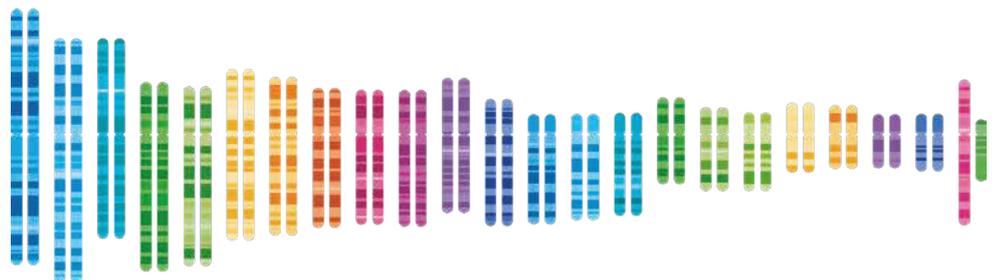


Genie Solution

The Genie Solution provides comprehensive genetic analysis of pre-implantation embryo biopsy samples including the detection of aneuploidy, structural rearrangements and linkage analysis to detect the inheritance of single gene disorders. End-to-end solution where, sample libraires are prepared with **Genie Kits**, sequenced on the **Genie Sequencer** followed by data analysis and reporting on the **Genie Data Analysis Software**.



GENIE-A SOLUTION

Genie-A is a solution for detecting CNV, ploidy, Sibling QC and ROH (regions of homozygosity) in preimplantation embryo biopsy samples. We offer two approaches: **Genie-A Fast** and **Genie-A Advanced**.

KEY FEATURES



COMPREHENSIVE DETECTION.

A single solution to detect CNV, Ploidy, Sibling QC and ROH (regions of homozygosity).



DEFAULT SNP ANALYSIS.

Genie-A uses genome-wide SNP information to detect Ploidy, Sibling QC and ROH.



HIGH RESOLUTION.

$\geq 4\text{Mb}$ CNVs (unknown), and $\geq 1\text{ Mb}$ inherited CNVs (known).



HIGH THROUGHPUT AND SCALABILITY.

5M reads/samples, 1x100 bp reads, Up to 48, 96 samples/run.

| Genie-A Kits | Analysis Methods | Reporting Content |
|-------------------------|--|--|
| Genie-A Fast | <ul style="list-style-type: none">• CBS Algorithm• Log Likelihood Ratio (LLR)• SNPs BAF Analysis | <ul style="list-style-type: none">• Aneuploidy, Ploidy (Triploidy, Haploidy)• $\geq 4\text{Mb}$ CNVs in unknown regions• $\geq 1\text{Mb}$ known inherited CNVs• Mosaicism $\geq 30\%$ C $\geq 10\text{Mb}$• Sibling QC• Whole chromosome level ROH |
| Genie-A Advanced | | |

GENIE-A: TWO APPROACHES



GENIE-A FAST:

Fast workflow, combining WGA and library preparation to reduce cost and time.

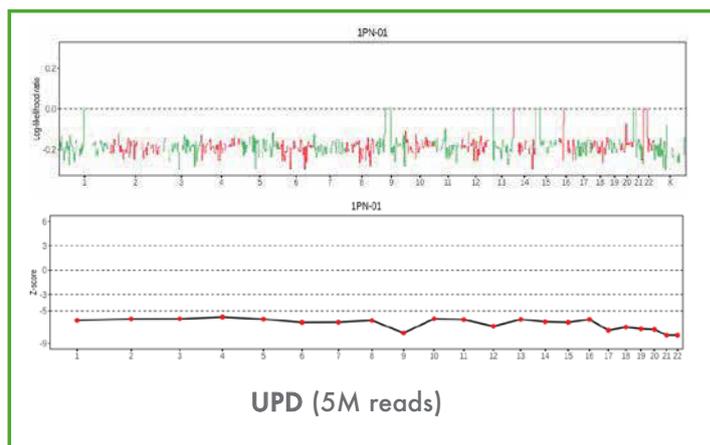
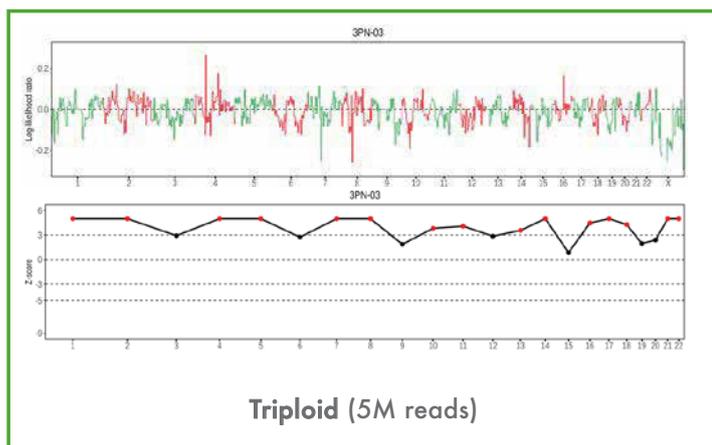
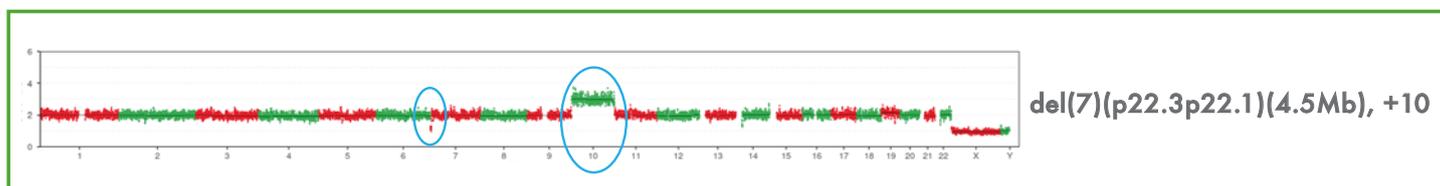


