



*I wish computer system validation
would be fully electronic...*

A fully electronic solution is available today!

Discover **QAvalid CSV™**

Clarmon
Compliance Management Solutions

automate your Computer System Validation (CSV) reduce the time and costs of updating documents

Automatically Generated Traceability Matrix

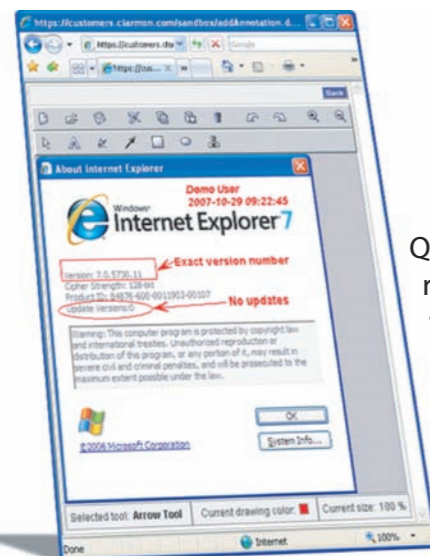
An up-to-date Traceability Matrix can be generated on-demand for any set of CSV requirements or verification documents, regardless of how many physically separate documents the information is distributed across.

Constantly up-to-date inventory and validation plan

QValid connects the inventory list with requirements, risk assessments or qualification documents. This ensures that the inventory is maintained up-to-date and that the compliance status of each equipment can be promptly justified.

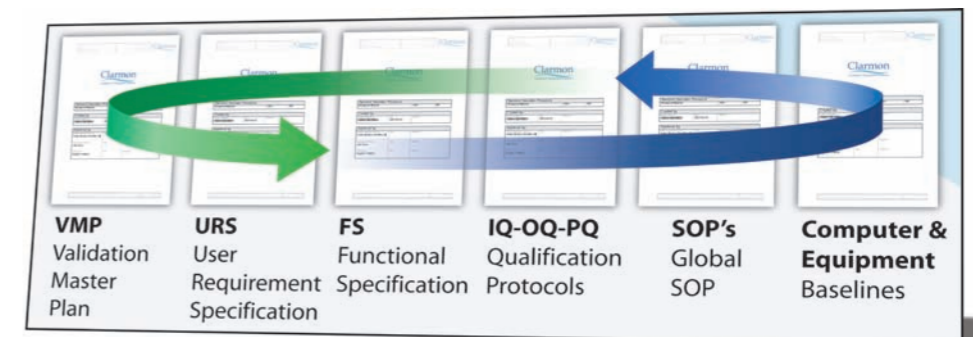
Requirements Management

QValid™ has built-in Requirements Management features to track requirements criticality and links to related Risk Assessments. In addition, QValid™ speeds up test script creation by generating and linking tests based on selected groups of requirements.



Quick and compliant test result evidence capturing with QValid™'s multi-layer annotation tool

QValid CSV™



Created with Microsoft Word, kept compliant with QValid™

Electronic Test Script Management and Execution

Test scripts are electronically managed within QValid™ with recording of test results, integrated capture or annotation of documented evidence, automatically generated deviation forms and automatic creation of phase reports.

Intelligent reuse based on templates and master documents

Information reuse is enhanced based on configurable GxP document and project templates as well as master documents. For example, users can select a set of requirements from a master document and have the possibility of automatically copying the related test scripts and traceability links.

Fully Integrated with Microsoft Word

Controlled document writing, editing, commenting and testing can be performed in Microsoft Word or using the QValid web interface. You can skip the "new software" training problems inherent with most new applications.



QValid™, Clarmon's quality and compliance management suite, provides a simple method to centrally manage and control documents without requiring users to reenter information.

Unique technology embedded in QValid™ allows quality professionals to connect and synchronize documents and regulations, drastically reducing the time needed to update documents.

QValid™ helps companies improve their efficiency in managing the following processes:

- Computerized System Validation
- Facility Commissioning and Qualification
- Corrective and Preventive Actions
- Equipment Inventory Management
- Change Management
- Audit Management

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 one of the most highly regulated environments.

www.clarmon.com