

Pseudomonas Selective Agar, Base (Cetrimide Agar)

CETRIMIDE Agar

A modification of the medium proposed by BROWN and LOWBURY (1965) for the isolation and differentiation of *Pseudomonas aeruginosa* from various materials.

General Information

This medium complies with the recommendations of the harmonised method in the European Pharmacopeia 5.6 and the United States Pharmacopeia 29 (2006).

Mode of Action

The use of cetrimide (cetyltrimethylammonium bromide) was recommended by LOWBURY (1951) and other authors; this compound largely inhibits the growth of the accompanying microbial flora. According to LOWBURY and COLLINS (1955), a concentration of 0.3 g/litre inhibits the accompanying organisms satisfactorily and minimizes interference with the growth of *Ps. aeruginosa*. The pigment production of *Ps. aeruginosa* is not inhibited when grown on this medium.

GOTO and ENOMOTO (1970) recommend the addition of 15 µg nalidixic acid/ml to improve the inhibition of the accompanying microbial flora.

Typical Composition (g/litre)

Peptone from gelatin 20.0; magnesium chloride 1.4; potassium sulfate 10.0; N-cetyl-N,N,N-trimethylammoniumbromide (cetrimide) 0.3; agar-agar 13.6.

Also be added:

Glycerol 10 ml.

Preparation

Suspend 45.3 g/litre, add 10 ml glycerol/litre, autoclave (15 min at 121 °C). Pour plates.

pH: 7.2 ± 0.2 at 25 °C.

The plates are turbid and light brown.

Experimental Procedure and Evaluation

Inoculate by spreading the sample on the surface of the plates.

Incubation: *Pseudomonas aeruginosa* 18 h at 35 °C, others 72 h.

Ps. aeruginosa colonies produce a yellow-green pigment (pyocyanin) and fluoresce under UV light. Further tests should be performed for further identification (HUGH and LEIFSON 1953, KOVACS 1956, THORNLEY 1960, BÜHLMANN et al. 1961 etc).

Literature

- DIN Deutsches Institut für Normung e.V.: Deutsche Einheitsverfahren zur Wasser-, Abwasser- und Schlammuntersuchung. Mikrobiologische Verfahren. Nachweis von *Pseudomonas aeruginosa*. - DIN 38411-8 (1982).
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- European Pharmacopeia 5.6, Chapter 2.6.13 B (Harmonised Method), Microbial examination of nonsterile products: Tests for specified microorganisms, 2006.
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- HUGH, R., a. LEIFSON, E.: The taxonomic significance of fermentative versus oxidative metabolism of carbohydrates by various gram-negative bacteria. - J. Bact., 66; 24-26 (1953).
- KOVACS, N.: Identification of *Pseudomonas pyocyanea* by the oxidase reaction. - Nature (Lond.), 178; 703 (1956).
- LOWBURY, E.J.L.: Improved culture methods for the detection of *Ps. pyocyanea*. - J. Clin. Pathol., 4; 66-72 (1951).
- LOWBURY, E.J.L., a. COLLINS, A.G.: The use of a new cetrimide product in a selective medium for *Pseudomonas pyocyanea*. - J. Clin. Pathol., 8; 47-48 (1955).
- THORNLEY, M.J.: The differentiation of *Pseudomonas* from other gram-negative bacteria on the basis of arginine metabolism. - J. Appl. Bact., 23; 37-52 (1960).
- United States Pharmacopeia 29 - NF 24, Chapter 62, Microbial examination of nonsterile products: Tests for specified microorganisms, 2006.

Ordering Information

Product	Ordering No.	Pack size
Pseudomonas Selective Agar, Base (Cetrimide Agar)	1.05284.0500	500 g
Glycerol (about 87 %)	1.04094.0500	500 ml
UV Lamp (366nm)	1.13203.0001	1 ea



Pseudomonas aeruginosa ATCC 9027

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Quality control (spiral plating method)

Test strains	Inoculum (CFU)	Recovery (%)	Yellow-green pigment
<i>Pseudomonas aeruginosa</i> ATCC 9027	10 - 100	≥ 30	+
<i>Pseudomonas aeruginosa</i> ATCC 25668	10 - 100	≥ 30	+
<i>Pseudomonas aeruginosa</i> ATCC 27853	10 - 100	≥ 30	+
<i>Escherichia coli</i> ATCC 8739	> 10 ⁴	no growth	-
<i>Proteus mirabilis</i> ATCC 29906	> 10 ⁴	no growth	-
<i>Salmonella typhimurium</i> ATCC 14028	> 10 ⁴	no growth	-
<i>Staphylococcus aureus</i> ATCC 6538	> 10 ⁴	no growth	-