Cognizant® Shared Investigator Platform

One clinical trial operations platform across sponsors and studies.

Cognizant® Shared Investigator Platform makes it easy for sites to collaborate with sponsors.

Cognizant Shared Investigator Platform (SIP) provides a single point of access for multiple clinical trial resources across sponsors and investigation sites. This collaborative platform enables investigators and site staff to work together with sponsors in one central workspace, so the management of studies is more efficient, and information is easily shared–even as trials become increasingly decentralized and sites, patients and sponsors must collaborate remotely.

Cognizant SIP helps reduce many of the redundant tasks associated with site selection and qualification. By tracking study startup tasks and routing key communications, such as notices of study changes and safety notifications, the system allows investigators and their staff to spend more time with their patients, doing the work that matters most.

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How it works

Cognizant SIP is a site-centric, cloud-based SaaS solution which provides operational workflows and capabilities, enabling clinical trial sites to streamline the management of multiple trials for multiple sponsors.

With its single sign-on, site users log on just once to access multiple studies, reducing the hassle of keeping track of different sponsors with different systems. Investigators and their staff can spend less time logging in and out. Feasibility surveys aren't as time consuming. What's more, because Cognizant SIP is an open system, investigators and site staff can use it to seamlessly navigate dozens of other clinical trial technologies used by their sponsors.

The result is less time spent on duplicate paperwork for each study and more time with patients.

Benefits for sites

Cognizant SIP makes life easier, according to our users, with benefits that are unmatched.

Gain the ease of single sign-on

Time consumed by logging in and out of multiple systems as work is done within different studies can be significantly reduced. Cognizant SIP is designed to provide sites access to clinical trials across participating sponsors with a single sign-on. It can also speed workflow by allowing sites to share credentials with underlying clinical trial systems including Electronic Data Capture (EDC), Interactive Response Technology (IRT) and where available, it can link to electronic Investigator Site File (eISF) vendors.

Work more efficiently

Cognizant SIP helps simplify and reduce the burden of survey responses, document requests and training requirements by allowing you to share your site's profile in a central location. Cognizant SIP allows site users the flexibility and control to configure which sponsors can view their user profile during the registration process or update their choices at any point after they have registered. Consistent with GDPR privacy principles, sponsors can search and view site user profile information for only those users who have consented to share their profile with that sponsor.

Promote your site's capabilities

Cognizant SIP provides a two-way information exchange: sites upload details on their facilities and investigators' qualifications, while sponsors search for sites based on the needs of their protocols. The result? New partnerships and enhanced access for sites to participate in new and cutting-edge clinical trials.

Focus on patients

With fewer administrative burdens, clinical sites can better focus on patients' needs.

Why Cognizant?

Cognizant SIP offers a streamlined process and features that accelerate study startup and reduce administrative burdens by offering a single platform for managing key communications.

1. Create a single sign-on. Sites can access all of their clinical trials across participating sponsors in one place.
2. Build a user profile. Enter data once and maintain credentials centrally for platform sponsors and studies. Includes sponsor-searchable, electronically-signed digital CVs for investigators.
3. Manage facility and department profiles. Sites need only to complete their profiles and upload once for them to be accessible to platform sponsors and their future studies. User profiles can also be associated with multiple facilities and departments.
4. Work digitally. Sites, facilities and investigators are interconnected, enabling all to complete feasibility surveys and manage startup and day-to-day tasks for clinical studies.
5. Participate in dedicated study workspaces. Each study has its own designated work area where sponsors and study teams can post, share and retrieve documents safely and securely.

6. Prioritize and manage work. Investigators have a consolidated view of tasks across studies and sponsors.
7. Access and acknowledge receipt of safety notification reports. Sites have access to a centralized dashboard view of their clinical trial safety reports.
8. Track training. Investigators can complete training courses for one sponsor and receive credit across all participating sponsors via mutually-recognized GCP training.
9. Facilitate and maintain site regulatory binders. Cognizant SIP streamlines access to eISF, which enables clinical trial sites to electronically manage their essential regulatory study documents and data. This capability also enables both sites and sponsors to seamlessly exchange documents and information.

For more information,
visit **cognizant.com/shared-investigator-platform.**

Less is More

- Helps reduce redundant system logins through SSO
- Integrated site survey information reduces time
- Helps reduce duplicate CVs and autogenerates 1572s
- Mutually-recognized training reduces duplicate training requirements

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Cognizant (Nasdaq-100: CTSH) engineers modern businesses. We help our clients modernize technology, reimagine processes and transform experiences so they can stay ahead in our fast-changing world. Together, we're improving everyday life. See how at at **www.cognizant.com** or follow us **@Cognizant.**

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