# Validated Electronic Record Approval

VERA is a versatile workflow engine and electronic signature solution that enables 21 CFR Part 11 compliance through integrations with many of today's leading DevOps tools. VERA's end-to-end approval system helps life sciences companies achieve regulatory compliance in their GxP systems. VERA can be deployed in your own environment or in the Tx3 VERA SaaS-qualified Cloud Platform.

## Leading Quality and Compliance Management Platform

- Standardize workflow and best practices
- Test automation enables continuous testing
- Enables use of Agile and DevOps tools in regulated environments
- Data-driven (not document-dependent) computer system validation

## Complete End-to-End Approval System

- Common UI for records approval across leading DevOps tools
- Approve or reject records on the fly, electronically
- Configurable approval routes
- End-to-end traceability

## 21 CFR Part 11 Compliant

- Maintain record versions and audit history
- Eliminates signing and managing paper documents
- Versatile system of record
- Verifies validity of electronic records and signatures

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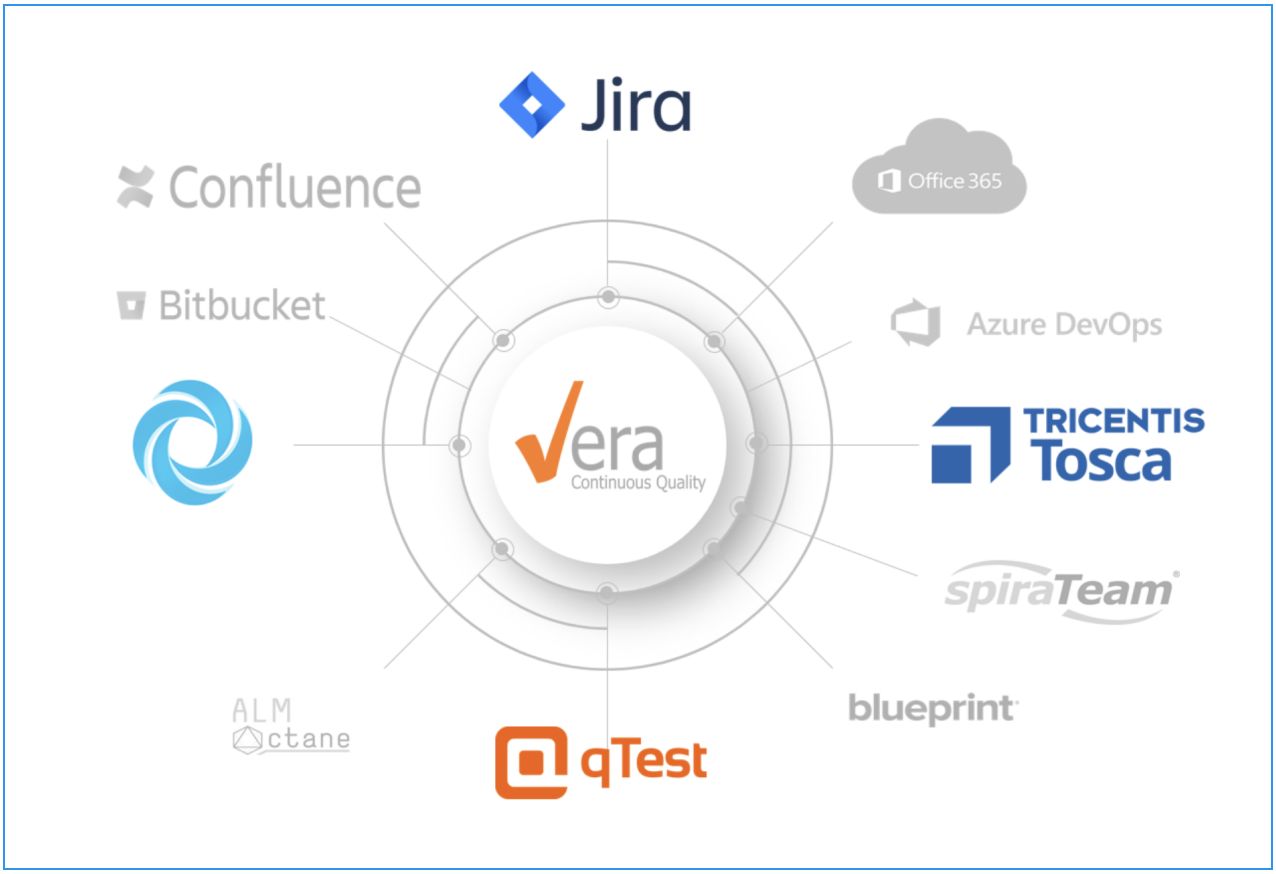
## Integrate VERA with Best-of-Breed DevOps Tools

The strength and adaptability of VERA is demonstrated by its ability to effortlessly integrate with many of the leading DevOps tools on the market today. These tools include MicroFocus ALM, Atlassian Jira and Tricentis qTest, among others. The benefits of these integrations include faster fixes, greater collaborative working and operational support, improved processes and automation across the IT department and teams, as well as increased flexibility and agility.




VERA is designed to help your organization move through the FDA approval process faster, more efficiently and with less expenditure.
Discover the advantages of VERA.
Ask for a demo today.

Tx3 Services LLC, 980 Harvest Drive, Suite 205, Blue Bell, PA 19422
www.tx3services.com