BIOMARK Laboratories-INDIA

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TECHNICAL SHEET

B1475 UREA IN	DOLE MEDIUM						
Formula							
Ingredients:	gms/lit.						
L- Tryptophan	3.00						
Sodium chloride 5.00							
Dipotassium phosphate 1.00							
Monopotassium phosphate	1.00						
Urea	20.00						
Phenol red	0.025						
Final pH (at 25°C):6.8+ 0.2							
Directions:							
Suspend 30.00 grams in 1000 ml distilled water. Mix well. Add 10 ml of 95% ethanol. Dissolve the medium							
completely. Dispense in 1 - 5 ml amounts into sterile tubes. AVOID OVERHEATING. DO NOT AUTOCLAVE.							
Principle:							
L-Tryptophan is an essential amino acid and is converted to skatole and indole. Sodium chloride maintains the osmotic balance. Potassium phosphates act as a buffer system. Urea is a source of nitrogen for those organisms producing urease. Phenol red is the pH indicator.Indole production is determined by adding a few drops of Kovacs Reagent (BA068). A positive test is indicated by the development of a red color in the reagent layer. Tryptophan deaminase (TDA) is demonstrated by adding to a 24 hours culture a few drops of a 30% solution, diluted 1:3. of iron perchloride. The appearance of a brown or red-brown color indicates a positive TDA.							
QC Tests – (I)Dehydrated Medium							
Colour:		Light pink					
Appearance:		Homogeneous Free Flowing powder					
(II)Rehydrated medium		5 51					

QC	Tests - (I)De	hydrated Medium						
	Colour:		Light pink	Light pink				
	Appearance:		Homogene	Homogeneous Free Flowing powder				
(II)Rehydrated n	nedium						
	pH (post autoclaving/heating) :		6.8 ± 0.2	6.8 ± 0.2				
	Colour (post	autoclaving/heating):	Orange	Orange				
	Clarity (post	autoclaving/heating) :	Clear	Clear				
(II	I)Q.C. Test M	licrobiological						
	Cultural char	acteristics observed after :	18 –24 hrs at 3	5-37°C.				
	MICROORGAN	CROORGANISM (ATCC)		Urease	Indole			
	Escherichia coli (25922)		luxuriant	-	+			
	Klebsiella pneumoniae (13883)		luxuriant	+	-			
	Proteus vulgaris(13315)		luxuriant	+	+			
	Salmonella typhimurium(14028)		luxuriant	-	-			
	*Yersinia enterocolitica (23715)		luxuriant	+	±			
	*Incubate at 30°C for 24 hours							
Pre	ecautions :	 For Laboratory Use. Follow proper, establis materials. 	shed laboratory	procedures in	n handling and	disposing of infectious		

Refer disclaimer Overleaf

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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:	To differentiate micro-organisms especially Enterobacteriaceae on the basis of their abili							
	to hydrolyze urea and indoleproduction. This medium is used for the presumptive							
	identification of Yersinia enterocolitica by ISO 10273.							
Storage:	Dehydrated medium-at 2to 8 ° C Prepared mediums– Between 2 to 8°C.							
Packing:	500 gm. bottle							
Product profile:	Reconstitution	Quantity on	pH (25°C)	Supplement	Sterilization			
		Preparation (500g)						
B1475	30g/l	16.66 L	6.8 ± 0.2	NIL	Heating			
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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. The information contained in this publication is based on our in-house studies and market performance and is to the best of our knowledge true and accurate. BIOMARK LABORATORIES reserves the right to make changes to specifications and information related to the products at any time. Products are not intended for human or animal or therapeutic use but for laboratory, diagnostic, research or further manufacturing use only, unless otherwise specified. Statements contained herein should not be considered as a warranty of any kind, expressed or implied, and no liability is accepted for infringement of any patents.

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