

Technical Information

MR-VP Medium (Glucose Phosphate Broth)

Product Code: DM 1070S

Application: - MR-VP Medium (Glucose Phosphate Broth) is recommended for studying Methyl Red and Voges-Proskauer tests to differentiation amongst coli-aerogenes group.

Composition*

Ingredients	Gms / Litre	Gms / Litre			
Peptic digest of animal tissue	5.000				
Dextrose	5.000				
Dipotassium phosphate	5.000				
Final pH (at 25°C)	7.5±0.1				
**Formula adjusted, standardized to suit performance	e				
narameters					

Principle & Interpretation

Clark and Lubs ⁽¹⁾ showed that the addition of methyl red to cultures of *Escherichia coli* resulted in a red colour due to high acidity produced during dextrose fermentation. Voges-Proskauer ⁽²⁾ found red colouration after addition of potassium hydroxide to specific culture media with organisms grown in it. The investigators developed MR-VP Broth in which both tests could be performed in same medium in different tubes. The red colour produced by the addition of potassium hydroxide to cultures is due to the ability of organisms to produce a neutral product acetoin (acetyl methyl carbinol) from dextrose ⁽³⁾. The acetoin is oxidized in the presence of oxygen and alkali to produce diacetyl which reacts with creatine to produce a red colour. This formulation is also recommended by BIS ⁽⁵⁾ and ISO committee ⁽⁸⁾ for the detection of coli-aerogenes group. A slightly modified formulation (DM1070S) is recommended by BIS ^(4,6,7) for the detection of *E. coli*, *Vibrio parahaemolyticus* and

Bacillus cereus responsible for food poisoning. To test *V.parahaemolyticus* for VP, addition of 2-3% Sodium chloride to the medium is required.

The Methyl Red (MR) test is performed after maximum of 5 days of incubation at 30°C ⁽⁹⁾ and Voges-Proskauer test (VP) cultures are incubated at 30°C for 24-48 hours ⁽¹⁰⁾. Various other tests have been suggested by Werkman ⁽¹¹⁾, OMeara ⁽¹²⁾ Levine, Epstein and Voughn ⁽¹³⁾ and Voughn, Mitchell and Levine ⁽⁹⁾. Werkmans Test ⁽⁸⁾: Add 2 drops of a 2% solution of ferric chloride to 50 ml culture and 5 ml of 10% sodium hydroxide. Shake the tube to mix well. Stable copper colour developing in a few minutes is positive reaction. OMeara Test ⁽⁸⁾: Add of 25 mg of solid creatine to 5ml culture and then add 5 ml concentrated (40%) sodium hydroxide. Red colour development in a few minutes after shaking the tube well, is a positive reaction. Levine, Epstein and Voughn ⁽¹³⁾ modified OMeara technique by dissolving the creatine in a concentrated solution of potassium hydroxide. Voughn, Mitchell and Levine ⁽⁹⁾ recommended the method of Barritt ⁽¹⁴⁾ as, addition of 1 ml of 40% potassium hydroxide and 3 ml of 5% a - naphthol in absolute ethanol to 5 ml culture. Positive test is indicated by eosine pink colour within 2-5 minutes.

Methodology

Suspend 15 grams of powder media in 1000 ml of distilled water. Shake well & distribute in test tubes in 3 ml amounts or as desired and sterilize by autoclaving at 15 lbs pressure (121°C) for 15 minutes.

Quality Control

Physical Appearance

Cream coloured homogeneous free flowing powder

Colour and Clarity of prepared medium

Light yellow coloured clear solution without any precipitate.

Reaction

Reaction of 1.5% w/v aqueous solutions at 25°C. pH: 7.5±0.1

pH range: 7.40-7.60

Cultural Response/Characteristics

DM 1070S: Cultural characteristics observed after an incubation at 30°C for 48 hours.





1				i
Organism	Inoculum (CFU)	Growth	MR Test	VP Test
Bacillus cereus ATCC 10876	50-100	Luxuriant	Negative reaction, yellow colour	Positive reaction, eosin pink/ red colour within 2-5 minutes
Enterobacter aerogenes ATCC 13048	50-100	luxuriant	Negative reaction, yellow colour	Positive reaction, eosin pink/ red colour within 2-5 minute
Escherichia coli ATCC 25922	50-100	Luxuriant	Positive reaction, bright red colour	Negative reaction, no colour change
Klebsiella pneumoniae ATCC 23357	50-100	Luxuriant	Negative reaction, yellow colour	Positive reaction, eosin pink/ red colour within 2-5 minute
Salmonella Typhi ATCC 6539	50-100	Luxuriant	Positive reaction, bright red colour	Negative reaction, no colour change
Vibrio parahaemolyticus ATCC 17802	50-100	Poor	Negative reaction, yellow colour	Negative reaction, no colour change

Storage and Shelf Life

Dried media: Store below 30°C in tightly closed container and use before expiry date as mentioned on the label.

Prepared Media: 2-8⁰ in sealable plastic bags for 2-5 days.

Further Reading

- 1. Clark and Lubs, 1915, J. Inf. Dis., 17: 160.
- 2. Voges and Proskauer, 1898, Zeit, Hyg., 28: 20.,,
- 3. MacFaddin, J.F., 1985, Media for Isolation-Cultivation-Identification-Maintenance of

Medical Bacteria, Vol. 1, Williams and Wilkins, Baltimore.

- 4. Bureau of Indian Standards, IS: 5887 (Part I) 1976, reaffirmed 1986.
- 5. Bureau of Indian Standards, IS: 5887 (Part III) 1976.
- 6. Bureau of Indian Standards, IS: 5887 (Part IV) 1976.
- 7. Bureau of Indian Standards, IS: 5887 (Part V) 1976, reaffirmed 1986.
- 8. International Organization for Standardization (ISO), 1993, Draft ISO/DIS 6597.
- 9. Vaughn, Mitchell and Levine, 1939, J. Am. Water Works Association, 31:993.
- 10. Kallas, Chinn and Coulter, 1931, J. Bact., 22: 125.
- 11. Werkman, 1930, J. Bact., 20: 121.
- 12. OMeara, 1931, J. Path. Bacteriol., 34: 401.
- 13. Levine, Epstein and Voughn, 1934, Am. J. of Publ. Health, 24: 505.
- 14. Barritt, 1936, J. Path. Bacteriol., 42 : 441.

Disclaimer :

- User must ensure suitability of the product(s) in their application prior to use.
- The product conforms solely to the technical information provided in this booklet and to the best of knowledge research and development work carried at **CDH** is true and accurate.
- Central Drug House 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 diagnostic or therapeutic use but for laboratory, research or further manufacturing of diagnostic reagents extra.
- 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.
- Do not use the products if it fails to meet specifications for identity and performens parameters.





