

## Fraser Broth (NCM0050)

### Intended Use

Fraser Broth is used with ferric ammonium citrate for the selective enrichment of *Listeria* species and is not intended for use in the diagnosis of disease or other conditions in humans.

### Description

Fraser Broth is a secondary enrichment broth for the isolation of *Listeria* spp. formulated according to ISO 11290. The selective components acriflavine and nalidixic acid are blended into the base powder and the ferric ammonium citrate (X211 or 7984) is added to the tempered broth after sterilization.

### Typical Formulation

Peptone	15.0 g/L
Yeast Extract	5.0 g/L
Esculin	1.0 g/L
Disodium Hydrogen Phosphate	9.6 g/L*
Potassium Dihydrogen Phosphate	1.35 g/L
Sodium Chloride	20.0 g/L
Lithium Chloride	3.0 g/L
Acriflavine	0.025 g/L
Nalidixic Acid	0.02 g/L

pH: 7.2 ± 0.2 at 25°C

\*equivalent to Disodium Hydrogen Phosphate Dihydrate 12.0 g/l

Formula may be adjusted and/or supplemented as required to meet performance specifications.

Ferric Ammonium Citrate Supplement (X211) or Fraser broth base supplement (7984) may be used

### Precaution

Refer to SDS

### Preparation

1. Dissolve 55 grams of the medium in 1 liter of purified water.
2. Mix thoroughly
3. Autoclave at 121°C for 15 minutes.
4. Cool to 45-50°C.
5. Aseptically add 2 vials of Ferric Ammonium Citrate Supplement (X211) or 2 vials of Fraser Broth Base Supplement (7984) and mix thoroughly before dispensing.

### Test Procedure

To isolate *Listeria monocytogenes* and other *Listeria* spp., refer to appropriate references

### Quality Control Specifications

**Dehydrated Appearance:** Powder is homogeneous, free flowing and beige.

**Prepared Appearance:** Prepared medium is yellow, clear, with none to slight precipitate.

# Technical Specification Sheet



## Expected Cultural Response: (Fertility and Selectivity Testing)

The medium was prepared according to label directions and 10mL volumes inoculated with the organisms listed below. Cultures were incubated aerobically at 37 ± 1°C and examined for growth at 22 to 26 hours. Following incubation, 10µL of *E. faecalis* and *E. coli* cultures were subcultured onto Tryptone Soy Agar (NCM0020) at 37 ± 1°C and examined for growth at 22 to 26 hours

MICROORGANISM	APPROX. INOCULUM (CFU)	Expected Results		
		Growth in FB	Reaction in FB	Recovery on TSA
<i>Enterococcus faecalis</i> ATCC® 19433	> 10 <sup>3</sup>	Partial to Complete Inhibition	---	< 100 cfu
<i>Enterococcus faecalis</i> ATCC® 29212	> 10 <sup>3</sup>	Partial to Complete Inhibition	---	< 100 cfu
<i>Escherichia coli</i> ATCC® 25922	> 10 <sup>3</sup>	Complete Inhibition	---	< 10 cfu
<i>Escherichia coli</i> ATCC® 8739	> 10 <sup>3</sup>	Complete Inhibition	---	< 10 cfu
<i>Listeria innocua</i> ATCC® 33090	> 10 <sup>3</sup>	Growth	Blackening	NA
<i>Listeria monocytogenes</i> ATCC® 35152	10-100	Growth	Blackening	NA
<i>Listeria monocytogenes</i> ATCC® 13932	10-100	Growth	Blackening	NA
<i>Staphylococcus aureus</i> ATCC® 25923	> 10 <sup>4</sup>	Inhibited	---	NA

## (Productivity Testing)

The medium was prepared according to label directions and 10mL volumes inoculated as a mixed culture using the organisms listed below. Cultures were incubated aerobically at 37 ± 1°C, examined for growth at 22 to 26 hours and subcultured onto Harlequin™ *Listeria* Chromogenic Agar according to Ottaviani and Agosti (NCM1004) at 37 ± 1°C and plates examined for growth at 22-48 hours

Mixed Culture Testing	ATCC	APPROX. INOCULUM (CFU)	EXPECTED RESULTS	
			Recovery on <i>Listeria</i> Agar*	Reaction on <i>Listeria</i> Agar*
<i>Enterococcus faecalis</i> + <i>Escherichia coli</i> + <i>Listeria monocytogenes</i>	29212 25922 35152	> 10 <sup>3</sup> >10 <sup>3</sup> 10-100	Inhibited Inhibited >10 cfu	--- --- Blue colonies with opaque halo
<i>Enterococcus faecalis</i> + <i>Escherichia coli</i> + <i>Listeria monocytogenes</i>	19433 8739 13932	> 10 <sup>3</sup> >10 <sup>3</sup> 10-100	Inhibited Inhibited >10 cfu	--- --- Blue colonies with opaque halo

The organism listed are minimum that should be used for quality control testing.



# Technical Specification Sheet



## **Results**

For further identification and confirmation of *Listeria* species, consult appropriate references. Rapid slide and macroscopic tube tests can be used for definitive serological identification.

## **Expiration**

Refer to expiration date stamped on the container. The dehydrated medium should be discarded if not free flowing, or if appearance has changed from the original color. Expiry applies to medium in its intact container when stored as directed.

## **Limitations of the Procedure**

Due to nutritional variation, some strains may grow poorly or fail to grow on this medium.

## **Storage**

Store dehydrated culture media at 2 – 30°C away from direct sunlight. Once opened and recapped, place the container in a low humidity environment at the same storage temperature. Protect from moisture and light by keeping container tightly closed.

## **References**

1. Fraser J.A. and Sperber W.H. (1988). Rapid detection of *Listeria* spp in food and environmental samples by esculin hydrolysis. *J. Food Protect.* 51, No.10, 762-765.
2. McClain D. and Lee W.H. (1989). FSIS method for isolation of *L. monocytogenes* from processed meat and poultry products. Lab.Comm.No.57, Revised May 24, (1989). US Dept of Agric.FSIS, Microbiol. Div.
3. ISO 11290-1:2017 (Microbiology of food and animal feeding stuffs - Horizontal method for the detection and enumeration of *Listeria monocytogenes* and of *Listeria* spp. -- Part 1: Detection method

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