## **BIOMARK Laboratories-INDIA**

### www.biomarklabs.com

## **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 <u>+</u> 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 ( $121^{\circ}$ C) for 15 minutes. Cool to 45 -  $50^{\circ}$ 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.

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QC Tests - (I)Dehydrated Medium									
	Colour:		Crea	Cream to yellow					
	Appearance :		Homogeneous Free Flowing powder						
(II)Rehydrated medium									
	pH (post autoclaving/heating) :			$7.4 \pm 0.2$					
				A) Basal medium : Light amber to yellow					
	3, 3, 3,			B) After addition of 7% sterile defibrinated blood: Cherry red.					
	Clarity (post autoclaving/heating):			A) Clear to slightly opalescent					
	, (1			B) Opaque					
(III)Q.C. Test Microbiological									
	Cultural chara	characteristics observed with added 5-7% sterile defibrinated blood, after an incubation at 35-							
	37°C for 18-48								
	MICROORGANISM (ATCC )			GROWTH	HAEMOLYSIS				
	Neisseria meningitides (13090)			Good to luxuriant	none				
	Streptococcus pneumoniae (6303)			Good to luxuriant	alpha				
	Streptococcus pyogenes (19615)			Good to luxuriant	beta				
	Staphylococcus aureus (25923)			Good to luxuriant	beta				
<b>Precautions:</b> 1. For Laboratory Use		€.	•	<u> </u>					
		2. Follow proper, established laboratory procedures in handling and disposing of							
infectious materials.				, ,	J				
Lim	<b>Limitations</b> : 1. Since the nutritional			Il requirements of organisms vary, some strains may be					
	encountered that fail to grow or grow poorly on this medium.								
Use: Specially devised to per				ermit maximum recovery of fastidious pathogenic microorganisms					
	without interfering with their haemolytic reactions after addition of blood.								

Refer disclaimer Overleaf

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Storage :	Dehydrated medium- below 30°C Prepared medium- Between 2 to 8°C.								
Packing:	500 gm. bottle								
Product profile: Reconstitution Qu		Quantity on	pH (25°C)	Supplement	Sterilization				
		Preparation (500g)							
B117	39.5 g/l	12.658 L	7.4 <u>+</u> 0.2	7% v/v sterile	121°C / 15 minutes				
				defribinated					
				blood.					

#### 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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