ForPatients
by Roche

Alzheimer's Disease (AD)

A Study of Crenezumab Versus Placebo in Preclinical Presenilin1 (PSEN1) E280A Mutation Carriers to Evaluate Efficacy and Safety in the Treatment of Autosomal-Dominant Alzheimer's Disease (AD), Including a Placebo-Treated Non-Carrier Cohort

Trial Status	Trial Runs In	Trial Identifier
Completed	1 Countries	NCT01998841 GN28352

The source of the below information is the publicly available website ClinicalTrials.gov. It has been summarised and edited into simpler language.

Trial Summary:

This study consists of 2 periods: [1] Study Period A - evaluating the efficacy and safety of Crenezumab versus Placebo in participants who carry the PSEN1 E280A autosomal-dominant mutation and do not meet the criteria for mild cognitive impairment due to AD or dementia due to AD and are thus, in a preclinical phase of AD. Participants will be randomised in a 1:1 ratio to receive either Crenezumab or Placebo subcutaneously (every 2 weeks) or intravenously (every 4 weeks) for at least 260 weeks. A cohort of participants (non-mutation carriers) will also be enrolled and will be dosed solely on Placebo and [2] Study Period B - Participants will be offered the opportunity to continue to receive study drug until the results of the study are known and post trial access to Crenezumab is started or development of Crenezumab is discontinued.

Genentech, Inc. Sponsor	**Phase 2** Phase

NCT01998841 GN28352
Trial Identifiers

Eligibility Criteria:

Gender	Age	Healthy Volunteers
All	>=30 Years & <= 60 Years	No