

Supercharge Your MedTech Client Portfolio

QuickVault offers a tailored solution for service providers working with MedTech companies of all sizes.

With QuickVault Hub, you can collaborate with clients on a single platform that unifies product design, regulatory submissions, quality, and operational workflows in a scalable environment.

Why Join the QuickVault Partner Program?

By partnering with QuickVault, you can:

- Enhance operational efficiency with structured, automated workflows
- Serve more clients without adding headcount
- Differentiate your services by aligning with Veeva, the trusted leader in life sciences

Proven Business Impact

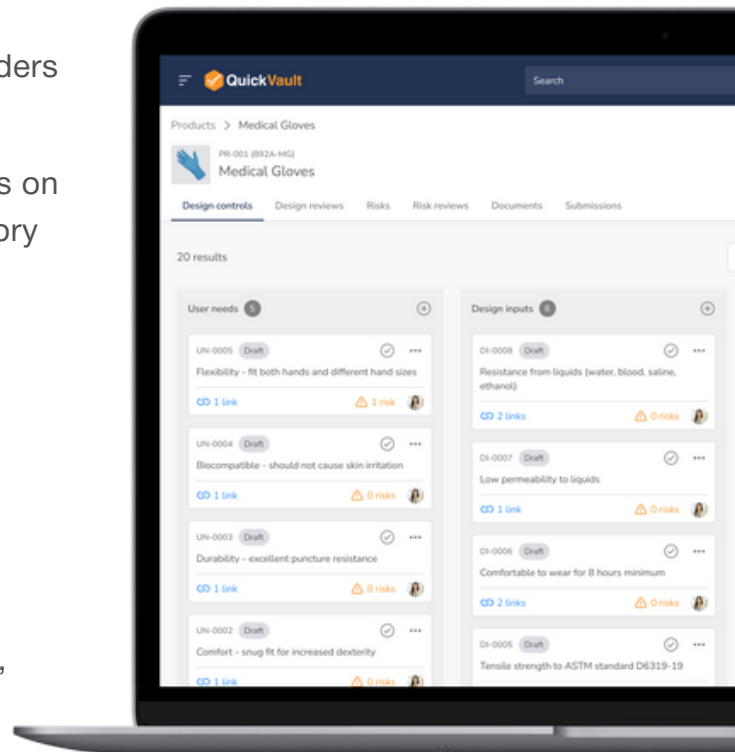
Partners using QuickVault Hub are seeing measurable results — including reducing document preparation time by up to 50%, cutting submission readiness from weeks to hours, and delivering faster audit confidence across their client base.

These outcomes directly translate into:

- More billable capacity
- Faster client project execution
- Higher client satisfaction and loyalty

The Opportunity

QuickVault Hub transforms how you work with MedTech clients, giving you the tools to elevate your business, deliver greater client value, and stay ahead in a market that is rapidly moving toward Veeva solutions.



Scaling Compliance with Speed

How CS Lifesciences Cut Prep Time in Half and Gained Instant Audit Confidence with QuickVault Hub



CS Lifesciences, a leading consultancy serving MedTech companies in the UK, Ireland, and the United States, needed a better way to manage client documentation and regulatory submissions. Disconnected tools, fragmented file shares, and manual submission prep were slowing projects and creating compliance risk.

KEY METRICS

CS Lifesciences' transformation with QuickVault Hub wasn't just qualitative, it was measurable. The team unlocked significant time savings, improved readiness, and eliminated the burden of manual tasks.



100%

Centralized, version-controlled documentation



DAYS TO HOURS

Submission readiness



50%

Reduction in document prep time



A powerful solution to help your clients navigate compliance, achieve approvals, and scale efficiently.

BECOME A QUICKVAULT PARTNER

CHALLENGES

CS Lifesciences and their MedTech clients were slowed by fragmented systems of emails, file shares, and single-use tools. Manual and error-prone preparation for FDA and ISO 13485 submissions consumed valuable time. Rogue document copies and inconsistent version control created ongoing audit risks.

SOLUTIONS

QuickVault Hub gave CS Lifesciences a centralized, audit-ready platform for managing client documentation and regulatory submissions. QuickVault's secure workspaces with version control and audit trails made collaboration seamless, while its Regulatory Submission functionality auto-organizes dossiers, enables one-click, pre-validated exports, and cuts turnaround time in half.

“

QuickVault Hub pays for itself the moment a client or auditor asks for a document and you can produce the latest, fully-tracked version in seconds.

”

BENEFITS

1

Faster Document Preparation

50% reduction in prep and review cycles for audits and submissions.

2

Accelerated Submissions

Submission-ready packages can now be assembled in hours instead of days.

3

Always-Ready Compliance

Eliminates “document hunting” and provides immediate audit confidence for clients and regulators.