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B1296	M E AGAR						
Formula							
Ingredients :	gms/lit.						
Peptone	10.00						
Sodium chloride	15.00						
Yeast extract	30.00						
Esculin	1.00						
Actidione	0.05						
Sodium azide	0.15						
Agar	gar 15.00						
Final pH (at 25°C) : Self							
Directions :							
Suspend 71.2 gms. in 1000 ml. distilled water. Boil to dissolve the medium completely. Sterilize by							
autoclaving at 15 lbs pressure (121° C) for 15 minutes.							
Principle :							
Peptone and Yeast extract provide essential nutrients Sodium chloride maintains osmotic equilibrium.							
Sodium azide inhibits growth of gram negative organisms and permits the growth of faecal Streptococci.							
Esculin hydrolyzers hydrolyze the glycoside esculin to esculetin and dextrose.							
Following modifications make the medium more selective ;							
1. Add 0.75 gm of indoxyl- β -D-glucoside to 1000ml of distilled water containing 71.2 gm of dehydrated							
Basal medium (M E Agar). Autoclave at 15lbs pressure (121°C) for 15 minutes.							
2. Add 0.24 gm Nalidixic acid in 5ml sterile distilled water (Add a few drops of 0.1 N NaOH to dissolve if							
required) Add this solution to autoclaved medium.							
3. Add 0.02 gm triphenyl tetrazolium chloride to the medium and mix.							
The MEI Medium is similar to the ME Medium except that it contains a reduced amount of							
triphenyltetrazaol	ium chloride (TT	C) and co	ontains a su	bstrate, indox	yl β-D-glucoside	that turns blue when	
cleaved by an enz	yme present in	enterocod	cci (β-glucos	sidase). All co	olonies with any b	ue halo are recorded	
as enterococci, regardless of colony color. Magnification with a dissecting microscope is used for counting							
to give maximum visibility of colonies. It gives faster results (within 24 hrs.) at 41°C instead of 48 hrs. at							
37°C for ME Agar.		-		-	-		
QC Tests - (I)Deh	ydrated Medium						
Colour :			Cream to light yellow				
Appearance :			Homogeneous Free Flowing powder				
(II)Rehvdrated m							
nH (nost autoclaving/heating)			Self				
Colour (post autoclaving/heating) :			Cream to vellow				
Clarity (post autoclaving/heating) :			Clear to slightly opalescent				
(III) C Test Microbiological							
Cultural characteristics observed after 48 brs at 25, 2790							
				<u>57 C.</u>			
Enterococcus faccalis (20212)		Luxuriant					
	aecalis (29212)		Luxunani				
Dragoutiona	1 Farlaharata	m / 1100					
recautions: 1. For Laboratory USE.							
2. Follow proper, established laboratory procedures in handling and disposing of							
Limitations : 1. Since the nutritional requirements of organisms vary, some strains may be encountered that fail to grow or grow poorly on this medium. Use : For isolation and enumeration of enterococci from potable, estuarine, marine and							
						ns may be	
						e, marine and	
•	shellfish growing waters.						
Storage :	Dehydrated medium- below 8°C Prepared medium – Between 2 to 8°C.						
Packing :	500 gm. bottle				1		
Product profile:	Reconstitution	Quantity	on	pH (25°C)	Supplement	Sterilization	
		Preparati	ion (500g)				
B1296	71.2g/l	7.	022L	Self	NIL	121°C / 15 minutes	

Refer Disclaimer overleaf

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Disclaimer:

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