



LABORATORY REPORT

Example Client, XYZ123
1234 Warde Road
Ann Arbor MI 48108

EXAMPLE, REPORT W
WX0000003826 F 12/05/1988 34 Y

Referral Testing

Collected: 09/06/2023 11:42 Received: 09/06/2023 11:42

Table with 6 columns: Test Name, Result, Flag, Ref-Ranges, Units, Site. Row 1: Rickettsia typhi (Typhus Fever) IgG and IgM Ab, Typhus Fever Antibody, IgG, <1:64, <1:64, ARRL

INTERPRETIVE INFORMATION: Typhus Fever Antibody, IgG

- Less than 1:64 Negative - No significant level of IgG antibody detected.
1:64 - 1:128 Equivocal - Questionable presence of IgG antibody detected. Repeat testing in 10-14 days may be helpful.
1:256 or greater Positive - Presence of IgG antibody to detected, suggestive of current or past infection.

Antibody reactivity to Rickettsia typhi antigen should be considered group-reactive for the Typhus Fever group, which includes Rickettsia prowazekii.

Seroconversion between acute and convalescent sera is considered strong evidence of recent infection. The best evidence for infection is a significant change (fourfold difference in titer) on two appropriately timed specimens, where both tests are done in the same laboratory at the same time.

Table with 6 columns: Test Name, Result, Flag, Ref-Ranges, Units, Site. Row 1: Typhus Fever Antibody, IgM, <1:64, <1:64, ARRL

INTERPRETIVE INFORMATION: Typhus Fever Antibody, IgM

- Less than 1:64 Negative-No significant level of IgM antibody detected.
1:64 or greater Positive-Presence of IgM antibody detected, which may indicate a current or recent infection; however, low levels of IgM

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



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Table with 6 columns: Test Name, Result, Flag, Ref-Ranges, Units, Site. Content: antibodies may occasionally persist for more than 12 months post-infection.

Antibody reactivity to Rickettsia typhi antigen should be considered group-reactive for the Typhus Fever group, which includes Rickettsia prowazekii.

Seroconversion between acute and convalescent sera is considered strong evidence of recent infection. The best evidence is a significant change (fourfold difference in titer) on two appropriately timed specimens, where both tests are done in the same laboratory at the same time. Acute-phase specimens are collected during the first week of illness and convalescent-phase samples are generally obtained 2-4 weeks after resolution of illness. Ideally these samples should be tested simultaneously at the same facility. If the sample submitted was collected during the acute-phase of illness, submit a marked convalescent sample within 25 days for paired testing.

Performed By: ARUP Laboratories
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Salt Lake City, UT 84108
Laboratory Director: Jonathan R. Genzen, MD, PhD
CLIA Number: 46D0523979

Performing Site:
ARRL: ARUP REFERENCE LAB 500 Chipeta Way Salt Lake City UT 841081221
Reported Date: 2023.09.06 11:42 TYPGM

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