Labwise® XD

Labwise® XD Provides a single platform in the cloud for all processes and demands of the laboratory Workplace

Key Benefits:

- Automated Reporting and document control for critical regulated files.
- Enterprise Risk Reduction and improved regulatory compliance.
- Cost reduction due to IT system consolidation
- Streamlined compliance monitoring and reporting
- Improved visibility into key laboratory workflows and reduction in cycle times.
- Configurable workflows driven by conditional business rules.
- Task triggered workflows allowing collaborative work.

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Labwise® XD is a cloud-based total business process management solution designed for the laboratory environment. Leveraging a highly configurable low code technology stack, Labwise® XD is the only solution which can fully integrate all laboratory operations management areas with a single cloud platform, facilitating full 21 CFR Part 11 compliance. Whether you are subject to GLP, GMP, or GCP, Labwise® XD provides a digital workplace for integrating critical workflow processes and data across the enterprise, interconnected on a single platform & database.

Labwise® XD is a unique, comprehensive solution for Quality, Governance, Risk and Compliance designed to optimize laboratory business process flows. By consolidating the functionality normally found in disparate systems, such as LIMS, Electronic Laboratory Notebooks (ELNs), Quality Management Systems (QMS), Electronic Data Capture Systems, Learning Management Systems, and others in to one platform, Labwise® XD provides substantial gains in productivity while minimizing IT costs.

Labwise® XD Features:

- Integration of Laboratory Environment. Labwise XD creates an integrat-ed lab environment in which lab processes are automated, methods are managed electronically, instruments are connected and results are generated through automated workflows.
- Link scientific data to critical controlled documents. Seamless integration allows users to access key documents such as SOPs, Material Safety Data Sheets (MSDS) documents, and Certificate of Analysis (C of A) documents just in time.
- Manage Deviations and Corrective Actions/Preventive Actions (CAPA). The system delivers workflow processes that enable proactive manage-ment of Corrective Actions/ Preventive Actions in a timely manner.

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ONE platform UNIQUELY designed to support the COMPLETE demands of Laboratory Departments.

