PERFORMANCE VALIDATION | Moving Quality Forward

THREE DIGITAL VALIDATION OPTIONS. **ONE SUCCESSFUL OUTCOME.**

The life sciences industry is under increasing pressure to go digital — driven by FDA requirements and ISPE's Pharma 4.0 vision. But making the transition is not without challenges. Cost, time, operational disruption, managing change, and the need for validation expertise often stand in the way. That's why PV provides 3 pathways to help you jump to the front of digital transformation.

DIGITAL VALIDATION AS A SERVICE (dVaaS)

- PV provides a 3rd generation Digital Validation Tool (DVT) as part of its CQV services.
- Get started with minimal investment and effort yet migrate to your own DVT at any time.

INVEST IN YOUR OWN DVT

PV helps you define requirements, supports vendor selection, and provides implementation services.

SYSTEM SUPPORT

Already invested in a DVT?
PV generates and executes validation documents in the three industry-leading DVT platforms.

PERFORMANCE VALIDATION

GO DIGITAL TODAY, NOT TOMORROW!

BECOME A LEADER IN DIGITAL VALIDATION

Digital transformation is now within reach for all FDA-regulated environments — regardless of company size or cGMP experience. PV's three solutions are built on Performance Validation's in-depth experience with regulatory frameworks and a 37+ year track record of delivering audit-ready CQV documentation.

DIGITAL VALIDATION AS A SERVICE (dVaaS)

Ideal for companies new to U.S. cGMP regulatory frameworks, including late-stage clinical start-ups, smaller CDMOs seeking greater impact, overseas manufacturers building their first U.S. facility, new suppliers to the medical device industry, or 3PL logistics facilities.

PV provides a 3rd generation DVT as part of our comprehensive CQV services, with no software licensing, hosting, or DVT validation costs. Our experts configure workflows aligned with your QMS, VMP, and SOPs, while you focus only on review and approval.

When you are ready to scale, your digital validation footprint can transition seamlessly to your own licensed system. dVaaS is a great, low-involvement way of jumping to the forefront of the digital revolution.

INVEST IN YOUR OWN DVT

Ideal for companies where digital transformation is an urgent priority and who have the software licensing budget and time to transform an existing paper-based system.

PV provides the depth to make that transformation a rapid reality. With expertise across the top three industry-leading DVT platforms, we help author URS requirements, guide system selection, and evaluate each platform's attributes to your needs.

Our team can lead end-to-end implementation, from system setup and user management to SOP development, template configuration, training, and ongoing support.

SYSTEM SUPPORT

Ideal for industry leaders who have already made the move to a digital validation platform.

Every day, the majority of our validation engineers use a DVT from one of the three leading vendors. We digitize paper documents, support document review and editing, and execute test scripts with accuracy and quality.

PV brings digital validation expertise but also decades of experience in commissioning, qualification, and validation of equipment, systems, automation, utilities, and facilities across the full pharmaceutical and medical device workflow.

COMPREHENSIVE DIGITALLY ENABLED SERVICES

DIGITAL VALIDATION TOOL

- Our preferred partner is Valkit
- 3rd generation, true SaaS solution
- Enterprise-grade security
- Easy to validate, easy to configure, easy to support
- Rapid document cloning
- AI augmentation options for even faster content generation

RELATED CQV SERVICES

- Consulting on best-in-class digital transformation practices
- Audit preparation and remediation
- Program and project management
- Paper to digital document transformation
- Digital document authoring and test execution
- Change control
- Library of Golden Validation Packages
- User and admin training on DVT

LET'S TALK ABOUT HOW PV CAN HELP YOU GO DIGITAL

with efficiency, confidence, and compliance built in.

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