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:

01 Products are novel discoveries representing significant research and development investments

02 Resources are limited and teams serve many functions

03 Deadlines are tight and extremely consequential

04 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.

COMPREHENSIVE TECHNOLOGY, OPTIMIZED TO YOUR STUDY

Our clinical trial experts tailor technology to your specific study needs—from our Unified Platform EDC with add-on modules for eCOA, eConsent, and Randomization, to our specialized TrialMax, Rater Station, and SmartSignals RTSM and eConsent solutions for complex protocols.

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.

EASY-TO-USE TECHNOLOGIES

Our suite of solutions are easy to use for sites and sponsors. Our EDC provides a unified experience with add-on modules for eCOA, eConsent, Randomization and TeleVisits –enabling users to move effortlessly between applications.

Why Choose Signant Biotech?

01 ENSURE RELIABLE DATA FOR ACCURATE DECISION MAKING

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

02 MEET CRITICAL TIMELINES & MILESTONES

Our attentive study implementation specialists get studies setup quickly to meet FPI targets and implement mid-study changes efficiently.

03 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.

04 SIMPLIFY SITE, CRA, AND SPONSOR EXPERIENCE

Simplify working using multiple solutions together with single sign on, and converged workflow, using our comprehensive Unified Platform EDC with fully integrated add-on modules. Its single database enables efficient, consolidated oversight and reporting.

05 GLOBAL EXECUTION, LOCAL TEAMS

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

SIGNANT BIOTECH

WE'RE COMMITTED TO YOUR SUCCESS

PROVEN EXPERTISE THAT DELIVERS

As 25+ year veterans in clinical research with comprehensive digital-enablement, scientific, and operational solutions, Signant is fully resourced to help you achieve research milestones on time and on budget.

POWERING REGULATORY SUCCESS

Signant Heath's solutions and services have contributed to 28% of all recent novel drug approvals by the FDA and EMA.

ACCELERATING YOUR INNOVATION JOURNEY

Your discoveries advance healthcare, so let us help you optimize the development process with targeted applications of our solutions to accelerate processes as well as reduce risks and costs.

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.