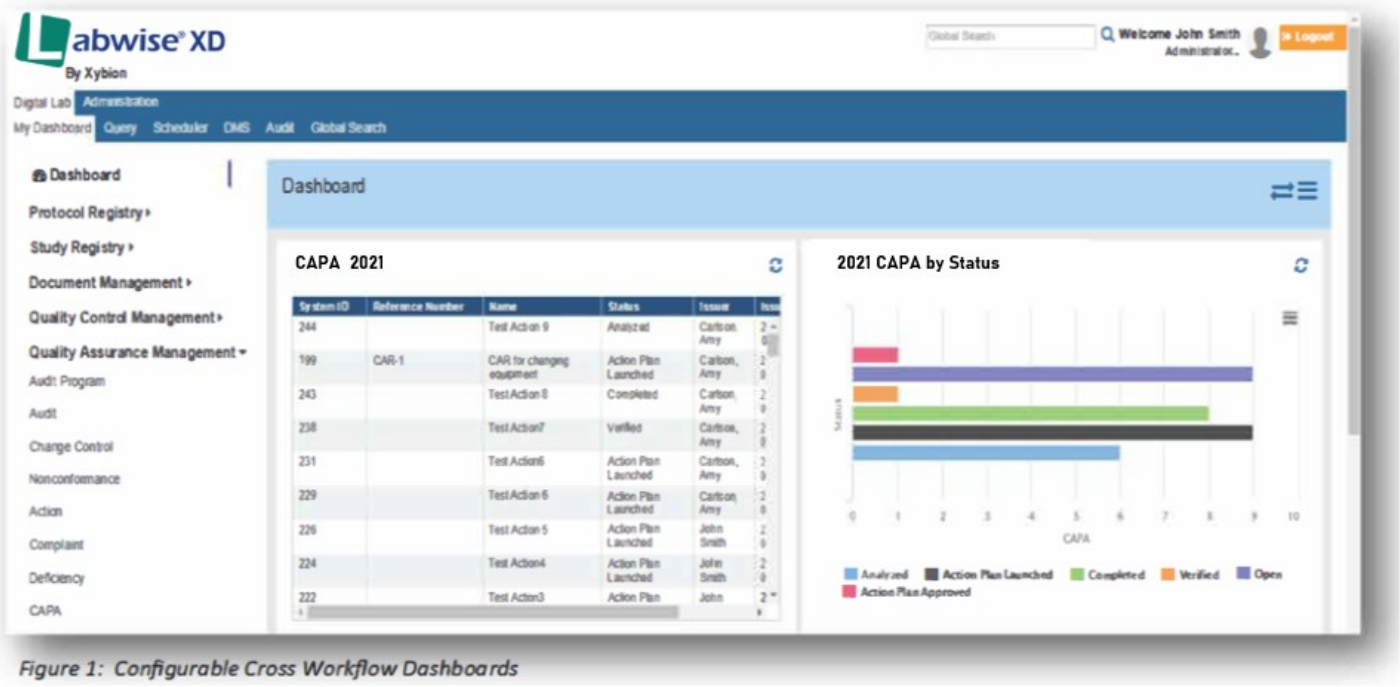


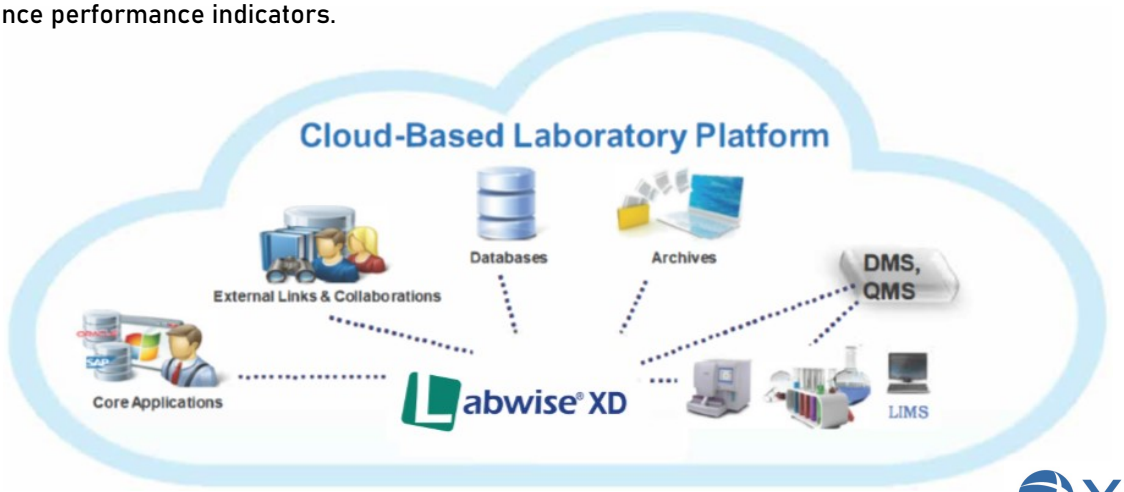
Figure 1: Configurable Cross Workflow Dashboards

Labwise® XD Features Continued:

- Manage Controlled Documents. The system provides out-of-the box capabilities for full versioning, document approvals, PDF rendering, document training records, connections to other CMIS compliant systems, and more.
- Conduct audits and track non-conformance. A robust feature set empowers your team to plan, prepare, and execute internal and external audits and accomplish day-to-day operational tasks related to all non-conformances.
- Streamline compliance monitoring and reporting. Easy-to-configure dashboards, scorecards and reports allow you to track compliance monitoring and report against key compliance performance indicators.

Xybion - Your Expert Partner

- Xybion has over 20 years of experience managing laboratory workflows and helping clients stream-line compliance and mitigate risk.
- Xybion has extensive global experience and works with companies of all sizes from multiple industries, including 150+ blue chip enterprise customers in 15 countries.
- Xybion offers deep domain knowledge of FDA regulations and has a long track record of helping clients build effective risk mitigation strategies.



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