

Labwise® XD is a powerful, flexible total business process management solution designed for the laboratory environment. Whether you are subject to GLP, GMP, or GCP, Labwise® XD provides a framework for integrating business critical workflow processes and data across the enterprise. With flexible deployment options, Labwise® XD can be installed on premise or implemented as a hosted service.

Labwise® XD Overview

Labwise® XD consolidates the functionality normally found in many disparate systems into one platform.

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.
- Business process enablement and enhanced efficiencies.
- Improved visibility into key laboratory workflows and reduction in cycle times

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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 (OMS), 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.** Can be used as a standalone laboratory operations management system or integrated with Pristima® XD Core, the industry standard for total preclinical data management.
- **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 management of Corrective Actions/ Preventive Actions in a timely manner.
- **Manage controlled documents** with full versioning, PDF rendering, user access controls, and storage and retrieval of all documents including lab results and instrument data across all sites, locations and work areas.

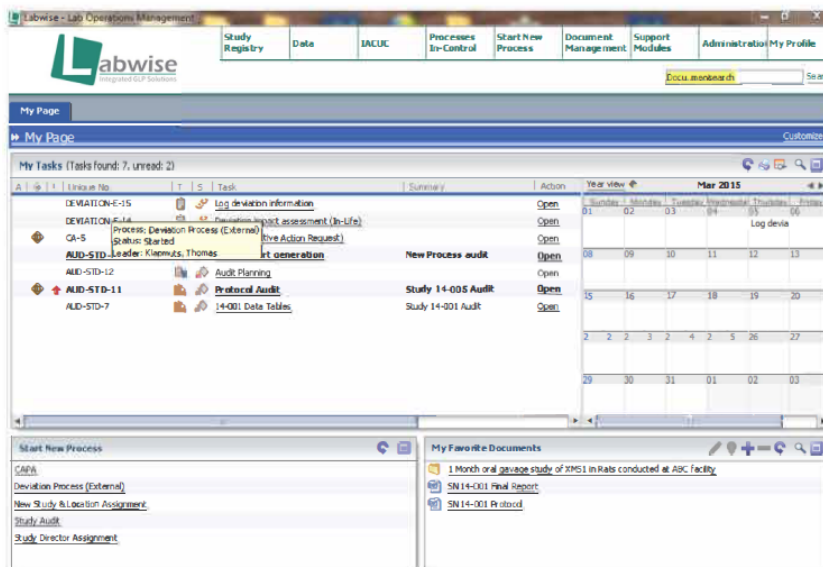


Figure 1. Labwise home page provides easy access to key tasks, processes, and documents.

- **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

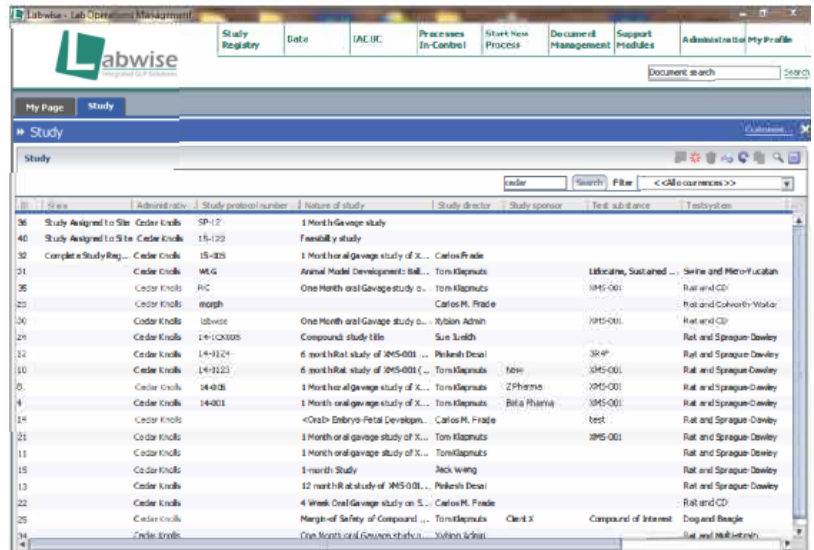


Figure 2. Manage the Master Schedule.

Xybion - Your Expert Partner

- Xybion has over 20 years of experience managing laboratory workflows and helping clients streamline compliance and mitigate risk.
- Xybion has extensive global experience and works with companies of all sizes from multiple industries.
- Xybion offers deep domain knowledge of FDA regulations and has a long track record of helping clients build effective risk mitigation strategies.

