Subawards for Study Sites

Subawards will be distributed after the site start-up surveys are completed. For those that are ready to proceed, you will hear from the CLARiTI team and NACC to review your budget and begin collecting required documents. To learn more, please view the February 28 webinar below.

Site Initiation Meetings

Thank you to everyone who has met with us to begin on-boarding discussions at site initiation meetings in the past few months. We’ve already met with 26 ADRC sites! We'll schedule the remaining meetings in the weeks to come.
CLARiTI Concierge

Each site is paired with one of our site project coordinators. Find out which coordinator is assigned to your site on our website. Meet our coordinators below.

Kelsey Shuda is a Clinical Research Project Coordinator for CLARiTI and is eager to begin collaborating with sites to implement CLARiTI. She previously worked as an Outreach Specialist for Wisconsin ADRC's ORE Core where she served as a liaison for research participants. Kelsey has worked closely with the Clinical Core and DVCID studies to fulfill study milestones. She also coordinated WI ADRC's ancillary studies to meet study and recruitment goals. Kelsey enjoys a good cup of coffee, exercising, and traveling.

Email: klshuda@medicine.wisc.edu

Kelsey Shuda
Hanzhe Gao is a Clinical Research Project Coordinator with a background in research and medical sciences. Previously, he worked as a coordinator for the NIH "All of Us" research program. He has skills in coordinating collaborative, multi-site research projects, as well as insight into the many tenants of successful community engagement initiatives. With the CLARiTI project, he will help coordinate the consortium of ADRCs across the US. Outside of work, he enjoys hiking the trails with his husky, Riceball, or looking for unique cuisines to try with his wife of three years, Cynthia.

Email: hgao@medicine.wisc.edu

Upcoming CLARiTI Events

**CLARiTI WEBINAR**

*Study Startup & Inclusion Processes*

*Wednesday, March 27, 2024
2:30 - 3:30pm ET*

**CLARiTI - Recruiting for Inclusion**

*Wednesday, March 27, 2024, at 2:30-3:30 pm ET*

Register for the Webinar

*Special thank you to Dorothy Edwards, PhD, Chair of the ORE Core Steering Committee for offering valuable feedback on behalf of the committee.*
We look forward to connecting at the upcoming Spring ADRC meeting May 6 & 7, 2024 in Austin, Texas! Invite for CLARiTI Luncheon to follow.

Register for the Spring ADRC Meeting

CLARiTI Received IRB Approval - Sites May Initiate Cede Application with Local IRB

The CLARiTI protocol received IRB approval on January 9, 2024. Your site may now initiate a cede application with your local IRB to cede review to our single IRB, WCG. The approved documents, which include two consent templates (one for cognitively impaired, one for cognitively unimpaired), and the study protocol, are filed here. Additional materials will be forthcoming.

Updates on Radioligand Contracts

Contracts are either in place or nearly complete with the companies who supply the several radioligands used in this study. Our team submitted the Investigational New Drug (IND) application to the FDA for review on February 6.

The CLARiTI Administrative team will be in touch once our IND is approved and will notify sites of the onboarding logistics of ordering the radioligand tracers and training involved with each company. Please do not reach out to the radioligand companies regarding CLARiTI as we want to avoid duplication of efforts.
Visit the CLARiTI Website

Our website is live and holds lots of pertinent information for your study teams. We continue to update the FAQ page. Visit the website here for:

- Webinars & events, including recordings
- Links to study documents
- FAQs
- News
- About CLARiTI and team

Contact Us:

Email: clariti@medicine.wisc.edu
Web: https://naccdata.org/nacc-collaborations/clariti