ClinTrak® SITE SOURCE STUDY PORTAL

INCREASING MONITORING EFFICIENCIES WITH REMOTE REVIEW OF DATA

The ClinTrak Site Source Study Portal provides a solution to the ever-evolving needs of remote data review by providing a secure platform for sites to easily upload source documentation for remote CRA review. Through this platform, you can accelerate your study's timelines by offering a seamless source document collection and review process for remote monitoring visits.

Medpace's approach to clinical trials ensures patient safety and data quality are met. The ClinTrak Site Source Study Portal complies with the General Data Protection Regulation (EU) 679/2016 ("GDPR") with regards to the processing of personal data of clinical trial participants. Internal technical and organizational resources and processes ensure a level of security appropriate to the risk, aligned with NIST Cybersecurity Framework.

MEDPACE REMOTE SOURCE REVIEW

175
STUDIES SET-UP IN UNDER A YEAR

< 2 weeks
AVERAGE TIME TO GO-LIVE

>16,000
PATIENT VISIT SOURCE DOCUMENTS UPLOADED FOR REVIEW

WHAT CAN THE CLINTRAK SITE SOURCE STUDY PORTAL DO FOR YOUR TRIAL?

MAINTAIN DATA QUALITY & SITE COMPLIANCE

- Critical data points accessed remotely in real time
- CRA ability to query directly in portal
- Reports available to track user upload, updates, and corrections
- Document certification confirmation built within system

SECURE PORTAL

- System controls protect documents during transit, upload, and maintenance
- Built in encryption code allows for optional site redaction, based on local regulations
- Protected access to assigned users
- Role based controls
- Antivirus, backups, disaster recovery, intrusion detection/protection, vulnerability management etc.

EASY TO USE

- Quick Reference Guide available for site training and reference
- Fully accessible on any browser
- Access provided via Medpace's website, which authorizes appropriate access and roles to Medpace systems (Access Request Management (ARM))
- Can be used in combination with Medpace's electronic data capture system (ClinTrak EDC)
- Upload within minutes directly from site computer
- Multiple pages may be included within one upload

STUDY TEAM & TECHNICAL SUPPORT

- Study team efficiencies to provide rapid role access/deactivation and usability
- Dedicated team to address technical issues
- CRA support and Quick Reference Guide (QRG) available

FULL-SERVICE CLINICAL DEVELOPMENT

Medpace is a scientifically-driven, global, full-service clinical contract research organization (CRO) providing Phase I-IV clinical development services to the biotechnology, pharmaceutical and medical device industries. Medpace's mission is to accelerate the global development of safe and effective medical therapeutics through its high-science and disciplined operating approach that leverages local regulatory and deep therapeutic expertise across all major areas including oncology, cardiology, metabolic disease, endocrinology, central nervous system and anti-viral and anti-infective.

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MAKING THE COMPLEX SEAMLESS™