Pioneering science, personalized service

Cost estimator
Wondering what your out-of-pocket costs may be? Visit sequenom.com/everymom, select a test name, enter your insurance info, and get an estimate in seconds.

Convenient blood draws
Getting your blood drawn is easier than ever. As a LabCorp company, we have a nationwide network of patient service centers, allowing for convenient access to sample collection. Visit www.LabCorp.com to find your nearest location.

Genetic counseling
Patients with a positive test result may be offered counseling, and Sequenom and Integrated Genetics offer the largest national commercial network of genetic counselors to help inform and support patients.

Every Mom Pledge
We believe every mom should have access to the best possible care. That’s why we work directly with every patient to make sure our testing services are both accessible and affordable.

Watch a short video to learn about the test: sequenom.com/videos

Sequenom Center for Molecular Medicine, LLC d/b/a Sequenom Laboratories, a wholly owned subsidiary of Sequenom, Inc., is a CAP-accredited and Clinical Laboratory Improvement Amendment (CLIA)-certified molecular diagnostics laboratory dedicated to improving patient outcomes by offering revolutionary laboratory-developed tests for a variety of prenatal conditions. Sequenom, Inc. is a wholly owned subsidiary of Laboratory Corporation of America Holdings.

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References
Did you know that information on your baby’s health can be found in your own bloodstream?

The MaterniT 21 PLUS test analyzes genetic information that enters your bloodstream from the placenta. It screens for certain chromosomal abnormalities that could affect your baby’s health and development—such as trisomy 21 (Down syndrome) and sex chromosome aneuploidies (SCAs, abnormal numbers of X or Y chromosomes)—and can also detect if you’re having a boy or a girl.

The MaterniT 21 PLUS test detects the following chromosomal abnormalities:

- **Trisomies**
  - Trisomy 21 (Down syndrome)
  - Trisomy 18 (Edwards syndrome)
  - Trisomy 13 (Patau syndrome)

- **SCAs***
  - 45,X (Turner syndrome)
  - 47,XXY (Klinefelter syndrome)
  - 47,XXX (Triple X syndrome)
  - 47,XYY (XYY syndrome)

There are many ways to get this information, including methods such as serum screens and diagnostic procedures such as amniocentesis.

As a noninvasive prenatal test, MaterniT 21 PLUS is different from both. It has higher detection rates than serum screening\(^1\) (determined to be 97.9% positive predictive value for trisomy 21 in a high-risk cohort\(^2\)), and requires only a blood draw from the mother; amniocentesis requires withdrawing fluid from around the developing baby.

Most women who get the MaterniT 21 PLUS will screen negative for chromosomal abnormalities and may not require further testing.

However, any patient with a positive test result may be offered genetic counseling and/or diagnostic testing for confirmation of test results.

\(^{*}\) Reported as an additional finding. Talk to your doctor about your options.

Accurate, easy-to-understand results

MaterniT 21 PLUS is the pioneering noninvasive prenatal test. Since its introduction in 2011, the test has been used by more than 600,000 women and has been validated in clinical studies.

And unlike some similar tests, MaterniT 21 PLUS delivers answers in terms you can easily understand, with clear positive or negative results for well known chromosomal abnormalities, such as trisomy 21 (Down syndrome), typically returned within five days.

Also, if you’re carrying twins, MaterniT 21 PLUS can detect common chromosomal abnormalities in multiple gestation pregnancies.

A noninvasive blood test

With a blood draw from you as early as nine weeks into your pregnancy, the MaterniT® 21 PLUS test can screen for certain chromosomal abnormalities that could affect your baby’s health and development, providing you with more information earlier in your pregnancy.