



# L. mono Differential Agar Base

L. mono Differential Agar Base has been recommended for the selective and differential isolation of *Listeria monocytogenes*.

# **Composition\*\***

Ingredients	Gms / Litre
Meat peptone	18.000
Casein enzymic hydrolysate	6.000
Yeast extract	10.000
Sodium pyruvate	2.000
Glucose	2.000
Magnesium glycerophosphate	1.000
Magnesium sulphate	0.500
Sodium chloride	5.000
Lithium chloride	10.000
Disodium hydrogen phosphate anhydrous	2.500
Chromogenic substrate	0.050
Agar	15.000
Final pH ( at 25°C)	7.2±0.2
**Formula adjusted standardized to suit performance parameters	

\*\*Formula adjusted, standardized to suit performance parameters

## Directions

Suspend 36.02 grams in 460 ml distilled water. Heat to boiling to dissolve the medium completely. Sterilize by autoclaving at15 lbs pressure (121°C) for 15 minutes. Cool to 45-50°C. Aseptically add sterile contents of 1 vial of L. mono Enrichment Supplement I (FD214) and sterile rehydrated contents of L .mono Selective Supplement I (FD212), L .mono Selective Supplement II (FD213). Mix well and pour into sterile Petri plates.

Warning : Lithium chloride is harmful. Avoid bodily contact and inhalation of vapours. On contact with skin wash with plenty of water immediately.

# **Principle And Interpretation**

*Listeria monocytogenes* is a gram-positive foodborne human pathogen responsible for serious infections in pregnant women that may ultimately result in abortion, stillbirth, birth of a child with neonatal listeriosis and meningitis or primary bacteremia in adults and juveniles. The pathogenicity of *Listeria ivanovii* for humans is uncertain. Since *L. monocytogenes* and *L. innocua* have similar biochemical properties, they cannot be differentiated on traditional media (PALCAM, Oxford). L. mono Differential Agar Base is based on the formulation of Ottoviani and Agosti (1, 2) for the selective and differential isolation of *Listeria monocytogenes* from food and animal feeds which is adopted by ISO Committee (3).

Meat peptone, casein enzymic hydrolysate, yeast extract and sodium pyruvate provide essential growth nutrients and nitrogenous substances. Glucose is the fermentable carbohydrate. Sodium chloride maintains osmotic equilibrium. Phosphate buffers the medium. Lithium chloride and added selective supplements (FD212 and FD213) inhibit accompanying microflora and allow the growth of *Listeria* species. *Listeria* species hydrolyse the chromogenic substrate which produces green coloured colonies. Differentiation of *Listeria monocytogenes* from other *Listeria* species is based on phosphatidylinositol-specific phospholipase C (PIPLC) activity. Phospholipase C enzyme hydrolyses the purified substrate (FD214) added to the medium resulting in an opaque halo around *Listeria monocytogenes* colonies.

# **Quality Control**

Appearance Cream to yellow homogeneous free flowing powder Gelling

# **M1540**

Firm, comparable with 1.5% Agar gel

#### Colour and Clarity of prepared medium

Light amber coloured, opalescent gel forms in Petri plates

## Reaction

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

#### pН

7.00-7.40

#### **Cultural Response**

Cultural characteristics observed with added sterile L. mono Selective supplement I (FD212), L. mono Selective Supplement II (FD213) and L.mono Enrichment supplement I (FD214) after an incubation at  $35 - 37^{\circ}$ C for 24 - 48 hours.

## **Cultural Response**

Organism	Inoculum (CFU)	Growth	Recovery	Colour of Colony	PIPLC activity
Cultural Response					
Candida albicans ATCC 10231	>=103	inhibited	0%		
Enterococcus faecalis ATCO 29212	C>=10 <sup>3</sup>	inhibited	0%		
Escherichia coli ATCC 25922	>=103	inhibited	0%		
Listeria innocua ATCC 33090	>=103	luxuriant	>=50%	greenish-blue	negative
Listeria grayi ATCC 19120 Listeria ivanovii ATCC 19119 Listeria monocytogenes ATCC 19112	50-100 50-100 50-100	luxuriant luxuriant luxuriant	>=50% >=50%	greenish-blue greenish-blue	negative positive, opaque halo around the colony exhibiting phophatidylinositol specific phospholipase acivity positive, opaque halo around the colony exhibiting phosphatidylinositol specific phospholipase
Listeria seeligeri ATCC 35967	50-100	luxuriant	>=50%	greenish-blue	activity negative
Listeria welshimeri ATCC 43549	50-100	luxuriant	>=50%	greenish-blue	negative
Pseudomonas aeruginosa ATCC 27853	>=103	inhibited	0%		

## **Storage and Shelf Life**

Store dehydrated powder and the prepared medium at 2-8° C in tightly closed container . Use before expiry date on the label.

### Reference

1. Ottaviani F., Ottaviani M., and Agosti M. (1997 a), Industrie Alimentari 36, 1-3.

2. Ottaviani F., Ottaviani M., and Agosti M. (1997 b), Quimper Froid Symposium Proceedings p. 6, A.D.R.I.A. Quimper, France, 16-18 June 1997.

3. Draft Amendment ISO 11290-2:1996/DAM 1.

#### Disclaimer :

User must ensure suitability of the product(s) in their application prior to use. Products conform solely to the information contained in this and other related HiMedia<sup>TM</sup> publications. The information contained in this publication is based on our research and development work and is to the best of our knowledge true and accurate. HiMedia<sup>TM</sup> Laboratories Pvt Ltd reserves the right to make changes to specifications and information related to the products at any time. Products are not intended for human or animal or therapeutic use but for laboratory, diagnostic, research or further manufacturing use only, unless otherwise specified. Statements contained herein should not be considered as a warranty of any kind, expressed or implied, and no liability is accepted for infringement of any patents.

HiMedia Laboratories Pvt. Ltd. A-516, Swastik Disha Business Park, Via Vadhani Ind. Est., LBS Marg, Mumbai-400086, India. Customer care No.: 022-6147 1919 Email: techhelp@himedialabs.com