

LABORATORY REPORT

Example Client, XYZ123 1234 Warde Road Ann Arbor MI 48108 **EXAMPLE, REPORT W**

WX0000003827 M 07/08/1978 45 Y

Referral Testing

Collected: 09/25/2023 08:24 Received: 09/25/2023 08:24

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

KIT D816 Mutation Analysis (Mastocytosis)

Specimen Source LAV EDTA WB QCRL
Block / Specimen Id N/A QCRL
Clinical Indication N/A QCRL
KIT D816 Mutation Detected AB QCRL

Reference range: Not Detected

Interpretation See Note QCRL

The D816V KIT mutation (c.2447A>T) is detected in extracted DNA by a PCR-based pyrosequencing method.

This mutation is characteristic of systemic mastocytosis, related disorders and certain AML subtypes.

This assay, which has an approximate sensitivity of 2.0% mutation-bearing cells, can also be used to monitor follow-up samples for involvement. KIT mutations outside of this codon will not be detected.

Renee Mohrmann, M.D. Medical Director, Molecular Oncology

This test was developed and its analytical performance characteristics have been determined by Quest Diagnostics Nichols Institute Chantilly, VA. It has not been cleared or approved by the U.S. Food and Drug Administration. This assay has been validated pursuant to the CLIA regulations and is used for clinical purposes.

Test Performed at: Quest Diagnostics Chantilly Nichols Institute, 14225 Newbrook Dr. Chantilly, VA 20153-0841 P W Mason MD, PhD

Performing Site:

QCRL: QUEST DIAGNOSTICS REFERENCE LAB CAPISTRANO 33608 Ortega Highway San Juan Capistrano CA 92675

Reported Date: 2023.09.25 8:24 D816M

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

F325000006 WX0000003827 Printed D&T: 09/25/23 08:25 Ordered By: KAJAL SITWALA, MD, PhD WX00000000002365

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