

# Cetrimide Agar

U.S.P.

# **LAB 133**

#### **Description**

A medium recommended by the United States Pharmacopoeia for the isolation of *Pseudomonas aeruginosa* from pharmacological preparations. Subculture is carried out onto the medium after enrichment in LAB 4 Tryptone Soy Broth. Cetrimide inhibits the growth of many micro organisms whilst allowing *P. aeruginosa* to develop typical colonies which will fluoresce in ultraviolet light and produce green pigment.

Formula	g/litre
Pancreatic Digest of Gelatin	20.0
Magnesium chloride	1.4
Potassium sulphate	10.0
Cetyl trimethylammonium bromide (cetrimide)	0.3
Agar	13.6

## **Method for reconstitution**

Weigh 45.3 grams of powder, disperse in 1 litre of deionised water. Add 10ml of glycerol, allow to soak for 10 minutes then swirl to mix. Sterilise at 121°C for 10 minutes.

Appearance: Opalescent, pale yellow agar.

**pH:**  $7.2 \pm 0.2$ 

Minimum Q.C. organisms: P. aeruginosa NCIMB 50067

E. coli (inhibition) NCIMB 50034

**Storage of Prepared Medium:** Plates – up to 7 days at 2-8°C in the dark.

Inoculation: Subculture from enrichment broth, streak out for single colonies.

Incubation: 30-35°C aerobically for 24-48 hours.

Growth Characteristics				
organism	colony size (mm)	shape & surface	colour	
P. aeruginosa	0.5-1.0	F.CR.D.	green pigment (non pigment) green/yellow fluorescence	
P. fluorescens	0.5	CV.R.E.G.	green/yellow fluorescence	
E. coli	N.G.			
S. aureus	N.G.			
Proteus spp.	N.G.			

### References

United States Pharmacopoeia XXI. 1985.

Brown V.I., Lowbury E.J.L. (1965). Use of an improved Cetrimide Agar Medium and other culture methods for *Pseudomonas aeruginosa*. J. Clin. Pathol. 18, 752-756.