

Technical Data

Phenol Red Maltose Broth

M276

Phenol Red Maltose Broth is used for maltose fermentation studies of microorganisms.

Composition**

Ingredients	Gms / Litre
Proteose peptone	10.000
Beef extract	1.000
Sodium chloride	5.000
Maltose	5.000
Phenol red	0.018
Final pH (at 25°C)	7.4±0.2

^{**}Formula adjusted, standardized to suit performance parameters

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 Maltose Broth is used to study maltose 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 maltose. 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 coloured 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

рH

7.20-7.60

Cultural Response

Cultural characteristics observed after an incubation at 35 - 37°C for 18 - 24 hours (longer if necessary)

Cultural Response

Organism	Inoculum (CFU)	Growth	Acid	Gas
Cultural Response				
Citrobacter freundii ATCC	50-100	luxuriant	Positive	Positive
8090			reaction, yellowreaction	
			colour	

HiMedia Laboratories Technical Data

Escherichia coli ATCC	50-100	luxuriant	Positive	Positive	
25922			reaction, yellowreaction		
			colour		
Enterobacter aerogenes	50-100	luxuriant	Positive	Positive	
ATCC 13048	50 100	Tuxurum	reaction, yellowreaction		
			colour	wicaction	
Klebsiella pneumoniae	50-100	luxuriant	Positive	Positive	
ATCC 13883	30-100	iuxuiiaiit			
AICC 13003			reaction, yellowreaction colour		
Proteus vulgaris ATCC	50-100	luxuriant	Positive	Positive	
13315			reaction, yellowreaction		
			colour		
Salmonella Typhi ATCC	50-100	luxuriant	Positive	Negative	
6539			reaction, yello	C	
			colour		
Salmonella Typhimurium ATCC 14028	50-100	luxuriant	Positive	Positive	
			reaction, yellowreaction		
			colour	W Tedetion	
Serratia marcescens ATCC	50-100	luxuriant	Positive	Negative	
8100	30-100	luxurum	reaction, yello	U	
			colour	wicaction	
Chi II Ch ATCC	50-100	luxuriant		NI4:	
Shigella flexneri ATCC	30-100	iuxuriani	Positive	Negative	
12022			reaction, yellowreaction		
			colour		

Storage and Shelf Life

Store below 30°C in tightly closed container and 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.

Revision: 2 / 2015

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 HiMedia™ publications. The information contained in this publication is based on our research and development work and is to the best of our knowledge true and accurate. HiMedia™ Laboratories Pvt Ltd reserves the right to make changes to specifications and information related to the products at any time. Products are not intended for human or animal or therapeutic use but for laboratory, diagnostic, research or further manufacturing use only, unless otherwise specified. Statements contained herein should not be considered as a warranty of any kind, expressed or implied, and no liability is accepted for infringement of any patents.