



LABORATORY REPORT

Example Client, XYZ123
1234 Warde Road
Ann Arbor MI 48108

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

Molecular

Collected: 09/25/2023 14:18 Received: 09/25/2023 14:18

Table with 6 columns: Test Name, Result, Flag, Ref-Ranges, Units, Site. Rows include Influenza Virus A and B PCR, Specimen Source (Nasopharyngeal Swab), Influenza A (DETECTED), and Influenza B (Not detected).

This test utilizes a real-time reverse-transcriptase polymerase chain reaction procedure to amplify and detect Influenza A and Influenza B virus. The Influenza A assay detects a conserved region of the matrix gene and the Influenza B assay detects a conserved region of the NS1 gene. Available nucleic acid sequence information suggests that the Influenza A test will detect all known Influenza A subtypes including avian influenzas. However, this method cannot distinguish avian from non-avian subtypes. The analytical sensitivity of this assay is 100 copies/mL. A "Not detected" result does not rule out infection.

This test uses commercial reagents that have not been approved or cleared by the FDA. The FDA has determined that such clearance or approval is not necessary. The performance characteristics of this procedure were determined by Warde Medical Laboratory.

This test is performed pursuant to a license agreement with Roche Molecular Systems, Inc.

Performing Site:

WMRL: WARDE MEDICAL LABORATORY 300 West Textile Road Ann Arbor MI 48108

Reported Date: 2023.09.25 14:19 FLPCR

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

F325000056
WX0000003826
Printed D&T: 09/25/23 14:19

Ordered By: KAJAL SITWALA, MD, PhD
WX00000000002353

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