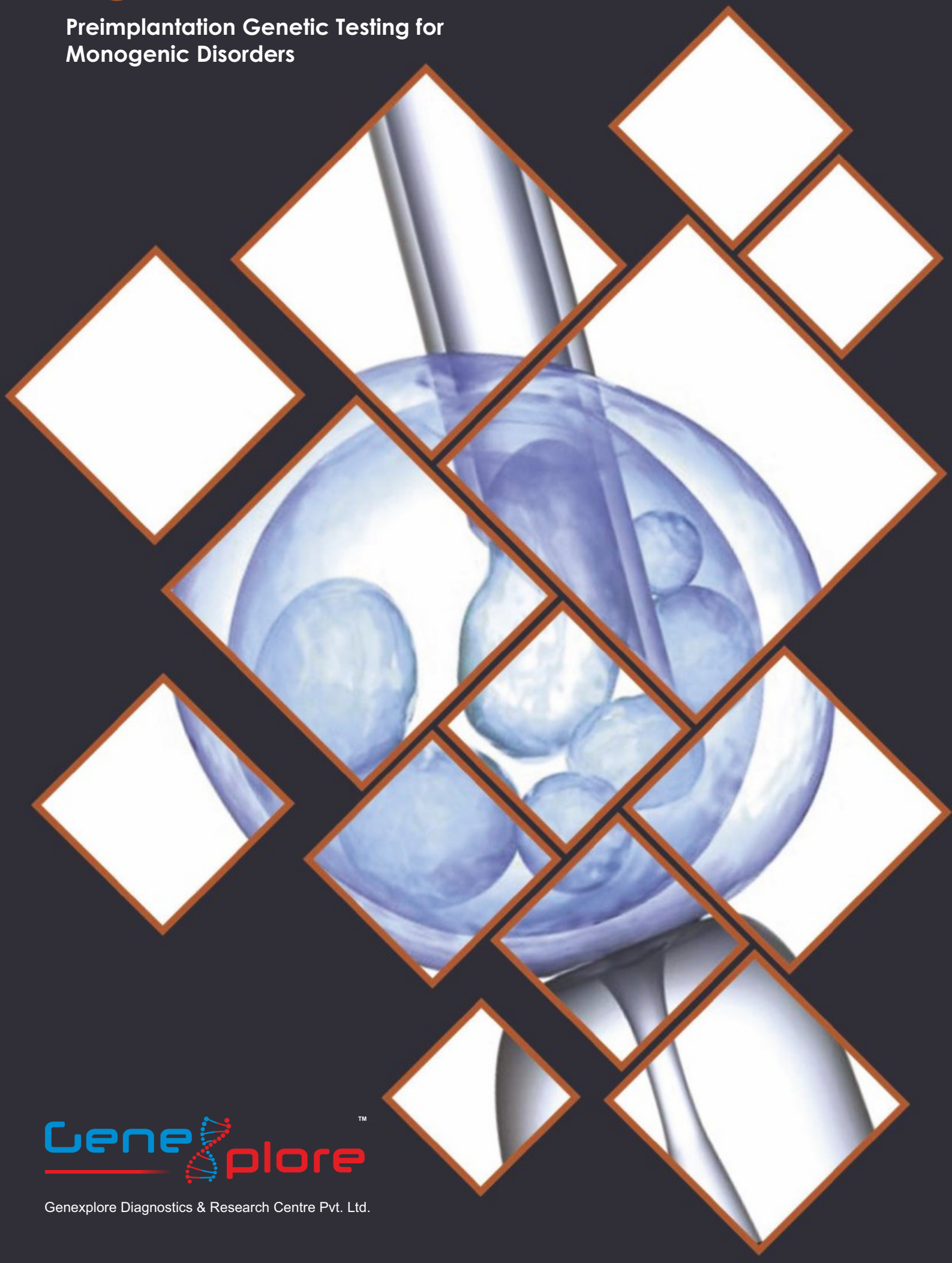
PGT - M

Preimplantation Genetic Testing for Monogenic Disorders



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PGT - M

- PGT-M enables detection of point mutation in single gene from embryo biopsy that can lead to Monogenic Disorders e.g. ß-thalassemia, Hemophilia, Cystic Fibrosis, Dystrophy.
- PGT-M helps to identify genetic mutation in embryo, ensuring healthy fetus.
- PGT-M from single/few cell/s of embryo will be analyzed for common identified monogenic disorders.

WHO SHOULD GO FOR PGT-M

- Both parents are the carriers of a single-gene Autosomal Recessive Genetic disorder.
- One of parent carries a mutation for a single-gene Autosomal Dominant Genetic disorder.
- One of the parent is a carrier of a chromosomal Microdeletion & Microduplications.

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