PLAN FOR A HEALTHY FAMILY

When you are looking for actionable information to manage your pregnancy, the Super Panel provides a comprehensive assessment of your risk of being a carrier for a genetic condition that you could pass on to your child. **The Super Panel provides equitable care across populations with a >99% detection rate for most conditions and ethnicities.**

CARRIER SCREENING CAN HELP INFORM LIFE-SAVING INTERVENTIONS FOR YOUR NEWBORN

NXGEN'S PHILOSOPHY IS ABOUT EMPOWERING YOUR REPRODUCTIVE DECISIONS WITH ACTIONABLE INFORMATION

NxGen MDx

NxGen MDx

SUPER PANEL

There are many unknowns in life

FAMILY PLANNING DOESN'T HAVE TO BE ONE OF THEM

Ask your doctor about NxGen MDx carrier screening today and feel confident about tomorrow.

NxGen MDx, LLC is a leading healthcare company delivering comprehensive genetic testing. NxGen MDx's history of next generation sequencing combined with advanced technology allows us to provide accessible, high-quality testing options to families as they plan for the future.

www.nxgenmdx.com
Customer Service: 855-776-9436

facebook.com/nxgenmdx
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@nxgenmdx

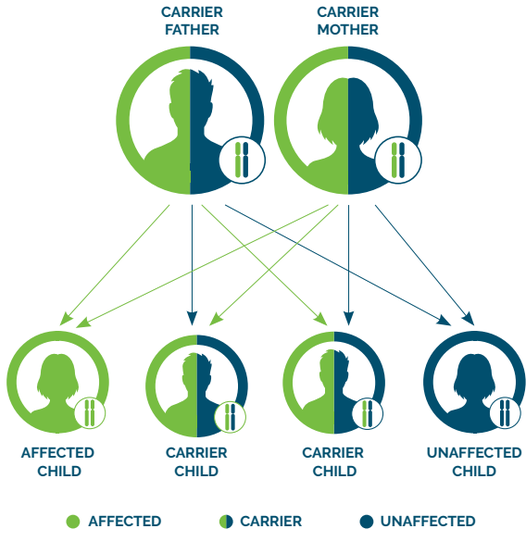
Revised (11/14/22)

NxGen MDx

What is Carrier Screening?

Carrier screening looks at your DNA to determine if you are a carrier of any autosomal recessive conditions (such as cystic fibrosis) or X-linked conditions (such as Fragile X). Carriers are generally healthy and typically don't show symptoms. However, if both reproductive partners are carriers of the same condition, there is a risk their baby could be affected by it.

How are Recessive Conditions Passed Down?



Two reproductive partners who both carry the same autosomal recessive condition have a **25% chance of having an affected child. With X-linked conditions, if the mother is a carrier, there is a 50% chance that male children will be affected.**

Why Choose the Super Panel?

1 in 300 pregnancies will be impacted by a genetic condition. The Super Panel screens for **conditions** that are common across multiple ethnicities including **35 conditions that are federally recommended for newborn screening (NBS).**

Minimize surprise NBS results by screening for all recommended NBS conditions

Maximize detection by screening for the most common conditions

If both you and your partner are carriers for the same condition, there are options available to you based on where you are in your reproductive journey:

Preconception	Pregnancy
Improved preconception planning	Prenatal diagnosis
In vitro fertilization (IVF)	Improved pregnancy and postnatal planning
Donor sperm/egg	Earlier diagnosis and timely treatment
Adoption	

See the complete list of Super Panel conditions at nxgenmdx.com

Want to know more about carrier screening? Scan this code to learn more.

What Do My Results Mean?

Negative means you are at a low risk of being a carrier

A negative result is reassuring because your chance of having a child with any of the conditions screened for is reduced. However no screen can detect all carriers. You still have a small remaining chance (known as your "residual risk") of being a carrier.

Positive means you are a carrier of a genetic condition

For those that screen positive, it is recommended that your reproductive partner also be screened for that condition. Our genetic counselors are available to review your results and discuss next steps.

Why NxGen MDx?

Simple & Convenient
Only requires a blood or saliva sample taken by your doctor.

Fast Turnaround Time
Results in approximately 2 weeks from sample receipt.

Accessible Genetic Counseling
Consultations with our board-certified genetic counselors to answer any questions you have about the screen or your results.

Personalized Customer Service
For questions, or to schedule an appointment with one of our genetic counselors, call us at **855-776-9436.**

Access for All
If you think you will have difficulty paying for the Super Panel, our Access for All Program may be able to help.

Visit **nxgenmdx.com/access-for-all** or call **855-776-9436** to learn more.

CareCredit®
Making care possible...today.

Special Financing Available*

For transactions of $200 or more made with your CareCredit credit card.

*Subject to credit approval. Minimum monthly payments required. Scan the QR code for details.

Access for All

The NxGen difference is making testing accessible for everyone

AFFORDABLE AND ACCESSIBLE GENETIC TESTING WITH NXGEN MDX

About NxGen MDx

NxGen MDx, LLC is a leading healthcare company delivering comprehensive genetic testing. NxGen MDx's history of next-generation sequencing combined with advanced technology allows us to provide accessible, high-quality testing options to families as they plan for the future.

Follow us on social media

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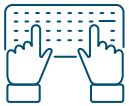
Updated: 8/10/22

NxGen MDx

What to Expect with NxGen MDx



1. Your healthcare provider orders a test and collects your sample to send to our lab.

2. We receive your sample and begin processing it.

3. We believe in access for all when it comes to genetic screening, and we will work with you to help you understand the available payment options.

4. If you think you will have difficulty paying for your testing, NxGen MDx offers an income-based Access for All Program.

Access for All Program

Our highest priority is ensuring testing is affordable and accessible. If you will have trouble paying for testing, you can apply for **the Access for All Program which can reduce the cost of testing for eligible applicants.**

The Access for All Program will determine your cost reduction based on **your income, your household size, and federally determined financial criteria.**

Note that only individuals using their health insurance or who are uninsured can qualify for the Access for All Program. Additionally, **patients who are covered under government-funded programs such as Medicaid, Medicare, Medicare Advantage, and TriCare are not eligible due to regulatory restrictions.**

How to Apply

To apply for the Access for All Program, you will need to provide the following:

- Insurance information
- Household income
- Household size
- Your most recent 1040 tax form

Apply for the Access for All Program at nxgenmdx.com/access-for-all

Contact the NxGen MDx Billing Department to:

- Ask questions related to your bill or insurance
- Get an estimated cost for testing
- Request a paper application for the Access for All Program

By phone at (855) 776-9436 or via email: cost@nxgenmdx.com