and held for a maximum of 72 hours.

10 - READING AND INTERPRETATION

After incubation, observe the bacterial growth and record the specific morphological and chromatic characteristics of the colonies.

N. gonorrhoeae colonies are variable in size, usually small (0,5-2 mm), moderately convex, raised, granular, glistening, moist, with entire to lobate margins, usually greyish-white to translucent; almost all strains become mucoid after 48 hours.

A Gram staining must be performed on suspected Neisseria colonies to confirm the presence of uniform Gram-negative diplococci. Performance of oxidase test is mandatory for colonies suspected to belong to Neisseria that shall be positive for N. gonorrhoeae.

11 - USER QUALITY CONTROL

All manufactured lots of the product are released for sale after the Quality Control has been performed to check the compliance with the specifications. However, the end user can perform its own Quality Control testing in accordance with the local applicable regulations, in compliance with accreditation requirements and the experience of the Laboratory. Here below are listed some test strains useful for the quality control.8

Control strains Modified Thayer-Martin Medium		INCUBATION T°/ T / ATM	EXPECTED RESULTS	
N.gonorrhoeae	ATCC	43069	35-36,5°C / 24-48H / CO ₂	good growth
P.mirabilis E.coli			35-36,5°C / 24-48H / CO ₂ 35-36.5°C / 24-48H / CO ₂	inhibited inhibited
N.sicca	ATCC		35-36,5°C / 24-48H / CO ₂	growth partially inhibited
S.epidermidis C.albicans		12228 60193	35-36,5°C / 24-48H / CO ₂ 35-36,5°C / 24-48H / CO ₂	inhibited growth partially inhibited

ATCC is a trademark of American Type Culture Collection

12- PERFORMANCES CHARACTERISTICS

Prior to release for sale a representative sample of all lots of VCNT Antimicrobic Supplement are tested for productivity and selectivity properties with Modified Thayer-Martin plates.

Productivity is tested by semi-quantitative ecometric technique with 2 gonococcal strains: N.gonorrhoeae ATCC 43069, N.gonorrhoeae ATCC 19424. After incubation at 35-36.5°C for 24-48 hours, with 3-7% of CO₂, the amount of growth is evaluated and recorded. All strains show a good growth with typical morphology. The selectivity is evaluated with modified Miles-Misra surface drop method by inoculating the plates with suitable decimal dilutions in saline of a 0.5 McFarland suspension of the non-target organisms N.sicca ATCC 9913, S.epidermidis ATCC 12228, E.coli ATCC 25922, P.mirabilis ATCC 43071, P.rettgeri ATCC 39944, S.aureus ATCC 25923, E.faecalis ATCC 19433 and C.albicans ATCC 60193. After incubation at 35-36.5°C for 24-48 hours, with 3-7% of CO2, the growth of non-target strains is inhibited.

13 - LIMITATIONS OF THE METHOD

- · Vancomycin sensitive strains of some auxotypes of N.gonorrhoeae which fail to grow on MTM, have been reported from 3% to 10% of the total isolates. 9,10 Some gonococci are susceptible to trimethoprim too. 11
- It is recommended that both a selective and a non-selective medium be used when isolating pathogenic Neisseria in order to avoid the loss of vancomycin and/or trimethoprim sensitive strains.7
- MTM is not useful for the isolation of Neisseria spp. from supposedly sterile sites as cerebrospinal fluid, conjunctival swab, skin biopsy, ioint fluid for which non-selective media are recommended.
- For the growth of N.gonorrhoeae it is necessary that the surface of the plates is moist; if it appears dry, humidify with a few drops of sterile distilled water. Place damp gauze or paper towels in the CO₂ container before incubation or use an incubator with humidifier.
- On MTM N. gonorrhoeae grows with smaller and more granular colonies than with non-selective chocolate agar.
- Some saprophytic non-target microorganisms, resistant to antimicrobials present in the media may grow. N.lactamica may grow on MTM with colonies smaller and less moist than gonococci, occasionally with a yellowish tint.7
- The gonococci are one of the most fragile Gram-negative bacteria. It is recommended that any suspected Neisseria containing specimen should be inoculated onto primary isolation medium immediately on collection to avoid any loss in viability and/or overgrowth of contaminants; if this is not possible *N.gonorrhoeae* swabs are better held at 4-6° C for not more than 3 hours.⁷
 • The incubator temperature should be set at 35-36,5°C¹² because many strains of *N.gonorrhoeae* will not grow well at 37°C. ^{7,12}
- Examine plates after 24 hours incubation. At 48 hours the Gram morphology may exhibit atypical forms.
- Many standard protocols¹³⁻¹⁵ describe the use of Modified Thayer-Martin medium for the detection of meningococcal carriage in oropharyngeal and nasopharyngeal swabs. This application is out the intended use of GC Medium base supplemented with Biovitex and VCNT supplements The end user should validate this application before routinely using those selective media for N.meningitidis detection in clinical specimens.
- · Use dacron or calcium alginate swabs for specimen collection, avoid cotton swabs since they contain fatty acids which are inhibitory for N.gonorrhoeae.7
- Incorrect specimen collection, incubation temperature, CO2 level, humidity and pH can adversely affect growth and viability of the microorganisms.
- Inactivation or deterioration of antibiotics into selective media can allow the growth of contaminants.
- It is recommended to measure the pH of complete media. GC Medium Base has sufficient buffering capability however sometimes it could be necessary to adjust the final pH.
- Even if the microbial colonies on the plates are differentiated on the basis of their morphological and chromatic characteristics, it is recommended that biochemical, immunological, molecular, or mass spectrometry testing be performed on isolates, from pure culture, for complete identification. If relevant, perform antimicrobial susceptibility testing.
- The VCNT supplement and the prepared media are intended as an aid in the diagnosis of infectious diseases; the interpretation of the results must be made considering the patient's clinical history, the origin of the sample and the results of other diagnostic tests.

