

Rapid preimplantation genetic testing (PGT) analysis from a single embryo biopsy

Combine PGT-A and PGT-M in one workflow

Simple integrated workflow:



Collect sample

Embryo biopsy



Perform library prep

- ✓ Ion Torrent[™] ReproSeq[™] PGS Kit
- PGD-SEQ™ kit







Prepare template and run sequence

- ✓ Ion Chef™ Instrument
- ✓ Ion GeneStudio™ S5 system



Analyze results

- ✓ Ion Reporter[™] Software
- PGD-SEQ[™] software

Key features include:



Comprehensive results from a single biopsy Perform PGT-A and PGT-M simultaneously



Easy to learn

Get started quickly with minimal training—no prior bioinformatics knowledge required



Fewer manual steps

Minimize staff time with automation and a combined workflow



Rapid turnaround

Reduce wait time by running samples in-house

Simultaneously identify an euploidy (A) and monogenic (M) disease biomarkers with a single embryo biopsy for reproductive health research:

PGT-A

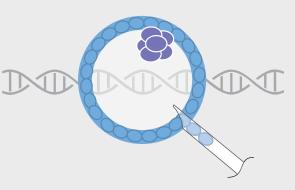
Identifies aneuploidies across 22 autosomes and X and Y chromosomes:

- Whole chromosome aneuploidy
- Mosaicism
- Segmental aneuploidy

PGT-M

Identifies relevant alleles for over 200 monogenic diseases:

- SMN1, spinal muscular atrophy
- CFTR, cystic fibrosis
- COL1A1, osteogenesis imperfecta
- FMR1, fragile X syndrome
- HBB, beta thalassemia
- HBA1 and HBA2, alpha thalassemia





Learn more at thermofisher.com/pgt

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