GENIE-A ADVANCED:

Separate WGA (MDA) followed by a patented library preparation method to support wider genome coverage.

TWO VERSIONS OF GENIE-A KITS WITH DIFFERENT SAMPLE PREPARATION METHODS FOR GREATER FLEXIBILITY, BUT THE SAME OUTCOME.

ACCURATE DETECTION OF CNV AND PLOIDY



GENIE-PLUS SOLUTION

Genie-Plus Solution is a genome-wide sequencing solution designed to detect the inheritance of monogenic disorders, structural rearrangements, in addition to CNV, ploidy, UPD, Sibling QC and regions of homozygosity (ROH) in preimplantation embryo biopsy samples.

KEY FEATURES



COMPREHENSIVE DETECTION.

A single solution to detect the inheritance of monogenic disorders, structural rearrangements, in addition to CNV, Ploidy, UPD, sibling QC and ROH.



GENOME-WIDE COVERAGE.

WGA by MDA and a patented library preparation method for better whole-genome coverage.



HIGH RESOLUTION.

$\geq 4\text{Mb}$ CNVs (unknown), and $\geq 1\text{ Mb}$ inherited CNVs (known).



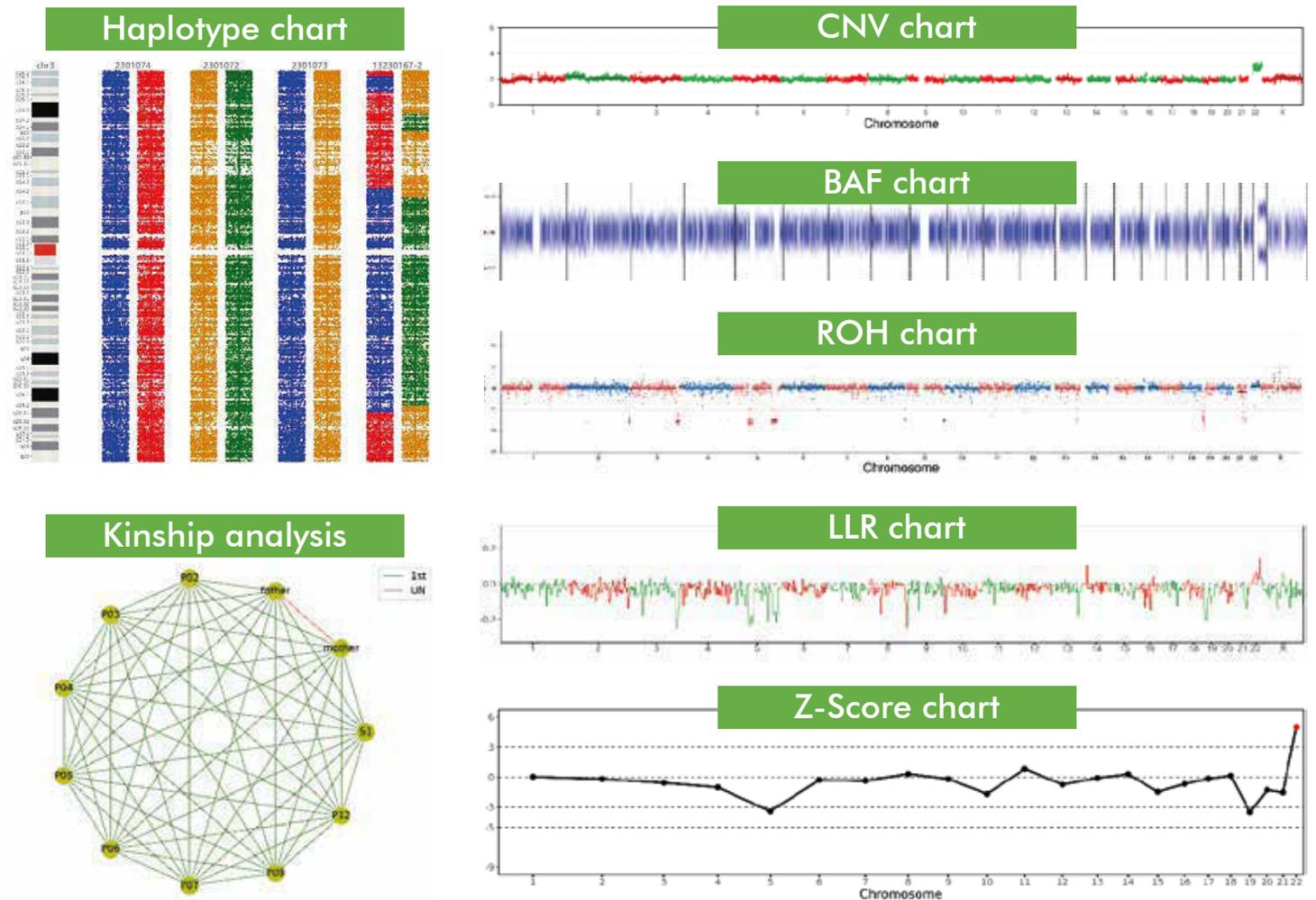
HIGH THROUGHPUT AND SCALABILITY.

40M reads/samples, 2x100 bp reads, Up to 24, 48 samples/run.



COMPLETE GENETIC ANALYSIS

The Genie-Plus solution performs comprehensive genetic analysis of embryo biopsy samples and provides results including, haplotype chart, CNV chart, BAF chart, ROH chart, LLR chart, Z-Score chart and Kinship analysis.



| Genie Kits | Analysis Methods | Reporting Content |
|-------------------|--|---|
