SIGNANT HEALTH
THE EVIDENCE GENERATION COMPANY™

Solutions for Hemophilia A

CONDUCT BETTER HEMOPHILIA A TRIALS FOR BETTER OUTCOMES

While the standard of care has advanced dramatically, new discoveries and approaches offer opportunities to develop better hemophilia A treatments. Signant Health can apply decades of deep therapeutic-area, pediatric, and rare disease expertise, as well as innovative clinical trial technologies within our Signant SmartSignals™ suite to help researchers support hemophilia A study endpoints.

STREAMLINE COA MANAGEMENT

With many types and versions of home- and site-based clinical outcome assessments involved in hemophilia A trials, capturing accurate, reliable COA data can be challenging. Leverage our comprehensive eCOA platform and our scientific expertise to simplify COA administration, reduce patient burdens, optimize data quality, and launch studies quickly.

MINIMIZE PARTICIPATION BURDENS

Make study participation more convenient for patients and caregivers. With eConsent, patients can take their time reviewing study requirements at home before making the effort to visit the clinic. Once enrolled, leverage video-assisted patient engagement to connect with patients and caregivers between visits for safety monitoring and regular checkups.

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 hemophilia A protocols will support your trial operations from study launch to database lock.

LEVERAGE OUR HEMATOLOGY RESEARCH EXPERTS

From consulting on outcome measure selection and implementation to reducing burdens on sites and patients, Signant's in-house experts will help you navigate study design and operational challenges throughout the study lifecycle. Talk through your protocol with our full-time clinical science and medicine experts experienced in hematology, rare disease, and pediatric studies.

DIGITALIZE THE PROCESS

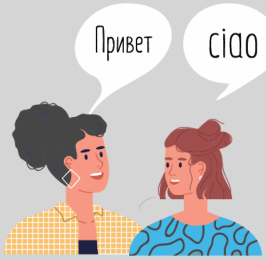
Each solution and service within the Signant SmartSignals evidence generation suite can be applied to a study independently. However, when combined, they optimize trial operations and improve the participation experience for sites and patients. Plus, every study is supported by a dedicated team of clinical science and operations experts.

With Signant, you gain a single-source partner for comprehensive evidence generation and trial optimization solutions that support the full clinical trial lifecycle.

SIGNANT'S HEMOPHILIA A CLINICAL TRIAL EXPERIENCE

Phases

I II III IV



40+
Languages

1,100
Sites

30
Countries

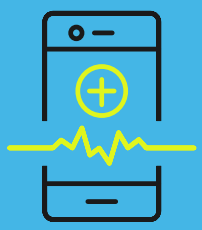
2,000+
Patients

20+
Protocols

SMARTSIGNALS SUITE

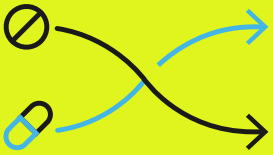
SmartSignals solutions can be used individually or integrated together for a seamless, end-to-end digital experience.

eCOA



Patient Engagment

RTSM



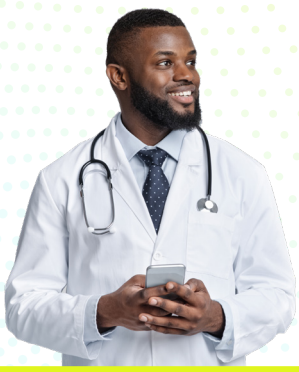
eConsent

Scientific Advisory

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