

## TM 1377 – LISTERIA MOTILITY MEDIUM

### INTENDED USE

For testing motility of *Listeria monocytogenes*.

### PRODUCT SUMMARY AND EXPLANATION

Bacterial motility is one of the important determinants in making a final species identification. Bacteria move by means of flagella, the number and location of which vary among different species. Semisolid media in tubes are most commonly employed for detecting motility. Motility media have agar concentration of 0.4% or less. The motility test is interpreted by making a macroscopic examination of medium for a diffused zone of growth flaring out from the line of inoculation. *Listeria monocytogenes* requires room temperature incubation before motility develops, since in some organisms; flagellar proteins develop more rapidly at lower temperatures (room temperature) such as in *L. monocytogenes* and *Yersinia enterocolitica*. Listeria Motility Medium is formulated in accordance with ISO Committee specification for the determination of motility by *L. monocytogenes*.

### COMPOSITION

Ingredients	Gms / Ltr
Casein enzymic hydrolysate	20.000
Peptic digest of animal tissue	6.100
Agar	3.500

### PRINCIPLE

This medium consists of Casein enzymic hydrolysate and peptic digest of animal tissue which act as source of growth nutrients. The motility of *L. monocytogenes* is best demonstrated by stab inoculating two tubes of semisolid medium and incubating one at room temperature (20 - 25°C) and the other at 35°C. Motility is better observed at room temperature. An umbrella-like zone of growth 2 to 5 mm below the surface of the medium is characteristic of *L. monocytogenes*. Motility at 35°C incubation is either absent or extremely sluggish.

### INSTRUCTION FOR USE

- Dissolve 29.6 grams in 1000 ml purified/distilled water.
- Heat to boiling to dissolve the medium completely.
- Dispense in tubes and sterilize by autoclaving at 15 psi pressure (121°C) for 15 minutes.
- Allow the tubed medium to cool in an upright position.

### QUALITY CONTROL SPECIFICATIONS

<b>Appearance of Powder</b>	: Cream to yellow homogeneous free flowing powder.
<b>Appearance of prepared medium</b>	: Light yellow coloured, clear to slightly opalescent gel forms in tubes as butts.
<b>pH (at 25°C)</b>	: 7.3 ± 0.2

### INTERPRETATION

Cultural characteristics observed after incubation.

Microorganism	ATCC	Inoculum (CFU/ml)	Growth	Motility	Incubation Temperature	Incubation Period



<i>Listeria monocytogenes</i>	19117	50-100	Luxuriant	Positive, growth away from stabline causing turbidity	25-30°C	24-48 Hours
<i>Listeria monocytogenes</i>	19111	50-100	Luxuriant	Positive, growth away from stabline causing turbidity	25-30°C	24-48 Hours
<i>Listeria monocytogenes</i>	19112	50-100	Luxuriant	Positive, growth away from stabline causing turbidity	25-30°C	24-48 Hours
<i>Staphylococcus aureus</i>	25923	50-100	Luxuriant	Negative, growth along the stabline, surrounding medium remains clear	25-30°C	24-48 Hours

**PACKAGING:**

In pack size 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. International Organization for Standardization (ISO), 1993, Draft ISO/DIS 10560.
2. Bailey and Scotts Diagnostic Microbiology, 1986, 7th Ed., The C.V. Mosby Co., St. Louis.

 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 <small>MedNet GmbH Borkstrasse 10, 48163 Muenster, Germany</small>	 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 Lab Use Only**  
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