



LABORATORY REPORT

Example Client, XYZ123
1234 Warde Road
Ann Arbor MI 48108

EXAMPLE, REPORT W
WX0000003827 M 07/08/1978 45 Y

Referral Testing

Collected: 09/21/2023 13:52 Received: 09/21/2023 13:52

Table with 6 columns: Test Name, Result, Flag, Ref-Ranges, Units, Site. Row 1: Listeria Antibody, CF, Serum, <1:8, QCR

REFERENCE RANGE: <1:8

INTERPRETIVE CRITERIA:
<1:8 Antibody Not Detected
> or = 1:8 Antibody Detected

The recommended laboratory method for diagnosing Listeria infection is culture. The intended use of this complement fixation assay for Listeria antibodies is to document infection when culture is not performed, or when antibiotic treatment precludes obtaining an accurate culture result.

Single titers of > or = 1:8 are suggestive of either recent or past Listeria infection. A four-fold or greater increase in titer between acute and convalescent specimens confirms a diagnosis of recent infection.

This test was developed and its analytical performance characteristics have been determined by Quest Diagnostics. It has not been cleared or approved by FDA. This assay has been validated pursuant to the CLIA regulations and is used for clinical purposes.

Test Performed at:
Quest Diagnostics Nichols Institute
33608 Ortega Highway
San Juan Capistrano, CA 92675-2042 I Maramica MD, PhD

Performing Site:
QCRL: QUEST DIAGNOSTICS REFERENCE LAB CAPISTRANO 33608 Ortega Highway San Juan Capistrano CA 92675

Reported Date: 2023.09.21 13:52

LAB: L - LOW, H - HIGH, AB - ABNORMAL, C - CRITICAL, . - NOT TESTED

F321000007 Ordered By: KAJAL SITWALA, MD, PhD
WX0000003827 WX00000000002365
Printed D&T: 09/21/23 13:52

Kajal V. Sitwala, MD, PhD - Medical Director
Form: MM RL1
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