Pioneering science, personalized service

Understand your cost options

Visit our website for your cost estimate and personalized payment options and the opportunity to learn about our *Moms Helping Moms of Tomorrow* initiative.
www.integratedgenetics.com/transparency
844.799.3243

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 Integrated Genetics offers 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 you to make sure our testing services are accessible and out-of-pocket costs are transparent.

We are ready to help!

www.integratedgenetics.com
askSQNMCS@labcorp.com

General questions?

Call toll-free within the US
877.821.7266

Billing and cost questions?

Call toll-free within the US
844.799.3243

Every Mom Pledge team is ready to answer questions about your insurance coverage and cost options
integratedgenetics.com/transparency

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

Sequenom Laboratories
3595 John Hopkins Court
San Diego, CA 92121

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. Sequenom®, Sequenom Laboratories™, and MaterniT® are trademarks of Sequenom, Inc. This brochure is provided by Integrated Genetics as an educational service for healthcare providers and their patients.

Note: This material is provided for general information purposes only. It is not intended as a substitute for medical advice and/or consultation with a physician or technical expert.

REFERENCES

1. Internal data.

Integrated GENETICS
LabCorp Specialty Testing Group

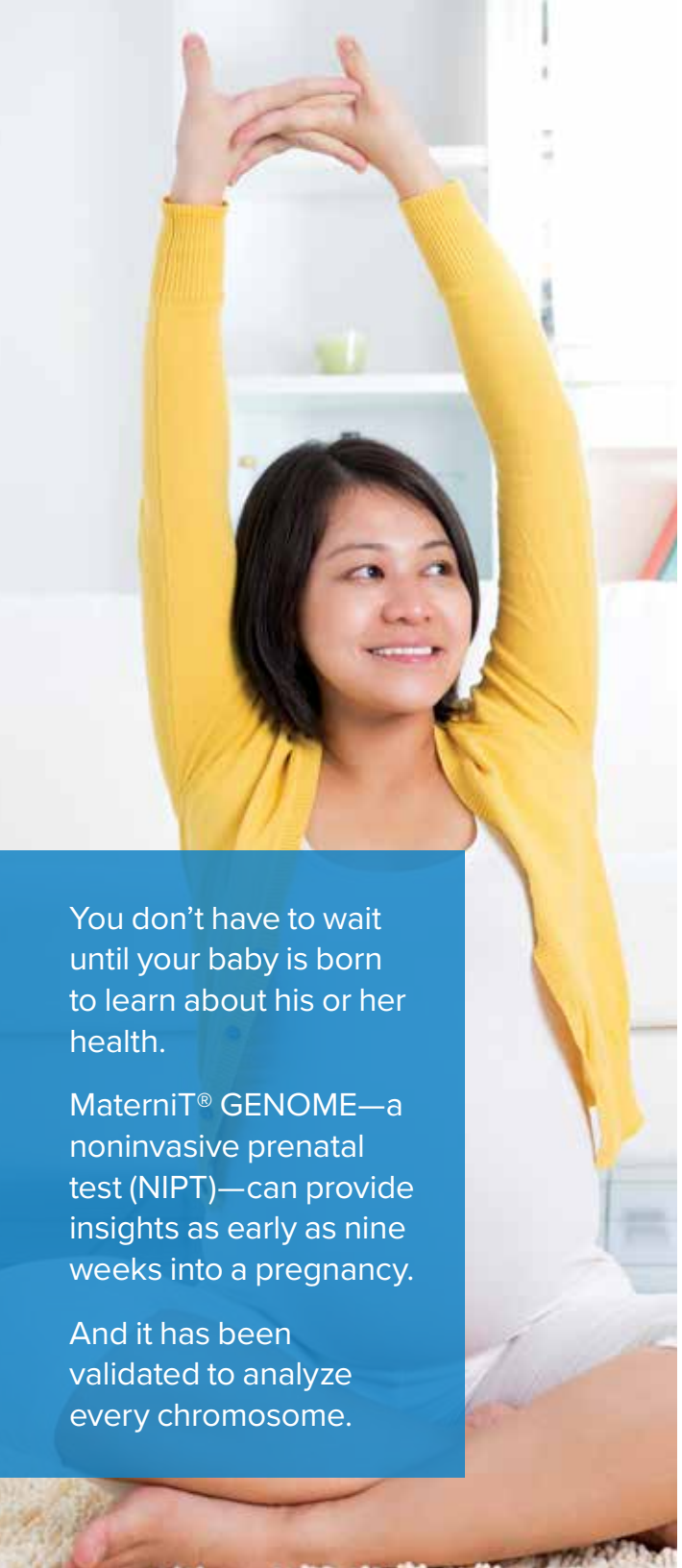
Rep-1038-v4-1019 | L17294-1019-4

MaterniT® GENOME

Screen for more than Down syndrome

A test that analyzes every chromosome to tell you more about your baby's health

Integrated GENETICS
LabCorp Specialty Testing Group



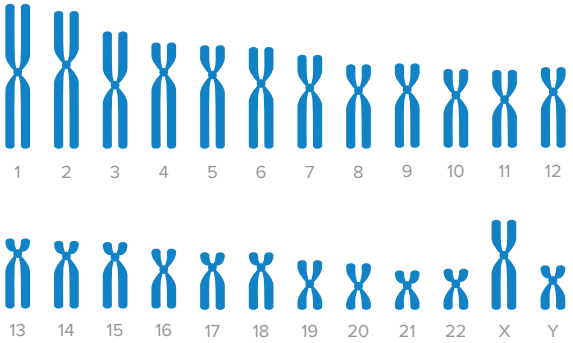
You don't have to wait until your baby is born to learn about his or her health.

MaterniT® GENOME—a noninvasive prenatal test (NIPT)—can provide insights as early as nine weeks into a pregnancy.

