



## Co-Enrollment Program

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### Suggested Recruitment Materials (Collaborator to Participant)

These materials are provided as reference for the collaborator. The collaborator can choose to use whatever materials work for their study. Please note that all recruitment materials will need to be submitted to in the collaborator's IRB submission.

#### ***Phone Script***

Hello, my name is \_\_\_\_\_. I'm calling from the <COLLABORATOR study name and information>.

We are excited to announce that the <COLLABORATOR study name> will be partnering with the Brain Health Registry. The Brain Health Registry (BHR) is welcoming <COLLABORATOR study name> participants to enhance their experience by completing online assessments! Participation in this study is completely optional and you can withdraw at any time.

Is now a good time to talk about the Brain Health Registry?

➤ *If "yes": continue with the below.*

First I have a couple of questions that will help me figure out if the Brain Health Registry is right for you.

- 1) Do you have regular access to a computer with the internet?
- 2) Do you have an email address?

➤ *If "yes": continue with the below.*

#### Version 1 (Short)

Great! I'd like to tell you a little bit more the study. The **Brain Health Registry** is an internet study conducted entirely online. Participation consists of going online to answer questions and take online brain function tests, which are similar to paper and pencil tests you might take in a doctor's office. It takes about <TIME> minutes/hour(s) total to participate but you don't have to complete all study tasks at once. You can return to the website to finish remaining tasks. Participants are emailed every <TIME> months/year and asked to log back in to answer follow-up questions and retake brain function tests. Your caregiver or care partner will also have the opportunity to participate, which will help us learn more about their caregiving experience and how it affects their health. In addition, the Brain Health Registry gives you the opportunity to learn about and participate in other research studies, including clinical trials.

Would you like to participate in the <COLLABORATOR study name> - Brain Health Registry?

➤ *If "yes"*

Great, then the next step is to <explain study specific registration process>

#### Version 2 (long)

The Brain Health Registry is a free internet study lead by researchers at the University of California, San Francisco. Volunteers who agree to participate in this study go to the Brain Health Registry website, answer questions about themselves, and take online brain tests—which are like brain games. Volunteers are emailed every <TIME> months/year

and asked to log back in to answer follow-up questions and retake brain tests. They receive a newsletter every few months that gives interesting information about brain health and Alzheimer’s disease. Volunteers may also be eligible to take part in other studies through the Brain Health Registry, such as clinical trials.

The goal of this collaboration is to <brief description of the BHR Collaboration goals>

**Optional if co-enrolled participants can be referred to another study:** *(In addition, the Brain Health Registry is designed to accelerate the recruitment process for clinical research, as recruitment is typically time-consuming and expensive. By helping speed the recruitment process for treatment trials, the Brain Health Registry hopes to speed the path to treatments for Alzheimer’s and other brain diseases.*

*The Brain Health Registry study staff use the information they collect from their volunteers to determine if there are other research studies volunteers can participate in. As a Brain Health Registry participant you have the option to learn about these research opportunities, or not.)*

Participation in the Brain Health Registry is entirely online. You can login from a desktop computer, laptop, tablet or smartphone to complete most tasks. Portions of the Brain Health cannot be completed on tablets or smartphones, and require login from a computer or laptop, but you can still participate if you don’t have access to a desktop or laptop computer.

For most participants, it takes about <TIME> minutes/hour(s) to complete all study tasks, but you don’t have to do them at once. You can leave the website and log back in to complete tasks at a convenient time. Participants receive emails, reminding them to log back in and finish remaining tasks.

Before completing any tasks, volunteers must agree to an Informed Consent Form. All information entered in the Brain Health Registry is secure and privacy is protected.

Volunteers who enroll in the Brain Health Registry will have the opportunity to invite a friend or loved one to participate as their Study Partner. A study partner is someone who can answer questions about your health, memory, and day-to-day activities. Having the pair of you participate in the Brain Health Registry together helps illuminate a more complete understanding of brain health. Study Partners also go to the Brain Health Registry website to answer questions and take brain tests. Study partners get newsletters and email reminders, and even have the opportunity to join the main Brain Health Registry study. Study Partners and participants will not have access to the information the other person provides.

Would you like to participate in the <COLLABORATOR study name> - Brain Health Registry?

➤ *If “yes”*

Great, then the next step is to <explain study specific registration process>