NewbornGene ID

Introducing NewbornGene ID™, a Comprehensive Genetic Carrier Test from GeneID Advanced Molecular Diagnostics, LLC

GeneID
The Future Today

What is carrier screening?

NewbornGene ID is a non-invasive carrier test for over 60 of the most prevalent and devastating hereditary diseases including:

Cystic Fibrosis	*Spinal Muscular Atrophy*	*Gaucher Disease*	*Alpha Thalassemia*
Fanconi Anemia	*Familial Dysautonomia*	*Bloom Syndrome*	*Beta Thalassemia*
Tay-Sachs	*Sickle Cell Disease*	*Usher Syndrome*	*Many others*

Why is carrier screening important?

Roughly 80% of all recessive occur in families with no known history of the disease and therefore have no indicator that they are carrying a disease causing mutation

In populations where testing is prevalent, incidence of disease has dropped by 90% or more (Source: ACOG Committee opinion 442)

How is carrier testing performed?

Test performed with a simple mouth rinse... no blood!

Who may be a carrier for a hereditary disease?

Anyone can be a carrier. Studies have shown that the likelihood of being a carrier can be as high as 1 in 4 in certain populations

Who is eligible for carrier screening?

Carrier screening is recommended by ACOG and ACMG for ALL patients of reproductive age who plan on having a child

What is the turn-around time and how are results reported?

Results returned within 8-10 days after insurance verification via web portal, hard copy and/or fax