

LABORATORY REPORT

Example Client, XYZ123 1234 Warde Road Ann Arbor MI 48108

EXAMPLE, REPORT

WX0000073111 F 02/15/1985 39 Y

Molecular

Collected: 08/01/2024 13:53 Received: 08/01/2024 13:53

<u>Test Name</u> <u>Result</u> <u>Flag Ref-Ranges</u> <u>Units</u> <u>Site</u>

Cystic Fibrosis Mutation Panel

Cystic Fibrosis Mutation Analysis See Below WMRL

Result: Normal Genotype

Interpretation:

This individual is negative for the 39 Cystic Fibrosis (CF) mutations included in this assay, indicating a reduced risk for CF. These results do not exclude all pathogenic CFTR mutations. An individual's residual risk after a negative test varies by ancestry (see table below).

Estimated carrier risk:

| | Detection Rate | Risk Before Test | Residual Risk After Negative Test |
|--------------------|-------------------|---------------------|---|
| Ethnic group | | | Negative lest |
| | | | |
| Ashkenazi Jewish | 94% | 1/24 | 1 in 400 |
| European Caucasian | 91% | 1/25 | 1 in 263 |
| African American | 68% | 1/61 | 1 in 187 |
| Hispanic American | 74% | 1/58 | 1 in 221 |
| Asian American | 49% | 1/94 | 1 in 184 |

NOTE:

Limited information is available for individuals from other ethnic populations. Residual carrier risk after a negative test is modified by the presence of a positive family history of CF (i.e., having a first, second, or third degree relative affected with CF) and/or by a mixture of various ethnic groups. For these specific situations, accurate risk assessment requires standard Bayesian analysis and genetic counseling.

Methodology: Multiplex targeted amplification and direct mutation analysis using the Luminex analyzer was performed to test for the presence of 39 mutations within the CFTR gene (GenBank accession number NM_00492). A full list of mutations tested for by this assay can be found at the Warde Laboratory website:

(https://wardelab.com/resources/forms)

under "CFMPL Cystic Fibrosis Mutation Panel List of Variants Targeted".

Limitations: This assay does not include all known CF-causing pathogenic variants. The absence of a variant does not rule out the possibility of this individual being

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

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



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or patient management decisions.

Reported Date: 08/01/2024 13:54 CFMPL

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

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