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Solution overview

Elevate site efficiency and engagement with the Cognizant Shared Investigator Platform mobile app

Accelerate clinical research site actions with fast, intuitive and secure access to essential study tasks—anytime, anywhere. Purpose-built for today's busy investigators and support staff, the SIP mobile app streamlines study responsibilities so site staff can stay connected, productive and responsive throughout the day.

Clinical research sites are operating under growing pressure to handle more studies while maintaining speed, accuracy and compliance. The Cognizant® Shared Investigator Platform (SIP) mobile app is designed to ease these demands by bringing key clinical operations to mobile devices. It offers rapid access to essential study tasks and mobile-friendly document workflows, helping site teams work efficiently, stay informed and remain fully engaged in their studies.

By extending the capabilities of the SIP platform to smartphones and tablets, the app enables site staff to manage responsibilities such as document acknowledgments, e-signatures, safety notifications and study invitations with far less effort. With secure biometric access, intelligent alerts and seamless transitions between mobile and desktop, clinical teams can complete meaningful operational work anytime. Built for real-world site environments, the app supports quick approvals and productive moments between patient visits, helping investigators and study coordinators stay organized and compliant throughout the day.

What can site users do on the go?

Log in securely: Instantly access studies with fingerprint or facial recognition; no passwords required.

Accept pre-study and study invitations: New study opportunity? Accept sponsor/CRO invitations on the go without delay; no logging into multiple systems or searching emails.

Preview, generate and print FDA Form 1572: No more waiting for your PI to get to a desktop. Investigators can preview, e-sign, generate and even print FDA Form 1572 directly from their phones.

View documents and tasks: Access study documents anytime to stay informed and up to date with essential information regarding their studies.

- **Acknowledge and e-sign study documents:** Whether it's a protocol amendment or an essential form, you can review and e-sign study documents instantly. Plus, acknowledgments are just a tap away.
- **Biometric e-signatures:** Sign documents with confidence using fingerprint or facial recognition. It's fast, secure and eliminates extra steps.

Review safety notifications: Receive real-time safety alerts on the go and acknowledge them instantly because patient safety always comes first.

Read study news: Stay informed on the go with a centralized feed of study announcements and updates. Access the latest developments, insights and reminders of upcoming milestones instantly; no more searching through emails. Your studies stay right at your fingertips.

Accept manage facility associations: Investigators and other site leadership roles can add/remove staff associated with their facility quickly and securely right from their mobile device.

Benefits for sites:

Work anywhere, anytime

- Complete essential tasks like document acknowledgments, e-signatures, safety notifications and study invitations from your mobile device, without needing to return to a desktop.

Reduced administrative burden

- Clear task prioritization, intuitive navigation and real-time alerts help staff stay focused and avoid missing urgent or high priority actions.

Faster, more secure access

- Biometric login eliminates the need to remember or manage passwords, reducing friction during busy clinical workflows.

Seamless document handling

- Preview, generate, print, review, acknowledge and e-sign documents including FDA Form 1572 directly from your phone.

Stay informed and connected

- Access the latest study news, updates and safety alerts in one place, ensuring teams remain aligned and aware of critical study changes.

Benefits for sponsors:

Accelerated study startup

- Faster site response to pre-study and study invitations from their mobile devices, helps sponsors keep study startup timelines on track.

Improved site engagement and responsiveness

- Ease of access to new safety notifications, essential documents/forms and study updates leads to quicker site action and stronger site-sponsor communication.

Enhanced compliance and data integrity

- Biometric e-signatures and secure access ensure regulatory requirements are met, while reducing risks associated with password fatigue or manual document handling.

Higher productivity across the network

- By giving sites a simple, mobile-optimized way to complete tasks, sponsors benefit from more consistent site participation and smoother operational execution.

Better oversight and fewer bottlenecks

- With documents acknowledged and regulatory forms completed more quickly, sponsors experience fewer administrative delays throughout the study lifecycle.

About Cognizant Life Sciences

Cognizant's Life Sciences business unit partners with biopharmaceutical and medtech companies to develop strategies and solutions for healthcare challenges across the value chain. Our services and products, including SIP, digitize interactions between sponsors and investigators, helping the industry subtract time from clinical development and add it to improving patient outcomes.

Learn more about SIP: www.cognizant.com/life-sciences/shared-investigator-platform

For more information, visit: www.cognizant.com/life-sciences-technology-solutions

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