| Genie-Plus | <ul style="list-style-type: none"> CBS Algorithm Log Likelihood Ratio (LLR) SNPs BAF Analysis | <ul style="list-style-type: none"> Aneuploidy, Ploidy (Triploidy, Haploidy) $\geq 4\text{Mb}$ CNVs in unknown regions $\geq 1\text{Mb}$ known inherited CNVs Mosaicism $\geq 30\%$ $C \geq 10\text{Mb}$ Sibling QC Whole chromosome level ROH |
| | <ul style="list-style-type: none"> Linkage Analysis SNP analysis CNV Analysis | <ul style="list-style-type: none"> > 4000 monogenic Disorders Balanced translocations, Robertsonian translocations, inversion |

GENIE SECURE DATA ANALYSIS SYSTEM

Highest security level and privacy during data analysis, data generated by the Genie Sequencer is transferred to the secure local server, through a LAN connection for data analysis on the Genie Data Analysis Software.

- **MAXIMUM SECURITY:**

All data analysis happens locally on a secure local server, eliminating cloud risks and ensuring full data protection.

- **100% LOCAL, 100% PRIVATE:**

Your data never leaves your infrastructure, ensuring complete control and compliance with strict security standards.



WITH GENIE, YOU GET CUTTING-EDGE TECHNOLOGY WITH UNMATCHED SECURITY



LOCAL PC

Genie result
(web)



LOCAL SERVER

Raw data
(rsync)



SEQUENCER

LOCAL LAB

GENIE SEQUENCER

The Genie Sequencer supports the sequencing of sample libraries prepared through the Genie Kits.

- Compact benchtop sequencer designed for ease of use and integration.
- High-throughput sequencer generates 500M reads/run, offering the ability to be used for various applications, including carrier screening, whole exome sequencing, and whole genome sequencing.
- Patterned flow cell facilitates highly accurate sequencing with Q30 >85%, decreased duplicates, and reduced index hopping.

