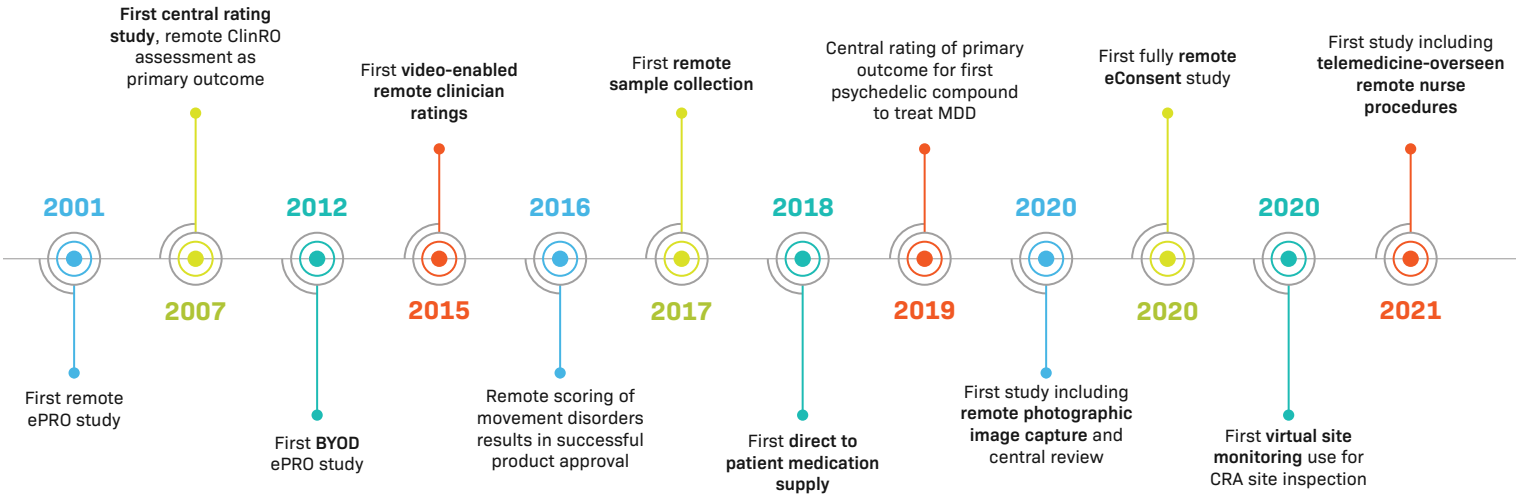
OPTIMIZING THE TRIALS OF THE FUTURE TODAY

At Signant Health, we understand that every clinical trial requires accurate, reliable measures of efficacy and safety as well as methods to simplify participation and conduct. In addition, each trial involves unique study characteristics such as study design, the disease, the concepts to be measured, the patient characteristics, and the geographies involved.

To ensure reliable evidence generation and simplified participation, **our solutions leverage thoughtful application of our technology suite and in-house clinical science expertise to design and deliver optimized clinical trials for any model of conduct.**

