## **IOMARK Laboratories-INDIA**

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TECHNICAL SHEET								
B580 DNASE TEST AGAR BASE W/ METHYL GREEN								
Formula								
Ingredients:	gm	s/lit.						
Tryptose 20.00								
Deoxyribonucleic	00							
Sodium chloride								
Methyl green	en 0.05							
Agar	15.	00						
Final pH (at 25°C) : 7.3 <u>+</u> 0.2								
Directions:								
Suspend 42.05 grams in 1000 ml distilled water. Heat to boiling to dissolve the medium completely.								
Sterilize by autoclaving at 15 lbs pressure (121°C) for 15 minutes. Mix well and pour into sterile Petri								
plates.								
Principle :								
Tryptose serves as nitrogenous source for the organisms. DNase produced by microorganisms								
depolymerizes the DNA substrate in the medium. Methyl green fades into a colourless compound								
producing distinct clear zones surrounding colonies (or band/ spot inocula) in an otherwise green coloured								
medium. Methyl green requires a highly polymerized DNA substrate and it combines with polymerized DNA								
forming a stable, green complex at pH 7.5. As hydrolysis progresses, methyl green is released and when								
not combined at this pH it fades and becomes a colourless compound. Therefore clear zones are observed								
QC Tests – (I)Dehydrated Medium								
Colour :			Light yellow to greeenish yellow					
Appearance :			Homogeneous Free Flowing powder					
(II)Rehvdrated medium								
pH (post autoclaving/heating) :			7.3 ± 0.2					
Colour (post autoclaving/heating) :			Green					
Clarity (post autoclaving/heating) :			Clear to slightly opalescent					
(III)O.C. Test Microbiological								
Cultural characteristics observed after 18 –24 hrs at 35-37°C								
MICROORGANISM (ATCC.)			GROWTH		DNASE ACTIVITY			
Staphylococcus aureus (25923)			Luxuriant		positive, clear halo around the growth			
Staphylococcus enidermidis (12228)			Luxuriant		negative	ative reaction		
Streptococcus pyogenes (19615)			Luxuriant		positive, clear halo around the growth			
Serratia marcescens (8100)			Luxuriant		positive, clear halo around the growth			
Precautions :	ns · 1 For Laboratory Lise							
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	2. Follow proper, established iduoratory procedures in indimining and disposing						ig and disposing of	
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Ennitations . If Since the nutritional requirements of organisms vary, some strains may						ne strains may be		
It is recommended for the detection of deevuribenuclease activity of bacteria						of bactoria and fungi		
lit is recommended for the detection of deoxymbonucledse activity of Dacteria and land especially for identification of nathogonic Stanbylococci						o bacteria anu iuliyi,		
Storago I	torage I Debydrated medium- below 30°C Propared medium- Between 2 to 9°C						<u>۹۹</u>	
Dacking :	Engurateu medium- below 50°C Prepareu medium- belween 2 lo 8°C.							
Product profile: Deconstitution Quantity on Large (2500)				(2500)	Cupplement	Ctorilization		
Product profile:	Reconstitution	Quantity on		рн	(25°C)	Supplement	Stermzation	
DEOO		Preparati		_	2 . 0 2	NU	11000 += 12100) (	
6280	42.05 g/l	11	.89 L	7.	3 <u>+</u> 0.2	INII	118°C to 121°C) for	
							15 minutes.	

## Disclaimer:

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