How to Integrate Your Clinical Trial Document Ecosystem

Protocols. Amendments. Investigator brochures. CVs. Informed-consent forms. Your site manages thousands of documents for each trial you run. Now there's an easy and efficient way to make sure you have the right documents—and the right versions—where you need them.

Complion integrates with WCG solutions to automate clinical trial document flow.

Benefits of integration include:

Faster study startup and easier management

- Upload documents once rather than to multiple individual systems
- Shared documents in WCG IRB Connexus™ flow to Complion eRegulatory/eISF
- Shared documents in Complion eRegulatory/eISF flow to Velos CTMS
- Flexible Complion platform enables you to choose the documents your site shares across systems

Less potential for error

- Versions update across all systems automatically—so users always have the correct version
- Ensure that clinicians always access the correct protocol documents when seeing patients for greater patient safety
- Eliminate confusion that can lead to frustrating patient call-backs
- Automated system performs regardless of staff tenure or staffing levels

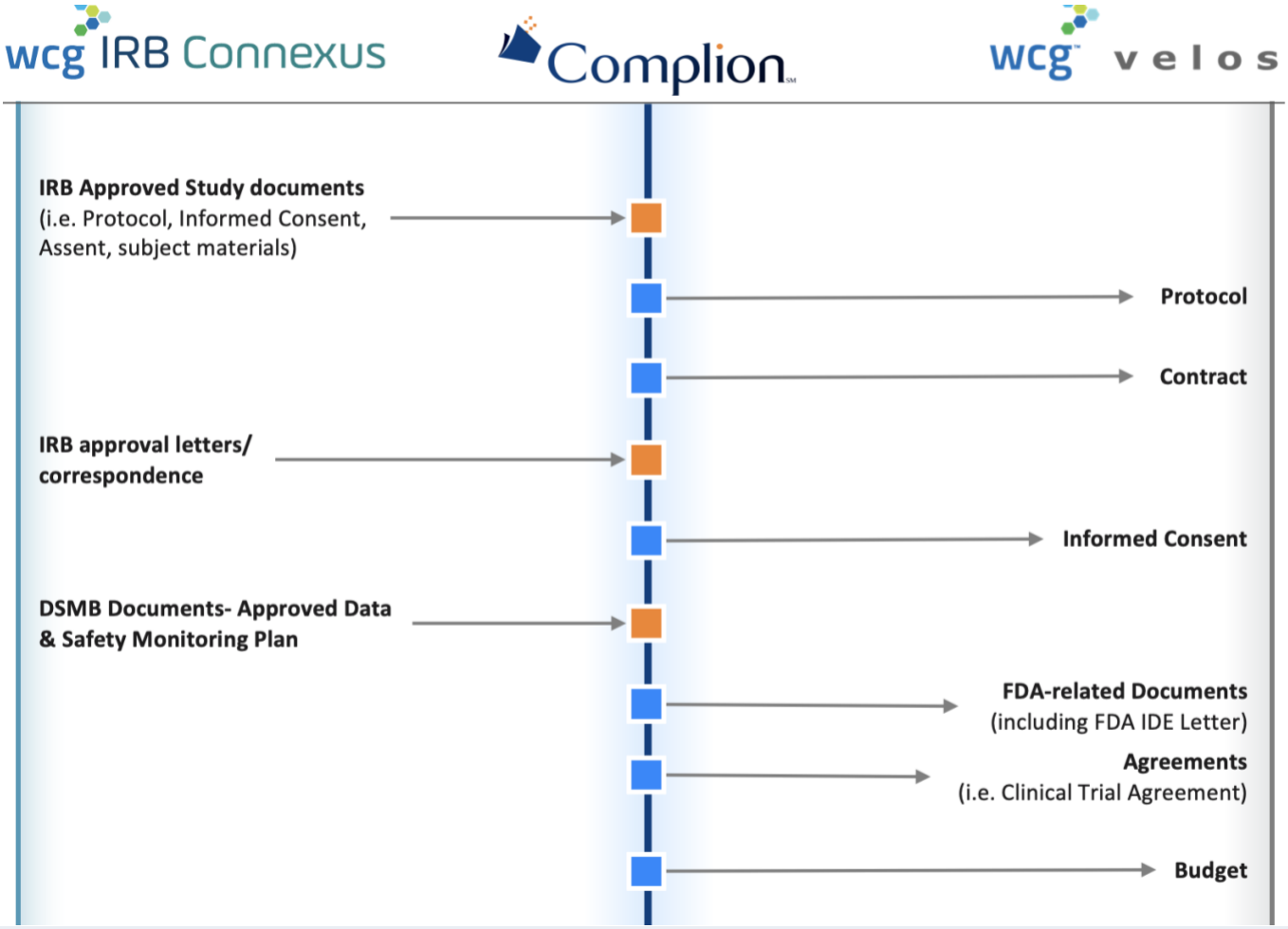
Readiness for audits and FDA submissions

- Always know the status and location of key documents
- Quickly identify documents that are missing or require signatures
- Ensure that auditors and reviewers always see the latest versions

Managing trial documents in a single place increases efficiency for both regulatory and clinical teams. WCG and Complion have partnered to help sites streamline document management.

The right version. The right place. In real time.

A typical study shares dozens of documents across multiple systems. With Complion integration, you decide which documents you want to share. The system does the rest.



A powerful partnership for your site's success

- **The WIRB-Copernicus Group (WCG™) is the world's leading provider** of solutions that measurably improve the quality and efficiency of clinical research. Those solutions include:
- **MyXConnexus** enables WIRB-Copernicus Group IRB customers to securely submit and track research at any time throughout the course of a study. The portal offers guidelines that outline the submission process for your protocol with step-by-step instructions for submitting and tracking your review.
- **Velos eResearch** clinical trial management system platform simplifies the management of institution research by linking study status, patient enrollment, calendars, budgets, billing and electronic data capture (EDC) in a cloud-based system.

Complion is a leading provider of eRegulatory and eISF solutions for Sites, Sponsors and CROs. Founded by a clinical researcher participating in an NIH-funded MD/PhD training program, Complion's unique cloud-based purposebuilt software enables collaboration of site regulatory workflows and the management of essential documents to ensure clinical trials are conducted in an audit ready manner.

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