

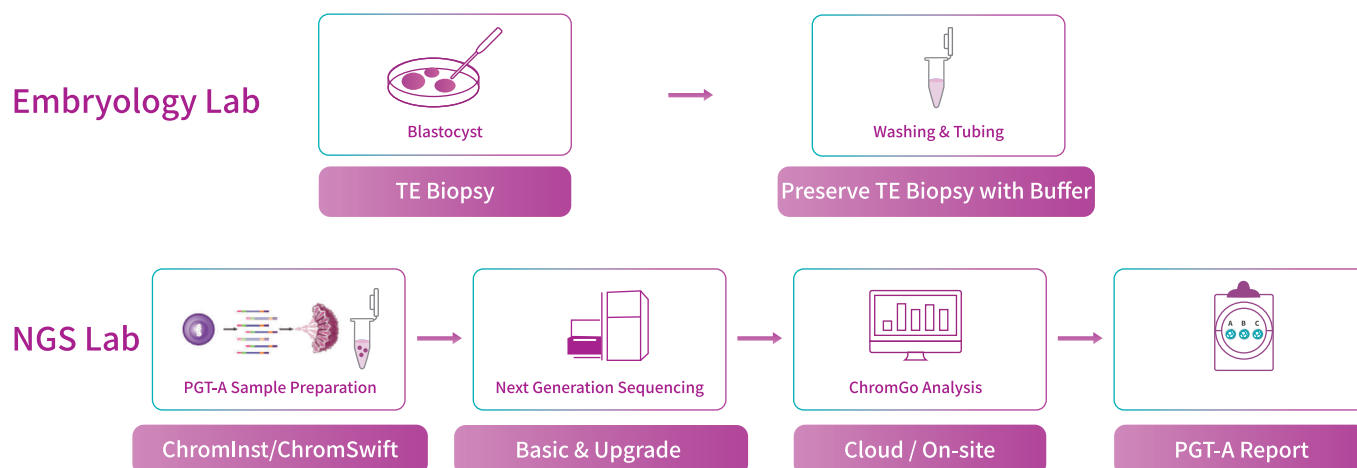
ChromInst®/ChromSwift®

Most Advanced PGT-A Test Solutions



	PGT-A Basic	PGT-A Upgrade (NGS Based SNP Analysis)
	ChromInst/ChromSwift	ChromInst/ChromSwift
Aneuploidy	•	•
Mosaicism	•	•
Segment Duplication and Deletion	•	•
Sibling QC	•	•
Genetic PN Check/Ploidy		•
UPD		•
mtDNA		•
Chromosomal Syndromes		• (40+)
NGS/TGS Sequencing Reads	200K (Chr resolution) -1M(Mb resolution),single end	3M, single end
Target Group	<ul style="list-style-type: none"> ✓ Advanced maternal age ✓ Infertility concerns ✓ Recurrent miscarriages ✓ Multiple IVF failed cycles 	<ul style="list-style-type: none"> ✓ History of chromosomal abnormalities ✓ 3PN embryo ✓ Reproductive history of triploidy/haploidy ✓ Previous molar pregnancy

Workflow



Advantages

One Set of
Reagents

One Analysis
Platform

One Sample
Needed

Two-Level PGT-A to Meet the Needs of Basic and Suspected Abnormal Embryos

Basic

- ✓ High-resolution for precise segmental abnormality detection
- ✓ Validated technology detects aneuploidies and mosaicism
- ✓ Sibling QC ensures genetic relatedness of tested embryos

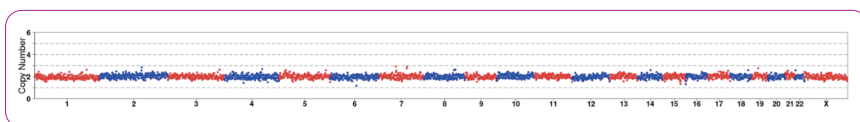
Upgrade

- ✓ Identifying true 2PN embryos from morphological 1PN and 3PN embryos
- ✓ UPD detection eliminates the occurrence of chromosome pair inherited from one parent

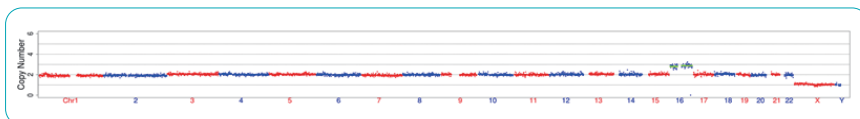
Who Is PGT-A For?

- Advanced maternal age.
- Couples with infertility concerns.
- Couples with recurrent miscarriages.
- Couples with many failed IVF cycles.
- Couples with a family history of chromosomal abnormalities.
- Couples undergoing IVF treatment for other reasons.

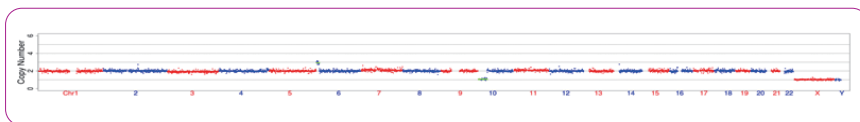
Reliable PGT-A Results



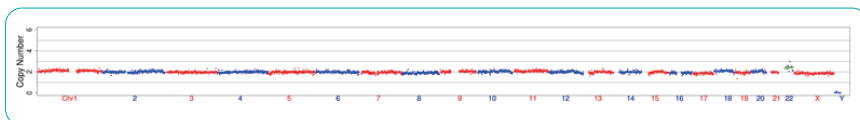
CNV Result: Paternal Triploidy 23X



CNV Result:46,XN,+16(X3)



CNV Result:46,XN,
+6p(p25.3→p24.2,~11Mb,X3),
-10p(p15.3→p11.22,~34Mb,X1)



CNV Result:46,XN,+22(X3,mos,~39%)