SIGNANT BIOTECH

CLINICAL TRIAL TECHNOLOGY SOLUTIONS AND SERVICES TAILORED FOR EMERGING BIOPHARMA

Biotech and emerging biopharma innovations are critical for advancing healthcare, but resources, expertise, and speed are vital for effective processes that fuel novel drug discovery and development. Signant transforms clinical trials so emerging biopharma organizations can transform medicine.

WE UNDERSTAND THE CHALLENGES INHERENT IN DEVELOPING NEW DISCOVERIES INTO COMMERCIALLY VIABLE HEALTHCARE SOLUTIONS:

- Products are novel discoveries representing significant research and development investments
- Resources are limited and teams serve many functions
- Deadlines are tight and extremely consequential
- Trials can be as complex or even more so than those of larger pharmaceutical companies

SIGNANT BIOTECH SOLUTIONS & SERVICES ADDRESS CLINICAL DEVELOPMENT PRIORITIES:

ON-DEMAND EXPERTISE

Supplement your team with clinical, scientific, and operational experts available in your time zone, reducing burdens on your team and facilitating smooth collaborations.

COLLABORATIVE PROTOCOL DESIGN

Leverage our medical and clinical experts to assist in early protocol development and design, ensuring protocols are optimized to generate reliable data for any indication while simplifying participation.

COMPREHENSIVE TECHNOLOGY, OPTIMIZED TO YOUR STUDY

Drive accurate and reliable data, optimal site and patient experience, and simplified study management using eCaseLink, our fully integrated, unified platform of eClinical solutions, customized to meet your study design quickly and cost-effectively.

EASY TO USE TECHNOLOGIES

Easy to use for sites and sponsors, our unified eClinical platform means single sign on, integrated data, and converged, guided workflows enabling users to move effortlessly between applications.

WHY CHOOSE SIGNANT BIOTECH?

1. ENSURE RELIABLE DATA FOR ACCURATE DECISION MAKING

Capture high-quality data from anyone, anywhere with our comprehensive data capture solutions, including EDC/DDC, eCOA and eConsent.

2. MEET CRITICAL TIMELINES & MILESTONES

Get studies setup quickly to meet FPI targets and implement mid-study changes efficiently with our eCaseLink unified platform.

3. SUPPLEMENT YOUR TEAM WITH OUR SCIENTIFIC EXPERTISE

Leverage more than 50 full-time clinical and digital health science experts experienced in all therapeutic areas who proactively collaborate with your team in early protocol development and to optimize study solutions to meet study goals.

4. SIMPLIFY SITE, CRA, AND SPONSOR EXPERIENCE

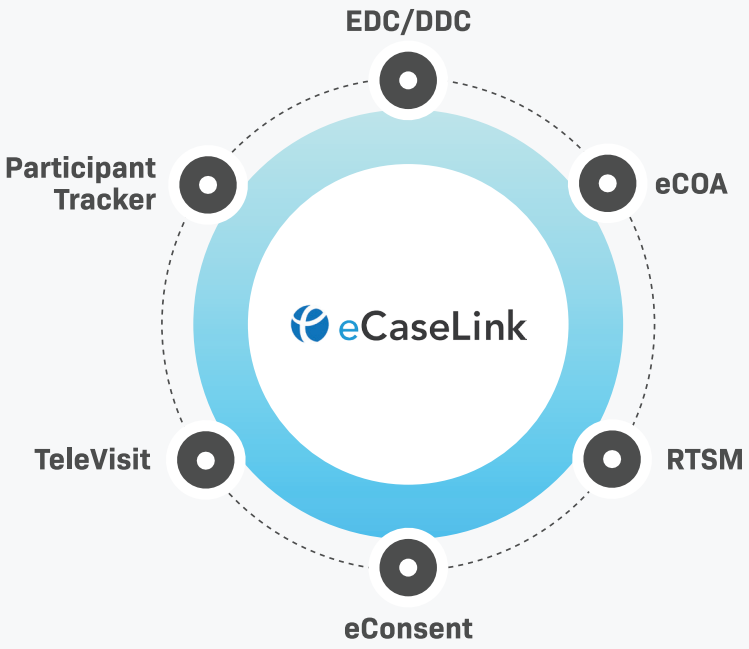
Simplify working using multiple solutions together with single sign on, converged workflow across individual platform components, and a single database enabling efficient, consolidated oversight and reporting.

5. GLOBAL EXECUTION, LOCAL TEAMS

Work with project teams in your time-zone, to simplify and streamline execution for local and global studies.

eCASELINK

UNIFIED PLATFORM



OUR SOLUTIONS & RESOURCES, YOUR INNOVATIONS

When the stakes are high, rely on our full suite of evidence generation and trial optimization solutions. Signant is fully resourced with the infrastructure and expertise biotechs and emerging biopharma companies need to achieve mission-critical milestones on time and on budget.