



LABORATORY REPORT

Example Client, XYZ123
1234 Warde Road
Ann Arbor MI 48108

EXAMPLE, REPORT W
WX0000003826 F 12/05/1988

Collected: 12/28/2023 14:52

Received: 12/28/2023 14:52

First Trimester Screen

Test Name	Result	Flag	Ref-Ranges	Units	Site
Screen Result	Negative		Negative		WMRL
Age at EDD (years)	37				WMRL
GA (weeks)	13				WMRL
GA (days)	2				WMRL
Ultrasound Date	12/20/2023				WMRL
CRL	55.1			milimetre	WMRL
Nasal Bone	N/A				WMRL
Weight (lbs)	150				WMRL
Multiple Gestation	Single				WMRL
Ethnic Origin	Other				WMRL
IDDM	None				WMRL
Smoker	No				WMRL
NT	1.0			milimetre	WMRL
PAPP-A	1,234.5			ng/mL	WMRL
hCG	55.0			IU/mL	WMRL
NT MOM	1.00				WMRL
PAPP-A MOM	1.00				WMRL
HCG MOM	1.00				WMRL
Down Syndrome Risk	1:19000				WMRL
Maternal Age Risk for Down	1:1000				WMRL
Trisomy 18 Risk	1:11000				WMRL
Interpretation	SeeBelow				WMRL

This is the initial sample received at Warde Laboratory for First Trimester Screening.

SCREEN NEGATIVE FOR DOWN SYNDROME AND TRISOMY 18.

This test does not screen for open neural tube defects (NTD). Please collect blood draw for MSAFP in the second trimester.

Additional Test Information:
The First Trimester Screen provides risk estimates, 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 1:270 or when the risk for Trisomy 18 is greater than or equal to 1:100.

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

F628000024

Ordered By: KAJAL V SITWALA, MD

WMB-23-1410

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PAGE 1 OF 2

Printed D&T: 12/28/2023 2:56 PM

Kajal V. Sitwala, MD, PhD - Medical Director



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Table header with columns: Test Name, Result, Flag, Ref-Ranges, Units, Site

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.

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.

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