

INTRALINKS**STUDY SPACE**

Intralinks Studyspace™

For Study Startup

Expedite

Instant Access to Study Start-Up (SSU) Documents

Facilitate Communication among Stakeholders:

- Sponsors
- Sites
- CROs
- Ethics Committees

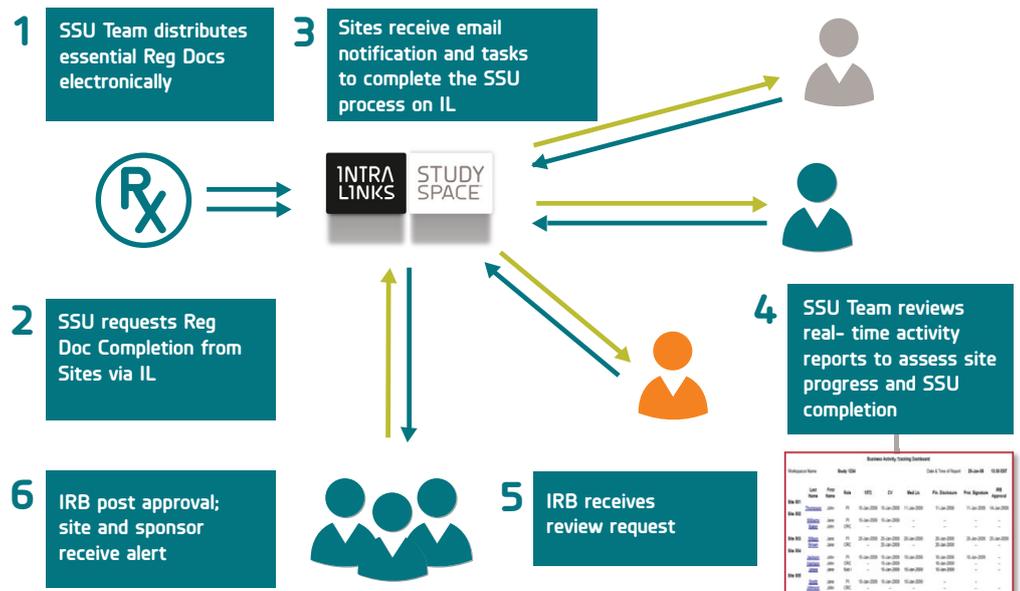
Decrease Costs

- Eliminate overnight mail shipping costs
- Reduce number of man-hours associated with creating SSU packets

Start Studies Faster

Intralinks Studyspace™ accelerates study start up time by automating the exchange and tracking of all necessary documents related to the site activation process. Documents posted to the Intralinks Platform are automatically and securely routed to the appropriate parties, and alert notifications prompt site personnel, institutional review boards (IRBs) and ethics committees (ECs) to review and respond to any requests. Real-time activity reports ensure study start up teams understand what needs to be accomplished across investigator sites and geographies.

Site Activation: The New Way





Who we work with:

THE TOP 20

pharmaceutical companies

THE TOP 10

biotech companies

THE TOP 5

CROs

17

IRBs

60,000

investigative sites around the globe

OVER 200

biopharmaceutical companies

Benefits

- Reduces average study start-up (SSU) cycle time by 25% - 50%, depending on study size and IRBs (saving 15 days = \$450,000)
- Eliminates the need for overnight mail, reducing costs by \$36,000 per study
- Reduces man hours associated with manual site management processes by 80%
- Saves \$35,000 with every day shaved off of the clinical trial process
- Enables 21 CFR 11 Compliance

Intralinks Studyspace removes the manual tracking and follow-up of study start up packages prevalent in the industry today and provides deeper insight into the progress of sites and IRBs/ECs. The rich metadata captured by the Intralinks Platform provides a complete audit trail and the fully validated system enables 21 CFR Part 11 compliance.