



Labwise® XD is a powerful, flexible life science laboratory execution platform and quality management system. Integrated with the industry standard Pristima preclinical data suite, it allows organizations to manage study operations, business processes, controlled documents, non-conformance/deviations, and more, all within one seamless platform.

## Labwise® XD for Life Science Labs

Labwise® XD for Life Science Labs is a unique, comprehensive solution for Quality, Governance, Risk and Compliance designed for GLP laboratory environments. It can be used as a stand-alone laboratory compliance system or integrated with Pristima, the industry standard for total preclinical data management.

### Key Benefits

Labwise® XD for Life Sciences Labs maximizes the efficiency of the preclinical laboratory workflow by allowing you to:

- Link preclinical 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 Corrective Actions/Preventive Actions (CAPA). The system delivers workflow processes that enable proactive management of Corrective Actions I Preventive Actions in a timely manner.
- 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.
- Manage business processes, including task notifications, powerful queries, and reports to users.

### Why Labwise?

*Labwise® XD for Life Science Labs consolidates the functionality normally found in disparate systems, such as LIMS, Quality Management, Document Management, and BPM tools in one platform.*

*Converging on the Labwise platform lowers your total cost of ownership, reduces your IT infrastructure complexity, and streamlines cross-department collaboration.*

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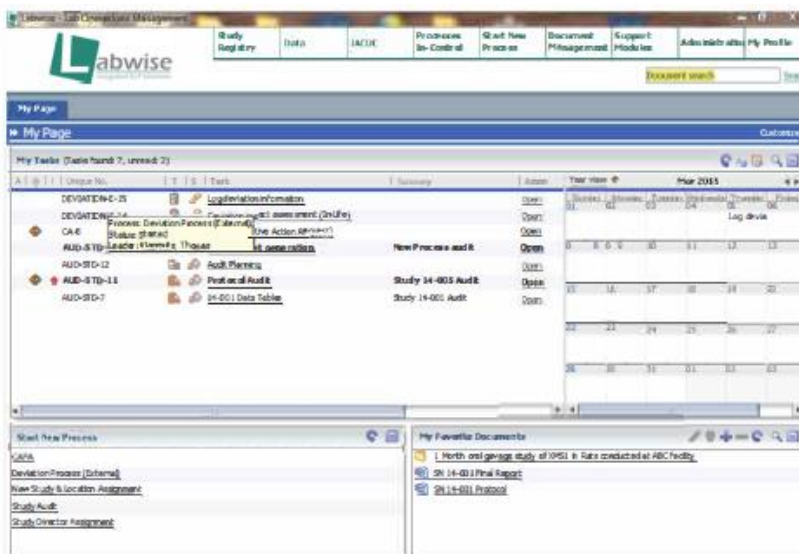


Figure 1. Labwise® XD for Life Science Labs home page provides easy access to key tasks, processes, and documents.



- **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 performance indicators.

### Xybio-Your Expert Partner

- Over 20 years of experience managing preclinical workflows and helping clients streamline compliance and mitigate risk.
- Consultants have extensive global experience and work with companies of all sizes from multiple industries.
- Offer deep domain knowledge of FDA regulations and long track record of helping clients build effective risk mitigation strategies

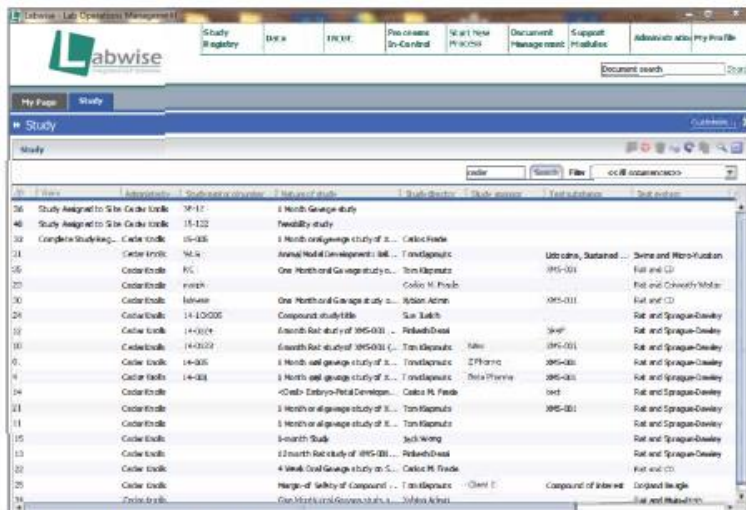


Figure 2. Manage the Study Registry.

