



Technical Data

Pseudomonas Agar Base (CN Agar)

M085

Intended use :

For selective isolation of *Pseudomonas* species. The composition and performance criteria of this medium are as per the specifications laid down in ISO 16266-1:2006

Composition**

ISO specification - *Pseudomonas* agar base/CN-agar

Ingredients	g/ L
Gelatin peptone	16.000
Casein hydrolysate	10.000
Potassium sulphate, anhydrous (K ₂ SO ₄)	10.000
Magnesium chloride, anhydrous (MgCl ₂)	1.400
Agar	11.00 - 18.00
Final pH (of solidified, complete medium)	7.1±0.2

Supplement to be added after autoclaving g / L

Hexadecyltrimethyl ammonium bromide (Cetrimide)	0.200
Nalidixic acid	0.150

Pseudomonas Agar Base

Ingredients	g/ L
Gelatin peptone	16.000
Tryptone	10.000
Potassium sulphate	10.000
Magnesium chloride	1.400
Agar	11.000
Final pH (at 25°C)	7.1±0.2

Cetrinix Supplement - FD029

Cetrimide	100mg
Nalidixic acid	7.500mg

**Formula adjusted, standardized to suit performance parameters

Directions

Suspend 24.2 grams in 500 ml purified/distilled water containing 5 ml glycerol. Heat to boiling to dissolve the medium completely. Sterilize by autoclaving at 15 lbs pressure (121°C) for 15 minutes. Cool to 45-50°C and aseptically add 1 vial of sterile rehydrated contents of Cetrinix Supplement (FD029). Mix well and pour into sterile Petri plates.

Note : Do not keep the molten agar for longer than 4 hours.

Principle And Interpretation

Pseudomonas Agar Base is as per the specification laid down in ISO 16266 for testing water quality by MPN method (1). It is also a modification of Kings A medium (2) which contains magnesium chloride and potassium sulphate to enhance pigment production. Cetrinix supplement is specified for the selective isolation of *Pseudomonas aeruginosa* from water samples, it suppresses the growth of *Klebsiella*, *Proteus* and *Providencia* species. Lowbury and Collins (3) studied cetrimide as a selective agent.

Tryptone and gelatin peptone supplies nitrogenous and carbonaceous compounds, long chain amino acids, and other essential growth nutrients.

Examine inoculated plates after incubation at UV light 360 ± 20 nm. The presence of blue-green colonies with fluorescence may be considered as presumptive evidence of *Pseudomonas aeruginosa*.

Type of specimen

Clinical samples - pus, urine, Water samples.

Specimen Collection and Handling:

ISO 16266-1:2006

Preparation of test sample: Prepare tenfold dilutions of water samples (1,4). After use, contaminated materials must be sterilized by autoclaving before discarding.

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

Limitations :

1. Individual organisms differ in their growth requirement and may show variable growth patterns on the medium.
2. Each lot of the medium has been tested for the organisms specified on the COA. It is recommended to users to validate the medium for any specific microorganism other than mentioned in the COA based on the user's unique requirement.

Performance and Evaluation

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

Quality Control

Appearance

Cream to yellow homogeneous free flowing powder

Gelling

Firm, comparable with 1.1% Agar gel.

Colour and Clarity of prepared medium

Yellow coloured clear to slightly opalescent gel forms in Petri plates

Reaction

Reaction of 4.84% w/v aqueous solution containing 1% v/v glycerol at 25°C. pH : 7.1±0.2

pH

6.90-7.30

Cultural Response

Productivity : Cultural response was observed after an incubation at $36 \pm 2^\circ\text{C}$ for 44 ± 4 hours, with added sterile Cetrinix Supplement (FD029). Recovery rate is considered as 100% for bacteria growth on Reference medium - Soyabean Casein Digest Agar (Tryptone Soya Agar).

Selectivity : Cultural response was observed after an incubation at $36 \pm 2^\circ\text{C}$ for 44 ± 4 hours, with added sterile Cetrinix Supplement (FD029).

Organisms	Inoculum (CFU)	Growth	Recovery#	Fluorescence under UV 360± 20 nm
Productivity				
<i>Pseudomonas aeruginosa</i> ATCC 27853 (00025*)	50-100	good-luxuriant	$\geq 50\%$	blue-green with Fluorescence
^ <i>Pseudomonas paraaeruginosa</i> ATCC 9027 (00026*)	50-100	good-luxuriant	$\geq 50\%$	blue-green with Fluorescence
<i>Pseudomonas aeruginosa</i> ATCC 10145 (00024*)	50-100	good-luxuriant	$\geq 50\%$	blue-green with Fluorescence
Selectivity				
<i>Enterococcus faecalis</i> ATCC 29212 (00087*)	$\geq 10^4$	inhibited	0%	
<i>Enterococcus faecalis</i> ATCC 19433 (00009*)	$\geq 10^4$	inhibited	0%	
<i>Escherichia coli</i> ATCC 25922 (00013*)	$\geq 10^4$	inhibited	0%	
<i>Escherichia coli</i> ATCC 8739 (00012*)	$\geq 10^4$	inhibited	0%	

Key : (*) Corresponding WDCM numbers,

^ Formerly known as *Pseudomonas aeruginosa*

- Recovery obtained for productivity is $\geq 70\%$ when compared to a previously validated batch of *Pseudomonas* Agar Base (CN Agar) is used.

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.

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).

References

1. Water quality — Detection and enumeration of *Pseudomonas aeruginosa*-- Method by membrane filtration ; ISO 16266-1:2006
2. King E.O., Ward M.K. and Raney D.E., 1954, J. Lab and Clin. Med., 44:301.
3. Lowbury E.J. and Collins A.G., 1955, Clin. Path., 8:47.
4. Lipps WC, Braun-Howland EB, Baxter TE, eds. Standard methods for the Examination of Water and Wastewater, 24th ed. Washington DC:APHA Press; 2023.
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.

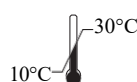
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