

The ever-changing regulatory landscape makes it challenging to create an environment that fosters compliance and efficiency. Professionals must create and share documentation, but are faced with time-consuming and costly workflows as they pull information from multiple sources and locations.

Patients are waiting, and 88% of medical writing professionals believe they could complete submissions faster by using technology to speed up the document review process.*

ENHANCE DOCUMENT WORKFLOW & PRODUCTION

Content Companion™ delivers the most efficient path to compliance. Document creation professionals can create and store approved copy and information in a centralized library where content can be easily added, updated and distributed throughout the organization to ensure consistency.

MAINTAIN COMPLIANCE

Build new or update existing documentation by easily accessing approved content to maintain consistency and keep regulators happy.

INCREASE OPERATIONAL EFFICIENCY

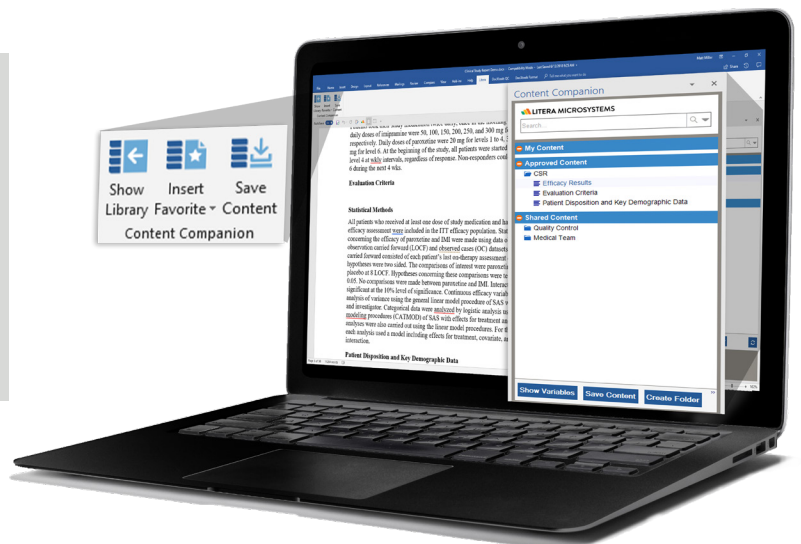
Optimize the workflow by reducing the time it takes to find and pull content from multiple locations and enable the team to focus their attention on higher value activities.

SIMPLIFY CONTENT MANAGEMENT

Decrease hassle among anyone who leverages relevant regulatory information by storing, retrieving, and sharing standardized content from one central location.

“I needed a better way to share and manage my standardized regulatory document content. Content Companion does everything I need it to—stores my content in an approved location and makes it easy to access through Microsoft Word tools.”

**Director, Regulatory Operations
Global Pharmaceutical Company**



Contact Litera Microsystems for a demo today at litera.com/request-a-demo

*Microsystems Market Segmentation Survey, 2017

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