



# **Phenol Red Rhamnose Broth**

M1183

Phenol Red Rhamnose Broth is used for rhamnose fermentation studies of microorganisms.

Composition**	
Ingredients	Gms / Litre
Proteose peptone	10.000
Beef extract	1.000
Sodium chloride	5.000
Rhamnose	5.000
Phenol red	0.018
Final pH ( at 25°C)	$7.4\pm0.2$
**Formula adjusted, standardized to suit performance parameters	5

## Directions

Suspend 21 grams in 1000 ml distilled water and mix well. Heat if necessary to ensure complete dissolution. Distribute in fermentation tubes (tubes containing inverted Durham's tubes). Sterilize by autoclaving at 15 lbs pressure (121°C) for 15 minutes.

## **Principle And Interpretation**

Phenol Red Broth Medium is formulated as per Vera (2) and is recommended to determine the fermentation reaction of carbohydrates for the differentiation of microorganisms (3, 4, 5). Phenol Red Broth Medium with various carbohydrates serves as a differential medium by aiding in differentiation of various species and genera by their ability to ferment the specific carbohydrate, with the production of acid or acid and gas (6). Phenol Red Rhamnose Broth is used to study rhamnose fermentation in various bacteria.

Proteose peptone and beef extract serve as sources for carbon and nitrogen. Sodium chloride is the osmotic stabilizer. Phenol red is the pH indicator, which turns yellow at acidic pH i.e. on fermentation of rhamnose. Gas formation is seen in Durhams tubes. All of the *Enterobacteriaceae* grow well in this medium. In addition to producing a pH colour shift, the production of mixed acids, notably butyric acids, often results in a pungent, foul odour from the culture medium (1).

#### **Quality Control** Appearance Light yellow to pink homogeneous free flowing powder Colour and Clarity of prepared medium Red coloured clear solution without any precipitate Reaction Reaction of 2.1% w/v aqueous solution at 25°C. pH : 7.4±0.2 pН 7.20-7.60 **Cultural Response** M1183: Cultural characteristics observed after an incubation at 35 - 37°C for 18 - 24 hours. Inoculum Growth Acid Organism Gas (CFU) **Cultural Response** Citrobacter freundii ATCC 50-100 luxuriant Positive Positive 8090 reaction, yellowreaction colour Escherichia coli ATCC 50-100 luxuriant Positive Positive 25922 reaction, yellowreaction colour

Enterobacter aerogenes ATCC 13048	50-100	luxuriant	Positive reaction, yello colour	Positive wreaction
Klebsiella pneumoniae ATCC 13883	50-100	luxuriant	Positive reaction, yello colour	Positive wreaction
Proteus vulgaris ATCC 13315	50-100	luxuriant	Negative reaction, no colour change	Negative reaction
Salmonella Typhi ATCC 6539	50-100	luxuriant	Negative reaction, no colour change	Negative reaction
Salmonella Typhimurium ATCC 14028	50-100	luxuriant	Positive reaction, yello colour	Positive wreaction
Serratia marcescens ATCC 8100	50-100	luxuriant	Negative reaction, no colour change	Negative reaction
Shigella flexneri ATCC 12022	50-100	luxuriant	Negative reaction, no colour change	Negative reaction

## **Storage and Shelf Life**

Store below 30°C in tightly closed containerand the prepared medium at 2-8°C. Use before expiry date on the label.

### Reference

1. Koneman E. W., Allen S. D., Janda W.M., Schreckenberger P.C., Winn W.C. Jr., 1992, Colour Atlas and Textbook of Diagnostic Microbiology, 4th Ed., J. B. Lippinccott Company

2. Vera H. D., 1950, Am. J. Public Health, 40, 1267

3. MacFaddin J. F., 1985, Media for Isolation-Cultivation-Identification -Maintenanceof Medical Bacteria, Vol. I, Williams and Wilkins, Baltimore.

4. Finegold S. M. and Baron E. J., 1986, Bailey and Scotts Diagnostic Microbiology, 7th Ed., The C.V. Mosby Co., St. Louis.

5. Ewing W. H., 1986, Edwards and Ewings Identification of Enterobacteriaceae, 4th ed., Elsevier Science Publishing Co., Inc., New York.

6. MacFaddin J. F., 2000, Biochemical tests for Identification of Medical Bacteria, 3rd edi., Lippincott, Williams and Wilkins, Baltimore.

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