

Technical Information

Clostridium Difficile Agar Base

Product Code: DM 1836

Application: - Clostridium Difficile Agar Base with supplement is used for cultivation of *Clostridium difficile* from food and certain pathological specimens.

Composition**

Ingredients	Gms / Litre
Proteose peptone	40.000
Disodium phosphate	5.000
Monopotassium phosphate	1.000
Magnesium sulphate	0.100
Sodium chloride	2.000
Fructose	6.000
Agar	15.000
Final pH (at 25°C)	7.4±0.2

**Formula adjusted, standardized to suit performance parameters

Principle & Interpretation

The spectrum of disease caused by *Clostridium difficile* (a pathogenic *Clostridium* affecting the bowel) ranges from pseudomembranous colitis (PMC) an antibiotic associated colitis (AAC) to chronic inflammatory bowel diseases, post-operative diarrhoea and non-antibiotic associated diarrhoea ⁽¹⁾. First report of *C. difficile* in human infections was published by Smith and King ⁽²⁾ George et al ⁽³⁾ recommended the use of a fructose-containing medium with egg yolk for the isolation of *C. difficile* from faecal specimens. The medium was made inhibitory to the accompanying flora by the addition of the selective agents namely, D-cycloserine and cefoxitin.

This medium does not contain neutral red indicator, as in the original formulation, but recommend the use of sheep or horse blood that increases the size of colony and recovery of *C. difficile* also ⁽³⁾. The medium composition is designed so as to obtain luxuriant growth of *C. difficile*. The selective agents D-cycloserine and cefoxitin used in the medium inhibit the growth of majority of *Enterobacteriaceae* and also *Enterococcus faecalis*, Staphylococci, gram-negative anaerobic bacilli and *Clostridium* species other than *C. difficile*, which may be found abundantly in faecal samples.

Spread a part of the faecal sample on the medium to obtain isolated colonies. Incubate the plates anaerobically at 37°C for 18 - 48 hours. *C. difficile* forms grayish white, irregular, raised and opaque colonies, 4-6 mm in diameter, after 48 hours. Typical gram stain morphology of *C. difficile* may not be seen in colonies taken from this medium due to the presence of antibiotics. Subculture on Blood Agar (DM1073) is required to obtain characteristic morphology. *C. difficile* colonies will not show the typical fluorescence and colour of colony on this medium whereas other Clostridia can give fluorescence. Therefore, for complete identification and confirmation, other tests like gram staining, morphology, biochemicals, specific cytotoxin and clinical observation should be carried out.

Methodology

Suspend 34.55 grams of powder media in 500 ml distilled water. Shake well & heat to dissolve the medium completely. Sterilize by autoclaving at 15 lbs pressure (121°C) for 15 minutes. Cool to 50°C. Aseptically add rehydrated contents of 1 vial of Clostridium Difficile Supplement (MS2010) together with 7% (v/v) defibrinated Horse blood or Sheep blood. Mix well and pour into sterile Petri plates.

Quality Control

Physical Appearance

Cream to yellow homogeneous free flowing powder

Gelling

Firm, comparable with 1.5% Agar gel

Colour and Clarity of prepared medium

Basal medium: Light amber coloured clear to slightly opalescent gel. After addition 7% v/v defibrinated horse blood: Cherry red coloured, opaque gel forms in Petri plates.

Reaction

Reaction of 6.9 1% w/v aqueous solution at 25°C. pH : 7.4±0.2

pH range 7.20-7.60

Cultural Response/ characteristics

DM1836: Cultural characteristics observed under anaerobic condition with added Clostridium Difficile Supplement (MS2010) and 7% v/v defibrinated horse blood, after an incubation at 35-37°C for 48 hours.

Organism	Inoculum (CFU)	Growth	Recovery	Colour of Colony
Clostridium difficile ATCC 11204	50-100	good-luxuriant	≥50%	greyish-white
Shigella flexneri ATCC 12022	≥10 ³	inhibited	0%	
Escherichia coli ATCC 25922	≥10 ³	inhibited	0%	
Staphylococcus aureus ATCC 25923	≥10 ³	inhibited	0%	

Storage and Shelf Life

Dried Media: Store below 30°C in tightly closed container and use before expiry date as mentioned on the label.

Prepared Media: 2-8° in sealable plastic bags for 2-5 days.

Further Reading

1. Collee J. G., Fraser A. G., Marmion B. P., Simmons A., (Eds.), Mackie and McCartney, Practical Medical Microbiology, 14th Ed., Churchill Livingstone.
2. Smith L. D. S. and King E. O., 1962, J. Bacteriol., 84:65.
3. George W. L., Sutter V. L., Citron D., and Finegold S. M., 1979, J. Clin. Microbiol., 9:214

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