

83339 Mac Conkey-Sorbitol ChromoSelect Agar

A selective agar for the direct isolation and differentiation of E. coli 0157:H7 strains from foodstuffs and clinical specimen.

Composition:

Ingredients	Grams/Litre
Casein enzymic hydrolysate	17.0
Proteose peptone	3.0
Sorbitol	10.0
Bile salts mixture	1.5
Sodium chloride	5.0
Crystal violet	0.001
Neutral red	0.03
5-Bromo-4-Chloro-3-Indolyl-ß-D-glucuronide Sodium salt	0.1
Agar	13.5
Final pH (at 25 °C) 7.1 ± 0.3	

Store prepared media below 8°C, protected from direct light. Store dehydrated powder, in a dry place, in tightly-sealed containers at 2-25°C.

Directions:

Suspend 50.1 grams in 1000 ml distilled water. Boil gently to dissolve the medium compleatly. **DO** NOT AUTOCLAVE. Cool to 50°C. Mix well and pour into sterile petri plates. If desired add Tellurite-Cefixime Supplement to the molten and cooled medium (50°C) before pouring into sterile petri plates.

Principle and Interpretation:

Sorbitol MacConkey Agar is based on the formulation described by Rappaport and Henigh (1). The medium contains sorbitol instead of lactose and it is recommanded for the detection of enteropathogenic strains of E. coli 0157:H7 which ferments lactose but does not ferment sorbitol (2) and hence produce colorless colonies. Sorbitol fermenting strains of *E. coli* produce pink-red colonies. The red colour is due to production of acid from sorbitol, absorption of neutral red and a subsequent colour change of the dye when pH of the medium falls below 6.8. E. coli 0157:H7 has been recognized as a cause of haemorrhagic colitis (3). March and Ratnam (2) reported that the detection of E.coli 0157:H7 had a sensitivity of 100% and specificity of 85% on Sorbitol Mac Conkey Agar and they recommended this medium as reliable means of screening E. coli 0157:H7.

B.C. indicator is added to detect the presence of an enzyme β -D-glucuronidase which is specific for E. coli (4). Strains of E. coli possessing β -D-glucuronidase appear as blue coloured colonies on the medium. Enteropathogenic strains of *E.coli 0157:H7* do not possess β -D-glucuronidase activity (5) and thus produce colorless colonies.

E. coli fermenting Sorbitol and possessing β -D-glucuronidase activity produce purple colored colonies. Casein enzymic hydrolysate and proteose peptone provide carbonaceous, nitrogenous and other essential growth nutrients. Most of the gram positive organisms are inhibited by crystal violet and bile salts. Sodium chloride maintains the osmotic equilibrium.

Addition of Tellurite-Cefixime Supplement makes the medium selective (6). Potassium tellurite selects the serogroups and inhibits Aeromonas species and Providencia species. Cefixime inhibits Proteus species. Pseudomonas if present produces colorless colonies on this medium. For confirmation oxidase test may be performed with suspect colonies and result should be noted within 5-10 seconds.

Cultural characteristics after 24 hours (48h if necessary) at 37°C.



Organisms (ATCC)	Color of Colony *	Sorbitol	β- glucuronidase	Oxidase	
Escherichia coli 0157:H7	colorless	_	-	_	
		_	_		
Escherichia coli (25922)	purple	+	+	-	
Pseudomonas aeruginosa (27853)	colorless	-	-	+	
* Color of the colors without addition of Tallwite Cofiving Complement					

^{*} Color of the colony without addition of Tellurite-Cefixime Supplement

References:

- 1. Rappaport F. and Henigh E., 1952, J. Clin. Path., 5:361.
- 2. March S.B., and Ratnam S. (1986), J. Clin. Microbiol. 23, 869-872.
- 3. Karmali M.A., Petric M., Lim C., et al, 1985, J. infect. Dis., 151-775.
- 4. Hansen W. und Yourassawsky E., 1984, J. Clin. Microbiol., 20:1177,
- 5. Thompson (1990), J. Clin. Microbiol. 29, 2165-2168.
- 6. Zadik P.M., Chapman P.A., and Siddons C.A. (1993), J. Med. Microbiol., 39, 155-158

Precautions and Disclaimer

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