## **BIOMARK Laboratories-INDIA**

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## **TECHNICAL SHEET**

BH1273 SABOURAUD DEXTROSE AGAR							
Formula							
Ingredients:			gms/lit.				
Mixture of Peptone &Tryptone (1:1)			10.00				
Dextrose (Glucose	40.00						
Agar			15.00				
Final pH (at 25°C): 5.6 <u>+</u> 0.2							
Directions:							
Suspend 65.0 grams in 1000 ml purified/ distilled water. Heat to boiling to dissolve the medium							
completely. Sterilize by autoclaving at 15 lbs pressure (121°C) for 15 minutes or as per validated							
cycle. Cool to 45-50°C. Mix well and pour into sterile Petri plates.							
Principle:							
Peptone and Tryptone provides carbonaceous, nitrogenous compounds, long chain amino acids,							
vitamins and other essential growth nutrients. Dextrose (Glucose) provides an energy source. High							
dextrose concentration and low pH favors fungal growth and inhibits contaminating bacteria from							
clinical specimens.							
QC Tests - (I)Dehydrated Medium  Colour :			Croom to light vollow				
			Cream to light yellow Homogeneous Free Flowing powder				
Appearance :			nomogeneous riee riowing powder				
(II)Rehydrated medium			5.6 ± 0.2				
pH (post autoclaving/heating) :			Cream to light amber				
Colour (post autoclaving/heating):			Clear to slightly opalescent				
Clarity (post autoclaving/heating) :			Clear to slightly opalescent				
Cultural characteristics observed after 24 – 48 hrs. at 30-35°C.							
Aspergillus niger (16404) Luxuria					<=5 Da	•	
` '		Luxuria			24 -48		
		Luxuria			24 -48		
Trychophyton rubrum (28191) Go		Good	20 -2	5 °C	<=5 Da	ıys	
		Luxuria			24 -48	hrs	
Escherichia coli (25922)		Good*	30 -3		24 -48	hrs	
Escherichia coli (8739)		Good*	30 -3		24 -48	24 -48 hrs	
Escherichia coli (NCTC9002)		Good*	30 -3		24 -48 hrs		
Lactobacillus casei (334) Luxu		Luxuria	nt 30 -3	85 °C	24 -48	hrs	
Key * = inhibited on media with lower pH.							
Precautions: 1. For Laboratory Use.							
2. Follow proper, established laboratory procedures in handling and disposir							
infectious materials. Wear protective gloves/protective clothing/eye						е	
	protection/face pro						
<b>Limitations:</b> 1. For heavily contaminated samples, the media must be supplemented w						ented with	
		bitory agents for inhibiting bacterial growth with lower pH					
2. Avoid overheating a medium with an acidic pH because this often of						en causes a	
	soft medium.  3. Some pathogenic fungi may produce infective spores which are easily						
				•			
	dispersed in air, so examination should be carried out in safety cabinet.						
	4. Further biochemical tests should be carried out for confirmation.						
<b>Use:</b> For the cultivation of yeasts, moulds and aciduric bacteria from phareducts in accordance with the microbial limit testing by harmonic							
products in accordance with the microbial limit testing by harmonized						zea	
Storage :	methodology of USP/EP/BP/JP .  Dehydrated medium - below 30°C & Prepared medium - Between 2 to 8°C						
Storage :	Dehydrated medium- below 30°C & Prepared medium – Between 2 to 8°C. 500 gm. bottle						
Product profile	<b>0</b> 0	рЦ (ЭЕОС)	Cupplant	Ctoriliantian			
Product profile:		Quantity		pH (25°C)	Supplement	Sterilization	
DU1272			ion (500g)	E 6 + 0 2	Nil	1210C/1Emin	
BH1273	65.0 g/l  7	'.69 L		$5.6 \pm 0.2$	INII	121°C/15min	

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