



LABORATORY REPORT

Example Client, XYZ123
1234 Warde Road
Ann Arbor MI 48108

EXAMPLE, REPORT W
WX0000003826 F 12/05/1988

Collected: 12/28/2023 15:02

Received: 12/28/2023 15:02

Sequential Screen Part 1

Table with 6 columns: Test Name, Result, Flag, Ref-Ranges, Units, Site. Rows include Screen Result (Negative), Age at EDD (years) (37), GA (weeks) (12), GA (days) (1), Ultrasound Date (12/20/2023), CRL (55.5), Nasal Bone (N/A), Weight (lbs) (150), Multiple Gestation (Single), Ethnic Origin (Other), IDDM (None), Smoker (No), NT (1.0), PAPP-A (1,234.5), hCG (123.4), NT MOM (N/A), PAPP-A MOM (N/A), HCG MOM (N/A), Down Syndrome Risk (1:N/A), Maternal Age Risk for Down (1:N/A), Trisomy 18 Risk (1:N/A), Interpretation (SeeBelow).

*** NOT AT INCREASED RISK FOR DOWN SYNDROME ***

This is the initial sample received at Warde Laboratory for Sequential Screening Part One. To complete the Sequential Screen for the final risk calculation, a second sample is needed on 12/30/2023 (15 weeks gestation) or as soon as possible after this date.

Additional Test Information:

The Sequential Screen Part One only provides risk estimates when positive and is not a diagnosis. Incorrect or missing information may considerably alter results.

A positive report is indicated when the Down syndrome risk is greater than or equal to 1:25.

LAB: L - LOW, H - HIGH, AB - ABNORMAL, C - CRITICAL,

F628000027

Ordered By: KAJAL V SITWALA, MD

WMB-23-1413

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PAGE 1 OF 2

Printed D&T: 12/28/2023 3:07 PM

Kajal V. Sitwala, MD, PhD - Medical Director



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<u>Test Name</u>	<u>Result</u>	<u>Flag</u>	<u>Ref-Ranges</u>	<u>Units</u>	<u>Site</u>
<p>Risks in twins cannot be precisely estimated because the relative contribution of each twin to serum marker levels cannot be determined. Assessment is adjusted for IVF, donor eggs, frozen embryos, insulin-dependent diabetic status, weight, race and smoking. Previous pregnancies affected with a neural tube defect or Down syndrome may significantly affect current results.</p> <p>The PAPP-A test uses a kit designated by the manufacturer as "for research use, not for clinical use". The performance characteristics of this test were validated by Warde Medical Laboratory. The U.S. Food and Drug Administration (FDA) has not approved or cleared this test. The results are not intended to be used as the sole means for clinical diagnosis or patient management decisions. Warde Medical Laboratory is authorized under Clinical Laboratory Improvement Amendments (CLIA) to perform high-complexity testing.</p>					

Performing Site:

WMRL: Warde Medical Laboratory 300 West Textile Road Ann Arbor MI 48108 (800)876-6522

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PAGE 2 OF 2

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Kajal V. Sitwala, MD, PhD - Medical Director