BIOMARK Laboratories-INDIA

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TECHNICAL SHEET

TEGITATO TEET						
B1080	KARMALI CAMPYLOBACTER AGAR BASE					
Formula						
Ingredients:	gms/lit.					
Peptone, special	23.00					
Corn starch 1.00						
Sodium chloride 5.00						
Charcoal 4.00						
Agar 12.00						
Final pH (at 25°0	C): 7.4 <u>+</u> 0.2					
Directions:						
Suspend 22.5 grams in 490 ml distilled water. Heat to boiling to dissolve the medium completely. Sterilize by autoclaving at 15 lbs pressure (121°C) for 15 minutes. Cool to 45-50°C. Aseptically add 5 mlof Hemin solution (16 mg/5ml)& rehydrated contents of 1vial of Campylobacter selective supplement Karmali(BF112). Alternatively, 1 vial of Campylobacter selective supplement w/Hemin(BF113) can be added instead of BF112 and Hemin. Mix well and pour into sterile Petri plates.						
Principle:						

Peptone special, cornstarch and hemin, serve as sources of essential nutrients required for bacterial metabolism. Presence of charcoal in the medium helps to neutralize the toxic metabolic products formed in the medium. Sodium pyruvate (present in Supplement) enhances, the aerotolerance of microaerophilic Campylobacter by quenching the toxic forms of oxygen. The antibiotics included in the selective supplement are Vancomycin, Amphotericin Band Cefoperazone. Vancomycin suppresses gram-positive organisms while Amphotericin inhibits the fungal flora. Cefoperazone has inhibitory action on gram-negative organisms other than Campylobacter. The inoculated plates are incubated in an atmosphere consisting of approximately 5-6% O2, 10% CO2 and 84-85% N2 at 42°C.

QC Tests - (I)Dehydrated Medium				
	Colour:	Greyish-black		
	Appearance :	Homogeneous Free Flowing powder		
(II)	Rehydrated medium			
	pH (post autoclaving/heating):	7.4 ± 0.2		
	Colour (post autoclaving/heating):	Black		
	Clarity (post autoclaving/heating):	Opalescent gel		
(III)Q.C. Test Microbiological				

Cultural characteristics observed with added Hemin solution and Campylobacter Selective Supplement Karmali (BF112) or Campylobacter selective supplement w/Hemin (BF113), after an incubation at 42°C for 42-48 hours.

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MICROORGANISM (ATCC)	GROWTH					
Campylobacter jejuni (29428)	Good-luxuriant					
Campylobacter coli (33559)	Good-luxuriant					
Escherichia coli (25922)	None-poor					

Precautions: 1. For Laboratory Use.

2. Follow proper, established laboratory procedures in handling and disposing of infectious materials.

Refer disclaimer Overleaf

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Limitations :	1. Since the nutritional requirements of organisms vary, some strains may be encountered that fail to grow or grow poorly on this medium.							
	2. Campylobacter Agar prepared with either Campylobacter Antimicrobic Supplement S or Campylobacter Antimicrobic Supplement B is selective primarily for Campylobacter species. Biochemical testing using a pure culture is necessary for complete identification. Consult appropriate references for further information. 3. Growth of Campylobacter fetus subsp. Intestinalis may be dramatically inhibited on Campylobacter Agar Blaser due to the prsence of cephalothin. The use of Campylobacter Agar Skirrow and incubation at 35°C is suggested when isolating this orgnisms from mixed populations.							
	4. Some strains may show poor growth due to strain variability. 5. Some strains of normal enteric organissm may be encountered that are not inhibited or only partially inhibited on Campylobacter Agar.							
Use :	For selective isolation and cultivation of thermotolerant Campylobacter species from food and animal feeds.							
Storage :	Dehydrated medium- below 30°C Prepared medium- Between 2 to 8°C.							
Packing:	500 gm bottle							
Product profile:	Reconstitution	Quantity on Preparation (500g)	pH (25°C)	Supplement	Sterilization			
B1080	45.0 g/l	11.111L	7.4 ± 0.2	Hemin solution(16mg/5ml) & rehydrated contents of 1vial Campylobacter selective supplement Karmali (BF112) or Campylobacter selective supplement w/Hemin (BF113)				

Disclaimer:

User must ensure suitability of the product(s) in their application prior to use. Products conform solely to the information contained in this and other related BIOMARKLABORATORIES publications.

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Rev: December 2020