



LABORATORY REPORT

Example Client, XYZ123
1234 Warde Road
Ann Arbor MI 48108

EXAMPLE, REPORT

WX0000073111 F 02/15/1985 35 Y

Molecular

Collected: 03/17/2020 09:21

Received: 03/17/2020 09:21

Table with 6 columns: Test Name, Result, Flag, Ref-Ranges, Units, Site. Row 1: Norovirus Group 1 and 2 PCR. Row 2: Specimen Source Feces WMRL. Row 3: Norovirus Group 1 Not detected Not detected WMRL. Row 4: Norovirus Group 2 Not detected Not detected WMRL.

This test utilizes a real-time reverse-transcriptase polymerase chain reaction procedure to amplify and detect RNA polymerase gene sequences of norovirus genogroup 1 and VP1 gene sequences from norovirus genogroup 2. The analytical sensitivity of this assay is approximately 100 virus copies/assay. A "Not detected" result does not rule out infection. All results should be correlated with clinical presentation as Norovirus are excreted in the stool for 2-3 weeks after clinical symptoms resolve.

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

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

6317000356
WX0000073111
Printed D&T: 03/26/20 11:09

Ordered By: CLIENT CLIENT
WX00000000409391

William G. Finn, M.D. - Medical Director
Form: MM RL1
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