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## Clarmon introduces QValid™ v1.5, the first Compliance Management Application with offline traceability and compliant test execution in Microsoft® Office

Clarmon Corporation announced today the introduction of version 1.5 of QValid, its main software solution for creating and managing documented evidence within regulated industries. The new version includes a number of added functionalities, the most important being the electronic testing which can now be executed in Microsoft Word.

QValid saves more than 40% of the time spent documenting compliance by providing unique knowledge capture capabilities, automatically generated traceability matrices, automatically generated and linked documents or total control over the information inside each document or status of each equipment. The features needed to accelerate and automate documentation and to meet requirements for electronic signatures and electronic records are delivered through efficient collaboration, using a simple Web interface or a QValid toolbar in the Microsoft Office package.

The Microsoft Office integration enables QValid users to work in the same simple manner as they do today. The QValid toolbar makes the integration possible and ensures the appropriate level of control over the information. The new toolbar enables the use of Microsoft Word to write documents, review or approve them, make risk assessments or track system or equipment inventory as compliant electronic records. Unique Linksense™ traceability references are now available offline to connect content across physically separate documents directly in Microsoft Word.

The most important new benefit of the Office integration is controlled execution of test scripts or audits to verify compliance. The test results are captured in electronic format and submitted to the QValid repository with 21 CFR Part 11 compliant electronic signatures, access controls and audit trails. Performing test execution in Microsoft Word provides great flexibility, enabling users to carry portable computers and electronically execute tests anywhere they like without sacrificing regulatory compliance.

### About Clarmon Corporation

Clarmon Corporation provides integrated software applications for the management of structured content within a number of target industries. Most of Clarmon's clients to date have been in the Pharmaceutical, Biotechnology and Medical Device Industries.

Through its QValid™ application, Clarmon provides a powerful tool to allow companies to achieve regulatory compliance faster and easier. By removing tedious manual tasks for Quality Assurance, QValid™ drives costs down, and diminishes the risk of non-compliance.

Clarmon Corporation is headquartered in London, with a development and consultancy team in Bucharest Romania, sales staff in Dubai and Singapore and worldwide sales and distribution partners. For more info about Clarmon Corporation, please visit [www.clarmon.com](http://www.clarmon.com).