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TECHNICAL SHEET

B1298	CARY - BLAIR MEDIUM
Formula	
Ingredients:	gms/lit.
Disodium phosphate	1.10
Sodium thioglycollate	1.50
Sodium chloride	5.00
Agar	5.00
Final pH (at 25°C): 8.	4 <u>+</u> 0.2

Directions:

Suspend 12.6 grams in 991 ml distilled water. Heat to boiling to dissolve the medium completely. Cool to 50° C and aseptically add 9 ml of 1% aqueous calcium chloride solution. Adjust pH to 8.4 if necessary. Distribute in 7 ml amounts in screw-capped tubes. Steam for 15 minutes. Cool and tighten the caps.

Principle:

The medium is prepared with minimal nutrients to increase the survival of the organisms without multiplying. Sodium thioglycollate is incorporated in the medium to provide a low oxidation – reduction potential. The pH of the medium is relatively alkaline which minimizes the bacterial destruction due to the formation of acid. Medium can maintain viability of fastidious microorganisms for only a short period of time. It is recommended that best results are obtained by direct inoculation of isolate and inoculation of enriched medium at the same time specimen is inoculated into transport medium.

For collection of the specimen, use sterile cotton tipped swabs on wooden sticks. Push the swabs down to one third of the medium depth and cut the stick so that when the cap is screwed down, the swab is forced to the bottom of the medium. Tighten the cap firmly on the bottle. The specimen will be preserved and the viability of the organisms will be also maintained during transport, but over the time it will diminish. Therefore direct inoculation of the specimen is advised. Some growth of accompanying contaminants may also occur during longer period of transit. The specimen should be inoculated into a proper medium as soon as possible.

	or transit. The specimen	siloulu de illoculateu	into a prope	er medium as soon a	as possible.		
	ehydrated Medium						
Colour :			Cream to yellow				
Appearance :		Homogeneous	Homogeneous Free Flowing powder				
(II)Rehydrated							
pH (post autoclaving/heating) :		8.4 ± 0.2					
Colour (post autoclaving/heating):			Light amber				
Clarity (post autoclaving/heating):		: Slightly opale	Slightly opalescent				
	: Microbiological						
Cultural c B039	haracteristics observed a	after 18-24 hrs at 35	-37°C when	subcultured on Tryp	otone Soya Agar		
MICROORG	MICROORGANISM (ATCC)						
Enterobacter aerogenes (13048)		Good-luxuriar	nt				
Escherichia coli (25922)		Good-luxuriar	Good-luxuriant				
Klebsiella pneumoniae (13883)		Good-luxuriar	nt				
Salmonella typhimurium (14028)		Good-luxuriar	Good-luxuriant				
Shigella flexneri (12022)		Good-luxuriar	Good-luxuriant				
Vibrio cholerae (15748)		Good-luxuriar	Good-luxuriant				
Vibrio parahaemolyticus (11344)		Good-luxuriar	Good-luxuriant				
Neisseria meningitidis (13090)		Good-luxuriar	Good-luxuriant				
Precautions: 1. For Laboratory Use.							
	in handling and di	sposing of infectious					
	materials. 3. IRRITANT: Irritating to eyes, respiratory system and skin. Avoid contact with skin and						
	eyes. Do not breathe dust. Wear suitable protective clothing. Keep container tightly closed.						
Limitations :							
	fail to grow or grow poorly on this medium.						
Use :	For collection and shipment of clinical specimens.						
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		
B1298	12.6 g/l	39.682 L	8.4 ± 0.2	-	Steam for 15 minutes		

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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