BIOMARK Laboratories-INDIA

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TECHNICAL SHEET

B342	TRYPTOSE BLOOD AGAR BASE							
Formula								
Ingredients :			gms/lit.					
Tryptose			10.00					
Meat Extract B #			3.00					
Sodium chloride			5.00					
Agar			15.00					
# Equvivalent to Beef extract			15.00					
Final pH (at 25°	•	7.2 <u>+</u> 0.2						
Directions :								
Suspend 33 grams in 950 ml purified / distilled water. Heat to boiling to dissolve the medium								
completely. Sterilize by autoclaving at 15 lbs pressure (121°C) for 15 minutes. Cool the autoclaved								
medium to 45 - 50°C and aseptically add 5% v/v sterile defibrinated blood. Mix thoroughly, avoiding								
air bubbles and pour into sterile Petri plates.								
Principle:								
Tryptose is the source of nitrogen, carbon and amino acids in Tryptose Blood Agar Base. Meat extract								
B provides additional nitrogen, Agar is the solidifying agent. Sodium chloride maintains osmotic								
balance. Supplementation with 5-10% blood provides additional growth factors for fastidious								
microorganisms, and is used to determine hemolytic patterns of bacteria. QC Tests – (I)Dehydrated Medium								
	d Medium							
Colour :		Cream to yellow						
Appearance		Homogeneous Free Flowing powder						
(II)Rehydrated medium								
pH (post autoclaving/heating) : 7.2 ± 0.2								
Colour (post autoclaving/heating): a) Basal medium: Yellow								
b) With addition of 5% v/v defibrinated sterile blood :Cherry red.								
Clarity (post autoclaving/heating) : a) Clear to slightly opalascent b) Opaque								
(III)Q.C. Test Microbiological Cultural characteristics observed after 18 – 48 hrs.at 35-37°C.								
MICROORGANISM (ATCC)			GROWTH W/O		W BLOOD	Haemolysis		
			BLOOD		W BLOOD			
Neisseria meningitidis (13090)			Good -		Luxuriant	None		
			luxuriant		Luxuriani	INOTIE		
Stanhylococo	ıc (25023)			Luvuriant	Beta/Gamma			
Staphylococcus aureus (25923			luxuriant		Luxuriant	Deta/Gamma		
Staphylococcus epidermidis			Good -		Luxuriant	Gamma		
(12228)			luxuriant		Laxuriant	Garrina		
Streptococcus pneumoniae			Fair-Good		Good	Alpha		
	(6303)			۳	2004	, uprid		
Streptococcus pyogenes			Fair-Good	d	Good	Beta		
(19615)			=== ====	_				
Precautions :	1. For L	aboratory l	Jse.					
2. Follow proper, established la						procedures in handlin	g and disposing of	
	infectious materials.							
Limitations: 1. Since the nutritional requirements of organisms vary, some strains may be								
	encountered that fail to grow or grow poorly on this medium.							
Use:			_					
haemolytic reactions. Storage: Dehydrated medium- below 30°C Prepared medium- Be						madium - Rotwoon 2 to	8°C	
	500 gm. bottle							
	Reconsti Quantity		on I		H (2E0C)	Cupplement	Sterilization	
				l	H (25°C)	Supplement	Stermzation	
	tution	Preparatio		7 .	2 + 0 2	5% v/v sterile	1210C /1E min	
B342	33g/l	15.1	LJL	/	2 ± 0.2		121ºC /15 min.	
1						defibrinated blood.		

Refer disclaimer Overleaf

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Disclaimer:

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