



Sequenom Laboratories
 3595 John Hopkins Court
 San Diego, CA 92121
 CLIA #: 05D2015356 CAP #: 7527138
 Lab Director: Phillip Cacheris, MD, PhD

MaterniT® 21 PLUS (Core) + ESS
Triplet Gestation

FINAL REPORT

877.821.7266

Ordering Provider:	SanDiego, Testdoc	Patient:	Last, First
Provider Location:	Sequenom SD	DOB:	11/17/1999
Provider Phone:		Specimen:	252094001
Date Ordered:	07/26/2025	Fetal Fraction:	9%
Date Collected:	07/26/2025	Gestational Age ≥ 9w:	Yes
Date Received:	07/28/2025	External Accession:	
Order ID:	ORD25209-90002	Referral Clinician:	
Patient ID:	pid20925000043	Date Reported:	07/28/2025 06:26 PM PT

Test Result	<h1 style="margin: 0;">Positive</h1> <h2 style="margin: 0;">Trisomy 18</h2>
<p>Lab Director Comments</p> <p>This specimen showed an increased representation of chromosome 18, suggestive of trisomy 18 (Edwards syndrome). Genetic counseling, confirmatory diagnostic testing, and clinical correlation are recommended.</p> <p>This is a reported multifetal gestation that was found to be positive, and the Y chromosome was detected. This assay is unable to determine if one or more of the fetuses are affected and/or male.</p>	

Result Table

Content	Result
FETAL SEX	Consistent with Male
AUTOSOMAL ANEUPLOIDIES	
Trisomy 21 (Down syndrome)	Negative
Trisomy 18 (Edwards syndrome)	Positive T18 PPV*: 35.5%
Trisomy 13 (Patau syndrome)	Negative
SELECT WHOLE CHROMOSOMES	
Trisomy 16	Not Detected
Trisomy 22	Not Detected
SELECT MICRODELETIONS	
22q11 deletion (associated with DiGeorge syndrome)	Not Detected
15q11 deletion (associated with Prader-Willi / Angelman syndrome)	Not Detected
11q23 deletion (associated with Jacobsen syndrome)	Not Detected
8q24 deletion (associated with Langer-Giedion syndrome)	Not Detected
5p15 deletion (associated with Cri-du-chat syndrome)	Not Detected
4p16 deletion (associated with Wolf-Hirschhorn syndrome)	Not Detected
1p36 deletion syndrome	Not Detected

Positive Predictive Value

* Positive Predictive Value (PPV) estimates the probability that a pregnancy with a positive test result is in fact an affected pregnancy. The PPV for this patient was calculated only using maternal age and gestational age[1], test performance[2] and the standard PPV formula.

For a more accurate and individualized PPV calculation, include additional clinical information from the patient's clinical history (which may include serum screen results, personal/family history, ultrasound findings, etc.), and refer to the table below.

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A Priori Risk (1:X)	10	20	30	40	50	100	200	300	400	500	1000	1500	2000	2500	3000	5000
PPV (%) TRISOMY 18	96.5	92.9	89.6	86.5	83.6	71.6	55.7	45.5	38.5	33.4	20.0	14.3	11.1	9.1	7.7	4.8

Negative Predictive Value

The Negative Predictive Value (NPV) for trisomy 21, 18, and 13 is greater than 99%. The NPV for SCA and ESS cannot be calculated as SCA and ESS are only reported when an abnormality is detected.

About the Test

The MaterniT® 21 PLUS laboratory-developed test (LDT) analyzes circulating cell-free DNA from a maternal blood sample. This test is used for screening purposes and not diagnostic. Clinical correlation is recommended. Validation data on twin pregnancies is limited and the ability of this test to detect aneuploidy in higher multiple gestations has not yet been validated.

Test Method

Circulating cell-free DNA was purified from the plasma component of maternal blood. The extracted DNA was then converted into a genomic DNA library for aneuploidy analysis of chromosomes 21, 18, and 13 via next generation sequencing.[3] Optional findings based on the test order include sex chromosome aneuploidy (SCA)[2], and enhanced sequencing series (ESS)[4], which will only be reported on as an additional finding when an abnormality is detected. SCA testing includes information on X and Y representation, while ESS testing includes deletions in selected regions (22q, 15q, 11q, 8q, 5p, 4p, 1p) and trisomy of chromosomes 16 and 22.

