# eReg Overview

## Streamline Your Regulatory Process

### Think beyond the binder

Managing essential clinical trial documents at academic medical centers, cancer centers and health systems is often an exhaustive, extremely manual process that requires both physical storage space and site resources. The Advarra eRegulatory Management System (Advarra eReg) provides integrated protocol, staff and institution documentation to streamline regulatory process and enhance compliance.

Advarra eReg takes a templated approach to creating and maintaining electronic binders. This is ideal for academic institutions managing a large number of protocols. Moving key regulatory tasks to the system level—instead of the protocol level—eliminates redundant workflows, boosts compliance and saves valuable staff time. Combined with 21 CFR Part 11-compliant electronic signatures for protocol documents, delegation of authority, and more, the system significantly reduces the regulatory burden for your staff.

## Improve compliance, maximize efficiency, and ensure return on your investment

Store all your essential protocol documents, track owners and expiration dates and more, all in a 21 CFR Part 11 compliant system

Leverage multi-site management, allowing a coordinating center to manage essential documents for participating sites

Efficiently route documents and manage electronic signature notifications

Integrate Advarra eReg with your OnCore Enterprise Research System

Allow regulatory leadership to standardize delegated tasks at the system level, dramatically reducing the amount of time spent routing documents

Utilize Advarra's data migration services to get up to speed quickly within the eReg system, and validation services to achieve and maintain 21 CFR Part 11 compliance

Easily and securely allow access to sponsors and monitors

Build new protocols using easy, intuitive templates

NEW

## Save staff time and boost compliance with technology integrations

Advarra eReg offers interfaces to connect with technology across your enterprise. Turnkey integration with Advarra's Center for IRB Intelligence (CIRBI) Platform and your local eIRB system allows regulatory staff to automatically upload documents into eReg. This unprecedented integration allows your institution to avoid the burden of manually downloading and uploading files and reduces the possibility of errors during the document upload process. Also, eReg provides email integration, allowing your staff to file correspondence with your IRB or sponsor/CRO directly into the eReg system, and send other documents straight to your system for filing.

To learn more, contact our Customer Relationships Team at CustomerRelationships@advarra.com.

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