BIOMARK Laboratories-INDIA

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TECHNICAL SHEET

B1047 FUR	UNCULOSIS AGAR						
Formula							
Ingredients : gr		gms	/lit.				
Casein enzymichydrolysate 1			.00				
Yeast extract)						
Tyrosine	1.00						
Sodium chloride 2.50							
Agar 15.0			0				
Final pH (at 25°C) : Self							
Directions :							
Suspend 33.5 gms in 1000 ml distilled water. Boil to dissolve the medium completely. Dispense in							
tubes and sterilize by autoclaving at 15 lbs pressure (121°C) for 15 minutes. DO NOT OVERHEAT. Cool							
the tubes to cool in slanted position.							
Principle :							
Furunculosis Agar is formulated as per Griffin et al for detection of Aeromonassalmonicida on the basis							
of production of brownish red pigment. Casein enzymichydrolysate and yeast extract provide							
essential nutrients for growth. Sodium chlrodie maintains osmotic equilibrium. Brownish red							
pigmentation within two to three days of incubation at 22°C is a positive presumptive evidence. For							
the more rapid presumptive test, 0.5 ml of 1% aqueous solution of paraphenylenediamine can be							
applied to the colonies of a 24 hours old culture growing on the surface of the agar slants. Contact							
the reagent with all the growth. After application of the reagent, the tubes should be tipped and							
rotated to spread the reagent to cover the growth on slant, a deep purple colour is seen within 45 to							
90 seconds.							
OC Tests – (I)Dehydrated Medium							
Colour :			Light yellow				
Appearance :			Homogeneous Free Flowing powder				
(II)Rehydrated medium							
pH (post autoclaving/heating) :			Self				
Colour (post autoclaving/heating) :			Light amber				
Clarity (post autoclaving/heating) :			Clear to slightly opalescent				
(III)O.C. Test Microbiological							
Cultural characteristics observed after 18 – 48 hrs at 35-37°C							
MICROORGANISM (ATCC.)			ROWTH COLOUR OF COLONY				
Aeromonassalmonicida (33658)			ood - luxuriant Brownish red				
/ cromonassam							
Precautions '	1 For Laborator	v llse					1
i i ccuations i	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						
Linitations .	encountered that fail to grow or grow poorly on this medium						
Ise For detection of Agromonassalmonicida by means of its brownish re						ed niament	
036.	nor detection of Actomonassamonicida by means of its brownish red pigment						
Storage :	Debudrated medium- below 30°C Prenared medium- Retween 2 to 8°C						
Packing : 500 am bottle							
Product profile	Poconstitution	nstitution Quantity o		nH (250C)	Supplement	Starilization	
Froduct profile:	Reconstitution	Droparat	$\frac{1}{100}$	pri (25°C)	Supplement	Stermzation	
B1047	22 Ea/l	1/			NTI	12100	/ 15
D1047	33.5g/I	14	+.92L	SELF			1 12
Disclaimory						minute	:5

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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