

CM 20942 – LEGIONELLA AGAR BASE

INTENDED USE

For cultivation of Legionella species.

PRODUCT SUMMARY AND EXPLANATION

Legionella Agar initially called as F-G agar was modified by Feely et al by replacing Starch with charcoal and casein hydrolysate with yeast extract which resulted in better recovery of Legionella pneumophila. Pasculle et al reported that the addition of ACES (N-2-acetamido-2-amino ethane sulphonic acid) buffer improved the nutritive value of medium. Edelstein suggested addition of α -Ketoglutarate to increase the sensitivity of this medium.

Legionella species have an absolute nutritional requirement for L-Cysteine. Presumptive Legionella species colonies can be sub-cultured onto both Legionella Agar Base with Legionella Growth Supplement and with Legionella Growth Supplement w/o L-Cysteine. All plates are incubated at 35°C. Colonies which grow on Legionella Agar Base with Legionella Growth Supplement, with L-Cysteine, but not on Legionella Agar Base with Legionella Growth Supplement w/o L-Cysteine, can be regarded as presumptive Legionella species.

COMPOSITION

Ingredients	Gms / Ltr
Charcoal activated	2.000
Yeast extract	10.000
Agar	13.000

PRINCIPLE

This medium consists of yeast extract to provide the necessary nitrogenous nutrients for Legionella growth. Activated charcoal nullifies toxic compounds that either accumulate in the medium during growth or develop during sterilization of medium. Addition of ACES buffer helps in maintaining proper pH of the medium for the optimal growth of Legionella. Antibiotics in the supplement inhibit the growth of various contaminating bacteria and fungi.

INSTRUCTION FOR USE

Dissolve 12.5 grams in 440 ml purified / distilled water.

Heat to boiling to dissolve the medium completely.

Sterilize by autoclaving at 15 psi pressure (121°C) for 15 minutes.

Cool to 45-50°C and aseptically add contents of 1 vial of Legionella Growth Supplement (BCYE). In case of non-incorporation of Legionella (GVPC) Selective Supplement, add aseptically 10 ml sterile distilled water to bring the total volume to 500 ml of medium.

Mix well and pour into sterile petri plates.

Stir the medium while dispensing to prevent the settling of charcoal particles. If desired, the medium can be made selective by aseptically adding rehydrated contents of 1 vial of either Legionella BIPA Selective Supplement or Legionella (GVPC) Selective Supplement, along with 1 vial of Legionella Growth Supplement (BCYE) to 440 ml sterile molten, cooled Legionella Agar Base. Simultaneously, a medium without L-Cysteine may be prepared by adding aseptically contents of 1 vial of Legionella Growth Supplement w/o L-Cysteine.

QUALITY CONTROL SPECIFICATIONS



Appearance of Powder	: Grey to black coloured homogeneous free flowing powder.
Appearance of prepared medium	: Black coloured opaque gel forms in Petri plates.
pH (at 25°C)	: 6.9 ± 0.2

INTERPRETATION

Cultural characteristics observed with added Sterile Legionella Growth Supplement (BCYE) and Legionella (GVPC) Selective Supplement or Legionella Growth Supplement w/o L-Cysteine after incubation.

Microorganism	ATCC	Inoculum (CFU/ml)	Growth (with TS 114 & TS 115)	Recovery	Growth (With TS 195)	Recovery	Incubation Temperature	Incubation Period
Escherichia coli	25922	$\geq 10^3$	Inhibited	0%	Good	40-50%	35-37°C	48-72 Hours
Legionella dumoffii	33343	50-100	Good-luxuriant	$\geq 50\%$	Inhibited	0%	35-37°C	48-72 Hours
Legionella pneumophila	33153	50-100	Good-luxuriant	$\geq 50\%$	Inhibited	0%	35-37°C	48-72 Hours

PACKAGING:

Inpacksize of 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. Baird R.B., Eaton A.D., and Rice E.W., (Eds.), 2015, Standard Methods for the Examination of Water and Wastewater, 23rd ed., APHA, Washington, D.C.
2. Dennis et al, 1984, Proceeding of the 2nd International Symposium, Washington D.C. Am. Soc. Microbiol. PP 294-296.
3. Edelstein, 1981, J. Clin. Microbiol., 14:298.
4. Feely J. C., et al, 1978, J. Clin. Microbiol., 8(3):320.
5. Feely, Gibson, Gorman, et al, 1979, J. Clin. Microbiol., 10(4):437.
6. Isenberg, H.D. Clinical Microbiology Procedures Handbook 2nd Edition.
7. Jorgensen, J.H., Pfaller, M.A., Carroll, K.C., Funke, G., Landry, M.L., Richter, S.S and Warnock, D.W. (2015) Manual of Clinical Microbiology, 11th Edition. Vol. 1.
8. Pasculle, Feely, Gibson et al, 1980, J. Infect. Dis., 141:72.



 GMP Good Manufacturing Practices Certified	 IVD For In Vitro Diagnostic Use	 QTY. Quantity	 LOT/ B. NO. Lot / Batch Number	 REF Catalogue Number	 Manufacturer
 Temperature Unit	 EC REP Authorized Representative MedNet GmbH Buckstraße 10, 48153 Münster, Germany	 European Conformity	 QR Code	 Consults Instructions for Use	 Best Before

NOTE: Please consult the Material Safety Data Sheet for information regarding hazards and safe handling Practices.

*For LabUse Only