A Leader in High Quality Pharmacogenomics Services

Gentris is a global provider of applied pharmacogenomic services. As pioneers in the field of pharmacogenomics, Gentris helps pharmaceutical companies and clinical research organizations effectively integrate pharmacogenomics into their drug development programs to deliver safer, more effective compounds to the market more quickly.

Quality, expertise, and customer satisfaction drive Gentris’ core business model in Innovative Genomic Biomarker Solutions. We provide seamless and dynamic coupling of comprehensive project life cycle management and full logistical support with the power of tested platforms for sample extraction and purification, genotyping, gene expression analysis, and highly customized pharmacogenomics (PGx) applications and expertise.

Gentris is transforming global drug development with clinical pharmacogenomic solutions that will help identify more qualified target populations for clinical studies and optimize safety and efficacy of therapeutic compounds. The result is an overall improvement in Return on Investment, an important consideration in these days of escalating drug discovery and development costs.

Comprehensive Quality Solutions

Genomics Services
- Genotyping
- Gene Expression
- Next Generation Sequencing

Clinical Applications
- Full-scale PGx-enabled Therapeutic Programs
- Patient Inclusion, Exclusion, and Stratification
- Support Phase I-IV, Endpoint, and Retrospective Analysis

Consulting Services
- Study Design and IRB Development
- Sample Acquisition and Collection
- Regulatory Data Submission Support

Genomics Capabilities
- Taqman® Allelic Discrimination
- Taqman® qRT-PCR
- Sanger Sequencing
- Affymetrix® Microarray
- Ion Torrent™ Personal Genome Machine
- DMET™ Plus Premier
- Sequenom MassARRAY Analyzer 4

Therapeutic Areas
- ADME
- Oncology
- Pain
- Immunology
- Inflammation
- CNS
- Custom Panels

Impact of Pharmacogenomics
- Minimize the occurrence & severity of adverse drug reactions
- Accelerate the drug discovery & development process
- Reduce cost of clinical trials

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www.gentris.com
The Gentris Advantage

Gentris’ clinical pharmacogenomics services are performed in a CLIA-certified, GCLP-compliant laboratory environment that exceeds the standards for quality, reliability, and consistency. Our clients have the assurance that the data generated to support their clinical trials is robust and will meet regulatory standards. Extensive validation procedures and a comprehensive Quality Assurance program, support our success rate in repeatedly passing rigorous on-site client audits.

Gentris at a Glance

- Founded in 2001
- >1 million sample biorepository
- 25,000 sq. ft. facility in Morrisville, NC
- CLIA, GLP, GCP, Safe Harbor, CAP (Biorepository)
- Supported >1,000 clinical studies globally
- Designed and validated >400 assays de novo

For more information, contact:

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