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TECHNICAL SHEET

B1106 B.C.P. – D.C.L.S. AGAR							
Formula							
Ingredients:	gm	s/lit.					
Peptic digest of animal tissue 5.00							
Casein enzymichy							
Yeast extract	Yeast extract 3.00						
Meat extract B# 3.00							
Lactose 7.50							
Sucrose 7.50							
Sodium citrate 10.00							
Sodium chloride 5.00							
Sodium thiosulphate 5.00							
Sodium deoxycholate 2.50							
Bromo cresol purple 0.02							
Agar	14	1.00					
# Equivalent to be	eef Extract						
Final pH (at 25°C): 7.2 <u>+</u> 0.2							
Directions :							
Suspend 67.5 gms. in 1000 ml. distilled water. Boil with frequent agitation to dissolve the medium							
completely. DO NOT AUTOCLAVE or OVERHEAT. Cool to 45°C and pour into sterile petri plates.							
Principle:							
Medium constituents like peptic digest of animal tissue, casein enzymichydrolysate, beef extract and yeast							
extract supply essential nutrients for the growth of the bacteria. Citrate and deoxycholate suppress							
coliforms and gram-positive bacteria. Inclusion of two sugars lactose and sucrose permits the formation of							
yellow colonies by the organisms that ferment either lactose or sucrose or both. Bromo cresol purple is							
the pH indicator. Proteus, Salmonella and Shigella species develop colourless colonies on this medium. QC Tests – (I)Dehydrated Medium							
Colour :	Beige						
Appearance :	Homogeneous Free Flowing powder						
(II)Rehydrated me	liomogen	iomogeneous rice norming powder					
pH (post autocla	7.2 ± 0.2						
	Purple						
(III)Q.C. Test Microbiological Cultural characteristics observed after 18 - 24 hrs at 45°C.							
MICROORGANISM (ATCC)						UR OF COLONY	
Salmonella typhimurium (14028)			Luxuriant			colourless	
			Luxuriant			colourless	
Shigellaflexneri (12022)			Luxuriant			colourless	
Shigellasonnei (25931)			Luxuriant			colourless	
			ood			colourless	
			or			-	
Escherichia coli (25922)			None – poor			Yellow	
Proteus vulgaris (13315)			None – poor		N	o swarming	
			•			-	
Precautions: 1. For Laboratory Use.							
	2. Follow proper, established laboratory procedures in handling 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.							ins may be
							- ,
Use: For isolation of Salmonella, Shigella and Arizona species.							
Storage :	Dehydrated medium-below 30°C Prepared medium- Between 2 to 8°C.						
Packing:	500 gm. bottle						
						Sterilization	
r roudet prome:	Neconstitution	Preparatio		Pii (2.	<i>J</i> ()	Supplement	Stermzation
B1106	67.5g/l	7.4		7.2 ±	U 3	NIL	DO NOT AUTOCLAVE
P1100	07.3g/i	7.4	U/L	/.2 =	0.2	INTL	or OVERHEAT.
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Rev: December 2020