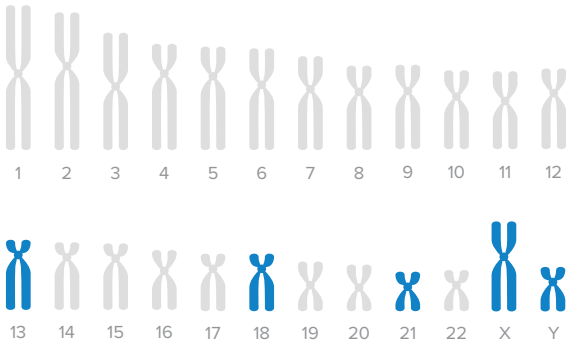
And it has been validated to analyze every chromosome.

Why does every chromosome matter?

Chromosomes are how cells transfer genetic information as a baby develops, and extra or missing parts of chromosomes, or whole chromosome changes, can severely impact the health of a baby. Most NIPTs analyze information from select chromosomes. But changes can be found in all chromosomes—which is why MaterniT GENOME analyzes them all.



Whole chromosomes analyzed by MaterniT GENOME



Whole chromosomes analyzed by most NIPTs

What makes MaterniT GENOME different?

After more than 70,000 tests resulted by Integrated Genetics, up to 30% of all positive findings could only be detected by MaterniT GENOME methodology.[1] Because most other NIPTs don't analyze for that 30%, they don't report on it. But that doesn't mean there's nothing to report.

What will MaterniT GENOME tell me?

Like most NIPTs, MaterniT GENOME can tell you if you screen positive or negative for trisomies 21 (Down syndrome), 18 (Edwards syndrome), and 13 (Patau syndrome), and if you're having a boy or a girl. But it can also find other chromosomal changes that may go undiagnosed at birth. Having information about these chromosomal changes before birth can help ensure your baby receives the proper and necessary support.

MaterniT GENOME reports on:

Any trisomy or monosomy	Trisomy: extra copy of a chromosome is present (3 instead of 2) Monosomy: missing copy of a chromosome (1 instead of 2)
Sex chromosome abnormalities	Atypical number of X and/or Y chromosomes beyond typical female (XX) or male (XY) complement
Partial chromosome abnormalities	Very small part of the chromosome is extra or missing

Results delivered clearly and quickly

Results from the MaterniT GENOME test are typically available within 5–10 days after your sample has been received in the laboratory. And while some NIPTs give you a risk score, MaterniT GENOME ensures screening results are communicated clearly—as positives or negatives.