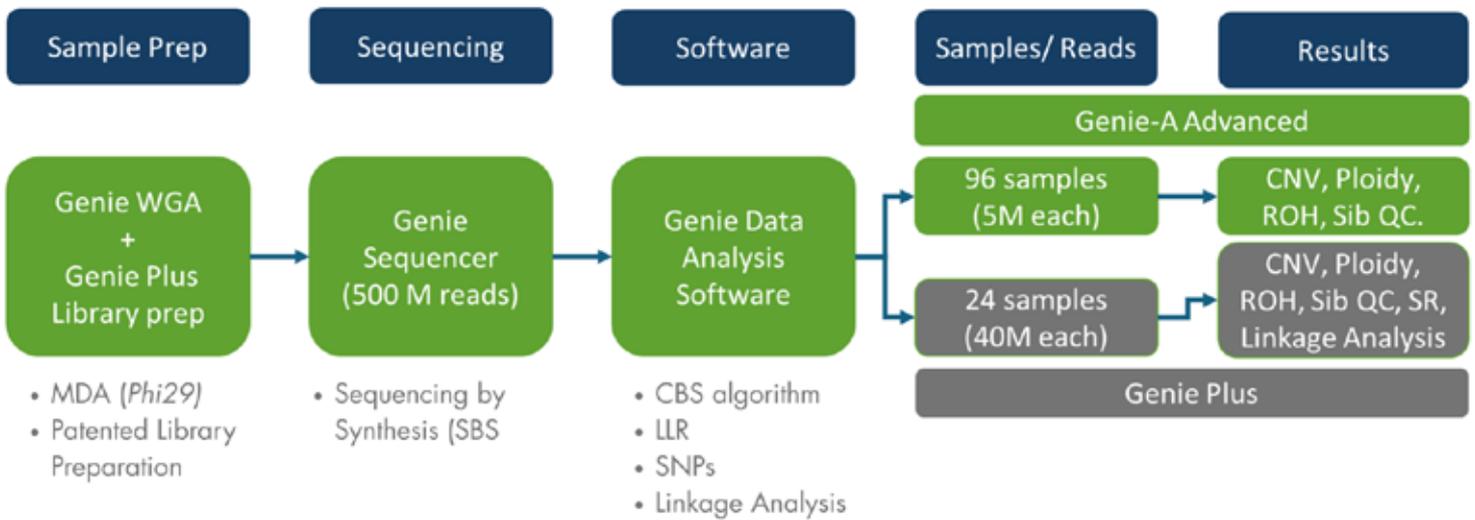


| Genie Solution | No. of Samples/run | Reads/sample | Read Length | Application |
|-------------------------|--------------------|--------------|-------------|--|
| Genie-A Fast | Up to 48 & 96 | 5M | 1 x 100 bp | CNV, Ploidy, Sibling QC, ROH |
| Genie-A Advanced | Up to 48 & 96 | 5M | 1 x 100 bp | CNV, Ploidy, Sibling QC, ROH |
| Genie-Plus | Up to 24 & 48 | 40M | 2 x 100 bp | Linkage analysis, SR, CNV, Ploidy, Sibling QC, ROH |