Performance

The performance characteristics of the MaterniT® 21 PLUS laboratory-developed test (LDT) have been determined in a clinical validation study with pregnant women at increased risk for fetal chromosomal aneuploidy.[2-5]

Fetal Sex	Accuracy: 99.4%	
Region (associated syndrome)	Estimated Sensitivity**	Estimated Specificity
Trisomy 21 (Down Syndrome)	99.1%	99.9%
Trisomy 18 (Edwards Syndrome)	>99.9%	99.6%
Trisomy 13 (Patau Syndrome)	91.7%	99.7%
Sex Chromosome Aneuploidies (singleton gestation only)	96.2%	99.7%

* As reported in ISCA database nstd37 [https://www.ncbi.nlm.nih.gov/dbvar/studies/nstd37/]

** Sensitivity estimated across the observed size distribution of each syndrome [per ISCA database nstd37] and across the range of fetal fractions observed in routine clinical NIPT. Actual sensitivity can also be influenced by other factors such as the size of the event, total sequence counts, amplification bias, or sequence bias.



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Limitations of the Test

While the results of these tests are highly reliable, discordant results, including inaccurate fetal sex prediction, may occur due to placental, maternal, or fetal mosaicism or neoplasm; vanishing twin; prior maternal organ transplant; or other causes. These tests are screening tests and not diagnostic; they do not replace the accuracy and precision of prenatal diagnosis with CVS or amniocentesis. A patient with a positive test result should be referred for genetic counseling and offered invasive prenatal diagnosis for confirmation of test results. [6] The results of this testing, including the benefits and limitations, should be discussed with a qualified healthcare provider. Pregnancy management decisions, including termination of the pregnancy, should not be based on the results of these tests alone. The healthcare provider is responsible for the use of this information in the management of their patient. Sex chromosomal aneuploidies are not reportable for known multiple gestations. A negative result does not ensure an unaffected pregnancy nor does it exclude the possibility of other chromosomal abnormalities or birth defects which are not a part of these tests. An uninformative result may be reported, the causes of which may include, but are not limited to, insufficient sequencing coverage, noise or artifacts in the region, amplification or sequencing bias, or insufficient fetal fraction. These tests are not intended to identify pregnancies at risk for neural tube defects or ventral wall defects. Testing for whole chromosome abnormalities (including sex chromosomes) and for subchromosomal abnormalities could lead to the potential discovery of both fetal and maternal genomic abnormalities that could have major, minor, or no, clinical significance. Evaluating the significance of a positive or a non-reportable result may involve both invasive testing and additional studies on the mother. Such investigations may lead to a diagnosis of maternal chromosomal or subchromosomal abnormalities, which on occasion may be associated with benign or malignant maternal neoplasms. These tests may not accurately identify fetal triploidy, balanced rearrangements, or the precise location of subchromosomal duplications or deletions; these may be detected by prenatal diagnosis with CVS or amniocentesis. The ability to report results may be impacted by maternal BMI, maternal weight, maternal systemic lupus erythematosus (SLE) and/or by certain pharmaceutical agents such as low molecular weight heparin (for example: Lovenox®, Xaparin®, Clexane® and Fragmin®).

Note

Sequenom, Inc. is a subsidiary of Laboratory Corporation of America Holdings, using the brand Labcorp. This test was developed and its performance characteristics determined by Labcorp. It has not been cleared or approved by the Food and Drug Administration. This laboratory is certified under the Clinical Laboratory Improvement Amendments (CLIA) as qualified to perform high complexity clinical laboratory testing and accredited by the College of American Pathologists (CAP).

References

1. Snijders RJ, et al. *Fetal Diag.* 1995;10(6):356-367.
2. Mazloom AR, et al. *Prenat Diag.* 2013;33(6):591-597.
3. Palomaki GE, et al. *Genet Med.* 2012;14(3):296-305.
4. Zhao C, et al. *Clin Chem.* 2015 Apr;61(4):608-616.
5. Palomaki GE, et al. *Genet Med.* 2011;13(11):913-920.
6. ACOG/SFM Practice Bulletin No. 226, Oct 2020.

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