



LABORATORY REPORT

Example Client, XYZ123
1234 Warde Road
Ann Arbor MI 48108

EXAMPLE, REPORT
WX0000073111 F 02/15/1985 38 Y

Molecular

Collected: 03/17/2020 09:22 Received: 03/17/2020 09:22

Table with 6 columns: Test Name, Result, Flag, Ref-Ranges, Units, Site. Rows include Parainfluenza Virus PCR and three sub-rows for Parainfluenza 1, 2, and 3.

This test utilizes a real-time reverse-transcriptase polymerase chain reaction procedure to amplify and detect type-specific areas within the HN gene of human parainfluenza viruses. The analytical sensitivity of this assay is 200 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.

Reported Date: 04/01/2020 10:01 PIPCR

Performing Site:

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

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

6317000357 Ordered By: CLIENT CLIENT
WX0000073111 WX00000000409391
Printed D&T: 12/01/23 10:01

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