



Technical Data

Phenol Red Adonitol Broth

M1200

Phenol Red Adonitol Broth is used for detection of adonitol fermenting bacteria.

Composition**

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

**Formula adjusted, standardized to suit performance parameters

Equivalent to Beef extract B

Directions

Suspend 21.02 grams in 1000 ml distilled water and mix well. Heat if necessary to ensure complete solution. 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 Adonitol Broth is used to study adonitol fermentation in various bacteria.

Proteose peptone and meat extract B 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 adonitol. 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. 7.4±0.2 pH : 7.4±0.2

pH

7.20-7.60

Cultural Response

M1200: Cultural characteristics observed after an incubation at 35-37°C for 18-24 hours.

Organism	Inoculum (CFU)	Growth	Acid production	Gas
Cultural Response				
<i>Citrobacter freundii</i> ATCC 8090	50-100	luxuriant	Negative reaction, no colour change	Negative reaction
<i>Escherichia coli</i> ATCC 25922	50-100	luxuriant	Negative reaction, no colour change	Negative reaction

<i>Enterobacter aerogenes</i> ATCC 13048	50-100	luxuriant	Positive reaction, yellow colour	Positive reaction
<i>Klebsiella pneumoniae</i> ATCC 13883	50-100	luxuriant	Positive reaction, yellow colour	Positive reaction
<i>Proteus vulgaris</i> ATCC 13315	50-100	luxuriant	Negative reaction, no colour change	Negative reaction
<i>Serratia marcescens</i> ATCC 8100	50-00	luxuriant	Negative reaction, no colour change	Negative reaction
<i>Salmonella Typhi</i> ATCC 6539	50-100	luxuriant	Negative reaction, no colour change	Negative reaction
<i>Salmonella Typhimurium</i> ATCC 14028	50-100	luxuriant	Negative reaction, no colour change	Negative reaction
<i>Shigella flexneri</i> ATCC 12022	50-100	luxuriant	Negative reaction, no colour change	Negative reaction

Storage and Shelf Life

Store below 30°C in tightly closed container and 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. Lippincott Company
2. Vera H. D., 1950, Am. J. Public Health, 40, 1267
3. MacFaddin J. F., 1985, Media for Isolation-Cultivation-Identification -Maintenance of 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 ed., Lippincott, Williams and Wilkins, Baltimore.

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