



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: 08/24/2023 15:57 Received: 08/24/2023 15:57

Table with 6 columns: Test Name, Result, Flag, Ref-Ranges, Units, Site. Contains data for Strep pneumoniae IgG Abs, 23 Serotypes, Ser and Interpretation.

Evaluation of the immune response following pneumococcal vaccination can be assessed by measuring serotype-specific Streptococcus pneumonia IgG antibodies. Either of the following conditions is consistent with a normal response to Streptococcus pneumonia vaccination: 1. When comparing pre and post-vaccination samples, antibody concentrations increased by at least 2-fold for either >50% of serotypes in children <6 years of age or >70% of serotypes for individuals >6 years of age. 2. In either a pre- or post-vaccination sample, antibody concentrations >=1.0 mcg/mL for either >50% of serotypes for children <6 years of age or >70% of serotypes for individuals >6 years of age. Results >=1.0 mcg/mL or those showing a >=2-fold change are consistent with an immune response, but are not necessarily sufficient to provide protection against

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

F224000029
WX0000003827
Printed D&T: 08/24/23 15:58

Ordered By: KAJAL SITWALA, MD, PhD
WX0000000002365

Kajal V. Sitwala, MD, PhD - Medical Director
Form: MM RL1
PAGE 1 OF 2



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Test Name Result Flag Ref-Ranges Units Site

infection.

-----ADDITIONAL INFORMATION-----

On 08/03/2023 Mayo Clinic Laboratories implemented a modified Streptococcal pneumoniae IgG antibody method. If patient was tested on previous method and is undergoing serial monitoring, particularly in the context of pre- and postvaccination responses, rebaselining may be indicated. Rebaselining, or testing the current sample on the previous method, is available at no charge, subject to reagent availability. The rebaseline result will be reported as an add-on test to this order under test code PN23. Contact Mayo Clinic Laboratories at 1-800-533-1710 within 7 days of initial report issuance to request this service. For Mayo Clinic patients, call (77)4-4403. This test was developed and its performance characteristics determined by Mayo Clinic in a manner consistent with CLIA requirements. This test has not been cleared or approved by the U.S. Food and Drug Administration.

Test Performed by:
Mayo Clinic Laboratories - Rochester Superior Drive
3050 Superior Drive NW, Rochester, MN 55905
Lab Director: William G. Morice M.D. Ph.D.; CLIA# 24D1040592

Performing Site:
MMRL: MAYO MEDICAL REFERENCE LAB 3050 Superior Drive NW Rochester MN 55901

Reported Date: 2023.08.24 15:57 PN23M

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