

NAHA Health provides an end-to-end solution for clinical trials that enables real-time sharing of data between the physician, nurse and study coordinator. Our paperless clinical trial platform reduces data errors, streamlines regulatory workflows, accelerates timelines and is HIIPAA-compliant. We can support a clinical trial model in which a patient participates in the trial from home or in a controlled clinical environment. Our team can provide assessments to determine the clinical and behavioral health status of the patient prior to starting the clinical trial. Any patients with increased risk will be more closely monitored for their safety throughout the trial by their dedicated NAHA Health nurse. The platform development and data automation is driven by your clinical trial requirements.

- Examples of Capabilities:

   Eligibility call preformed by a nurse

   Physician data is electronically captured

   Paper records can be digitized

   Dedicated nurse for each participant

   On-going quality of life assessments through completion of trial

   Clinical health risk assessments to determine risk level of participants

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- Physician assessments/ checklists
- Participant questionnaires / surveys





## The user interface is designed based on your clinical trial's needs

trial's needs
As your clinical trial partner, we will develop a
customized nurse monitoring program that will
maximize the safety of your study participants.
Together we will create a customized end-to-end
interface to monitor participants, collect real-time data
and digitize reporting for seamless execution.



Our nurse care team can facilitate questionnaires and quality of life checks throughout the trial. The NAHA team is trained in the field of medical research. We can oversee medical questionnaires, surveys or quality of life assessments before, during, after or as-needed during the trial. Additionally, we can monitor the behavioral health status of participants throughout the trial for safety.

