



HOW YOU BENEFIT

- Enhanced internal delivery capability for eCTD
- Improved organizational compliance
- Improved submission quality with data standardization
- Reduced compliance risk

WHAT SMARTDESK DOES

- Verify and correct legacy submission content
- Author content to meet eCTD standards
- Create define.pdf and define.xml content
- Create consistent navigational tools
- Meet and maintain CDISC standards

GET YOUR PUBLISHING OFF TO THE RIGHT START **LIQUENT *SmartDesk*[®]**

A COMPREHENSIVE SUITE OF PRODUCTS FOR MANAGING SUBMISSION CONTENT

LIQUENT[®] *SmartDesk*[™] is a suite of comprehensive desktop tools designed to enable users to complete their tasks at hand, in an easy, fast and efficient manner. Liquent *SmartDesk* tools aid in standardizing data for common technical documents, automating of publishing for PDF creation and in the preparation of Case Report Tabulations (CRT).

Each of the three components of LIQUENT[®] *SmartDesk*[™] can work independently, but are much more beneficial when integrated and deployed across your enterprise. LIQUENT[®] *SmartDesk*[™] consists of :

- LIQUENT[®] *SmartDesk*[™] for PDF
- LIQUENT[®] *SmartDesk*[™] for Authoring
- LIQUENT[®] *SmartDesk*[™] for CRT

Used together or separately LIQUENT[®] *SmartDesk*[™] products provide tremendous value to Life Sciences organizations and their employees.

LIQUENT[®] *SmartDesk*[™] Family





LIQUENT® *SmartDesk™ for PDF*

KEY BENEFITS

- ➔ Automated functions for consistency and error reduction
- ➔ Batch processing for increased efficiency
- ➔ Compliant with agency requirements and industry standards
- ➔ Intuitive and easy-to-use interface for immediate ROI

LIQUENT® *SmartDesk™ for PDF* automates the verification of submission content for compliance, enables a process for bookmarking & hyper-text linking and ensures that PDF documents are agency compliant and ready for submission within a tight timeframe.

LIQUENT® *SmartDesk™ for PDF* allows users to focus on building and viewing the electronic submission structure, rather than spending valuable time manually fixing broken links and missing bookmarks. If your legacy data isn't perfect, Liquent SmartDesk for PDF works directly in the PDF, removing the need for original Word documents.

LIQUENT® *SmartDesk™ for PDF* eliminates many of the manual processes associated with standard Adobe® offerings while providing the flexibility and intelligence to manage the entire publishing process.

LIQUENT® *SmartDesk™ for Authoring*

KEY BENEFITS

- ➔ Improved submission quality by standardization of Word content
- ➔ Decreased time when training authors and creating files
- ➔ Reduced time required to meet submission standards

LIQUENT® *SmartDesk™ for Authoring* delivers the perfect starting point to write structured, compliant submission content.

With LIQUENT® *SmartDesk™ for Authoring*, medical writers and regulatory affairs professionals can immediately start correctly authoring by using our standardized templates designed with formats based on the Common Technical Document Module 1 through Module 5, Investigational New Drug, and the Clinical Trial Application.

Designed by experienced regulatory professionals, LIQUENT® *SmartDesk™ for Authoring* reduces the time manual formatting requires, allowing end-users to create consistent and compliant documents without the need for extensive in-house training.

LIQUENT® *SmartDesk™ for CRT*

KEY BENEFITS

- ➔ Improve quality of CRT section by automation
- ➔ Reduce re-entry and rebuilding of metadata
- ➔ Meet and maintain CDISC standards

LIQUENT® *SmartDesk™ for CRT* is your complete solution for Electronic Clinical Data Submission to the FDA.

LIQUENT® *SmartDesk™ for CRT* automatically creates the entire CRT section of an electronic submission along with the define.pdf and/or define.xml files, SAS transports files and table of content files, including all of the necessary links and bookmarks. As a Certified CDISC Solutions Provider, LIQUENT designed LIQUENT *SmartDesk™ for CRT* according to the Case Report Tabulation Data Definition Specification (CRT-DDS) developed by CDISC define.xml team.

LIQUENT® *SmartDesk™ for CRT* helps ensure our clients are compliant with the FDA regulatory requirements that clearly stipulate that all study data must be delivered in an electronic format - even if the marketing application is in paper format.