

Clinical Trial Sample and Consent Tracking (CTST)

IQVIA Laboratories Digital Technologies offers **Labmatrix®**, a comprehensive, cloud-based sample management system designed to streamline clinical trial operations and maximize the value of biospecimens. With over 20 years of experience, IQVIA Laboratories Digital Technologies has a proven track record of successfully implementing and supporting Labmatrix for a wide range of clients, including top pharmaceutical companies, biotech firms, and academic institutions.

Why Labmatrix?



Enhanced sample oversight

- Centralized platform for tracking and managing samples throughout their lifecycle
- Ensures protocol adherence and reduces risk of lost or mishandled samples



Improved consent management

- Holds detailed consent metadata and consolidates allowable use parameters for compliant use of samples in future research



Real-time data insights

- Offers robust reporting, dashboards, and email alerts for real-time visibility facilitates proactive decision-making and timely identification of issues



Seamless integrations

- Integrates with various data sources, including Electronic Data Capture (EDC) systems, central labs, testing labs, LIMS, and third-party storage systems
- Ensures data consistency and eliminates manual data entry, saving time and reducing errors

KEY FEATURES

- Configurable data model, workflows, and user interfaces to meet specific client requirements
- Detailed chain of custody tracking for regulatory compliance
- Integration with Qigram, a patented graphical query and analysis tool for advanced data interrogation and visualization
- Customizable dashboards and reports for data-driven insights

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