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

Labwise™ 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 provides substantial gains in productivity while minimizing IT costs.

**Labwise™ Features**

- **Integration of Laboratory Environment.** Can be used as a stand-alone laboratory operations management system or integrated with Pristima, 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** The system provides out-of-the-box capabilities for full versioning, document approvals, PDF render-

![Labwise screenshot](image.png)

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

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**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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- 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 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.

Figures 2-5. Example Nonconformance (Deviation) Reports grouped by Deviation Type or Study.