BIOMARK Laboratories-INDIA

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TECHNICAL SHEET

B253 MC	CLUNG TOABE AGAR BASE						
Formula							
Ingredients :			/lit.				
Proteose peptone			0				
Dextrose							
Disodium hydrogen phosphate							
Monopotassium phosphate 1							
Sodium chloride 2							
Magnesium sulphate 0							
Agar 2			0				
Final pH (at 25°C) : 7.6 + 0.2							
Directions :							
Suspend 75.1 gms. in 900ml distilled water. Heat to boiling to dissolve the medium completely. Sterilize							
by autoclaving at 15 lbs pressure (121°C) for 20 minutes. Cool to 50°C and aseptically add 100ml of							
sterile Egg volk emulsion (BF003). Mix well and pour into sterile petri plates.							
Principle :							
McClung Toabe Agar Base is prepared the differntiation of Clostridium species on the basis of their							
lecithinase and lipase activity. Lecithinase enzyme lyses eag volk lecithin, producing an opague zone of							
precipitation surrounding the slightly raised colonies. Proteose peptone provides nitrogenous growth							
nutrients. Dextrose is the fermentable carbohydrate. Phosphates form good buffering system.							
OC Tests – (I)Dehvdrated Medium							
Colour :			Cream to Yellow				
Appearance '			Homogeneous Free Flowing powder				
(II)Rehvdrated medium							
nH (post autocl		7.6 ± 0.2					
Colour (post autoclaving/neuting) :			a) Basal medium : Amber				
b) After addition of equivalent vellow							
Clarity (post autoclaving/heating) :			a)Clear to slightly opalascent b) Opague				
(III)O.C. Test Microbiological							
Cultural characteristics observed after 18 – 24 hrs. at 35-37°C incubated anaerobically							
							SF
Clostridium porfringons (12010)							JL
Clostridium sporogonos (11/37)							
Staphylococcus aurous (25022)			uxuriant		-	<u> </u>	
Staphylococcu	s aureus (23923) [uxunanı		т —	Ŧ	
Kov Lasithingson L positive reaction apague range range range the							
rey : Leciumase : + = positive reaction, opaque Zone around the							
$t_{\text{Linaso}} + - \text{positive reaction}$ independent shoop on growth surface							
Procentions :	1 For Laborato						
2. Follow proper established laboratory precedures in handling and dispessing of						chocing of	
2. Follow proper, established aboratory procedures in nandling and disposing of							sposing of
IIIIectious IIIderidis.							
Limitations : 1. Since the nutritional requirements of organisms vary, some strains may l						may be	
Encountered that fail to grow of grow poorly on this medium.							
Sterage Debudrated medium, helew 2000 Prepared medium, Detween 2 to 0					S. 2 + - 00	<u> </u>	
Torage : Denyarated medium- below 30°C Prepared medium- Between 2 to 8°C.							
Packing: Dought boulde Provide the second file Dependition on the second file							
Product profile:	Reconstitution	Quantity c	n (FCC)	рн (25°С)	Supplemen	It	Sterilization
		Preparatio	n (500g)	7.6 . 0.5			
B253	/5.1g/l	6.6	06L	7.6 ± 0.2	sterile Egg		21°C / 15 minutes
	1			1	emulsion(BF0	03)	

Refer disclaimer Overleaf

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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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