SIGNANT HEALTH
THE EVIDENCE GENERATION COMPANY™

Cystic Fibrosis Clinical Trial Solutions

CLINICAL DEVELOPMENT SOLUTIONS TO IMPROVE CYSTIC FIBROSIS RESEARCH

Thanks to advancements in treatment, people with cystic fibrosis live (CF) longer than ever before, but work remains to develop safe and effective disease-modifying therapies. Signant Health can support the drug development process by applying our SmartSignals™ **clinical research solutions** as well as **scientific expertise** to help researchers improve trial outcomes.

CAPTURE BETTER ePRO DATA & KEEP PATIENTS ENGAGED

In one platform, Signant's eCOA solution provides clinicians and patients with tools to improve the assessment experience. Electronic administration of PROs such as CFQ/CFQ-R makes it easier to complete them at home or site, and produces more accurate data. Built-in patient engagement features also help keep participants on track and adherent.

CONDUCT STUDY ACTIVITIES REMOTELY

Reach the participants you need to meet enrollment goals by offering solutions that facilitate remote study activities. From RTSM to eConsent, eCOA, and Telemedicine, Signant's comprehensive range of solutions and global project teams will help you conduct trial activities anywhere and any way.

MINIMIZE PARTICIPATION BURDENS

Make study participation more convenient for patients and sites. With eConsent, patients can take their time reviewing study requirements at home before making the effort to visit the clinic. Once enrolled, your team can leverage our secure, compliant Telemedicine platform to connect with patients between visits or for appointments not requiring site-based assessments.

PARTNER WITH CF EXPERTS

With decades of clinical science experience, our CF and rare disease research leaders are prepared to serve as an extension of your study team. Rely on our in-house experts to help optimize protocols, implement clinically-appropriate instruments, and oversee data quality from launch through regulatory submission.

SIMPLIFY MEDICATION MANAGEMENT

Take control of inventory management for numerous medications and simplify randomization for multi-arm trial designs with Signant's global RTSM solution. Our agile design process will help you launch quickly and manage mid-study changes with ease. Plus, a dedicated team experienced in CF protocols will support your trial operations no matter how or where you conduct your trials.

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 cystic fibrosis patients everywhere.

SIGNANT'S CYSTIC FIBROSIS CLINICAL TRIAL EXPERIENCE

Phases

I II III IV

12 Languages

2,600+ Sites

18 Countries

2,600+ Patients

85 Protocols

SMARTSIGNALS SOLUTIONS

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

eCOA

Scientific Advisory

eConsent & Telemedicine

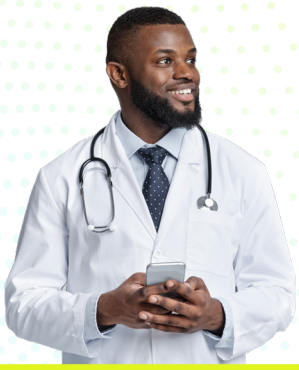
Decentralized Trial Activities

RTSM

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