

# GxP validation for leaders who see further.

See further. Move faster. Lead with confidence.

## Challenge

The life sciences industry is transforming at unprecedented speed, driven by AI breakthroughs, agentic frameworks, and emerging technologies. But validation hasn't kept pace:

- ✘ Emerging AI-driven and complex GxP systems demand a digital transformation of validation, but most **organizations lack the tools** to support them
- ✘ **Resource constraints** force teams to choose between speed and audit readiness
- ✘ Your experts **waste 30% of project costs** on manual validation instead of innovation
- ✘ Intelligent equipment is accelerating the shift to digital twins, turning validation into an **integrated software problem**

## Solution

Res\_Q delivers intelligent GxP validation that transforms compliance into competitive advantage. Our rigorous, workflow automation platform validates ANY system—from traditional enterprise to cutting-edge AI.



### With Res\_Q, you gain:



#### Centralize GxP validation control

Validate **software, infrastructure, AI/ML, and GxP processes** within a **single platform**. Establish consistent **oversight, risk-based execution, and end-to-end traceability** across validation activities—without relying on disconnected tools or manual handoffs.



#### Execute faster with CSA-aligned automation

Reduce validation timelines through **intelligent automation and CSA-ready workflows**. Streamline execution, approvals, and documentation to support **efficient, audit-ready validation** across global teams.



#### Validate AI and emerging technologies with traceability

Confidently validate non-deterministic **AI systems, agentic frameworks**, and emerging technologies with complete traceability. Support expectations from **FDA, EMA, GAMP 5, and Annex 11** while remaining adaptable as technologies evolve.



#### Deploy quickly and confidently

Implement Res\_Q in as little as **4 weeks** using a **proven approach** designed to align with **established validation practices and operating models**—supporting rapid adoption without altering how teams work today.

# Res\_Q is redefining GxP validation for the modern era.

**Turn validation into a catalyst for efficiency.  
Swap paperwork for a powerhouse of visibility and control.**

“*Our partnership with Sware offers Salesforce Life Sciences Cloud users a compelling way to embed robust, efficient, and scalable GxP validation directly into their cloud environment. [And] Sware doesn't just provide a platform; their leadership team has spent decades solving complex life sciences compliance challenges and building solutions with industry-wide impact.*”

—Frank D, SVP and General Manager, Salesforce

## Impact

- ✓ **Up to 80% faster validation cycles** with intelligent automation
- ✓ **60% reduction in total validation costs** across project lifecycle
- ✓ **4-week implementation** with turnkey deployment
- ✓ **100% audit readiness** with complete traceability and transparency

## Why Res\_Q

-  **AI-native by design, with custom agents, instant asset intelligence, video-to-script practice, and more**
-  **A single intelligent point of control for real-time insight and confident validation decisions**
-  **Turnkey connectivity with the ability to integrate with the apps and systems you care about**
-  **Built for what's next, combining regulatory trust with operational excellence at scale**

Box 2025 Life Sciences  
Partner of the Year

Salesforce Strategic Partner

SOC 2 Type II Certified  
(zero exceptions)

ISO 9001:2015



**Ready to build the future?**

Schedule a consultation with a GxP expert.