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B475 CR	RYSTAL VIOLET	LACTOSE	AGAF	2				
Formula								
Ingredients : gms/lit.								
Proteose peptone 5.00								
Meat extract B# 3.00								
Lactose 10.00								
Crystal violet 0.0033								
Agar 15.00								
# Equivalent to Beef extract								
Final pH (at 25°C) : 6.8 <u>+</u> 0.2								
Directions :								
Suspend 33 grams in 1000 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 to 45-50°C.								
Mix well and pour into sterile Petri plates.								
Principle :								
Crystal violet lactose agar was recommended for the differentiation of pure cultures of pathogenic								
from nonpathogenic strains of Staphylococci. Crystal violet is markedly inhibitory to Staphylococci.								
A fair growth can be obtained at a 1:300,000 concentration of the dye when the medium is								
inoculated heavily. So this medium is used for study of pure cultures where a mass inoculation can								
be used rather than for primary isolation. Staphylococcal colonies show different colours when								
cultured on Crystal violet Lactose Agar.								
OC Tests – (I)Dehvdrated Medium								
Colour :			Light tan					
Appearance :				Homogeneous Free Flowing powder				
(II)Rehvdrated medium								
pH (post autoclaving/heating) :				6.8 ± 0.2				
Colour (post autoclaving/heating) :				Purple				
Clarity (post autoclaving/heating) :				Clear to slightly opalescent				
(III)O.C. Test Microbiological					onghery open			
Cultural characteristics observed after 40 - 48 hrs. at 35- 37°C								
MICROORGANISM (ATCC)				th COLOUR OF COLONY				
Escherichia coli (25922)			Luxuriant		002	Purple		
Staphylococcus aureus (25923)			Fair – good			Light Yellow		
Staphylococcus enidermidis (12228)			Fair – good		Purnle	Purple /very slight vellow		
Streptococcus pyogenes (19615)			Nono-poor			Colourless		
Procentions 1	For Laborator							
	2. Follow proper, established laboratory procedures in handling and disposing o							
in								
imitations 1 Since the nutritional requirements of organisms yany some strains may be								nav ho
encountered that fail to grow or grow poorly on this medium								nay be
For differentiation of pure cultures of nathogenic and nonnathogenic								
Stanbylococcifrom clinical and food camples								
Storage : Debydrated medium, below 30°C Propared medium, Between 2 to 9°C								
Derivated medium- below 50 C Frepared medium- belween 2 to							L.	
Product Deconstitution Quantity on DH (250C) Supplement						Chault	Instian	
product Reconstitution Quant				0 ~)	рп (∠5°С)	Supplement	Sterili	zation
	220/			uy)	60100	NII	12100	/ 1 F
D4/J	339/T	1	5.15L		ο.o ± 0.2	INII	121°C	1 12
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Disclaimer:

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Rev: December 2020