



GENEWIZ Multiomics & Synthesis Solutions from Azenta Life Sciences is committed to providing reliable and scalable solutions across our clinical portfolio that produce the high-quality results required to support clinical studies. Our clinical lab follows Good Clinical Practices (GCP) guidelines and is CLIA certified by The Centers for Medicare & Medicaid Services (CMS) and accredited by the College of American Pathologists (CAP) for clinical NGS and Sanger sequencing. Whether you are a pharmaceutical or biotechnology company, our clinical services offer the flexibility to accommodate virtually any size project with the security of your data top of mind. When working with us, all your information, data, and intellectual property are treated with the utmost care and security so you can be sure your data is safe.

CLIA Capabilities

The following clinical laboratory assays are performed by CAP/CLIA-trained scientists on CAP/CLIA-qualified equipment, for research use only (RUO). For additional or customized CLIA capabilities, please contact us at clia@azenta.com.

Genomics

- Whole Genome Sequencing (WGS)
- Low-Pass WGS
- Whole Exome Sequencing (WES)
- Targeted Panels
- Amplicon Sequencing
- Metagenomics
- Genotyping

Transcriptomics

- Whole Transcriptome Sequencing
- mRNA Sequencing
- Small RNA Sequencing
- Immuno-Profiling
- qPCR/dPCR
- Spatial Profiling

Epigenomics

- Whole Genome Methylation Sequencing
- Targeted Methylation Sequencing

Proteomics

- Olink® Target Panels
- Olink® Explore Panels

Good Clinical Practice (GCP)

In addition to our CAP/CLIA certifications, our labs are now Good Clinical Practices (GCP) compliant (as applicable) in accordance with FDA 21 CFR Part 11 and 58 as well as adherence to the guidelines mentioned by the [European Medicines Agency \(EMA\)](#).

We are proud of our commitment to maintain the highest quality standards, ensuring regulatory compliance for our customers. Our labs meet clinical requirements with precision and excellence so that your data is accurate, and your future patients are safe.

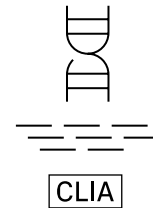


Clinical Services

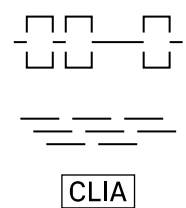


CLIA Registered Tests

GENEWIZ has the following CAP-accredited and CLIA-licensed validated tests available for commercial use; both assays are registered with CMS and the New Jersey Department of Health. Starting sample materials for these tests include genomic DNA, whole blood, and cell pellets (WES). Additional starting materials would necessitate a fit-for-purpose validation.



Whole Genome



Whole Exome

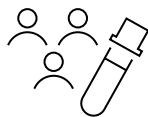
Features & Benefits

- CAP-accredited and CLIA-licensed laboratory and adherence to GCP guidelines with quality assurance oversight.
- Complete solution pipeline from nucleic acid extraction and sequencing to data analysis and sample storage.
- Dedicated Study Manager for proactive, transparent communication throughout the entire project.
- Superior data quality exceeding manufacturer benchmarks.
- Assay development expertise excelling in assay optimization and handling difficult templates.
- Industry-leading turnaround times with options for expedited assay development and sequencing.

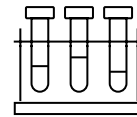
Applications



Clinical Trials



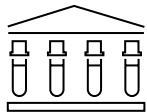
Clinical Research Studies



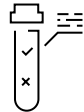
In Vitro Diagnostic Confirmation



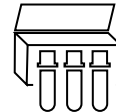
Diagnostic Assay Development



Sample Storage



Sample Sourcing



Sample Collection Kits



Explore our Clinical Service Solutions

