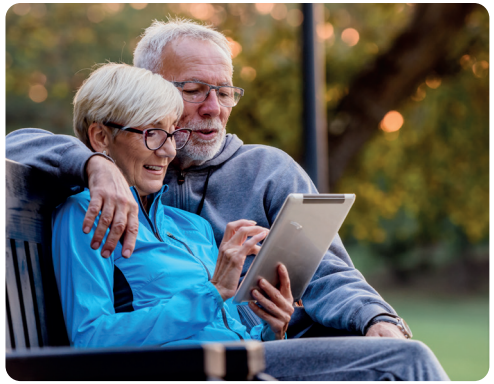
How do I get more information?

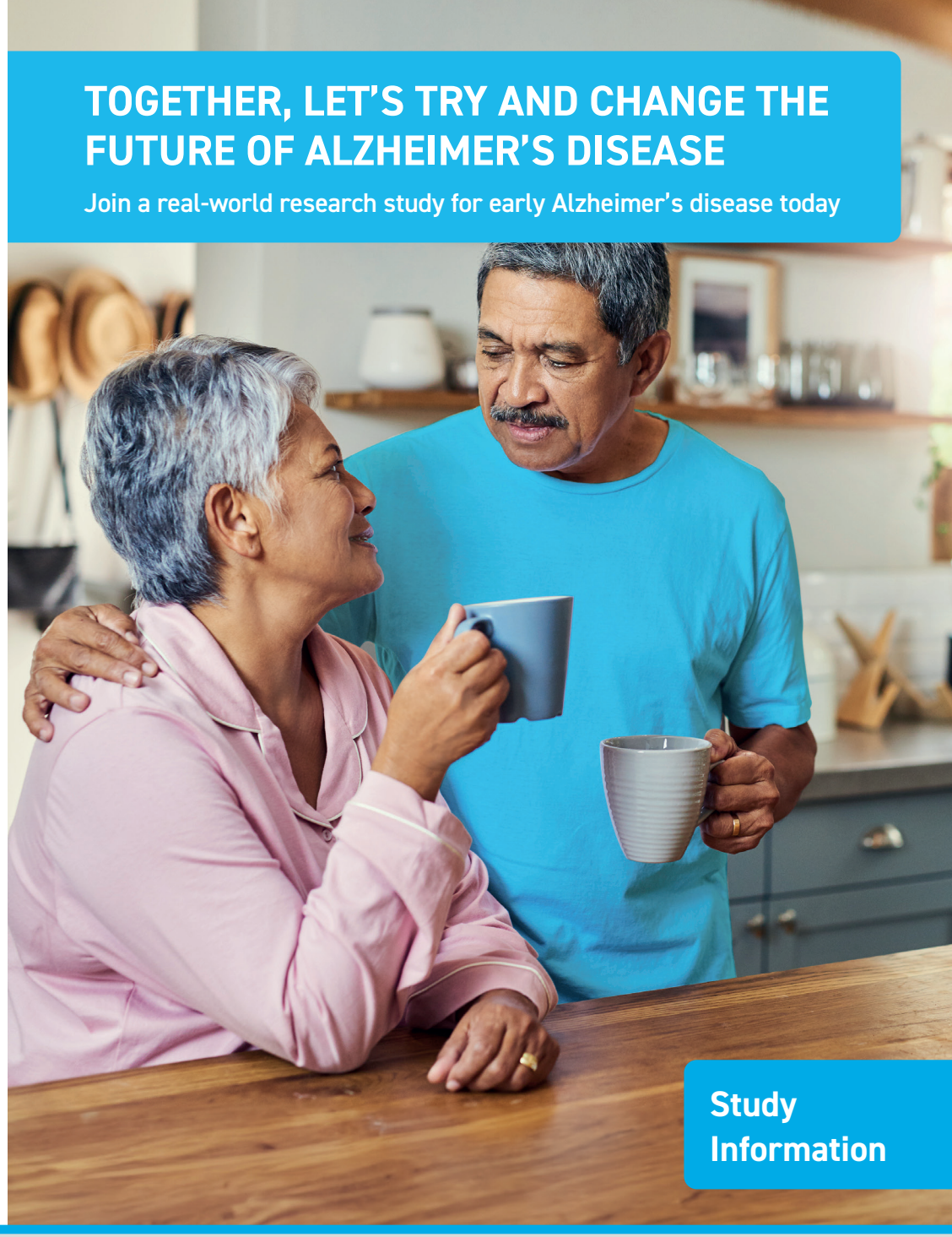
To find out more, please contact the study team using the information provided here. Study participation is voluntary. By contacting us, you are under no obligation to take part in the study.



Austin Regional Clinic
ARC Four Points
Visit ARCclinicalresearch.com
Call 737-247-7240

TOGETHER, LET'S TRY AND CHANGE THE FUTURE OF ALZHEIMER'S DISEASE

Join a real-world research study for early Alzheimer's disease today



Study Information

This brochure contains information about the TRAILBLAZER-REAL US Study. This information should help you decide whether you, or someone you know, may want to take part in the study.

What is a real-world evidence study?

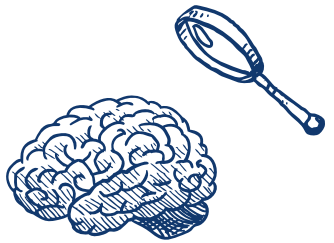
A real-world evidence study is done to gather more information about a new medication. Real-world evidence studies help doctors and scientists to learn more about the new medication's use in everyday life or in new situations.

Deciding to take part in a research study is an important decision. If you have any questions, you can contact the study team using the information provided in this brochure.

