INTRA LINKS STUDY SPACE

Intralinks Solutions for Life Sciences

33%
Reduced cycle time from final protocol to first subject first visit*

TOP 10
Pharmaceutical companies are Intralinks clients

TOP 10 Global CROs have conducted business on Intralinks

80,000+
Investigator sites are registered users of Intralins

*Source: Based on Intralinks exchange metrics for a top-five pharma client.

Accelerate Your Clinical Trials

Across the industry, companies are striving to find ways to improve operational efficiency. Intralinks® enables you to better manage the distribution, collection and tracking of critical information related to site recruitment, study start-up (SSU) and safety while also adding a layer of security to the process. Intralinks' Clinical Trial Portal improves collaboration and document exchange between everyone involved in a clinical trial — to accelerate the whole process.

Clinical Trials

Simplify Site Recruitment

Significantly shorten the site recruitment process by managing the distribution and collection of the feasibility survey online in a secure, centralized location. Track the status of site survey completion on Intralinks so you can focus on follow-up and get sites on-boarded quickly.

Shorten Study Start-Up Time

Distribute and collect regulatory documents such as 1572s, investigator brochures and study protocols online to shorten SSU. Manage and track site status by keeping tabs on the completion of key regulatory documents.

Enhance Study Conduct

Intralinks' Clinical Trial Portal is the single place of record for all study documents. Updating, tracking and discussing new trial information on Intralinks ensures everyone is always on the same page.

Easily Store, Manage, Review and Archive Site-Based Documentation

Enable sites to file and manage study documents with our Electronic Investigator-Controlled Site File (eISF). With document control everywhere and anytime, even the most sensitive content can be shared remotely, while secure, independent 3rd party hosted long-term archiving improves inspection-readiness after the completion of your study.

Manage Trial Information

Effortlessly exchange information among trial participants throughout the study. Create a single, secure repository for monitoring reports, data clarification forms, lab information, summary reports, patient diary data and adverse event reports.

intralinks.com

United States & Canada
+ 1 866 Intralinks
or 1 212 342 7684

Latin America
+ 55 11 4949 7700

Asia Pacific
+ 65 6232 2040

Europe, Middle East & Africa
+ 44(0) 20 7549 5200

INTRA LINKS STUDY SPACE

Safety Document Distribution

Centralize Information and Speed Up Reporting

Centralize SAE, SUSAR, IND, and ADR safety reporting on Intralinks so information is immediately available to all parties who need it. Principal Investigators, IRBs and other stakeholders can be notified via email or pager alerts when a document is posted so they have instant access to information.

Ensure Security and Reduce Risk

With Intralinks, you can control access to documents and reduce the risk of accidental disclosure to unauthorized individuals. Documents can be locked to prevent printing, saving and forwarding, and can be watermarked for additional security. And, Intralinks provides robust, real-time reporting capabilities to help with compliance.

Provide Global Access

Organize all documents in a centralized online location for instant global access by everyone who needs the information.

Intralinks

(NYSE: IL): The leading, global technology provider of inter-enterprise content management and collaboration solutions designed to enable the exchange, management control of documents and content between organizations securely and compliantly when working through the firewall. Thousands of companies and more than two million professionals use Intralinks for everything from ad hoc collaborations to complex, multi-business partnerships in financial services, life sciences, manufacturing, technology, and a wide range of other industries worldwide.