SIGNANT HEALTH
THE EVIDENCE GENERATION COMPANY™

SmartSignals™ Solutions for ADHD

CLINICAL DEVELOPMENT RESEARCH TO IMPROVE ADHD RESEARCH

While considerable advancements in the clinical understanding of ADHD have led to effective treatment options, there is still need for the development of new medications and improved therapeutic strategies. **Signant's technology solutions and unrivaled scientific expertise help sponsors and CROs capture high quality, reliable evidence while streamlining the clinical trial experience for patients and sites.**

CAPTURE BETTER COA DATA

Patient-reported and clinician-reported outcome assessments are vital to fully evaluate ADHD treatment risk-benefit profiles. Our eCOA solution can be tailored to sites' and patients' needs – guided assessments and built-in edit checks reduce COA errors, while alerts and reminders help improve adherence to medication dosing and assessment schedules.

TRAIN & CALIBRATE RATERS

Train and monitor your study's raters to ensure consistency and accuracy with Signant's Rater Training and Central Review services. Our experts ensure raters adhere to eligibility criteria and score diagnostic/efficacy measures correctly, and they can help mitigate excessive placebo response to improve your ADHD study's data reliability.

ENSURE COMPREHENSION

Signant's robust eConsent platform employs multimedia, managed interactions, content flagging, and virtual visits to help participants and caregivers comprehend your protocol requirements. Plus, automated version control ensures only the most recently approved document version is presented.

WORK WITH ADHD EXPERTS

From consulting on outcome measure selection and implementation to data quality management strategies, Signant's in-house experts will help you navigate common ADHD challenges throughout the study lifecycle. Talk through your protocol with our CNS and pediatric research experts well-versed in ADHD protocols.

OPTIMIZE SUPPLY MANAGEMENT

Take control of product management complexities and simplify randomization for multi-arm trial designs with Signant's global RTSM solution. Study teams trust our solution to not only launch studies faster but to handle patient cohorts, study supply forecasting, mid-study changes, and data transfers with efficiency and accuracy.

DIGITALIZE THE PROCESS

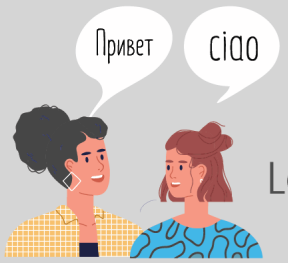
Each solution and service within our SmartSignals evidence generation platform can be applied to a study independently. However, when combined, they create an intuitive and powerful digital ecosystem for creating and managing complex global studies. Plus, every study is supported by a dedicated team of clinical science and operations experts.

At Signant, our focus is helping you develop and deliver treatments or therapies faster in order to improve the quality of life for people impacted with ADHD.

SIGNANT'S ADHD CLINICAL TRIAL EXPERIENCE

Phases

II III IV



16 Languages

900+ Sites

13 Countries

6,200+ Patients

56 Protocols

SMARTSIGNALS SUITE

SmartSignals solutions can be used individually or integrated together for a seamless, end-to-end digital experience.

eCOA

Scientific Advisory

RTSM

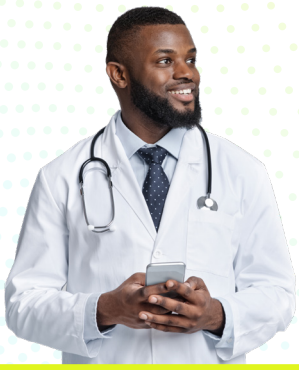
eConsent

Rater Training

DISCUSS YOUR NEXT STUDY WITH US

Our global team of therapeutic area experts advise on all areas of the clinical development process, including:

- Clinical science and medicine
- Data analysis
- Regulatory
- Operations and trial administration
- Global logistics



MEET THE EXPERTS