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TECHNICAL SHEET

B004	YEAST NITROGEN B	ASE W/O AMINO ACIDS AND AMMONIUM SULPHATE	
Formula			
Ingredients:		gms/lit.	
Biotin		0.000002	
Calcium pantothenate		0.0004	
Folic acid		0.000002	
Inositol		0.002	
Niacin		0.0004	
p-amino benzoic acid (PABA)		0.0002	
Pyridoxine h	ydrochloride	0.0004	
Riboflavin (vitamin B2)		0.0002	
Thiamine hydrochloride		0.0004	
Boric acid		0.0005	
Copper sulphate		0.00004	
Potassium io	dide	0.0001	
Ferric chloride		0.0002	
Manganese sulphate		0.0004	
Sodium moly	/bdate	0.0002	
Zinc sulphate		0.0004	
Monopotassium phosphate		1.00	
Magnesium sulphate		0.50	
Sodium chloride		0.10	
Calcium chlo	ride	0.10	
Final pH (at	25°C): 4.5 + 0.2		

Directions:

- A. For Carbon Assimilation tests, prepare the broth base in 10X concentration. Dissolve 1.7 gms. in 100 ml. distilled water. Add 5 gms. ammonium sulphate, 10 mg L-Histidine,20 mg DL-methionine and 20 mg DL-tryptophan. Carbon compounds for assimilation test are added in 10X concentration singly or in combination as required.
- B. For Nitrogen Assimilation tests, prepare the medium in 10X concentration. Dissolve 1.7 gms. in 100ml. distilled water. Add 1,0 gms. dextrose, 1 mg L-histidine, 2 mg DL-methionine and 2 mg DL-tryptophan. Add nitrogen compounds for assimilation test in 10X concentration singly or in combination as required. Wickerham employed the following nitrogen sources: ammonbium sulphate 1 gm. potassium nitrate 0.78 gms., urea 0.46 gms. aspargine 1 gm. peptone (gelatin) 1.32 gms.

For A and B, filter sterilize the 10X strength solution, Refrigerate and use as needed. Prepare final medium by aseptically pipetting 0.5 ml. of the 10X sterile medium into 4.5 ml. sterile distilled water. Mix well.

Principle :

Yeast Nitrogen Base without Amino Acids and Ammonium Sulphate is prepared as per Wickerham for classifying the yeasts on the basis of their carbon and nitrogen requirements. Wickerham used following nitrogen sources – ammonium sulphate 1.0 gm /litre, potassium nitrate 0.78 gm/litre, urea 0.46 gm/litre, asparagines 1.0 gm/litre, peptone (gelatin) 1.32 gm/litre.

Yeasts grown on rich medium may carry a reserve of nitrogen in the form of proteins and this may result in erroneous findings. To avoid this, 2 serial transfers in complete medium are recommended.

QC	Tests - (I)Dehydrated Medium	
	Colour:	Cream to off white
	Appearance :	Homogeneous Free Flowing powder
(II)	Rehydrated medium	
	pH (post autoclaving/heating):	4.5 ± 0.2
	Colour (post autoclaving/heating):	Very light yellow to colourless
	Clarity (post autoclaving/heating):	Clear

Refer disclaimer Overleaf

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TECHNICAL SHEET

(III)Q.C. Test M	icrobiological										
	Cultural characteristics observed after 6-7 days at 25 – 30°C.										
MICROORGANIS	MICROORGANISM (ATCC)			N) GRO	ROWTH WITH DEXTROSE						
Kloeckera apiculata (9774)			ne – poor	God	od						
Saccharomyces uvarum (9080)			one – poor	God	od						
Saccharomyce	es cerevisiae (9763) No	one – poor	God	od	d					
Precautions:	Precautions: 1. For Laboratory Use.										
2. Follow proper, estat			blished laboratory procedures in handling and disposing of								
	infectious materia	ctious materials.									
Limitations :	1. Since the nutritional requirements of organisms vary, some strains may be										
		at fail to grow or grow poorly on this medium.									
Use :	For classification of yeasts on the basis of their ability of assimilate carbon and										
	nitrogen compounds.										
Storage :	Dehydrated mediu	ehydrated medium and prepared medium- Between 2 to 8°C.									
Packing:	: 500 gm. bottle										
Product profile:	Reconstitution	Quantit	y on	pH (25	°C)	Supplement	Sterilization				
		Prepara	aration (500g)								
B004	1.7 g/l	294.11	L	4.5 <u>+</u> 0.2		Nil	FILTRATION				

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 BIOMARK LABORATORIES publications.

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