

Preimplantation Genetic Testing

There are 3 types of preimplantation genetic testing (PGT). All require in vitro fertilization (IVF).

PGT-M monogenic

formerly known as PGD for single gene disorders

EXAMPLES: cystic fibrosis, BRCA, Huntington's disease PGT-SR structural

formerly known as PCD for chromosome reastrangements

EXAMPLES: chromosome translocations, inversions

SEE OUR BLOG POST explaining the terminology in more detail

PGT-A aneuploidy

formerly known as PGS or CCS

screens for the presence of all 46 chromosomes



Preimplantation Genetic Testing



screens for the presence of all 46 chromosomes to identify embryos most likely to implant and result in a healthy pregnancy

FORMERLY KNOWN AS PGS (PREIMPLANTATION GENETIC SCREENING) OR CCS (COMPREHENSIVE CHROMOSOME SCREENING)



Preimplantation Genetic Testing

PGT-M monogenic

For patients who are at risk of passing on a known inherited genetic condition like cystic fibrosis, BRCA & Huntington's disease

FORMERLY KNOWN AS PGD (PREIMPLANTATION GENETIC DIAGNOSIS) FOR SINGLE GENE DISORDERS



Preimplantation Genetic Testing

PGT-SR

structural rearrangements

for patients who carry a known chromosome rearrangement and are at risk of passing this on

> FORMERLY KNOWN AS PGD FOR CHROMSOME REARRANGEMENTS