

INTRODUCING UNIFIED PLATFORM EDC. YOUR DATA. YOUR WAY. ON TIME.

Discover a comprehensive EDC solution that delivers high-quality data on time and addresses any study teams' goals:



EASE-OF USE

Sites and sponsors love our intuitive interface and guided workflow tools that simplify site and CRA tasks and streamline study management.



EFFICIENCY

Rapid setup tools help teams build studies in just 4-6 weeks, or less. Plus, roll out protocol amendments easily with no downtime.



FLEXIBILITY

Our no-coding EDC accommodates any study design from simple to highly complex.



SUPERIOR SERVICE

Every customer enjoys reliable, responsive, and attentive service. No project is too small, and no question unimportant. Our team is on your side.

Used on 3,000+ clinical trials, our EDC is the best choice for modern clinical trials

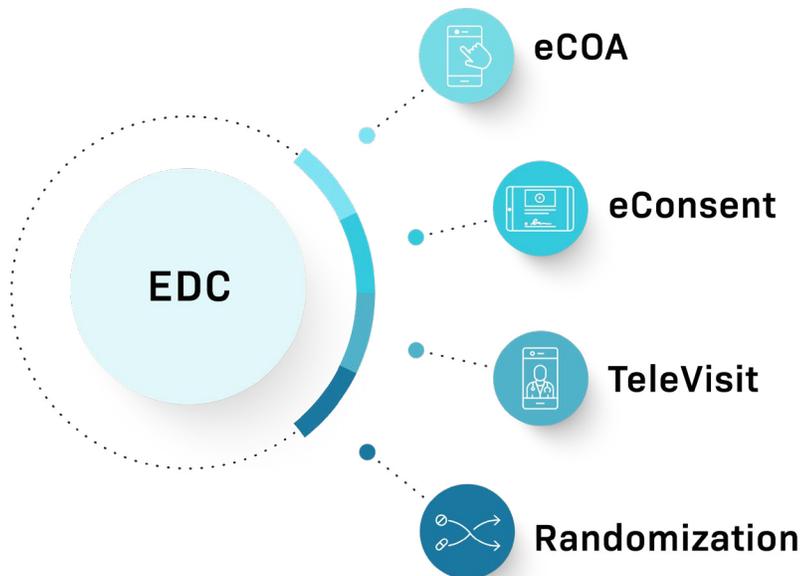
01 Comprehensive capabilities. Everything you need in an EDC system—from encoding and lab data imports to advanced edit checks and custom functions—all within a single, intuitive interface.

02 Optional eSource interface. Mobile-enabled responsive design allows investigators to capture data directly during patient consultations on tablets or other devices, with automatic identification of eSource data for streamlined SDV workflow.

03 Risk-based monitoring support. Advanced tools for targeted SDV and improved study oversight that optimize monitoring resources while maintaining data quality and regulatory compliance.

04 Modular add-ons. Start with our powerful EDC foundation and enhance as needed with eConsent, eCOA, Randomization, and TeleVisit modules through a single build and single database.

Our modular approach puts you in control, with a robust EDC foundation and optional add-ons tailored to your study needs.



eCOA

Capture patient-reported outcomes directly with our intuitive, device-agnostic interface using BYOD or provisioned devices.

eConsent

Streamline the consent process with digital forms, remote review options, and signature capture.

TeleVisit

Simplify patient follow-up with secure, compliant video visits.

Randomization

Easily implement patient randomization within the EDC workflow.

Why Choose Signant's EDC Solution?

With our EDC solution, you get much more than advanced technology. You benefit from Signant Health's deep operational, regulatory, and scientific experience, as well as our global scale and reach. Here's why customers love it:

① Scientific Expertise & Global Scale

- 25+ years of therapeutic and regulatory experience
- Deployed across 3,000+ clinical trials
- Used by 100,000+ sites in 83 countries
- 24/7 multilingual helpdesk support

③ Operational Excellence

- No downtime for mid-study changes
- Seamless mid-study changes with site-level versioning
- Attentive teams following mature operational processes

② Study Design & Build

- Rapid implementation (4-6 weeks)
- No-coding study design
- Comprehensive library of eCRFs, edit checks, and custom functions
- Flexible configuration for simple and complex studies alike

④ Technical Capabilities

- Direct data capture functionality
- Integration with other eClinical solutions
- Local and central lab data integration
- Central coding
- Standardized reporting
- Single database architecture across all modules

WHO IS SIGNANT HEALTH?



Signant Health is the evidence generation company. We are focused on leveraging software, deep therapeutic and scientific knowledge, and operational expertise to consistently capture, aggregate, and reveal quality evidence for clinical studies across traditional, virtual, and hybrid trial models. For more than 25 years, over 600 sponsors and CROs of all sizes – including all Top 20 pharma – have trusted Signant solutions for remote and site-based eCOA, EDC, eConsent, RTSM, supply chain management, and data quality analytics. Learn more at www.signanthealth.com.