14 - PRECAUTIONS AND WARNINGS

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- · VCNT Antimicrobic Supplement is a qualitative in vitro diagnostic, for professional use only; it must be used by adequately trained and qualified laboratory personnel, observing approved biohazard precautions and aseptic techniques.
- VCNT Antimicrobic Supplement is classified as dangerous according to current European legislation; consult the Safety Data Sheet before use.
- · The supplement and the medium base shall be used in association according to the directions described above. Apply Good Manufacturing Practice in the preparation process of plated media.
- VCNT Antimicrobic Supplement is sterilized by membrane filtration.
- Be careful when opening the metal ring to avoid injury.
- All laboratory specimens should be considered infectious.
- The laboratory area must be controlled to avoid contaminants such as medium powder and supplements or microbial agents.
- · Sterilize all biohazard waste before disposal. Dispose the unused supplements and the sterilized media inoculated with samples or microbial strains in accordance with current local legislation.
- Do not use VCNT Antimicrobic Supplement as active ingredients for pharmaceutical preparations or as production materials intended for human and animal consumption.
- The Certificates of Analysis and the Safety Data Sheets of the products are available on the website www.biolifeitaliana.it.
- · Notify Biolife Italiana Srl (complaint@biolifeitaliana.it) and the relevant Authorities of any serious incident occurring in connection with the use of the in vitro diagnostic.
- The information provided in this document has been defined to the best of our knowledge and ability and represents a guideline for the proper use of the product but without obligation or liability. In all cases existing local laws, regulations and standard procedures must be observed for the examination of samples collected from human and animal organic districts, for environmental samples and for products intended for human or animal consumption. Our information does not relieve our customers from their responsibility for checking the suitability of our product for the intended purpose.

15 - STORAGE CONDITIONS AND SHELF LIFE

Upon receipt, store the product in the original package at 2-8°C away from direct light. If properly stored, the product may be used up to the expiry date printed on the label; do not use beyond this date. Once the vial has been opened and the lyophilised product has been reconstituted, the resulting solution should be used immediately. Before use, examine the lyophilized and reconstituted product and discard if there are obvious signs of deterioration (e.g., contamination, atypical colour or other abnormal characteristics).

The user is responsible for the manufacturing and quality control processes of prepared media and the validation of their shelf life, according to the type (plates/tubes) and the applied storage conditions (temperature and packaging).

16 - REFERENCES

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VCNT ANTIMICROBIC SUPPLEMENT REF 4240008

Regulation (EU) 2020/878

Classification

The product is classified as hazardous. The product thus requires a safety data sheet that complies with the provisions of (EU) Regulation 2020/878

H302 Harmful if swallowed. Acute toxicity, category 4

Skin sensitization, category 1 H317 May cause an allergic skin reaction.

Labelling

Hazard pictograms:



Signal words: Warning



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Hazard statements:

H302 Harmful if swallowed.

H317 May cause an allergic skin reaction.

Precautionary statements:

P280 Wear protective gloves.

Avoid breathing dust / fume / gas / mist / vapours / spray.

If skin irritation or rash occurs: Get medical advice / attention.

Wash . . . thoroughly after handling. P261 P333+P313

P264

P362+P364 Take off contaminated clothing and wash it before reuse.

VANCOMYCIN HCL-COLISTIN Contains:

TABLE OF APPLICABLE SYMBOLS

REF or REF Catalogue number	LOT Batch code	IVD In vitro Diagnostic Medical Device	Manufacturer	This side up	
Temperature imitation	Content sufficient for <n> tests</n>	Consult Instructions for Use	Use by	Keep away from direct light	Fragile

REVISION HISTORY

Version	Description of changes	Date
Revision 1	Updated layout and content	2021/12
Revision 2	Removal of obsolete classification	2023/04

Note: minor typographical, grammatical, and formatting changes are not included in the revision history.