

## **BEHIND MEMORY LOSS:**

Disparities in the United States Alzheimer's Care System

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## The leeberg Model

- Inadequate insurance coverage for specific age groups.
- Patients without family support cannot provide the expenses for aftercare treatments.
- · Non-White populations in Alzheimer's are often underdiagnosed.
  - Patients' families are often overwhelmed on how to handle new symptoms.
    - Patients' medical insurance is not sufficient for their treatments (Medicare will often not cover expensive tests/treatments.)
    - Lack of guidance and information to assist patients and their families.
    - Many AD diagnostic tests have been developed in almost exclusively White AD population at early stage in the disease, these tests may not be representative of other populations.

**Insufficient funding from National Institutes of** 

Health (NIH) for high stake research programs.

U.S. Healthcare and insurance structures

usually do not include AD aftercare as part of what is covered for patients.

 Serious physical issues or comorbidities often overshadow AD at healthcare screenings.

 Knowledge gaps in minority communities regarding the progression of AD.

 Lack of awareness for seeking Alzheimer's testing, especially in younger populations.

• Cultural incompetence between doctors and patients.

 Stereotypes and biases against AD (feelings of shame) inhibit people from reaching out to seek treatment.

Physical ailments are often thought to be more important and are treated first compared to a more hidden disease like AD.

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 AD is viewed as a disease for the elderly when in reality younger populations are also susceptible.

 Physicians hold off testing for AD in younger people however treating AD early is the best way to combat the disease.





## Stakeholder Map

XC= 33X







Low Interest

Low Power

High Interest

## Journey Map



