

Post-Traumatic Stress Disorder (PTSD) is a psychiatric disorder that may occur in people who have experienced or witnessed a traumatic event. PTSD has largely been underserved in pharmaceutical research, which has made study teams less experienced in recognizing, diagnosing, and assessing changes in PTSD patients. Signant has the scientific expertise and innovative technology needed for sponsors run a successful PTSD study.



ADMINISTER SCALES CONSISTENTLY

We collaborated with internationally known PTSD experts to develop a comprehensive, electronic version of the CAPS-5 for our Electronic Clinician Ratings solution. With embedded training conventions and guidance, the electronic format collects reliable data to accurately diagnosis and detect changes in symptoms.



STANDARDIZE RATER TRAINING

Signant's Rater Training & Qualification solution standardizes PTSD assessment techniques through a customized and comprehensive program. The curriculum focuses on scale knowledge as well as applied skills training and assessment for a wide variety of gold standard scales such as the SCID, MINI, and CAPS-5.



GAIN EXPERT GUIDANCE

Signant's scientific team ensures the highest level of training at the outset and throughout the course of the trial. Our in-house experts will also provide real-time edit checks on the performed scales to identify potential errors for raters' consideration prior to data submission.



IMPROVE SIGNAL DETECTION

Select our Blinded Data Analytics to reduce data issues caused by inconsistent assessments, which will improve your ability to detect signals. Proprietary algorithms examine data throughout your study to identify trends, anomalies, inconsistencies, or discrepancies.

At Signant, our SmartSignals solution suite was designed specifically to meet the needs of PTSD sponsors, sites, and patients.

SIGNANT'S PSYCHIATRY CNS EXPERIENCE IN THE PAST 5 YEARS

Phases











24 Languages









47,300+ Patients



SMARTSIGNALS SOLUTIONS

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

eCOA



Telemedicine



Activity Monitoring





eConsent



Cognitive Assessments



DISCUSS YOUR NEXT STUDY WITH US

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

- Clinical science and medicine
- Data analysis
- Regulations

- Operations and trial administration
- Global logistics



MEET THE EXPERTS