

TM 1466 – RAPID-SENSITIVITY TEST BROTH

INTENDED USE

For antimicrobial susceptibility test.

PRODUCT SUMMARY AND EXPLANATION

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 microdilution, agar gradient and rapid automated instrument methods. Rapid Sensitivity Test Broth, which is used for antimicrobial susceptibility tests, is a semidefined 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 Rapid-Sensitivity Test Broth, due to very low levels of these compounds in the media as they are the naturally occurring antagonist of trimethoprim.

COMPOSITION

Ingredients	Gms / Ltr	
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	

PRINCIPLE

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

INSTRUCTION FOR USE

- Dissolve 23.4 grams in 1000 ml purified/distilled water.
- Heat to boiling to dissolve the medium completely.
- Sterilize by autoclaving at 15 psi pressure (121°C) for 15 minutes.
- Mix well and dispense as desired.

QUALITY CONTROL SPECIFICATIONS

Appearance of Powder : Cream to yellow homogeneous free flowing powder.

Appearance of prepared medium : 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.

pH (at 25°C) : 7.4 ± 0.2

INTERPRETATION

Cultural characteristics observed after incubation.

Microorganism	ATCC	Inoculum (CFU/ml)	Growth	Incubation Temperature	Incubation Period
Bacillus subtilis	6633	50-100	Good-luxuriant	35-37°C	18-24 Hours
Bacteroides vulgatus	8482	50-100	Good-luxuriant	35-37°C	18-24 Hours
Enterococcus faecalis	29212	50-100	Good-luxuriant	35-37°C	18-24 Hours









Salmonella Typhimurium	14028	50-100	Good-luxuriant	35-37°C	18-24 Hours
Staphylococcus aureus	25923	50-100	Good-luxuriant	35-37°C	18-24 Hours
Streptococcus pyogenes	19615	50-100	Good-luxuriant	35-37°C	18-24 Hours

PACKAGING:

In pack size of 100 gm and 500 gm bottles.

STORAGE

Dehydrated powder, hygroscopic in nature, store in a dry place, in tightly-sealed containers between 25-30°C and protect from direct sunlight. Under optimal conditions, the medium has a shelf life of 4 years. When the container is opened for the first time, note the time and date on the label space provided on the container. After the desired amount of medium has been taken out replace the cap tightly to protect from hydration.

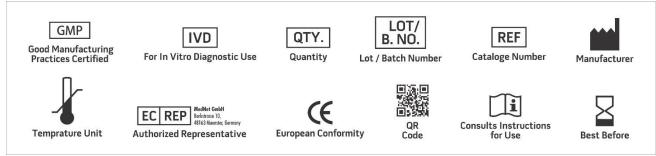
Product Deterioration: Do not use if they show evidence of microbial contamination, discoloration, drying or any other signs of deterioration.

DISPOSAL

After use, prepared plates, specimen/sample containers and other contaminated materials must be sterilized before discarding.

REFERENCES

- 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.
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- 5. Ericsson H. M. and Sherris J. C., 1971, Acta. Pathol. Microbiol Scand Suppl., 217:1.
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NOTE: Please consult the Material Safety Data Sheet for information regarding hazards and safe handling Practices.

*For Lab Use Only
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