

Bases / Media Supplements

# **Technical Information**

## **M-Bismuth Sulphite Broth**

#### Product Code: DM 2101

Application: - M-Bismuth Sulphite Broth is a selective medium used for the detection of Salmonellae by the membrane filter

Composition**		
Ingredients	Gms / Litre	
Peptic digest of animal tissue	20.000	
Beef extract	10.000	
Dextrose	10.000	
Disodium phosphate	8.000	
Ferrous sulphate	0.600	
Bismuth sulphite indicator	16.000	
Brilliant green	0.050	
Final pH (25°C)	7.7±0.2	
**Formula adjusted, standardized to suit performa	ince parameters	

#### Principle & Interpretation

Salmonella is a gram-negative, facultatively anaerobic, non-sporulating, these motile rods in the family Enterobacteriaceae. They are widely distributed in animals affecting mainly the stomach and the intestines some times these organisms are difficult to differentiate biochemically from Escherichia coli. M-Bismuth Sulphite Broth was formulated by Clark et al<sup>(1)</sup> and is recommended for detection of Salmonella Typhi from water and various clinical specimens by the membrane filtration technique. Preliminary enrichment of clinical sample on a non-selective medium is not necessary. M-Bismuth Sulphite Broth has a composition similar to Bismuth Sulphite Agar (DM1027), except in the broth medium, all the constituents are in double concentration & is without Agar.

Peptic digest of animal tissue, beef extract and dextrose provide essential growth nutrients. Ferrous sulphate and bismuth sulphite indicator together act as H<sub>2</sub>S indicators. Brilliant green acts as selective agent. Luxuriant growth of Salmonella Typhi is obtained after 30 hours incubation at 35°C but metallic sheen and brown-black halo is not developed before 40 hours. The importance of this medium has been repeatedly mentioned for detection of Salmonella Typhi by membrane filter technique has been reported by different workers <sup>(2-5)</sup>.

### Methodology

Suspend 64.65 grams of powder media in 1000 ml distilled water. Shake well & heat if necessary to dissolve the medium completely. Excessive heating destroys the selective properties of the medium. DO NOT AUTOCLAVE. The medium usually contains flocculent precipitate, which should be dispersed evenly by swirling the flask just before use. Cool to 35°C and saturate sterile absorbent cotton pad with 2 ml of the broth. The medium should be used within 24 hours of rehydration.

### **Quality Control**

Physical Appearance		
Light yellow to greenish yellow homogeneous free flowing p	owder	
Colour and Clarity of prepared medium		
Greenish yellow coloured opalescent solution with flocculen	t precipitate	
Reaction		
Reactionof 6.4% w/v aqueous solution at 25°C.pH:-7.7±0.2		
pH range 7.50-7.90		
Cultural Response/ characteristices		
DM 2101: Cultural characteristics observed in humid at	mosphere, after an incubation at	35-37°C for 40-48 hours.
Organism Inoculum (CFU)	Growth	Colour of Colony (on Membrane filter)
Escherichia coli ATCC 25922 50-100	none-poor	brown-green, if any





Dehydrated Culture Media Bases / Media Supplements

Salmonella Typhi ATCC 6539 50-100 Salmonella Typhimurium ATCC 14028 50-100 Staphylococcus aureus ATCC 25923 >=10<sup>3</sup> luxuriant luxuriant inhibited black with metallic sheen black with metallic sheen

### 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**: 2-8° in sealable plastic bags for 2-5 days.

#### Further Reading

1. Clark H. F., Geldreich E. E., Jeter M. L. and Kabler P. W., 1951, Pub I. Hlth. Reports, 66:95 1.

2. Goets A. and Tsuneishi N., 1951, J. Am. Water Works Assoc., 43:943.

3. Goets A. and Tsuneishi N., 1952, J. Am. Water Works Assoc., 44:471.

4. Goets A. and Tsuneishi N., 1953, J. Am. Water Works Assoc., 45 and 1196.

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

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