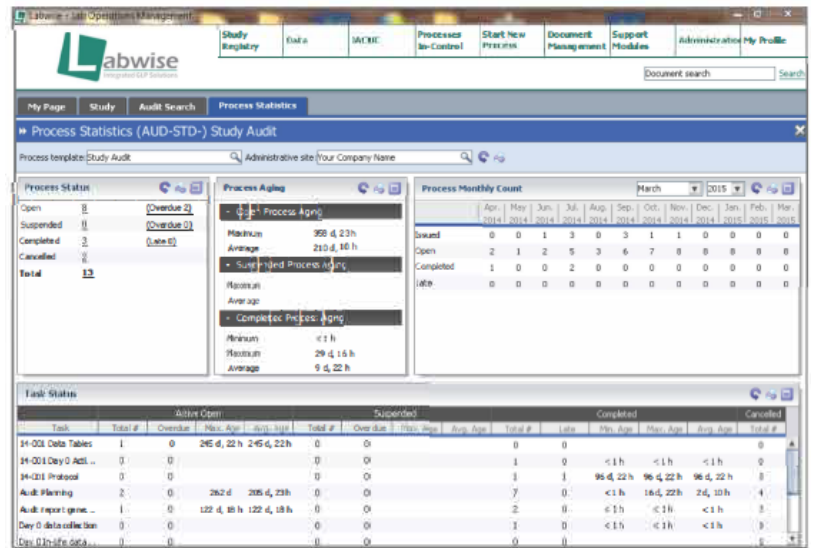
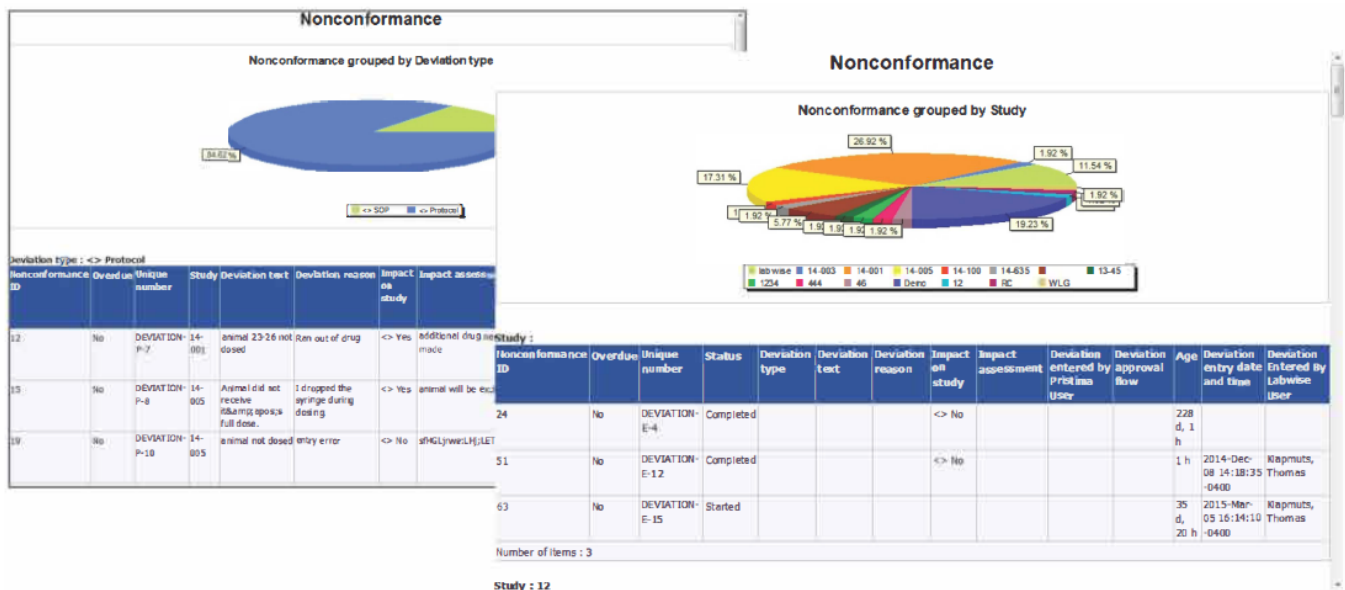


Figure 3. View Audit Process Statistics.



Figures 4-5. Example Nonconformance (Deviation) Reports grouped by Deviation Type or Study.

