



Fraser Broth w/ Supplements

Intended Use:

Recommended for the selective enrichment of Listeria species from food samples.

Composition**

Ingredients	Gms / Litre
Peptone	5.000
Tryptone	5.000
Yeast extract	5.000
HM peptone B #	5.000
Sodium chloride	20.000
Lithium chloride	3.000
Disodium hydrogen phosphate	9.600
Potassium dihydrogen phosphate	1.350
Esculin	1.000
Nalidixic acid	0.010
Acriflavin	0.0125
Ferric ammonium citrate	0.500
Final pH (at 25°C)	7.2±0.2
**Formula adjusted, standardized to suit performance parameters	
# Equivalent to Beef extract	

Directions

Suspend 55.47 grams of dehydrated medium in 1000 ml purified / distilled water. Heat to boiling to dissolve the medium completely. DO NOT AUTOCLAVE. Cool to 45-50°C. Mix well and dispense as desired in sterile tubes or flasks.

Principle And Interpretation

Listeria species are widely distributed and are isolated from soil, decaying vegetable matter, sewage, water, animal feed, fresh and frozen poultry, meats, raw milk, cheese and asymptomatic human and animal carriers (11). Only *Listeria monocytogenes* from the genus *Listeria*; causes infections in humans. *L. monocytogenes* primarily causes meningitis, encephalitis or septicemia in humans (9,12). In pregnant women, *Listeria monocytogenes* often causes an influenza like bacteremic illness that, if untreated, may lead to ammionitis and infection of the fetus, resulting in abortion, still birth or premature birth. Contaminated foods are the primary vehicles of transmission (8).

Fraser Broth w/ supplement is based on the formulation by Fraser and Sperber (2). It is recommended for selective enrichment of *Listeria* species from foods.

This medium contains peptone, HM peptone B, yeast extract and tryptone which provide essential nutrients like carbon and nitrogenous compounds including vitamins, amino acids and trace ingredients. Phosphates buffer the medium while sodium chloride maintains osmotic equilibrium. Nalidixic acid and Acriflavin inhibits the growth of gram-negative and gram-positive organisms respectively (5,6,7) except *Listeria* species (5,6,7). *Listeria* species hydrolyze esculin to glucose and esculetin. The latter combines with ferric ions of ferric ammonium citrate, resulting in the formation of 6-7 dihydroxycoumarin, a black brown complex. Ferric ammonium citrate also enhances the growth of *L. monocytogenes* (1). High salt tolerance due to sodium chloride of *Listeria* is used as means to inhibit the growth of Enterococci. Lithium chloride is also used to inhibit Enterococci, which also possess the ability to hydrolyze esculin.

Type of specimen

Food samples

M2002

Specimen Collection and Handling

For food samples, follow appropriate techniques for sample collection and processing as per guidelines (10). After use, contaminated materials must be sterilized by autoclaving before discarding.

Warning and Precautions

Read the label before opening the container. Wear protective gloves/protective clothing/eye protection/ face protection. Follow good microbiological lab practices while handling specimens and culture. Standard precautions as per established guidelines should be followed while handling specimens. Safety guidelines may be referred in individual safety data sheets.

Limitations

1. Due to nutritional variations, some strains may show poor growth.

- 2. Slight colour variation may be observed depending upon strains.
- 3. Further biochemical tests must be carried out for confirmation.

Quality Control

Appearance

Cream to yellow homogeneous free flowing powder

Colour and Clarity of prepared medium

Fluorescent yellow coloured clear solution.

Reaction

Reaction of 5.55% w/v aqueous solution at 25°C. pH : 7.2±0.2

pН

7.00-7.40

Cultural response

Cultural characteristics observed after an incubation at 35 - 37°C for 24-48 hours.

Organism	Inoculum (CFU)	Growth	Esculin Hydrolysis
Escherichia coli ATCC 25922 (00013*)	>=10 ⁴	inhibited	
Enterococcus faecalis ATCO 29212 (00087*)	C 50-100	none-poor	
Listeria monocytogenes subsp. serovar 1 ATCC 19111 (00020*)	50-100	good-luxuriant	positive reaction, blackening of medium
Listeria monocytogenes ATCC 19112	50-100	good-luxuriant	positive reaction, blackening of medium
Listeria monocytogenes ATCC 19117	50-100	good-luxuriant	positive reaction, blackening of medium
Listeria monocytogenes ATCC 19118	50-100	good-luxuriant	positive reaction, blackening of medium
Staphylococcus aureus subsp. aureus ATCC	50-100	none-poor	

25923 (00034*)

Key : (*) Corresponding WDCM numbers.

Storage and Shelf Life

Store dehydrated and the prepared medium at 2-8°C. Use before expiry date on the label. On opening, product should be properly stored dry, after tightly capping the bottle in order to prevent lump formation due to the hygroscopic nature of the product. Improper storage of the product may lead to lump formation. Store in dry ventilated area protected from extremes of temperature and sources of ignition Seal the container tightly after use.

Product performance is best if used within stated expiry period.

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User must ensure safe disposal by autoclaving and/or incineration of used or unusable preparations of this product. Follow established laboratory procedures in disposing of infectious materials and material that comes into contact with clinical sample must be decontaminated and disposed of in accordance with current laboratory techniques (3,4).

Reference

1.Cowart R. E. and Foster B. G., 1985, J. Infect. Dis.; 151:172.

2. Fraser, J., and W. Sperber. 1988. Rapid detection of *Listeria* in food and environmental samples by esculin hydrolysis. Journal of Food Protection 51: 762-765.

- ^{3.} Isenberg, H.D. Clinical Microbiology Procedures Handbook. 2nd Edition.
- 4. 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.
- 5. Lee W.K. and McClain D., 1986, Appl. Environ. Microbiol., 52:1215.
- 6. Lovette J., Francis D.W. and Hunt J.M., 1987, J. Food Prot., 50:188.
- 7. McClain D. and Lee W.H., 1988, J. Assoc. Off. Anal. Chem., 71:660.

8. Murray P. R., Baron E. J., Jorgensen J. H., Pfaller M. A., Yolken R. H., (Eds.), 8th Ed., 2003, Manual of Clinical Microbiology, ASM, Washington, D.C.

9. Nieman R. E., and Lorber B., 1980, Rev. Infect. Dis. 2 : 207-227.

- 10. Salfinger Y., and Tortorello M.L. Fifth (Ed.), 2015, Compendium of Methods for the Microbiological Examination of Foods, 5th Ed., American Public Health Association, Washington, D.C.
- Seeliger H. P. R., and Jones D., 1986, Bergeys Manual of Systematic Bacteriology, Vol. The Williams and Wilkins Co., Baltimore.
- 12. Schuchat A. B., Swaminathan and C. V. Broome, Clin. Microbiol. Rev. 4: 169-183.

Revision : 01 / 2019

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