

# **Technical Data**

## **Campylobacter Nitrate Broth**

M1240

#### **Intended Use:**

Recommended for identification of *Campylobacter* species on the basis of nitrate reduction obtained from clinical and non-clinical samples.

## Composition\*\*

Ingredients	g/L
HM infusion B from 500 g #	10.000
Tryptose	10.000
Sodium chloride	5.000
Potassium nitrate	2.000
Final pH (at 25°C)	$7.0 \pm 0.2$

<sup>\*\*</sup>Formula adjusted, standardized to suit performance parameters

#### **Directions**

Suspend 27.0 grams in 1000 ml purified/distilled water. Heat if necessary to dissolve the medium completely. Dispense into tubes or flasks as desired. Sterilize by autoclaving at 15 lbs pressure (121°C) for 15 mins.

## **Principle And Interpretation**

Campylobacter species are ubiquitous in the environment inhabiting a wide variety of ecological niches (1). Infection with Campylobacter species is one of the most common causes of human bacterial gastroenteritis (1). Most species are found in animals (cattle, swine) and cause infertility and abortion. Campylobacter species are non-fermentative and non-oxidative in their metabolism, deceiving energy from the use of amino acids (2). Also, they do not ferment or oxidize the usual carbohydrate substrates. Campylobacter Nitrate Broth is formulated as per APHA(3) and is used for identification of Campylobacter species on the basis of nitrate reduction (4). Campylobacter jejuni is oxidase positive and reduces nitrates. HM infusion B from and tryptose in the medium provide the essential nutrients including mainly the nitrogenous and a few carbon compounds to Campylobacter species. Sodium chloride maintains the osmotic balance of the medium. Potassium nitrate serves as the nitrate source. Biochemical reactions by which species may be differentiated are relatively few because of their inability to ferment or oxidize the usual carbohydrate substrates.

#### Preparation of Nitrate Test Reagents and Technique:

- 1. Sulphanilic acid: Dissolve 8 grams of sulphanilic acid in 1 litre 5 N acetic acid.
- 2. Alpha-naphthylamine reagent: Dissolve 5 grams of alpha-naphthylamine in 1 litre 5 N acetic acid.

For the test:

Put 2 - 3 drops of each reagent into the tube containing culture to be tested. A distinct red or pink colour indicates nitrate reduction. A control (uninoculated) tube should also be tested.

## Type of specimen

Clinical samples - Faeces; Food and dairy samples; Environmental samples.

#### **Specimen Collection and Handling:**

For clinical samples follow appropriate techniques for handling specimens as per established guidelines (5,6).

For food and dairy samples, follow appropriate techniques for sample collection and processing as per guidelines (3,4). After use, contaminated materials must be sterilized by autoclaving before discarding.

## **Warning and 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.

<sup># -</sup> Equivalent to Beef heart infusion from

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#### **Limitations:**

- 1. Individual organisms differ in their growth requirement and may show variable growth patterns on the medium
- 2. Further isolation and biochemical tests must be carried out for confirmation

#### **Performance and Evaluation**

Performance of the medium is expected when used as per the direction on the label within the expiry period recommended temperature.

### **Quality Control**

#### **Appearance**

Cream to yellow homogeneous free flowing powder

#### Colour and Clarity of prepared medium

Amber coloured, clear solution without any precipitate

#### Reaction

Reaction of 2.7% w/v aqueous solution at 25°C. pH: 7.0±0.2

pН

6.80 - 7.20

#### **Cultural Response**

Cultural characteristics observed after an incubation at 35-37°C for 18-24 hours.

Organism	Growth	Nitrate reduction
Acinetobacter calcoaceticus ATCC 23055	good-luxuriant	negative, no colour development
Campylobacter jejuni ATCC 29428 (00156*)	good-luxuriant	positive, red colour developed within 1-2
Escherichia coli ATCC 25922(00013*)	good-luxuriant	minutes positive, red colour developed within 1-2 minutes
#Klebsiella aerogenes ATCC 13048 (00175*)	good-luxuriant	positive, red colour developed within 1-2 minutes
Salmonella Typhimurium ATCC 14028 (00031*)	good-luxuriant	positive, red colour developed within 1-2 minutes

Key: \*Corresponding WDCM numbers.

# Formerly known as Enterobacter aerogenes

## **Storage and Shelf Life**

Store between 10-30°C in a tightly closed container and the prepared medium at 2-8°C. Use before expiry date on the label. On opening, product should be properly stored dry, after tightly capping the bottle in order to prevent lump formation due to the hygroscopic nature of the product. Improper storage of the product may lead to lump formation. Store in dry ventilated area protected from extremes of temperature and sources of ignition Seal the container tightly after use. Product performance is best if used within stated expiry period.

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## **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 (5,6).

#### Reference

- 1. Manning H., Duim B., Wassenaar T., Wagenaar A., Ridley A., Newell D. G., 2001, Appl. Environ. Microbiol., 67:1185
- 2. Koneman E. W., Allen S. D., Janda W. M, Schrenckenberger P. C., Winn W. C. Jr., 1992, Colour Atlas and Textbook of Diagnostic Microbiology, 4th Ed, J. B. Lippincott Company.
- 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.
- 4. Wehr H. M. and Frank J. H., 2004, Standard Methods for the Microbiological Examination of Dairy Products, 17th Ed., APHA Inc., Washington, D.C.
- 5. Isenberg, H.D. Clinical Microbiology Procedures Handbook. 2nd Edition
- 6. 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.

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In vitro diagnostic medical device



Storage temperature



CE Marking



Do not use if package is damaged

#### Disclaimer:

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