**DEMENTIA**

**Participant Inclusion Criteria**

**Neurotypical and Mild Cognitive Impairment**

Participants are >59 years old, not taking (or stable while taking) nootropic medication for 3 months or more, and able to read, speak, and understand spoken English. Participants will be excluded if they are unable to provide informed consent. This will be assessed with comprehension questions about the study embedded in the informed consent form.

**Participants will be excluded if they self-report any of the following health conditions on the general screening form:**

* major psychiatric disorder (e.g., schizophrenia)
* untreated major depression
* aphasia, apraxia of speech, dysarthria
* history of brain tumor, traumatic brain injury, or ≥ 3 concussions

**Dementia**

Participants are >59 years old, not taking (or stable while taking) nootropic medication for 3 months or more, and able to read, speak, and understand spoken English. Participants will be excluded if their LAR does not consent to the study or if the person with dementia does not provide verbal assent.

**Participants will be excluded if they have any of the following health conditions on the general screening form:**

* major psychiatric disorder (e.g., schizophrenia)
* untreated major depression
* aphasia, apraxia of speech, dysarthria
* history of brain tumor, traumatic brain injury, or ≥ 3 concussions

***Telehealth Inclusion Criteria:***

Participants who are completing session via telehealth must have access to Internet and an Internet-enabled device with a camera and have adequate hearing and vision to participate in the telecare platform. Adequate hearing will be determined by having the participant repeat words and phrases while turning up the volume on their headphones or computer speakers as needed. Adequate vision will be determined by the clinician projecting words and phrases onto the screen – as large as possible for the smallest font used in assessment materials.