

ENNNOV QUALITY SUITE

Ennov Quality Documents

Enterprise Quality Document Management Software

Manage and control GxP documents effectively and efficiently with **Ennov Quality Documents**. Streamline your operations, ensure compliance and increase overall efficiency in all quality processes.

The Quality Document Management Challenge

Managing controlled documents across sites, teams, and systems can quickly become difficult in regulated Life Sciences environments. When quality documents are stored across file shares, local drives, or disconnected systems, teams lose time, visibility, and confidence. That makes it harder to maintain version control, enforce document workflows, and support compliance with GxP and ISO requirements.

Structured Quality Document Management

Ennov Quality Documents helps Life Sciences organizations manage GxP documentation in a more structured, consistent way. The document inventory is preconfigured in alignment with the DIA GMP Reference Model, including document categories, groups, subgroups, and artifacts needed to support controlled quality documentation.

Available as either a preconfigured solution or a fully configurable system, **Ennov Quality Documents** gives organizations the flexibility to adopt proven best practices or tailor document management to their own structure and processes. Its metadata-based document model helps teams manage a wide range of GxP document types, including governance, procedures, manufacturing, quality, audit, validation, and packaging documents, while supporting 21 CFR Part 11 requirements such as electronic signatures, audit trails, and records management.

> CORE CAPABILITIES

- Pre-configured document inventory
- Advanced life cycle management
- Flexible rights management
- Automatic PDF rendering & tag management
- Full text and metadata-based search
- Controlled printing, copying and pasting
- Periodic review, expiration & archive management
- Office 365 and Google Drive integration

> KEY FEATURES

- Integrated worklist dashboard
- Configurable document types, workflows & views
- Automated email notifications
- Intuitive user interface
- Integrated PDF viewer
- Composite document support
- 21 CFR Part 11 compliant
- 100% web-based



Cloud Based or On Premises



Multi-Platform



ISO 9001 and 27001 Certified

Unified Document Management Across Compliance Functions

As part of Ennov's unified compliance platform, **Ennov Quality Documents** works within a common environment for documents, quality processes, data, training, and reporting. With a harmonized data model and shared user experience across applications, organizations can reduce redundancy, improve visibility, and manage controlled content more consistently across quality, regulatory, and operational teams.



Quality Documents - Part of the Ennov Quality Suite

QUALITY DOCUMENTS

QMS

TRAINING

Why Choose Ennov?

HUNDREDS OF COMPANIES TRUST ENNOV	PROVIDING YOU FREEDOM OF CHOICE
<p>Over 25 years' experience, and 450+ Life Science customers, with many more in other industries.</p>	<p>Available as cloud-based or on-premises deployment: You can switch between deployment options at any time.</p>
<p>Modern architecture and user interface: 100% web-based. Highly scalable. User-centric design.</p>	<p>No IT skills required for system configuration and management.</p>
<p>Our commitment to your success: Very high customer satisfaction. 98.5% of projects delivered on time and within budget.</p>	<p>Improved security and optimized performance: Data is hosted locally for total flexibility. Single tenancy minimizes business interruptions.</p>

Learn more about our **Unified Compliance Platform** to support the entire Life Sciences product development continuum at www.ennov.com

QUALITY	CLINICAL	REGULATORY	PHARMACOVIGILANCE	COMMERCIAL
<p>Our comprehensive QMS improves operational efficiency and ensures regulatory compliance</p>	<p>Our total solution for capturing and managing Clinical Trial information streamlines clinical operations</p>	<p>Our world-class Regulatory content and information management software accelerates HA approvals</p>	<p>Our end-to-end solution for collecting, reporting and analyzing human and vet PV data minimizes risk</p>	<p>Our complete management of professional events ensures DMOS, EFPIA, HCP and COI compliance</p>