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B511	EGG YOLK AGAR BASE							
Formula								
Ingredients :	gms/	lit.						
Proteose peptone 40.00								
Disodium phospha	phosphate 5.00							
Monopotassium p	Im phosphate 1.00							
Sodium chloride	2.00							
Magnesium sulpha	ate 0.10							
Glucose	2.00							
Hemin	0.005							
Agar	25.00							
Final pH (at 25°C) : 7.6 <u>+</u> 0.2								
Directions :						<u> </u>		
Suspend 75.10 grams in 900 ml distilled water. Heat to boiling to dissolve the medium completely. Dispense in 90 ml amounts and sterilize by autoclaving at 15 lbs pressure (121°C) for 15 minutes. Cool to 45-50°C and add 10 ml of sterile end volk emulsion (BE003) per 90 ml of medium. Mix well and pour into								
sterile Petri plates								
Principle :								
Proteose peptone	provide the essential nu	trients along with carbo	naceous and	d nitrog	enous	substances.		
Phosphates buffer the medium whereas sodium chloride maintains the osmotic equilibrium. Magnesium sulphate serves as a source of divalent cations along with sulphates. Glucose serves as a source of energy. Hemin help to enhance the growth of anaerobic organisms. Organisms producing lecithinase break down lecithin present in the egg yolk emulsion producing an insoluble opaque precipitate around the colonies. Lipase-producing organisms break down free fatty acids (in the egg yolk emulsion) forming an iridescent sheen on the surface of the colonies.								
QC Tests - (I)Deh	vdrated Medium							
Colour :	•	Cream to vellow	Cream to vellow					
Appearance :		Homogeneous Free Flov	Homogeneous Free Flowing, powder					
(II)Rehvdrated m	edium							
pH (post autocla	ving/heating):	7.6 ± 0.2						
Colour (post a	utoclaving/heating):	a) Basal medium : Cream to light yellow b) After addition Egg Yolk : Cream to yellow						
Clarity (post a	utoclaving/heating):	a) Clear to slightly opalescent b) Opaque						
(III)Q.C. Test M	icrobiological							
Cultural charac	cteristics observed with ad	ded Egg yolk emulsion (BF003), afte	r an inc	ubation	at 35-37°C		
for 48-72 hour	s when incubated anaerob	ically.						
MICROORGANIS	M (ATCC)	GROWTH	LEC	LIP	PRO			
Clostridium botu	llinum (25763)	Good-luxuriant	-	-	+			
Clostridium butyricum (13732)		Good-luxuriant	-	-	+			
Clostridium perfringens (13124)		Good-luxuriant	+	-	-			
Clostridium sp	orogenes (11437)	Good-luxuriant	-	+	+			
Bacteroides fra	Bacteroides fragilis (25285) Good-luxuriant							
Key: LEC = Lecithinase production, opaque precipitate around colonies.								
LIP = Lipase production, iridescent sheen on the surface colonies & medium.								
PRO = Proteolytic activity, clear zones surrounding colonies.								
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.							
Use :	For isolation and identification of Clostridia and other anaerobic microorganisms.							
Storage :	Dehydrated medium- belo medium – Use as fresh	n- below 30°C in cool dry place, away from bright light. Prepared fresh as possible.						
Packing :	500 gm. bottle							
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Product profile:	Reconstitution	Quantity on	pH (25°C)	Supplement	Sterilization
		Preparation (500g)			
B511	75.10 g/l	6.66L	7.6 ± 0.2	Nil	121°C / 15 minutes

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