

Technical Information

Sensitivity Test Broth

Product Code: DM 1486

Application: - Sensitivity Test Broth is recommended for antimicrobial susceptibility tests.

Composition**

Ingredients	Gms / Litre	
Casein enzymic hydrolysate	11.000	
Peptic digest of animal tissue	3.000	
Dextrose	2.000	
Sodium chloride	3.000	
Starch, soluble	1.000	
Disodium phosphate	2.000	
Sodium acetate	1.000	
Magnesium glycerophosphate	0.200	
Calcium gluconate	0.100	
Cobaltous sulphate	0.001	
Cupric sulphate	0.001	
Ferrous sulphate	0.001	
Zinc sulphate	0.001	
Manganous chloride	0.002	
Menadione	0.001	
Cyanocobalamin	0.001	
L-Cysteine hydrochloride	0.020	
L-Tryptophan	0.020	
Pyridoxine hydrochloride	0.003	
Calcium pantothenate	0.003	
Nicotinamide	0.003	
Biotin	0.0003	
Thiamine hydrochloride	0.00004	
Adenine	0.010	
Guanine	0.010	
Xanthine	0.010	
Uracil	0.010	
Final pH (at 25°C)	7.4±0.2	
**Formula adjusted, standardized to suit perform	ance parameters	

Principle & Interpretation

The goal of an antimicrobial susceptibility test is to predict through an in vitro assessment the likelihood of successfully treating an infection with a particular antimicrobial agent. There are several continual or novel methods for performing antibacterial susceptibility testing. These include the disk diffusion test, broth micro dilution, agar gradient and rapid automated instrument methods (1). Sensitivity Test Broth, which is used for antimicrobial susceptibility tests, is a semi-defined medium in which the mineral contents have been stabilized to give reproducible results. The thiamine and thymidine content is very low thus making it most suitable for testing antimicrobial activity of sulphonamides.





However some mutant strains which are totally dependent on thiamine and thymidine for their growth, will not grow in Sensitivity Test Broth, due to very low levels of these compounds in the media as they are the naturally occurring antagonist of trimethoprim. These strains should be carefully recognized (2, 3, 4).

Sensitivity Test Broth has been so designed to overcome the problems occurring in Mueller-Hinton Media that are as follows (5-11).

- 1. Mueller Hinton Agar and Mueller Hinton Broth give different MIC values.
- 2. Mueller Hinton Agar shows antagonistic effect towards tetracycline.
- 3. High levels of sulphonamide and trimethoprim antagonists.
- 4. Media prepared using peptone of different manufacturers give poor reproducibility.
- 5. Poor growth supporting ability for Streptococci and variable growth rates with gram-positive organisms.

Some pathogenic organisms are nutritionally dependent due to their intrinsic demands for special growth factors.

Casein enzymic hydrolysate, peptic digest of animal tissue, dextrose, and vitamins provides nitrogen, carbon compounds and other essential growth nutrients.

Methodology

Suspend 23.4 grams of dehydrated powder media in 1000 ml distilled water. Mix thoroughly & heat to boil to dissolve the medium completely. Sterilize by autoclaving at 15 lbs pressure (121°C) for 15 minutes. Shake well before dispense as desired.

Quality Control

Appearance

Cream to yellow homogeneous free flowing powder

Colour and Clarity

Basal medium: Light yellow; After addition of 5%v/v laked blood: Red to chocolate coloured, Basal medium: clear to slightly opalescent; After Addition: opalescent solution in tubes

Reaction

Reaction of 2.34% w/v aqueous solution at 25°C. pH: 7.4±0.2

pH Range

7.20-7.60

Cultural Response

DM 1486: Cultural characteristics observed after an incubation at 35-37°C for 18-24 hours.

Organism	Inoculum (CFU)	Growth
Salmonella Typhimurium ATCC 14028	50-100	good-luxuriant
Staphylococcus aureus ATCC 25923	50-100	good-luxuriant
Streptococcus pyogenes ATCC 19615	50-100	good-luxuriant
Enterococcus faecalis ATCC 29212	50-100	good-luxuriant
Bacillus subtilis ATCC 6633	50-100	good-luxuriant
Bacteroides vulgatus ATCC 8482	50-100	good-luxuriant





Storage and Shelf Life

Dried Media: Store dehydrated powder and prepared medium at 2-8°C in tightly closed container. Use before expiry period on the label. **Prepared Media**: 2-8° in sealable plastic bags for 2-5 days.

Further Reading

- 1. Murray P. R., Baron J. H., Pfaller M. A., Jorgensen J. H. and Yolken R. H., (Ed.). 2003, Manual of Clinical Microbiology, 8th Ed., American Society for Microbiology, Washington, D.C.
- 2. Tanner E. I. and Bullin C. H., 1974, J. Clin. Path., 27:565.
- 3. Thomas M. and Bond L., 1973, Med. Lab. Technol., 30:277.
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- 5. Ericsson H. M. and Sherris J. C., 1971, Acta. Pathol. Microbiol Scand Suppl., 217:1.
- 6. Garrod L. P. and Waterworth P. M., 1971, J. Clin. Path., 24:779.
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- 8. Duncan I. B. R., 1974, Antimicrob. Agents Chemother., 5:9.
- 9. Yourassowsky E., Vanderlinden M. P. and Schoutens E., 1974, J. Clin. Path., 27:897.
- 10. Neussil H., 1976, Chemother., Vol. 2:33.
- 11. Bridson E.Y., 1976, Arztl. Lab., 22:373.

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