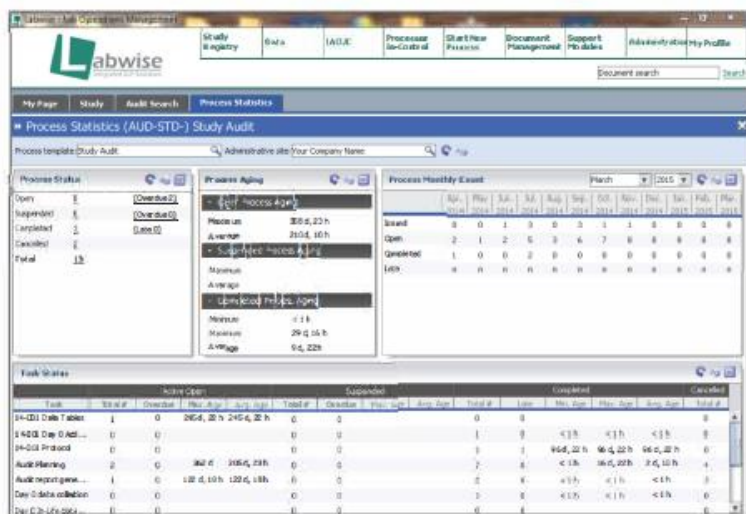
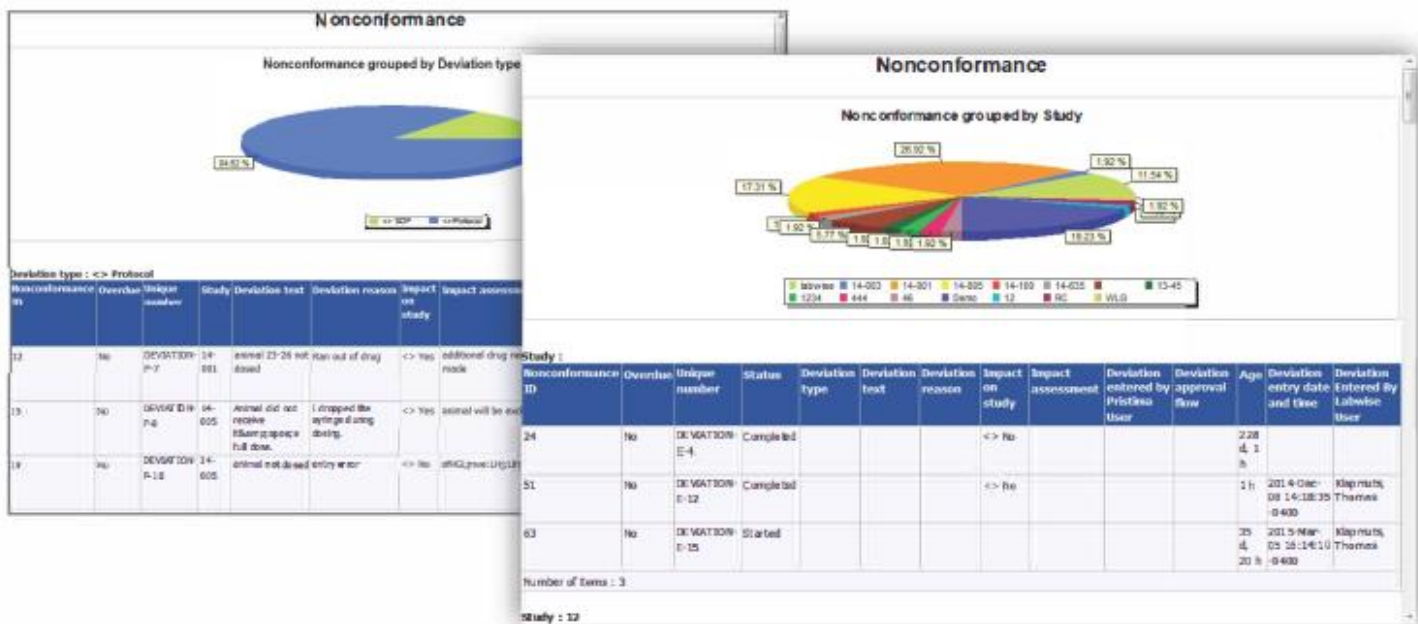


Figure 3. View Audit Process Statistics.



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

