

SAFEGUARD YOUR PROTOCOL AND ENDPOINT DATA INTEGRITY

With expert guidance from Signant's dedicated team of clinical science, medicine, and operations professionals, you can have confidence that your protocol and endpoint data quality will be safeguarded throughout the lifecycle of a study.

Guarantee the validity and accuracy of the evidence your study generates:



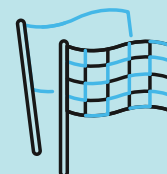
PAPER OR ELECTRONIC

Improve data quality regardless of how data are captured



EXTRA LAYER OF QUALITY CONTROL

Extend your team with the added support and oversight of our clinical science experts



START TO FINISH

Steer your study to success from design through regulatory submission

Why choose Signant's Advisory services?

Our in-house teams of experts are a second layer of support and oversight who guide your study from design through regulatory submission. With a range of services both included with our clinical trial technology solutions and available as stand-alone consulting, you can:

- 01** Increase confidence in data integrity and validity
- 02** Conduct paper-based or electronic studies, or a combination of the two
- 03** Decrease workloads for participants, sites, and sponsors/CROs
- 04** Reduce data variability and improve consistency
- 05** Rely on our experience in all therapeutic areas, phases, designs, and types of trial conduct
- 06** Minimize risks to data quality in every step of the development process

COMPREHENSIVE CLINICAL ADVISORY SERVICES INCLUDE:

Scientific Advisory for Clinical Trials



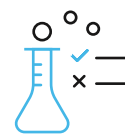
RATER TRAINING & QUALIFICATION

Ensure clinical raters score and administer study instruments accurately and consistently



DATA QUALITY MONITORING

Monitor and review clinical data throughout a trial to identify and remediate quality concerns



SCALE MANAGEMENT

Select, acquire, and implement study-appropriate scales in paper or electronic formats



LOCALIZATION & LANGUAGE MANAGEMENT

Preserve scale measurement properties by guaranteeing linguistic and cultural validity in any language



PROTOCOL ADVISORY

Leverage clinical and scientific experts to help develop an optimal measurement strategy to meet the objectives of your clinical trials

OUR SOLUTIONS ARE BETTER TOGETHER

UNLOCK THE FULL VALUE OF ADVISORY



eCOA



ELECTRONIC
CLINICIAN RATINGS



CENTRAL RATING



DATA ANALYTICS

Explore the Full Suite

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.

PROOF AT THE SPEED OF LIFE™

SIGNANTHEALTH.COM