Reference: 6040

Technical Data Sheet

Product: LISTERIA CHROMOGENIC SELECTIVE SUPPLEMENT

Specification

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Selective supplement for isolation and confirmation of Listeria monocytogenes formulated according to ISO 11290-1 and 2:1996 Amd 2004

Presentation

10 Freeze dried vials	Packaging Details	Shelf Life	Storage
Vial with: 3 ± 0.1 a	22±0.25 x 55±0.5 mm glass vials, tag labelled, White plastic cap - 10 vials per box.	49 months	2-25 ºC
	por uora		

Composition

Compositon (g/vial)

Polymyxin B	38350 IU
Cycloheximide	0.025
Ceftazidime	0.010
Nalidixic acid	. 0.010

NOTE : Each vial is sufficient to supplement 476 ml of Listeria selective Agar Base according to Ottavani and Agosti

Description /Technique

Description:

Completed with all its supplements the Agar Listeria Ottaviani & Agosti is a selective and differential medium for the detection of *Listeria* species and the presumptive identification of *Listeria* monocytogenes.

The selectivity is achieved by the high concentration of lithium chloride and the mixture of antimicrobics. The differential activity is due to the chromogenic substrate to detect the glucosidase, enzyme that is present in all *Listeria* species.

The specific identification is obtained by the L- α -phosphatidylinositol, that acts as substratre for a phospholipase C present only in *Listeria monocytogenes* and some strains of *Listeria ivanovii*. The combination of both substrates allows the differentiation *L. monocytogenes*, which produces colonies blue-green in colour and surrounded by an opaque zone, from the other *Listeria* species, which growth with blue-green colonies but without any halo. This differentiation is evident after incubate the plates for 24 ± 2 hours at 37 °C.

Sometimes, especially with highly contaminated samples, it is possible that some colonies, white in colour, that are not *Listeria* growth . In this case it is recommended an enrichment step previous to the plate inoculation.

Observations: Most Listeria ivanovii also produce an opaque halo around the colonies after 48 h of incubation. This presumptive evidence must be confirmed by performing the biochemical or serological identification tests (Rhamnose / Xylose sugar fermentation, hemolysis tests, CAMP test, etc.) or any test confirming the species without hesitation.

<u>Technique:</u>

Add 1 botlle Enrichment supplement Ottaviani & Agosti (L-alpha-phosphatidylinositol - 24ml) and 1 vial Selective supplemet Ottaviani & Agosti for complete 500ml medium. Homogenize by mixing and distribute in Petri dishes. The solidified cool medium appears homogeneously turbid.

There are a lot of standardized methodology (ISO, FDA-BAM, AOAC, AFNOR, etc.) The technician must follow the protocol validated in his laboratory.

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Quality control

Physical/Chemical control

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Color : White-Gray pH: at 25ºC

Microbiological control

Spiral Spreading: Practical range 100 ± 20 CFU. min. 50 CFU (productivity) / 10⁴-10⁶ CFU (selectivity). Microbiological control according to ISO 11133:2014/A1:2018.

Aerobiosis. Incubation at 35 ± 2 °C, reading at 24-48 hours.

Microorganism

Escherichia coli ATCC[®] 25922, WDCM 00013 Enterococcus faecalis ATCC[®] 29212, WDCM 00087 L. monocytogenes ATCC[®] 13932, WDCM 00021 Listeria innocua ATCC[®] 33090. WDCM 00017 Listeria monocytogenes ATCC[®] 35152

Sterility Control

Incubation 48 hours at 30-35 °C and 48 hours at 20-25 °C: NO GROWTH. Check at 7 days after incubation in same conditions.

Bibliography

Artault, S., j.L. Bind, Y. Delaval, N. Dureuil, N. Gallart (2000) AFNOR validation of the ALOA method for the detection of Listeria monocytogenes in foodstuffs. Coll. Soc. Fran. Microbiol. 19-20 Oct. Paris.

Bannerman, E.S. & J. Bille (1988) A new selective medium for isolating Listeria from heavily contaminated material. Appl.m Environm. Microbiol. 54:1:165-167.

·Greenwood, M., C. Willis, P. Dosweell, G. Allen & K. Pathak (2005) Evaluation of chromogenic media for the detection of Listeria species in food.

Hitchins, A.D. & K. Jinneman (1998) Listeria monocytogenes in FDA-BAM 8th edition Revision A. Updater January 2003. AOAC Intl. Gathersburg. MD. USA.

. ISO 11133:2014/ Adm 1:2018. Microbiology of food, animal feed and water. Preparation, production, storage and performance testing of culture media.

· ISO 11290-1:2017 Standard. Microbiology of the food chain. Horizontal method for the detection and enumeration of Listeria monocytogenes and for Listeria spp.- Part 1: Detection Method

· ISO 11290-2:2017 Standard. Microbiology of the food chain. Horizontal method for the detection and enumeration of Listeria monocytogenes and for Listeria spp.- Part 2: Enumeration Method -Jantzen, M.M., J. Navas, M. de Paz, B. Rodriguez, W.P. da Silva & M. Nuñez (2006) Evaluation of ALOA plating medium for its suitability to recover high pressure-injured Listeria monocytogenes from ground chicken meat. Letters Appl. Microbiol 43:313-317

Manafi, M. W. Kneifel & S. Bascomb (1991) Fluorogenic and chromogenic substrates used in bacterial diagnostics. Microbiol Rev. 55:3:335-348

Ottaviani, F., M. Ottaviani & M. Agosti (1997) Esperienza su un agar salettivo e differenziale per Listeria monocytogenes. Industrie Alimentari 36:1-3

-Victor Lachica, R. (1990) Selective plating medium for quantitative recovery of food-borne Listeria monocytogenes. Appl. Environm. Microbiol. 56:1:167-169

Watkins, J. & K.P. Sleath (1981) Isolation and enumeration of Listeria monocytogenes from sewage, sewage sludge and river water. J. Appl. Bacteriol. 50:1-9

. UNE-EN ISO 11133 (2014). Microbiología de los alimentos para consumo humano, alimentación animal y aqua.-Preparación, producción, conservación y ensayos de rendimiento de los medios de cultivo.

Inhibited Inhibited Good - Blue colonies with white halo Blue colonies without white halo Good - Blue colonies with white halo

Growth