BIOMARK Laboratories-INDIA

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TECHNICAL SHEET

B212 K	LIGLER IRON AGAR		
Formula			
Ingredients:	gms	s/lit.	
Peptone	15.0	00	
Meat Extract B#	3.00)	
Yeast extract	3.00)	
Proteose peptone	5.00)	
Lactose	10.0	00	
Dextrose	1.00)	
Ferrous sulphate	0.20)	
Sodium chloride	5.00)	
Sodium thiosulphate	0.30)	
Phenol red .	0.02	24	
Agar	15.0	00	
#- Equivalent to Bee	ef extract		
Final pH (at 25°C):	7.4 + 0.2		
Directions :			
Suspend 57.52 gra	ms in 1000 ml disti	lled water. Heat to boiling to dissolve the medium	
completely. Mix well	and distribute into tul	pes. Sterilize by autoclaving at 15 lbs pressure (121°C)	
		anted position to form slopes with about 1-inch butts.	
		ared medium. Do not use screw capped tubes or bottles.	
Principle :			
Kligler Iron Agar cor	mbines the principles of	of Russell double sugar agar and lead acetate agar into	
		differentiation of the gram-negative bacilli both by their	
		d to produce hydrogen sulfide. Meat Extract B, Yeast	
		ovide nitrogen, vitamins and minerals. Ferrous sulfate	
		of hydrogen sulfide production. Phenol red is the pH	
indicator. Sodium chloride maintains the osmotic balance of the medium. Agar is the solidifying			
agent.		, J , J	
QC Tests - (I)Dehydr	rated Medium		
Colour		Light vellow to nink	

QC	Tests - (I)Dehydrated Medium	
	Colour:	Light yellow to pink
	Appearance :	Homogeneous Free Flowing powder
(II	Rehydrated medium	
	pH (post autoclaving/heating):	7.4 ± 0.2
	Colour (post autoclaving/heating):	Reddish orange to red
	Clarity (post autoclaving/heating):	Clear to slightly opalescent
/TT	I)O C Test Microbiological	

(III)	0.C.	Test	Micro	bio	logical
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(III)Q.C. Test Microbiological					
Cultural characteristics observed after18 – 48 hrs at 35-37°C.					
MICROORGANISM (ATCC)	GROWTH	SLANT	BUTT	GAS	H ₂ S
Citrobacter freundii (8090)	Luxuriant	Α	Α	+	+
Escherichia coli (25922)	Luxuriant	Α	Α	+	ı
Enterobacter aerogenes (13048)	Luxuriant	Α	Α	+	ı
Klebsiella pneumoniae (13883)	Luxuriant	Α	Α	+	ı
Proteus vulgaris (6380)	Luxuriant	K	Α	-	+
Salmonella enteritidis (13076)	Luxuriant	K	Α	+	+
Salmonella paratyphi A (5006)	Luxuriant	K	Α	+	ı
Salmonella schottmuelleri (10719)	Luxuriant	K	Α	+	+
Salmonella typhi (6539)	Luxuriant	K	Α	-	+
Shigella flexneri (12022)	Luxuriant	K	Α	-	-
Pseudomonas aeruginosa (27853)	Luxuriant	K	K	-	-

Key: A = acid production (yellow)

K = alkaline reaction (red)

+ = positive or blacking - = negative reaction (no change)

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1. For Laboratory Use.					
2. Follow proper, established laboratory procedures in handling and disposing of					
infectious materials.					
1. Since the nutritional requirements of organisms vary, some strains may be					
encountered that fail to grow or grow poorly on this medium.					
For differential identification of gram-negative enteric bacilli on the basis of fermentation					
of dextrose, lactose and H ₂ S production.					
Dehydrated medium- below 30°C Prepared medium- Between 2 to 8°C.					
500 gm. bottle					
econstitution	Quantity on	pH (25°C)	Supplement	Sterilization	
	Preparation (500g)				
57.52 g/l	8.69L	7.4 ± 0.2	NIL	121°C / 15 minutes	
	Follow proper, of fectious material. Since the nutrition of the fection of the fe	Follow proper, established laborator fectious materials. Since the nutritional requirements of acountered that fail to grow or grow por differential identification of gram-net dextrose, lactose and H ₂ S production ehydrated medium- below 30°C Preparation (500g) Quantity on Preparation (500g)	Follow proper, established laboratory procedures if fectious materials. Since the nutritional requirements of organisms variountered that fail to grow or grow poorly on this report differential identification of gram-negative enterior dextrose, lactose and H ₂ S production. Sehydrated medium-below 30°C Prepared medium-below gm. bottle Econstitution Quantity on pH (25°C) Preparation (500g)	Follow proper, established laboratory procedures in handling and fectious materials. Since the nutritional requirements of organisms vary, some strain accountered that fail to grow or grow poorly on this medium. For differential identification of gram-negative enteric bacilli on the dextrose, lactose and H ₂ S production. Sehydrated medium- below 30°C Prepared medium- Between 2 to 20 gm. bottle Econstitution Quantity on pH (25°C) Supplement Preparation (500g)	

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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