Fact Sheet

Manage trials from end-to-end using the intuitive

IQVIA Investigator Site Portal

IQVIA Technologies' Investigator Site Portal offers sponsors, CROs, and sites the ability to easily and efficiently manage trials from beginning to end. Built by former investigators, coordinators, and study team personnel, the IQVIA Investigator Site Portal enables users to communicate effortlessly with sites, design and execute goals, provide reporting, trial metrics, and much more.

FEATURES

- Single sign-on
- Automatic reminders for upcoming tasks
- "Site-first" features such as patient visitation calculators, contact directories, robust FAQs and more
- Full-text searches
- Automatic audit trails
- Cross-trial safety notifications and training
- Comprehensive reporting

BENEFITS

- Faster, more efficient trials
- Ability to set, define, and execute trial goals with ease
- Better site relationships
- Complete audit coverage
- Reduction in trial costs

MODULES

SITE ACTIVATION

Activate sites up to 50% faster through reduction in cycle times, increased transparency, collaborative processes, and more by guiding sites, sponsors and CROs through a "to-do" checklist.

LEARNING MANAGEMENT SYSTEM

Save time, money, and reduce audit findings by providing sites and study teams with a single, efficient source for cross-trial education and learning, knowledge assessments, reporting and more.

SITE ENGAGEMENT

Set, execute, and measure trial goals, such as patient enrollment, protocol adherence, or clean trial data, all while offering tools that study teams can use to keep sites informed and appreciated.

DOCUMENT EXCHANGE

Access critical information the moment you need it, from anywhere in the platform. The 21 CFR Part 11-compliant system features fully searchable PDFs, dynamic distribution lists, and email notifications for a more streamlined way to manage trial documents.

SAFETY NOTIFICATIONS

Reduce the time, cost, and liability around safety notifications by disseminating, managing, and tracking SUSARs and other safety notifications with a centralized, simple cross-trial solution, all while providing 100% transparency.

CONTACT US
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