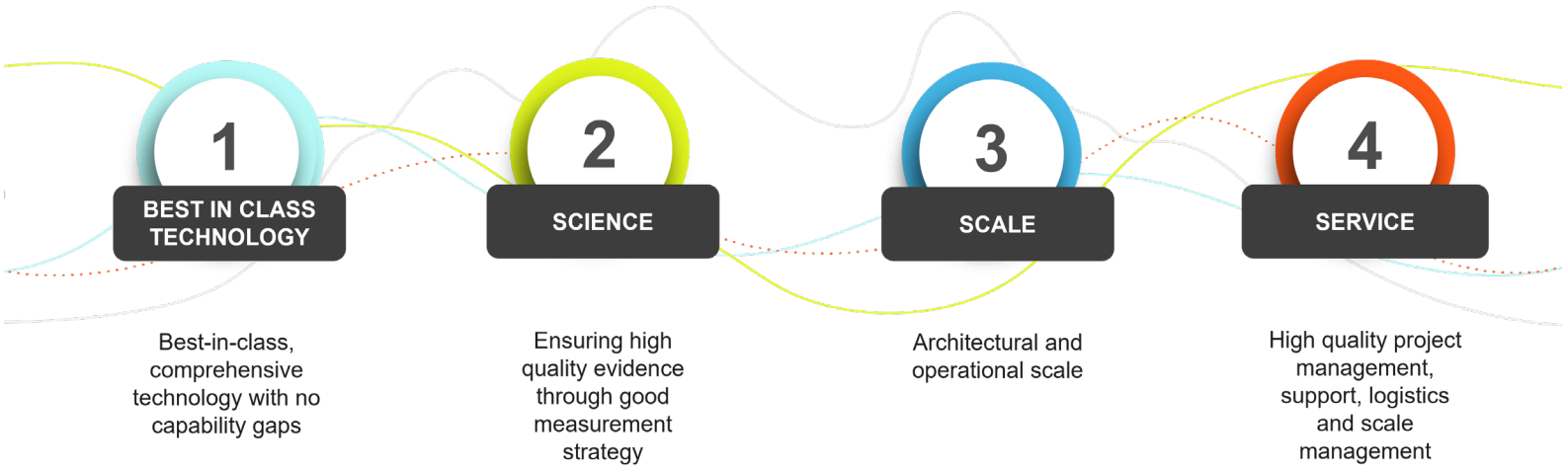
A RICH HERITAGE OF RESEARCH DECENTRALIZATION

For more than twenty years, Signant has pioneered the application of technology, scientific services, project delivery infrastructure, and scale to optimize clinical evidence generation and clinical trial conduct, regardless of how and where studies take place.



WHY CHOOSE SIGNANT SMARTSIGNALS?

Helping sponsors to optimize clinical trials involves thoughtful application of technology, in combination with clinical and scientific expertise. We focus on the four S's of decentralized clinical trial (DCT) success:



01

SOLUTIONS

Full-featured, pressure-tested software solutions optimize evidence generation, trial operation, and patient experience from enrollment to completion, without capability gaps. Our Signant SmartSignals suite also uniquely includes vital decentralized elements such as direct-to-patient medication supply, data aggregation solutions, and home nurse partnerships.

02

SCIENCE

Over 50 full-time clinicians oversee all aspects of study and solution design, ensuring optimal measurement strategy and measure validity.

03

SCALE

Extensive global reach with project teams in all time-zones and mature operational processes, combined with global architecture and infrastructure, facilitate hundreds of new trials per year, anywhere across the world.

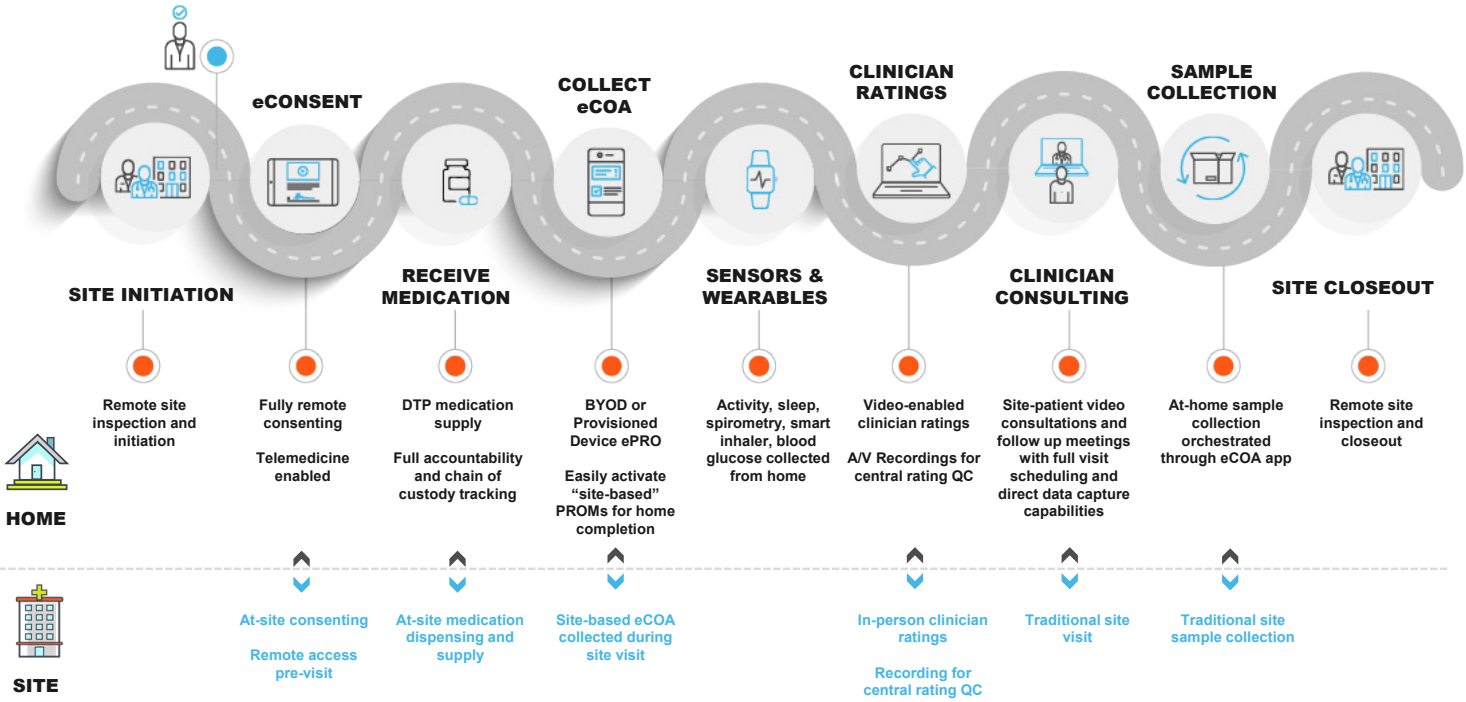
04

SERVICE

Experienced global project delivery teams, including end-to-end procurement and logistics, COA scale and license management, and 24/7 local language help desks drive hassle-free implementations.