About the TRAILBLAZER-REAL US Study

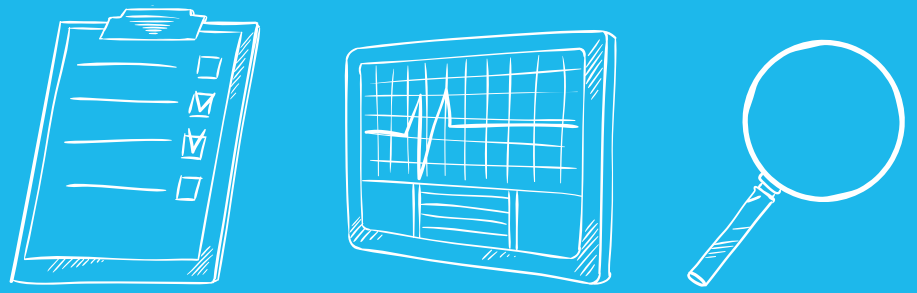
Alzheimer's disease (AD) causes changes in the structure and chemistry of the brain. This can cause symptoms such as memory loss, confusion, and trouble finding the right words or thoughts. There is currently no cure for AD, but researchers are looking for treatments to slow the disease or stop it from getting worse.

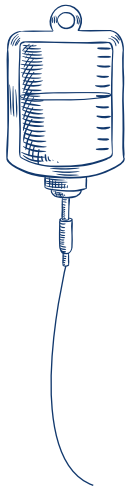
This study is looking to learn more about how well a new medication (study medication), when given alongside the usual care, works compared with usual care alone for people with early symptoms of AD in the real world.



Why is this study important?

Real-world studies are important because they can help doctors and scientists learn more about how a medication works when used in everyday care, rather than in a clinical trial. These studies can also help doctors and scientists understand how the medication affects outcomes that are especially important to patients and their caregivers.





What will this study involve?

If you take part in the TRAILBLAZER-REAL US Study, you will receive either the study medication along with your usual care, or you will receive usual care alone.

- Whether you will receive usual care alone or study medication plus usual care will be based on your research site.
- Usual care represents the standard of care that you would receive outside of a research setting.

If you receive the study medication, it will be given by intravenous (IV) infusion. This means the study medication is given directly into a vein in the arm.

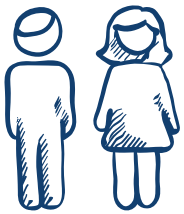
Screening period
(up to 90 days)

- You will visit the study center to see if the study is suitable for you.

Based on the study center you are participating in, you will be placed into 1 of 2 groups:

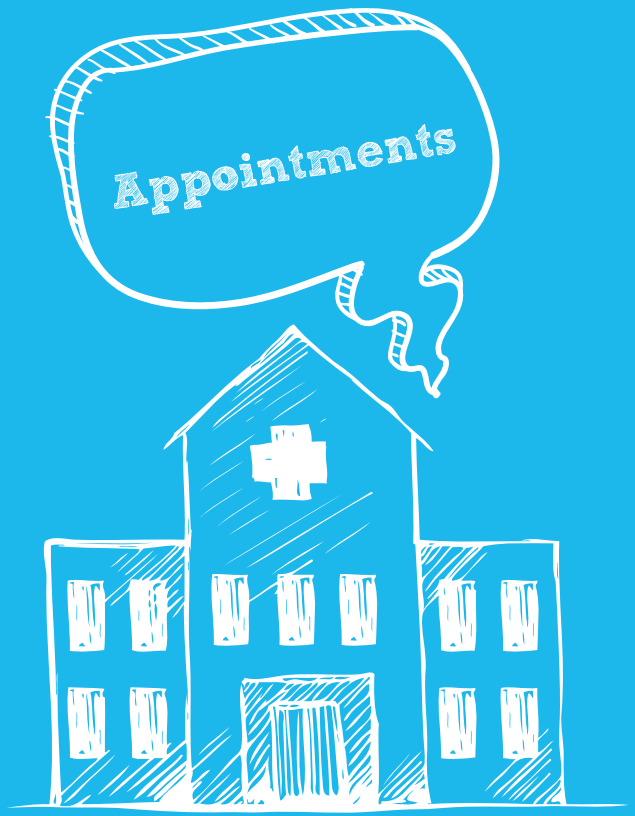
Usual care group
(up to 5 years)

- You will receive your usual care as directed by your healthcare team.
- A caregiver (your study partner) should be available to answer some questions by telephone about every 6 months during this time.



Treatment group
(up to 5 years)

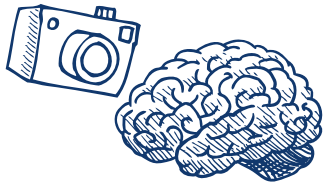
- You will receive the study medication for up to 18 months along with your usual care. Study medication is administered every 4 weeks. Administration of study medication after 6 months will be per investigator discretion.
- After you have completed the study medication, you will receive your usual care as directed by your healthcare team.
- A caregiver (your study partner) should be available to answer some questions by telephone about every 6 months during this time.



Who can take part?

You may be able to take part if you:

- are 60 years of age or older
- are under care for presumed mild cognitive impairment or mild dementia stage of AD
- have a reliable study partner who you are in frequent contact with (and who will be available by telephone at designated times).



What else do I need to consider?

- The study team will explain the possible benefits and risks of the study.
- You do not have to take part in the study if you do not want to.
- If you choose to take part in the study, you can stop participating at any time.
- Participation in the study treatment group is not the only way to get access to the study medication, but if you are in the study treatment group, treatment will be provided at no cost to you.
- Participation in the study usual care group does not prevent your physician prescribing and you receiving the treatment at a later time if you decide it is right for you.
- A team of doctors and nurses will monitor your health carefully during the study.
- An Institutional Review Board (IRB)/Ethics Committee (EC) has reviewed this study. An IRB/EC protects the rights, safety, and well-being of the participants.