

TECHNICAL SHEET

B582	LEAD ACETATE AGAR					
Formula						
Ingredients :		gms/lit.				
Peptic digest of animal tissue		15.00				
Proteose peptone		5.00				
Dextrose		1.00				
Lead acetate		0.20				
Sodium thiosulphate		0.08				
Agar		15.00				
Final pH (at 25°C) : 6.6 ± 0.2						
Directions :						
Suspend 36.28 gms. in 1000ml. distilled water. Boil to dissolve the medium completely. Dispense into test tubes sterilize by autoclaving at 15 lbs pressure (121°C) for 15 minutes. Allow to cool the tubes in a slanted position to obtain with generous butts. Inoculate pure culture by surface streaking slant and stabbing the butt.						
Principle :						
Peptic digest of animal tissue, proteose peptone, and dextrose provides all essential acts as an indicator for hydrogen sulphite production. Production of gas from dextrose is indicated by the presence of bubbles in the butt.						
QC Tests - (I) Dehydrated Medium						
Colour :		Light yellow				
Appearance :		Homogeneous Free Flowing powder				
(II) Rehydrated medium						
pH (post autoclaving/heating) :		6.6 ± 0.2				
Colour (post autoclaving/heating) :		Medium amber				
Clarity (post autoclaving/heating) :		Slightly opalescent				
(III) Q.C. Test Microbiological						
Cultural characteristics observed after 18 -24 hrs. at 35-37°C.						
MICROORGANISM (ATCC)	GROWTH	H ₂ S PRODUCTION	GAS PRODUCTION			
Enterobacter aerogenes (13048)	Luxuriant	-	+			
Escherichia coli (25922)	Luxuriant	-	+			
Salmonella typhi (6539)	Luxuriant	+	±			
Salmonella typhimurium (14028)	Luxuriant	+	-			
Shigella dysenteriae (13313)	Luxuriant	-	-			
Shigella flexneri (12022)	Luxuriant	-	-			
Salmonella paratyphi A	Luxuriant	-	-			
Salmonella paratyphi B	Luxuriant	+	-			
Key : H ₂ S + = browning of the medium						
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 detection of hydrogen sulphite producing enteric bacteria.				
Storage :		Dehydrated medium- below 30°C Prepared medium- Between 2 to 8°C.				
Packing :		500 gm bottle				
Product profile:		Reconstitution	Quantity on Preparation (500g)	pH (25°C)	Supplement	Sterilization
B582	36.28 g/l	13.78 lit	6.6 ± 0.2	Nil	121°C/15 min	

Refer disclaimer Overleaf

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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