BIOMARK Laboratories-INDIA

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TECHNICAL SHEET

B117	BLOOD AGAR BASE NO.2 (WITH 1.2% AGAR)						
Formula							
Ingredients:	gms/lit.						
Proteose peptone	15.00						
Liver extract	2.50						
Yeast extract	5.00						
Sodium chloride	5.00						
Agar	12.00						
Final pH (at 25°C)	: 7.4 + 0.2						

Directions:

Suspend 19.75 grams in 500 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 and aseptically add 7% v/v sterile defibrinated blood. Mix well and pour into sterile Petri plates

For Brucella species: Add rehydrated contents of 1 vial of Brucella Selective Supplement (BF012) to 500 ml sterile molten base.

For Campylobacter species: Add rehydrated contents of 1 vial of Campylobacter Supplement - I (BF013) or Campylobacter Supplement - II, Modified (BF014) or Campylobacter Supplement - III (Bf015) or Campylobacter Growth Supplement (BF016) to 500 ml sterile molten base.

For Streptococcus species: Add rehydrated contents of 1 vial of Strepto Supplement (BF017) to 500 ml sterile molten base.

Principle:

Proteose Peptone is the nitrogen source for Blood Agar Base No. 2 while Yeast Extract and Liver Digest provide essential carbon, vitamin, nitrogen and amino acids sources. Sodium Chloride to maintain osmotic balance.Blood Agar Bases are relatively free of reducing sugars, which have been reported to adversely influence the hemolytic reactions of beta-hemolytic streptococci.

Supplementation with blood (5-10%) provides additional growth factors for fastidious microorganisms and is the basis for determining hemolytic reactions. Hemolytic patterns may vary with the source of animal blood or type of base medium used.

QC Tests - (I)Dehydrated Medium											
Colour:			C	Cream to yellow							
Appearance :				Homogeneous Free Flowing powder							
(II)Rehydrated medium											
pH (post	pH (post autoclaving/heating):				7.4 ± 0.2						
Colour (Colour (post autoclaving/heating):				A) Basal medium : Light amber to yellow						
					B) After addition of 7% sterile defibrinated blood: Cherry red.						
Clarity (post autoclaving/heating):			٠,	A) Clear to slightly opalescent							
(III)Q.C. Test Microbiological				B) Opaque							
			nd with	added 5.70/ starile defibringted blood after an insubstitut at 25					scubation at 2E		
	Cultural characteristics observed with added 5-7% sterile defibrinated blood, after an incubation at 35-37°C for 18-48 hours.										
	MICROORGANISM (ATCC)				GROWTH		HAEMOLYSIS				
	Neisseria meningitides (13090)				Good to li	uxuriant	none				
	Staphylococcus pneumoniae (6303)				Good to luxuriant		alpha				
	Streptococcus pyogenes (19615)				Good to luxuriant		beta				
	Staphylococcus aureus (25923)				Good to li	uxuriant	beta				
Precautions: 1. For Laboratory Use.											
		2. Follow proper, established laboratory procedures in handling and disposing of									
	infectious materials.										
Limitations: 1. Since the nutritiona				I requirements of organisms vary, some strains may be							
				o grow or grow poorly on this medium.							
						nit maximum recovery of fastidious pathogenic microorganisms					
	without interfering with their haemolytic reactions after addition of blood.										
Storage :											
Packing:											
Product profile:		Reconstitution Quantity or		-			Supplement	Sterilization			
					(500g)				<u> </u>		
B117		39.5 g/l	1	2.65	8 L	7.4 <u>+</u> 0.2	7% v/v sterile	121	°C / 15 minutes		
							defribinated				
							blood.				