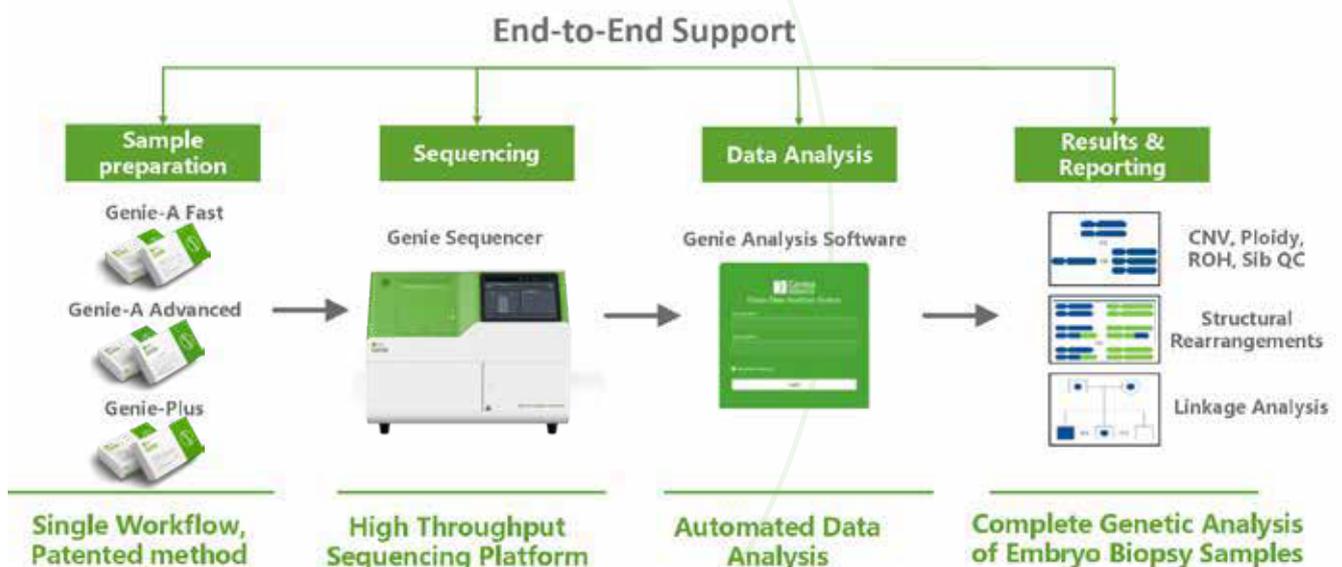
GENIE: SINGLE SOLUTION

A single patented workflow for the detection of CNV, ploidy, Sibling QC, ROH (Genie-A Advanced), structural rearrangements and linkage-based analysis to detect the inheritance of single gene disorders (Genie-Plus).



GENIE: END TO END SOLUTION

Genie provides an end-to-end solution for the genetic analysis of embryo biopsy samples. Sample libraires are prepared using the Genie kits, sequenced on the Genie Sequencer, followed by data analysis and reporting on the Genie Data Analysis Software.



GENIE KIT CATALOGUE

| Catalogue Number | Product Name | Packing List | Quantity |
|---------------------|--|---|----------|
| GEN-GAF-96-1 | Genie-A Fast for Genie Sequencer (96 samples/run) | Genie-A Sample Prep Kit (96 reactions) | 1 |
| | | Genie Sequencing kit | 1 |
| | | Genie Sequencing Flow Cell | 1 |
| GEN-GAF-48-2 | Genie-A Fast for Genie Sequencer (48 samples/run x2) | Genie-A Sample Prep Kit (96 reactions) | 1 |
| | | Genie Sequencing kit | 2 |
| | | Genie Sequencing Flow Cell | 2 |
| GEN-GAA-96-1 | Genie-A Advanced for Genie Sequencer (96 samples/run) | Genie WGA kit (48 reactions) | 2 |
| | | Genie-Plus Sample Prep Kit (50 reactions) | 2 |
| | | Genie-Plus Index Primers (96 indexes) | 1 |
| | | Genie Sequencing kit | 1 |
| | | Genie Sequencing Flow Cell | 1 |
| GEN-GAA-48-1 | Genie-A Advanced for Genie Sequencer (48 samples/run) | Genie WGA kit (48 reactions) | 1 |
| | | Genie-Plus Sample Prep Kit (50 reactions) | 1 |
| | | Genie-Plus Index Primers (96 indexes) | 1 |
| | | Genie Sequencing kit | 1 |
| | | Genie Sequencing Flow Cell | 1 |
| GEN-GAP-24-2 | Genie-Plus for Genie Sequencer (24 samples/run x2) | Genie WGA kit (48 reactions) | 1 |
| | | Genie-Plus Sample Prep Kit (50 reactions) | 1 |
| | | Genie-Plus Index Primers (96 indexes) | 1 |
| | | Genie Sequencing kit | 2 |
| | | Genie Sequencing Flow Cell | 2 |
| GEN-GAP-24-4 | Genie-Plus for Genie Sequencer (24 samples/run x4) | Genie WGA kit (48 reactions) | 2 |
| | | Genie-Plus Sample Prep Kit (50 reactions) | 2 |
| | | Genie-Plus Index Primers (96 indexes) | 1 |
| | | Genie Sequencing kit | 4 |
| | | Genie Sequencing Flow Cell | 4 |
| GEN-WGA-01* | Genie WGA kit | Genie WGA kit (48 reactions) | 1 |

*Common to both Genie-A Advanced and Genie-Plus

For detailed catalogue options to support GENIE-SEQ-01, please contact your Genea Biomedx representative.

Please refer to the instructions for use.

For further information, please contact your Service Representative or visit:

www.geneabiomedx.com