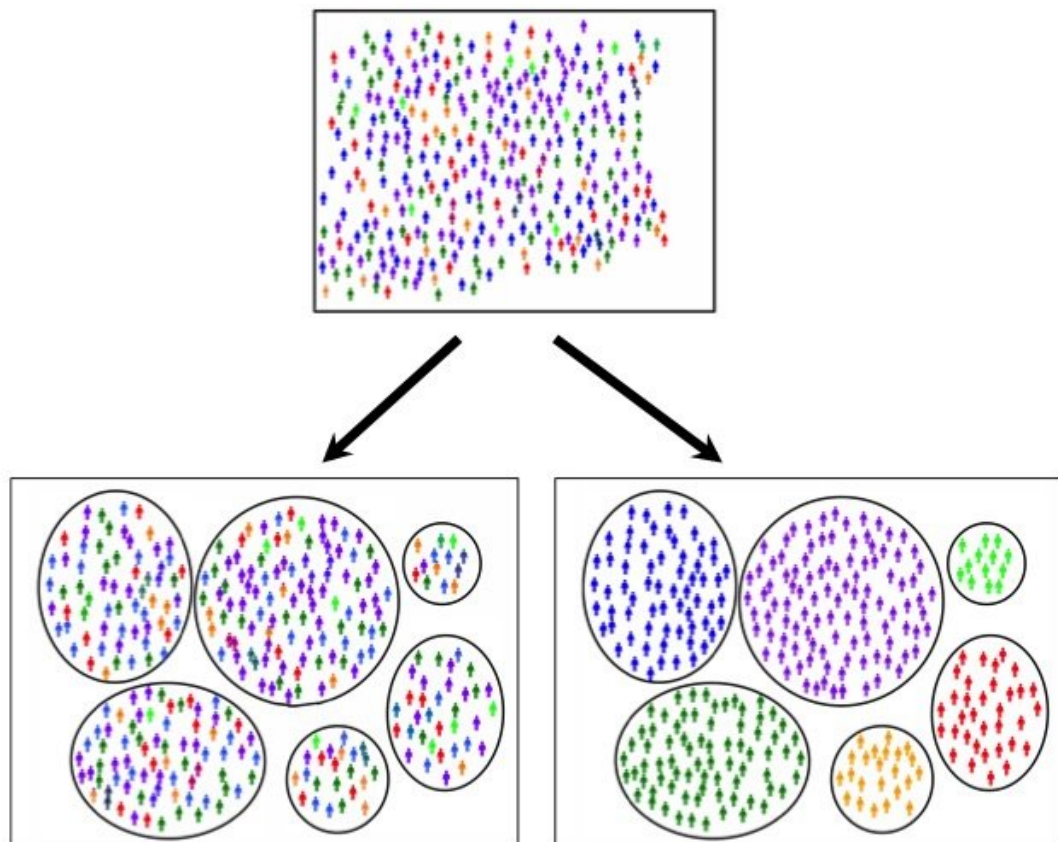


# Researchers classify Alzheimer's patients in six subgroups

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Genetic data supported the contention that a particular way of sorting people resulted in biologically coherent subgroups (pictured lower right). Credit: University of Washington School of Medicine

Researchers studying Alzheimer's disease have created an approach to classify patients with Alzheimer's disease, a finding that may open the door for personalized treatments.

"Alzheimer's, like breast cancer, is not one disease," said lead author Shubhabrata Mukherjee, research assistant professor in general internal [medicine](#) at the University of Washington School of Medicine. "I think a good drug might fail in a clinical trial because not all the subjects have the same kind of Alzheimer's."

This study, published in the recent issue of *Molecular Psychiatry*, involves 19 researchers from several institutions, including Boston University School of Medicine, the VA Puget Sound Health Care System and Indiana University School of Medicine.

The researchers put 4,050 people with late-onset Alzheimer's disease into six groups based on their cognitive functioning at the time of diagnosis and then used genetic data to find biological differences across these groups.

"The implications are exciting," said corresponding author Paul Crane, professor of [general internal medicine](#) at the University of Washington School of Medicine. "We have found substantial biological differences among cognitively defined subgroups of Alzheimer's patients."

Identification of cognitive subgroups related to [genetic differences](#) is an important step toward developing a precision medicine approach for Alzheimer's disease.

The participants received cognitive scores in four domains: memory, executive functioning, language, and visuospatial functioning.

The largest group (39%) had scores in all four domains that were fairly

close to each other. The next largest group (27%) had memory scores substantially lower than their other scores. Smaller groups had language scores substantially lower than their other scores (13%), visuospatial functioning scores substantially lower than their other scores (12%), and executive functioning scores substantially lower than their other scores (3%). There were 6% who had two domains that were substantially lower than their other scores.

The participants came from five studies, and it took more than two years to standardize the neuropsychological test scores across all the studies in order to detect meaningful patterns. The mean age was 80, 92 percent self-reported white race, and 61 percent were female.

The investigators used genome-wide [genetic data](#) to find out if the subgroups are biologically distinct.

Investigators found 33 [single nucleotide polymorphisms](#) (SNPs) - specific locations throughout the genome—where the genetic association was very strong for one of the subgroups. These genetic relationships were stronger than the strongest effects found by an earlier and much larger international consortium study where Alzheimer's disease was treated as a single homogeneous condition.

Several years ago, the International Genomics of Alzheimer's Project Consortium published the largest genome-wide association study of Alzheimer's disease and found about 20 SNPs associated with Alzheimer's disease risk.

This study found 33 additional SNPs with even stronger relationships with a single subgroup.

The study also found a particularly strong relationship between a particular variant of the APOE gene and risk for the memory subgroup.

The APOE e4 allele is a very strong risk factor for developing Alzheimer's disease for people with European ancestry, and it also appears to influence which cognitive subtype of Alzheimer's a person is likely to develop.

People can currently find out if they have an APOE e4 allele with direct-to-consumer testing; however, the researchers note that many people with an APOE e4 allele never develop Alzheimer's disease, and many who don't carry any known genetic risk factor nevertheless end up with the condition.

While world leaders want to find a cure for Alzheimer's by 2025, so far no one has been able to develop an effective treatment let alone a cure. But this study suggests that thinking of Alzheimer's [disease](#) as six distinct conditions may provide a way forward.

"This study is not the end, it's a start," said Mukherjee.

**More information:** et al, Genetic data and cognitively defined late-onset Alzheimer's disease subgroups, *Molecular Psychiatry* (2018). [DOI: 10.1038/s41380-018-0298-8](#)

Provided by University of Washington

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