

## Oxford Listeria Agar (NCM0056)

### Intended Use

Oxford Listeria Agar Base is used with antimicrobics for the selective isolation of *Listeria* spp. in a laboratory setting. Oxford Listeria Agar Base is not intended for use in the diagnosis of disease or other conditions in humans.

### Description

*Listeria monocytogenes*, described first in 1926 by Murray, Webb, and Swann, is an extensive problem in food industries. Epidemiological evidence from outbreaks of listeriosis has indicated that the principle route of transmission is via the consumption of foodstuffs contaminated with *Listeria monocytogenes*. Implicated vehicles of transmission included turkey frankfurters, coleslaw, pasteurized milk, Mexican style cheese, and pate. *Listeria* spp. are ubiquitous in nature, being present in a wide range of unprocessed foods as well as in soil, sewage and river water.

Oxford Listeria Agar is prepared according to the formulation of Curtis et al. *Listeria* spp. grow over a pH range of 5.0 - 9.6 and survive in food products with pH levels outside these parameters .

### Typical Formulation

Columbia Blood Agar Base	39.0 g/L
Esculin	1.0 g/L
Ferric Ammonium Citrate	0.5 g/L
Lithium Chloride	15.0 g/L
Agar	2.0 g/L

Final pH: 7.0 ± 0.2 at 25°C

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

### Precaution

Refer to SDS

### Preparation

#### **Oxford Listeria Agar**

1. Suspend 57.5 g of the medium in one liter of purified water.
2. Heat with frequent agitation and boil for one minute to completely dissolve the medium.
3. Autoclave at 121°C for 15 minutes.
4. Cool to 45 - 50°C.
5. Aseptically add 2 vials of Oxford Listeria Supplement (7986) or Oxford Supplement (X123) and mix well before dispensing.

#### **Modified Oxford Listeria Agar**

Perform steps 1-4 above and aseptically add a filtered sterilized aqueous solution of 10 mg colistin sulfate and 20 mg moxalactam. Mix well before dispensing.

### Test Procedure

The USDA method involves enrichment of the food sample in UVM Modified Listeria Enrichment Broth (one part sample to nine parts broth) at 30°C. After incubation, a portion of the enrichment mixture is plated onto Oxford or Modified Oxford Medium. The method according to FDA BAM involves adding 25 mL of liquid or 25 g of solid material to 225 mL Listeria Enrichment Broth and incubating at 30°C for two days. After enrichment, the broth is plated onto Oxford Listeria Agar. For further information consult appropriate references.

# Technical Specification Sheet



## Quality Control Specifications

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

**Prepared Appearance:** Prepared medium is light to medium amber and slightly hazy.

**Expected Cultural Response:** Cultural response in Oxford Listeria Agar and Modified Oxford Listeria Agar at 35 ± 2°C after 24 - 48 hours incubation.

Microorganism	Approx. Inoculum (CFU)	Expected Results	
		Oxford	Modified Oxford
<i>Candida albicans</i> ATCC® 10231	>10 <sup>5</sup>	Suppressed to complete inhibition	Suppressed to complete inhibition
<i>Escherichia coli</i> ATCC® 25922	>10 <sup>5</sup>	Complete inhibition	Complete inhibition
<i>Escherichia coli</i> ATCC® 8739	>10 <sup>5</sup>	Complete inhibition	Complete inhibition
<i>Staphylococcus aureus</i> ATCC® 25923	>10 <sup>5</sup>	Suppressed to complete inhibition	Complete inhibition
<i>Enterococcus faecalis</i> ATCC® 29212	>10 <sup>5</sup>	Suppressed to complete inhibition	Suppressed to complete inhibition
<i>Enterococcus faecalis</i> NCTC 775	>10 <sup>5</sup>	Suppressed to complete inhibition	Suppressed to complete inhibition
<i>Listeria innocua</i> ATCC® 33090	10-300	Growth; brown to black colonies	Growth; brown to black colonies
<i>Listeria monocytogenes</i> ATCC® 13932	10-300	Growth; brown to black colonies	Growth; brown to black colonies
<i>Listeria monocytogenes</i> ATCC® 19111	10-300	Growth; brown to black colonies	Growth; brown to black colonies
<i>Listeria monocytogenes</i> ATCC® 35152	10-300	Growth; brown to black colonies	Growth; brown to black colonies

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

## Results

Select esculin-positive colonies and confirm their identity through biochemical testing. Use macroscopic tube and rapid slide tests 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 the appearance has changed from the original color. Expiry applies to medium in its intact container when stored as directed.

## Limitation of the Procedure

1. An identification of *L. monocytogenes* must be confirmed through biochemical and serological testing.
2. Poor growth and a weak esculin reaction maybe seen after 40 hours incubation for some enterococci.

## 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.



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## References

1. Murray, E. G. D., R. A. Webb, and M. B. R. Swann. 1926. A disease of rabbits characterized by large mononuclear leucocytosis caused by a hitherto undescribed bacillus *Bacterium monocytogenes*. *J. Path. Bacteriol.* 29:407-439.
2. Bremer, P. J., and C. M. Osborne. 1995. Thermal-death times of *Listeria monocytogenes* in green shell mussels prepared for hot smoking. *J. Food Prot.* 58:604-608.
3. Graud, F. H., and P. B. Vanderlinde. 1992. Occurrence, numbers, and growth of *Listeria monocytogenes* on some vacuum-packaged processed meats. *J. Food Prot.* 55:4-7.
4. Patel, J. R., C. A. Hwang, L. R. Beuchat, M. P. Doyle, and R. E. Brackett. 1995. Comparison of oxygen scavengers for their ability to enhance resuscitation of heat-injured *Listeria monocytogenes*. *J. Food Prot.* 58: 244-250.
5. Curtis, G. D. W., R. G. Mitchell, A. F. King, and J. Emma. 1989. A selective differential medium for the isolation of *Listeria monocytogenes*. *Appl. Microbiol.* 8:95-98.
6. Vanderzant, C., and D. F. Splittstoesser (eds.). 2015. Compendium of methods for the microbiological examination of foods, 4<sup>th</sup> ed. American Public Health Association, Washington, D.C.
7. Fraser, J., and W. Sperber. 1988. Rapid detection of *Listeria* in food and environmental samples by esculin hydrolysis. *J. Food Prot.* 51:762-765.
8. Lee, W. H., and D. McClain. 1989. Laboratory Communication No. 57 (revised May 24, 1989). U.S.D.A., F.S.I.S. Microbiology Division, Beltsville, MD.
9. [www.fda.gov/Food/ScienceResearch/LaboratoryMethods/BacteriologicalAnalyticalmanualBAM/default.htm](http://www.fda.gov/Food/ScienceResearch/LaboratoryMethods/BacteriologicalAnalyticalmanualBAM/default.htm).

Effective Date: 3/1/2019

Revision: 0