EXPLORE SMARTSIGNALS SOLUTIONS FOR TRIALS WITH DECENTRALIZED ELEMENTS

Discover how our solutions simplify trial participation and decentralized trial operations. **Click** to explore the interactive patient data journey.



A CLOSER LOOK

SITE INITIATION, MONITORING, & CLOSEOUT

Enable CRAs to remotely inspect sites and facilities using proprietary video glasses worn by site staff.

- Save considerable time and money associated with CRA travel and on-site monitoring visits.

REMOTE eCONSENT

Our tiered eConsent solutions provide patients with remote access to study and consenting information alongside telemedicine to enable site-patient consultations and study participation discussion.

- Choose from rapidly implemented, PDF-based formats to rich multimedia solutions.
- Ensure well-informed patients to prevent unnecessary study dropout.
- Simplify consent/re-consent management and provide full traceability to limit inspection findings.

DIRECT-TO-PATIENT MEDICATION SUPPLY

Supply medication direct to the patient's home using our RTSM solution.

- Ship medication directly from site, central pharmacy, or country depot.
- Reduce unnecessary patient travel for dispensation when other on-site activities can be conducted remotely.
- Ensure full visibility of medication chain of custody and IP accountability when managing direct-to-patient supplies.

CLINICAL OUTCOME ASSESSMENTS (eCOA)

Collect patient-reported outcomes data at home using home diaries, quality-of-life measures, and other instruments.

- Select fully provisioned or bring-your-own-device models.
- High-quality COA data collection supports the simplest and most complex studies alike.
- Leverage our scientific, clinical, operational, and regulatory expertise to ensure endpoint data reliability.

SENSORS & WEARABLES

Collect a variety of objective clinical outcome assessments and biomarkers measured by sensors and wearables

- Gain richer insights into treatment effects with objective data captured by sensors and wearables.
- Ensure effective patient oversight through data access and reporting.
- Integrate with our eCOA to simplify usage for patients and sites.

CLINICIAN RATINGS (eCLINRO)

We train and enable sites to conduct complex clinician ratings remotely via video or telephone.

- Gain access to our large pool of qualified central raters for high-quality clinician ratings.
- Allow our clinical and scientific experts to assess each measurement scale for suitability for remote application.
- Ensure endpoint integrity and reliability when conducting clinician ratings remotely.

TELEMEDICINE CONSULTATIONS

Reduce the need for in-person visits by leveraging our leading telemedicine platform to offer remote, site-patient consultations as part of an engagement program or to replace regular follow-up assessments.

- Reduce patient burden by eliminating unnecessary on-site appointments.
- Interact with patients in an engaging manner throughout the study to retain a strong patient-site relationship.
- Accommodate caregivers or other health experts in the multi-person meetings.

SAMPLE COLLECTION

Enable patients to flag when samples are ready for collection and when new sample kits are needed through our eCOA solution.

- Ensure timely collection and delivery of sample kits to/from patients throughout the trial.
- Ensure sample pickups are convenient and quick for the patient.

PATIENT ENGAGEMENT

Provide patients with ongoing study participation information, including visit scheduling, site directions, and contact details.

- Guide patients through the study to limit missed visits and assessments.
- Increase engagement with study reminders, alerts, and study information.

HOME-HEALTH-SUPPORTED STUDY VISITS

Leverage Signant's established alliances with leading home health providers to enable a broader range of study activities to be conducted in patients' homes including:

- Conduct and oversee study assessments, and ensure accuracy and standardization through protocol-specific training
- Collect vital signs, EKG/ECG, blood draws, biologic sampling and preparation
- Administer medication and ensure adherence and accountability
- Support setup and use of at-home technologies such as telemedicine or devices

ASSESSMENT ADAPTATION

We ensure measurement validity when adapting measures from standard implementation.

- Migrate from face-to-face to sensor-, video-, or telephone-based assessments, ensuring measurement scientific validity.
- Train investigational and ancillary staff as well as provide training materials for patients on standardized assessment methodology for different settings.

LEARN MORE

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 20 years, over 400 sponsors and CROs of all sizes – including all Top 20 pharma – have trusted Signant solutions for remote and site-based eCOA, eConsent, RTSM, supply chain management, and data quality analytics. Learn more at www.signanthealth.com.