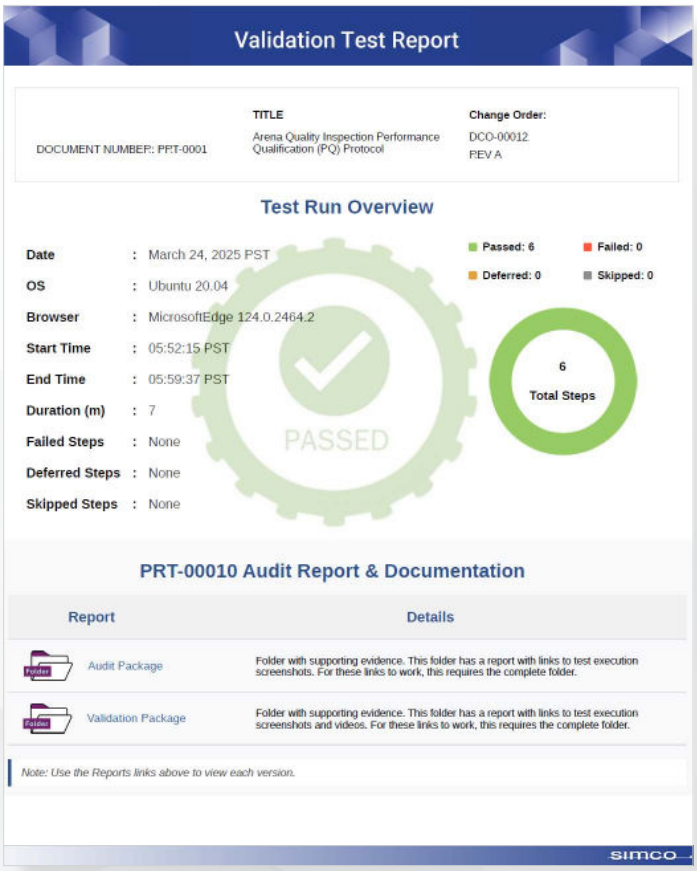
Automated Software Validation for Life Sciences

Ensure Compliance with Life Science Regulations Faster, More Efficient, and Fully Documented

Streamline and Accelerate Software Validation

Regulatory compliance requires rigorous validation of software used in critical operations such as Quality Management Systems (QMS), Laboratory Information Management Systems (LIMS), Manufacturing Execution Systems (MES), Enterprise Resource Planning (ERP), and more. Traditional validation methods are manual, time-consuming, and expensive. Our Automated Software Validation Service transforms this process - delivering faster, more reliable, validation documentation with minimal effort from your team.



How It Works:

1. **Share Your Validation Needs:** Kick off the automated testing process by submitting your validation test case document—enabling a thorough, efficient, and compliant evaluation of your software.
2. **SIMCO Automates the Test Cases:** Our system executes validation scripts tailored to your submitted test cases, ensuring full traceability and alignment with your compliance requirements.
3. **Generate Validation Test Reports:** Receive multiple reports automatically, including a concise auditor-friendly report with pass/fail results and digital signatures, as well as a comprehensive report detailing each test case with screenshots and video recordings.

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Key Benefits

Reduce Validation Time by 93%:
Automation removes manual effort and accelerates validation cycles.

Optimizing Workflow PQ:
Independent expertise to validate customized business workflow Performance Qualifications (PQ).

Complete Documentation:
Get fully structured, compliant validation documents for every test cycle.

Why Choose Our Automated Validation Service

Industry Expertise
SIMCO has decades of experience helping FDA-regulated life sciences companies meet complex validation and compliance requirements.

Supports Leading Software Solutions
Validates the critical systems used in life sciences—QMS, LIMS, MES, ERP, SharePoint, and more.

Rapid Validation for Software Updates
After the initial validation, automated re-testing makes every future update exponentially faster to approve.

Audit-Ready Documentation
Delivers clear, structured validation documentation with full traceability, screenshots, and video evidence that auditors prefer.

Validation Documentation Creation Services

In addition to automating test execution, SIMCO's validation experts can help you create essential software validation documents including User Requirements Specifications (URS), risk assessments, test cases, and traceability matrices—leveraging our regulatory expertise to deliver a complete, compliant, and audit-ready package.

Contact us today at SoftwareValidation@SIMCO.com to learn more.

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SIMCO is the leading provider of calibration and software services for technology organizations, bringing over 60 years of calibration industry leadership. Our experience enables us to develop exceptional solutions for service management.

Founded in 1962 to service NASA and high technology firms in Silicon Valley, SIMCO is committed to delivering life-saving quality leaner, by providing the highest level of quality and customer service. Learn more at **www.simco.com**