

Technical Information

Blood Free Campylobacter Selectivity Agar Base

Product Code: DM 1887

Application: - Blood Free Campylobacter Selectivity Agar Base is used for selective isolation and differentiation of Campylobacter species.

position**			
Gms / Litre			
10.000			
10.000 3.000			
5.000			
1.000			
0.250			
0.250			
4.000			
12.000 7.4±0.2			
	10.000 10.000 3.000 5.000 1.000 0.250 0.250 4.000 12.000		

Principle & Interpretation

As Campylobacters are carried in the intestinal tract of animal therefore they contaminate foods of animal origin ⁽¹⁾. *Campylobacter* causes intestinal upset or abortion in animals. It is also one of the most important causes of human gastroenteritis, particularly in children. Initially blood was used in the isolation of *Campylobacter*. Later it was reported that charcoal can be effectively used in place of blood ⁽²⁾ that rules out the variability noted during isolation of Compylobactor using blood. Blood Free Campylobacter Selectivity Agar Base ⁽³⁾ formulated as per APHA ⁽¹⁾ and recommended by the ISO Committee ⁽⁴⁾ is used for selective isolation of *Campylobacter* species. Cephalothin in the original formula was replaced by Cefoperazone as the latter gave better selectivity ⁽⁵⁾. *Campylobacter* species are highly resistant to cefoperazone, an antibiotic which effectively suppresses growth of *Pseudomonas* and *Enterobacteriaceae* ⁽⁷⁻⁹⁾ and increases the selectivity of the medium. Due to this addition, the medium is also known as Campylobacter Charcoal Differential Agar (CCDA). Charcoal, sodium pyruvate and ferrous sulphate reduces the aerotolerance of medium by quenching photochemically generated toxic oxygen derivatives ⁽⁹⁾. Peptic digest of animal tissue, casein enzymic hydrolysate and beef extract serve as sources of essential nutrients and amino acids. Casein is added to help grow certain strains of nalidixic acid resistant thermophilic environmental *Campylobacter* organisms ⁽⁶⁾. Additional Amphotericin B in Blood Free Campylobacter Broth Base suppresses the growth of yeast and mold contaminants. Colonies tend to swarm when initially isolated from clinical specimens.

Methodology

Suspend 22.75 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 and aseptically add rehydrated contents of 1 vial of Campylobacter Supplement V (MS2067). Alternatively to increase the selectivity of the medium, rehydrated content of one vial of CAT Selective Supplement (MS2145) may be added to 500 ml sterile molten base. Mix well and pour into sterile Petri plates.





Bases / Media Supplements

Quality Control

Physical Appearance

Grey to black homogeneous free flowing powder

Gelling

Firm, comparable with 1.2% Agar gel

Colour and Clarity of prepared medium

coloured, opaque gel forms in Petri plates

Reaction

Reaction of 4.55% w/v aqueous solution at 25°C. pH : 7.4±0.2

pH range 7.20-7.60

Cultural Response/Characteristics

DM1887: Cultural characteristics observed with added Campylobacter Supplement V (MS2067), after an incubation at 42°C for 24-48 hours.

Organism	Inoculum (CFU)	Growth	Recovery	Haemolysis
Campylobacter coli ATCC 33559	50-100	good-luxuriant	>=50%	creamy-grey
Campylobacter jejuni ATCC 29428	50-100	good-luxuriant	>=50%	Grey
Campylobacter laridis ATCC 35222	50-100	good-luxuriant	>=50%	varying type

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:** 2-8⁰ in sealable plastic bags for 2-5 days.

Further Reading

 Downes F. P. and Ito K., (Eds.), 2001, Compendium of Methods for the Microbiological Examination of Foods, 4th Ed., APHA, Washington, D.C.

- 2. Bolton F. J., Hutchinson D. N and Coates D., 1984, J. Clin. Microbiol., 19:169.
- 3. Atlas R. M., 2004, Handbook of Microbiological Media, 3rd Ed, CRC Press.
- 4. International Organization for Standardization (ISO), 1995, Draft ISO/DIS 10272.
- 5. Hutchinson D. N and Bolton F.J., 1984, J. Clin. Pathol., 34:956.
- 6. Koneman E. W., Allen S. D., Janda W. M., Schreckenberger P. C., Winn W. C. Jr., 1992, Colour Atlas and Textbook of Diagnostic Microbiology, 4 th Ed., J. B. Lippinccott Company
- 7. Ahonkai V. I., et al, 1981, Antimicrob. Agents. Chemother.,20:850
- 8. Jones R. N., et al, 1980, Antimicrob. Agents. Chemother.,17:743
- 9. Karmali M. A., et al, 1986, J. Clin. Microbiol., 23 :456

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- User must ensure suitability of the product(s) in their application prior to use.
- The product conform solely to the technical information provided in this booklet and to the best of knowledge research and development work carried at **CDH** is true and accurate
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