

Quality Documents Management

Create, manage, and distribute
Quality documentation

Quality Documents Management on the CARA Life Sciences Platform™ provides a modern, central DMS for all types of Quality documentation.

Beyond the standard functionality of content creation, review, and approval, CARA supports publishing, training, eSignature, controlled printing, and site-control to support enterprise needs.

Bring Quality documentation to your employees, with QR codes for laboratories and manufacturing, portals for rapid searching and viewing across the business, and the CARA Mobile App for on-the-go work.

Benefits

Designed to be Used

Different experiences for consumers and authors ensure that the system is intuitive and functional for all users.

Compliance & GxP

Visibility and automations across your processes keeps your content up to date, and your business compliant.

Integrate Quality into your Business

Quality Documents Management on the CARA Life Sciences Platform connects seamlessly not only with QMS, LMS, and Audits Management, but across Regulatory, Safety, and Clinical, creating a digital universe of quality embedded in everything you do.

Audit Readiness

Create certainty in audits with clear and flexible views of your audit trails, a network of connected quality information, and instant search.

Generate Content Automatically

From simple template-based authoring to PDF concatenation and full document generation, CARA leads the industry in automatic content creation.

Contact us for a demo or evaluation
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www.caralifesciences.generiscorp.com



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