MAST[®] Culture Media and Supplements

Technical Information Sheet

Product Code DM 160

Mannitol Salt Agar

A selective medium for the isolation of pathogenic staphylococci.

1. Description

Following the observation by Koch¹ that usually only staphylococci are able to grow on 7.5% salt agar, Chapman² described the use of Mannitol Salt Agar for the differential isolation of pathogenic staphylococci. It is worth noting that Gunn *et a^B* improved the medium by the addition of 2% egg yolk which identified 94.7% of coagulase producing strains. Recent work has shown that some strains of staphylococci may be partially inhibited by the high level of sodium chloride in Chapman's medium.^{4,5} To overcome this, the medium has been

2. Technical Formula*

Formula	grams per litre
Peptone	8.0
Yeast extract	2.0
Lactalbumin	3.0
Sodium chloride	30.0
Phenol red	0.0225
Lithium chloride	7.0
Glycine	1.0
Sodium pyruvate	3.0
Agar B	12.0
pH approx.7.4	

modified by reducing the sodium chloride concentration from 7.5 to 3.0%. Inhibition of Gram negative bacilli has been maintained by the inclusion of Lithium chloride. Pyruvate and glycine are included as growth promoters to enhance the recovery of *S.aureus* from samples containing small numbers of the organism.^{6,7} The medium is suitable for use with Oxacillin SelectatabTM (MS29) for the detection of Methicillin Resistant *Staphylococcus aureus* (MRSA) using the method described by Lally *et al*[§].

3. Directions

1. Suspend by swirling 76g of powder in 1 litre or the contents of the sachet in the stated volume of distilled or deionised water.

2. Autoclave at 121°C (15p.s.i.) for 15 minutes.

3. Mix well before pouring.





4. In Use

Inoculate plates by spreading the sample over the surface of the medium and incubate at 37°C for up to 48 hours. Pathogenic staphylococci grow well and ferment mannitol, indicated by yellow zones around the colonies, whereas non-pathogenic staphylococci produce smaller white colonies indicating non-fermentation of mannitol.

5. References

1. Koch FE. *Zentbl Bakt ParasitKde* (Abt.1) 1942; **149:** 122-124.

2. Chapman GH. J Bact. 1945; 50: 202-203.

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4. Van Enk RA, Thompson DK. *J Clin Microbiol.* 1992; **30:** 504-505.

5. Faiers M, George R, Jolly J, Wheat P. *Multipoint Methods in the Clinical Laboratory* - *A Handbook*. p41 PHLS, Lon 6.

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7. Giolloti C, Cantoni C. *J Appl Bact.* 1966; **29:** 395-398.

8. Lally RT, Ederer MN, Woolfrey BF. *J Clin Microbiol.* 1985; **22:** 501-504

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