

# ASSESS & OPTIMIZE RATINGS RELIABILITY



When research involves complex, clinician-reported outcome (ClinRO) assessments, endpoint reliability is a critical consideration. Signant's Central Ratings services help study teams **meet protocol requirements, optimize data quality, and reduce burdens on patients.**

Our global network of trained central raters can provide expert clinical ratings remotely as well as independent, expert scoring of assessments performed by sites. This enables sponsors and clinical research organizations to:



**Generate accurate, reliable ClinRO data**



**Assess eligibility and adjudication with independent experts**



**Offer remote assessments to patients to reduce study burden**

## WHY CHOOSE SIGNANT'S CENTRAL RATINGS?

- 01** Satisfy regulatory requirements for site-independent assessment of ClinROs for treatments that may have functional unblinding characteristics, such as psychedelics.
- 02** Broaden access to studies that require specialized clinical expertise, such as rare and orphan diseases.
- 03** Improve endpoint accuracy and reliability by performing in-study quality control, supported by over 50 full-time experts and our worldwide network of local language clinicians.
- 04** Work with a single partner that offers end-to-end technology solutions and scientific services to enhance signal detection and optimize evidence generation.

# B E T T E R T O G E T H E R

Combine Central Ratings with supporting technology solutions and scientific services to facilitate central ratings and gain additional in-trial quality assurances:



**PROTOCOL  
ADVISORY**



**SCALE  
MANAGEMENT**



**ELECTRONIC  
CLINICIAN  
RATINGS**



**RATER  
TRAINING**



**TELEMEDICINE**



**PURESIGNAL  
ANALYTICS**



## GET IN TOUCH

Contact our experts to review your protocol, submit an RFP, review our posters, or learn more about these endpoint reliability solutions.

**CONNECT TO EXPERTS**

## 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](http://www.signanthealth.com).