

PGT-M

Preimplantation Genetic
Testing for Monogenic
Diseases

by **Igenomix**[®]

Helping couples at
risk of passing on a
genetic condition
have a healthy baby



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What is PGT-M?

PGT-M (formerly PGD) helps significantly decrease the chance of having a child with an inherited genetic disorder.

By analyzing DNA from each embryo, unaffected embryos can be preferentially selected for transfer.

This test is indicated for couples at risk for passing on a single gene disorder such as cystic fibrosis, fragile X syndrome, muscular dystrophy, Huntington disease, and many others.

Benefits of PGT-M



Identifies embryos affected with a genetic disorder prior to transfer.



Unique probe, custom-designed for every couple.



In-depth genetic counseling sessions available at no extra cost.



Igenomix understands each patient and situation is unique. It is our promise to customize the process to each couple's individual needs.



How does it work?



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