
Clarmon introduces QValid v2.0, its new generation, web based validation and compliance management system

Clarmon Corporation announced today the release of QValid 2.0, the latest version of its leading validation and compliance management system.

As a real-life alternative to paper or manual systems, the QValid software suite is the optimal solution for Life Science companies focused on standardizing their validation and compliance documentation activities across multiple sites and reducing the cost of compliance. QValid ensures that all documents are maintained consistent and controlled and enables users to focus exclusively on critical tasks by automating recurring documentation and compliance processes.

Unlike traditional document management and compliance systems, QValid allows users to work directly in the familiar Microsoft Office® interface and thanks to its unique Linksense™ technology still ensures full traceability, drastically reducing the time needed to manage documents. The technology enables users to connect physically separate but related content such as requirements, risk assessments or tests across multiple documents, ensuring that the impact of any change is always visible. Furthermore, an up-to-date traceability matrix can be generated on demand for any set of validation or compliance documents in order to identify gaps or justify the extent of the documentation effort.

The enhancements in version 2.0 are strengthening QValid's position as the most efficient validation management system today and are extending the system's capabilities to automate all of the main quality management processes. QValid 2.0 ensures control and provides the optimal solution to demonstrate regulatory compliance without changing the user's way of working. Improved information reuse based on master templates, extended integration with 3rd party systems such as Crystal Reports and extended workflow support for quality system documentation processes increase the product's time and money saving potential even more.

About Clarmon Corporation

Clarmon Corporation provides advanced software solutions for quality and compliance management. The majority of Clarmon's customers are Life Science companies that operate inside highly regulated environments. Using Clarmon's Web and Microsoft Office® based solutions these companies have been able to replace paper and manual systems with a fully electronic, automated environment that reduces the efforts and costs of demonstrating compliance.

QValid, Clarmon's validation and compliance management suite, provides a simple method to centrally manage and control documents without requiring users to re-enter information. Unique technology embedded in QValid allows quality professionals to connect and synchronize documents and regulations, drastically reducing the time needed to update documents. Clarmon is a privately owned company established and sustained by people with extensive experience in Compliance Management, Quality Assurance and Validation. For more information about Clarmon Corporation please visit www.clarmon.com