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TECHNICAL SHEET

B172 DIAGNOSTIC SENSITIVITY TEST AGAR (D.S.T. AGAR)				
Formula				
Ingredients:	gms/lit.			
Proteose peptone	10.00			
Veal infusion solids	10.00			
Dextrose	2.00			
Sodium chloride	3.00			
Disodium phosphate	2.00			
Sodium acetate	1.00			
Adenine sulphate	0.01			
Guanine hydrochloride	0.01			
Uracil	0.01			
Xanthine	0.01			
Aneurine	0.00002			
Agar	15.00			
Final pH (at 25°C): 7.4 + 0.2				

Directions

Suspend 43.04 grams in 1000 ml distilled water. Heat to boiling to dissolve the medium completely. Sterilize by autoclaving at 15 lbs pressure (121°C) for 15 minutes. For blood agar, cool the base to 45-50°C and add 7% v/v sterile defibrinated horse blood aseptically. Mix well with gentle rotation and pour into sterile Petri plates.

Principle:

Peptone is the source of nitrogen. The medium is nutritionally rich due to presence of amino acid bases and glucose. The salts present, help in avoiding sudden pH shifts due to acid production which might affect the susceptibility test and haemolytic reactions and the MIC values of pH susceptible antimicrobials. Aneurine acts as vitamin source. Addition of the bases like adenine, guanine, uracil and xanthine improve the antibiotic testing performance of the medium. Agar is the solidifying agent.

solidifying agent.					
QC Tests – (I)Dehydrated Medium					
Colour:	Cream to yellow				
Appearance :	Homogeneous Free Flowing powder				
(II)Rehydrated medium					
PH (post autoclaving/heating):	7.4 ± 0.2				
Colour (post autoclaving/heating):	A) Basal medium: Cream to medium amber B) After addition of blood: Cherry red				
Clarity (post autoclaving/heating):	A) Clear to slightly opalescent B) Opaque				
(III)Q.C. Test Microbiological					
Cultural characteristics observed after	18-24 hrs.at 35-37°C.				
MICROORGANISM (ATCC)	GROWTH (WITHOUT ANTIBIOTICS)				
Escherichia coli (25922)	Luxuriant				
Micrococcus luteus (10240)	Luxuriant				
Neisseria meningitides (13090)	Luxuriant (with the addition of blood)				
Proteus mirabilis (25933)	Luxuriant				
Salmonella typhi (6539)	Luxuriant				
Shigella flexneri (12022)	Luxuriant				
Staphylococcus aureus (25923)	Luxuriant				
Streptococcus faecalis (29212)	Luxuriant				
Streptococcus pyogenes (19615)	Luxuriant (with the addition of blood)				
Streptococcus pneumoniae (6303)	Luxuriant(with the addition of blood)				

Refer disclaimer Overleaf

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TECHNICAL SHEET

1. For Laboratory Use.					
2. Follow proper, established laboratory procedures in handling and disposing of					
infectious materials.					
1. Since the nutritional requirements of organisms vary, some strains may be					
encountered that fail to grow or grow poorly on this medium.					
2.Inoculum density affects inhibtion zone. Heavy inoculum may result in smaller zones					
3. The salts present, helps in avoiding sudden pH shifts due to acid production, which					
might affect the susceptibility test and haemolytic reactions and the MIC values of pH					
susceptible antimicrobials.					
For antibiotic sensitivity testing of fastidious pathogens such as Neisseria, Streptococcus					
and Haemophillus species with blood enrichments.					
Dehydrated medium- below 30°C Prepared medium- Between 2 to 8°C.					
500 gm bottle					
Reconstitution	Quantity on	pH (25°C)	Supplement	Sterilization	
	Preparation (500g)				
	11.617 L	7.4 ± 0.2	7% v/v sterile	121°C / 15 minutes	
]	-		defibrinated	,	
			horse blood		
	2. Follow propinfectious mate 1. Since the encountered the 2.Inoculum der while scanty grows 3. The salts premight affect the susceptible anti For antibiotic so and Haemophill Dehydrated me 500 gm bottle Reconstitution	2. Follow proper, established lab infectious materials. 1. Since the nutritional requiremencountered that fail to grow or grow 2. Inoculum density affects inhibtion while scanty growth may result in eras. The salts present, helps in avoid might affect the susceptibility test susceptible antimicrobials. For antibiotic sensitivity testing of fand Haemophillus species with blood Dehydrated medium-below 30°C Practices of the property of the pro	2. Follow proper, established laboratory proceinfectious materials. 1. Since the nutritional requirements of organ encountered that fail to grow or grow poorly on the 2. Inoculum density affects inhibition zone. Heavy while scanty growth may result in enlarged zones. 3. The salts present, helps in avoiding sudden phromight affect the susceptibility test and haemolytis susceptible antimicrobials. For antibiotic sensitivity testing of fastidious path and Haemophillus species with blood enrichments. Dehydrated medium-below 30°C Prepared medium 500 gm bottle Reconstitution Quantity on pH (25°C) Preparation (500g)	2. Follow proper, established laboratory procedures in handling infectious materials. 1. Since the nutritional requirements of organisms vary, so encountered that fail to grow or grow poorly on this medium. 2. Inoculum density affects inhibition zone. Heavy inoculum may rewhile scanty growth may result in enlarged zones. 3. The salts present, helps in avoiding sudden pH shifts due to admight affect the susceptibility test and haemolytic reactions and the susceptible antimicrobials. For antibiotic sensitivity testing of fastidious pathogens such as Neand Haemophillus species with blood enrichments. Dehydrated medium- below 30°C Prepared medium- Between 2 to 500 gm bottle Reconstitution Quantity on pH (25°C) Supplement Preparation (500g) 43.04 g/l 11.617 L 7.4 ± 0.2 7% v/v sterile defibrinated	

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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