



Translational Genomics Research Outsourcing Services

Our translational genomics research outsourcing services bridge the gap between genomic discoveries and clinical applications, enabling biomarker validation, precision diagnostics, and personalized therapeutic strategies.

Our Translational Genomics Research Capabilities

Our genomic scientists and clinical researchers perform large-scale genomic analyses, integrate multi-omics data, and translate findings into actionable clinical insights.

Types of Translational Genomics Research We Handle

- Whole Genome and Exome Sequencing
- Clinical Variant Annotation
- Genotype-Phenotype Association Studies
- Biomarker Discovery and Validation
- Somatic and Germline Mutation Profiling
- Pharmacogenomics Analysis
- Multi-Omics Data Integration
- Pathway and Network-Based Disease Analysis
- Population Genomics for Disease Susceptibility
- Clinical Trial Genomics Support
- Tumor Genomics and Cancer Biomarkers
- Rare Disease Genomics
- Predictive and Prognostic Marker Identification
- Functional Genomics Validation
- Gene Expression Correlation Studies
- Regulatory and Submission-Ready Reports
- Publication-Ready Translational Genomics Data
- Cross-Lab Validation of Genomic Findings
- Custom Translational Genomics Research Projects

Key Research Outsourcing Services Offered

- Patient Sample Collection and QC
- Sequencing and Data Generation

- Clinical Grade Bioinformatics Analysis
- Biomarker Screening and Verification
- Functional Validation Studies
- Genotype-Phenotype Correlation Analysis
- Pathway Impact and Network Modeling
- Integration with Clinical Data
- Data Visualization and Interpretation
- Publication-Ready Reports and Graphs
- Stakeholder Presentation Preparation
- Confidential Data Handling and NDA
- Interim Technical Reports and Updates
- Workshops and Training in Translational Genomics
- Post-Project Technical Consulting
- IP and Patent Filing Support for Biomarkers
- Grant Proposal and Manuscript Assistance
- Secure Data Storage and Archiving
- Compliance with GLP, ISO, FAIR Standards
- Cross-Lab Validation of Workflows
- Collaboration with Hospitals and Clinical Labs
- Long-Term Translational Genomics Partnerships
- Custom SOP Development for Clinical Genomics
- Regulatory Dossier Preparation for Approvals
- Market-Ready Translational Genomics Reports

Why Choose Us for Translational Genomics Research Outsourcing?

Our deep experience in human genomics, robust data analysis pipelines, and strong clinical collaborations ensure reliable, clinically actionable genomic insights for precision medicine.

Industries & Sectors We Serve

- Clinical Research Organizations (CROs)
- Hospitals and Healthcare Systems
- Pharmaceutical and Biotech Companies
- Personalized Medicine Startups
- Academic Medical Research Centers
- Regulatory and Compliance Agencies

Customized Translational Genomics Solutions

We deliver validated genomic biomarkers, clinically annotated data, and interpretation reports tailored to your precision diagnostics and therapeutic development pipelines.

Quality Assurance & Regulatory Compliance

Our translational genomics workflows comply with GLP, ISO, and FAIR guidelines to ensure validated, reproducible, and regulator-ready deliverables for clinical use.

Case Studies & Client Success Stories

Learn how our translational genomics outsourcing has accelerated biomarker validation, improved patient stratification, and supported regulatory submissions. References available on request.

How It Works: Our Research Outsourcing Process

1. **Requirement Gathering:** Define clinical question, patient cohort, and desired outcomes.
2. **Proposal & Quotation:** Provide detailed plan, sequencing and analysis workflow, timeline, and cost.
3. **Data Generation and Analysis:** Perform sequencing, analyze variants, and validate biomarkers.
4. **Reporting:** Deliver clinical-grade reports, annotated variants, and submission-ready documents.
5. **Post-Project Support:** Offer follow-up consulting, manuscript drafting, and IP support.

Frequently Asked Questions (FAQs)

Q: Can you help with regulatory submissions?

A: Yes — we generate regulator-ready genomic data and reports for submission.

Q: Do you provide patient cohort analysis?

A: Absolutely — we handle cohort-level genotype-phenotype correlations and biomarker studies.

Q: How secure is my clinical genomic data?

A: We enforce strict NDAs, secure storage, and complete IP and data protection.

Get Started / Request a Quote

Contact us today to discuss your translational genomics project and receive a custom plan, timeline, and cost estimate aligned with your precision medicine goals.

Contact Us

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