SIGNANT BIOTECH

Signant SmartSignals® Unified Platform | EDC/DDC

A LEADING EDC/DDC SOLUTION FOR MODERN CLINICAL TRIALS

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/DDC is the best choice for modern clinical trials

01 **Mobile-enabled**. Use on any mobile device, with the responsive UI optimizing to any screen size. Investigators can use the system to record eSource data with the patient at site, or nurses can use it while conducting home visit assessments.

02 **Direct data capture**. No need to use different systems for eSource and EDC – our solution captures data genealogy so source data verification (SDV) and other data management activity can be determined accordingly.

03 **Remote SDV**. Reduce travel time and enable CRAs to conduct SDV remotely using our comprehensive remote SDV capabilities.

04 **Unified platform**. Incorporate eConsent, ePRO, RTSM, and televisits within the same unified platform; or integrate with third-party solutions.

Why choose Signant's EDC/DDC solution?

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

- ✓ Rapid implementation
- ✓ Flexible configuration for simple and complex studies alike
- ✓ No-coding study design
- ✓ Comprehensive library of eCRFs, edit checks, and custom functions
- ✓ Leading UI and workflow management
- ✓ No downtime
- ✓ Seamless mid-study changes without time-consuming data migration, with site-level versioning
- ✓ Remote monitoring/remote SDV
- ✓ Use with risk-based and targeted monitoring
- ✓ Fully mobile-enabled with responsive design
- ✓ Direct data capture functionality
- ✓ Ease of integration
- ✓ Local and central lab data integration
- ✓ Central coding
- ✓ Standard, custom, and ad hoc reporting
- ✓ Secure, clean data
- ✓ Experienced and attentive project team
- ✓ Available as full-service and self-service options
- ✓ Part of a single, unified platform including eConsent, Televisits, RTSM, and ePRO

Signant Biotech is part of Signant Health, 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/DDC, eConsent, RTSM, supply chain management, and data quality analytics.
Learn more at www.signanthealth.com

SIGNANT HEALTH
THE EVIDENCE GENERATION COMPANY™