

Plate count agar (Standard methods agar)

REF.	Pack size	
1410 001	100 gm	
1410 002	500 gm	

Intended Use

Plate count agar (Standard Methods Agar) is used for the determination of plate counts of microorganisms isolated from blood and urine.

Background

Plate count agar was developed by Buchbinder, Baris, and Goldstein in 1953 at the request of the American Public Health Association (APHA). This medium is recommended for the plate count of microorganisms in milk and other dairy products and may also be used to determine sanitary quality foods, water and other materials.

Principle

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Tryptone provides nitrogenous substances and other amino acids. Yeast extract provides B complex vitamins and dextrose is the energy source

Components	gm/Liter
Yeast Extract	2.5
Dextrose	1.0
Tryptone	5.0
Agar	15.0

Final pH (at 25°C) 7.0 ± 0.2

Preparation, Storage and Stability

Store the dehydrated medium at 10-30°C and use before the expiry date on the label.Store the prepared medium at 2-8°C. After the desired amount of medium is taken out, replace the cap tightly to protect from hydration.

Procedure

1. Suspend 23.5 g of the powder in 1 L distilled water and mix well.

2.Heat with frequent agitation to dissolve the powder completely.

3.Sterilize by autoclaving at 121°C for 15 minutes.

4. Cool to 45-50°C and pour into sterile petri plates.

SYMBOLS IN PRODUCT LABELLING

ECREP	Authorised Representative	Z	Use by/Expiration Date
IVD	For in-vitro diagnostic use	∕!∖	CAUTION. Consult instructions
LOT	Batch Code/Lot number		for use
REF	Catalogue Number	-	Manufactured by
i	Consult instructions for use	X	(Xi) - Irritant
°°	Temperature Limitation		

Quality control

Appearance

Organisms (ATCC)	Growth
3-Cultural Response	: after 18-24 hours at 35 ± 2°C for clinical specimens or 30-35 (as per pharmacopoeia)
2-Prepared Appearance	: Prepared medium is trace to slightly hazy, and light beige to medium amber.
1-Dehydrated Appearance	e :Powder is homogeneous, free flowing, and light beige.

	0.010
Escherichia coli	Good
Enterococcus faecalis	Good
Bacillus subtilis	Good
Lactobacillus rhamnosus	Good

Interpretation of the results

Count colonies on all plates containing 30 - 300 colonies. Calculate bacterial count per milliliter of sample by multiplying the average number of colonies per plate by the reciprocal of the dilution used. Report the count as CFU/mL.

Precautions

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 specimens. Safety guidelines may be referred in individual safety data sheets.

Bibliography

1. Marshall, R. T. (ed.). 1993. Standard methods for the microbiological examination of dairy products, 16th ed. American Public Health Association, Washington, D.C.

2. Cunnif, P. (ed.). 1995. Official methods of analysis AOAC International, 16th

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Egyptian Co for Biotechnology - Spectrum Diagnostics (S.A.E) Obour city industrial area. block 20008 piece 19 A. Cairo. Egypt. Tel: +202 4489 2248 - Fax: +202 4489 2247 www.spectrum-diagnostics.com E-mail:info@spectrum-diagnostics.com







Rev.(5), 16/7/2023