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### **TECHNICAL SHEET**

Phenol RedGalactoseBroth

B1104

Earmula										
Formula		ams/lit								
Ingredients:	-	gms/lit.								
Proteose peptone			10.00							
Meat extract B#			1.00							
Sodium chloride			5.00							
Phenol red		0	0.018							
Galactose5.00										
#- Equivalent to Beef extract										
Final pH (at 25°C	C): 7.4 <u>+</u> 0.2									
Directions :										
Suspend 21gms. in 1000 ml. distilled water Heat to dissolve the medium completely. Dispense in tubes										
containing invert	ed Durham's tubes and	d ste	erilize by autoc	laving at 15 lbs pr	essure (121°C) for	15 minutes.				
Principle :										
Proteose Peptone	e and Meat extract B p	rovi	de the carbon	and nitrogen sour	ces required for go	od growth of				
	organisms. Sodium (									
	icator, turning from re									
the added carbol			,		J					
	QC Tests - (I)Dehydrated Medium									
Colour :	-		Pink							
Appearance :			Homogeneous Free Flowing powder							
(II)Rehydrated n	nedium									
	laving/heating) :		$7.4 \pm 0.2$							
Colour (post autoclaving/heating):			Red to orange red							
Clarity (post autoclaving/heating):			Clear							
	(III)Q.C. Test Microbiological									
		or 1	0 24 brc at 2	E 270C						
Cultural characteristics observed after					CAC	T				
MICROORGANISM (ATCC )		G	ROWTH	ACID	GAS					
Citrobacterfreundii (8090)		L	uxuriant	+	+					
Enterobacteraerogenes (13048)		L	uxuriant	+	+					
Escherichia coli (25922)		- 11	uxuriant	+	+					
Klebsiellapneumoniae (13883)			uxuriant		+					
	<u> </u>			+						
Proteus vulgaris (13315)		Li	uxuriant	+	+					
Salmonella typhimurium (14028)		L	uxuriant	+	+					
Salmonella typhi ( 6539 )		L	uxuriant	+	-					
Serratiamarcescens (8100)			uxuriant	+	-					
Shigellaflexne			uxuriant	+	_					
	egative reaction, no		axariant	'						
colour change										
	eaction, yellow colour									
	eaction, yellow colour									
Precautions:	1. For Laboratory Us									
	2. Follow proper, established laboratory procedures in handling and disposing of									
	infectious materials.									
Limitations :	1. Since the nutrition	al re	quirements of	organisms vary, se	ome strains may b	е				
	encountered that fail	to g	row or grow po	oorly on this mediu	ım.					
	2. The addition of some carbohydrates to the basal medium may cause an acid reaction.									
To restore the original pH (and colour of the medium), add 0.1 N sodium hydroxide on										
drop - by - drop basis. Take care not to make the medium too alkaline, which wou										
prevent fermentation from occurring within the usual incubation period.										
	3. To ensure accuracy of interpretation, uninoculated control tubes and/or inoculate									
	Phenol Red Broth Base control tubes should be run in parallel with the fermentation tests									
	Refer disclaimer Overleaf				e 01 of 02					
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Use :	<b>B1104:</b> For Galactose fermentation studies of microorganisms.								
Storage :	Dehydrated medium- below 30°C Prepared medium- Between 2 to 8°C.								
Packing:	500 gm. bottle								
Product profile:		Quantity on Preparation (500g)	pH (25°C)	Supplement	Sterilization				
B1104	21.00 g/l	23.80 L	7.4 <u>+</u> 0.2	Nil	121 <sup>0</sup> C / 15 minutes				

#### 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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