

# Promising new methods for early detection of Alzheimer's disease

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New methods to examine the brain and spinal fluid heighten the chance of early diagnosis of Alzheimer's disease. Results from a large European study, led by researchers at the Karolinska Institutet, are now published in the medical journal *Brain*. These findings may have important implications for early detection of the disease, the choice of drug treatment and the inclusion of patients in clinical trials.

Despite many years of intensive research, no effective treatment currently exists for Alzheimer's disease, which is the most common form of dementia. It has become increasingly clear that, if the disease is to be treated successfully, it must be detected early, perhaps even before symptoms are evident. Thus, there is a great need for reliable diagnostic methods so that treatment to slow or prevent the disease can begin as early as possible.

A characteristic, pathological sign of Alzheimer's disease is the formation of insoluble amyloid plaques that accumulate in the brain. The presence of these plaques can be measured in the brain using [positron emission tomography](#) (PET camera) to visualize radioactive tracer molecules that bind to the amyloid plaques. Amyloid levels can also be measured in [spinal fluid](#). While amyloid accumulates in the brain in Alzheimer's disease, research has shown that levels of amyloid in the spinal fluid is instead reduced.

In the current study, researchers compared the amyloid-PET measurements in the brain with amyloid- $\beta$ 42 in the spinal fluid to see how well they align. The investigations were performed at seven European memory clinics on 230 patients who were examined for memory disorders. Patients received various diagnoses, such as mild cognitive impairment (MCI), Alzheimer's disease and various types of dementia. A small group of healthy individuals were control subjects.

PET studies on the patients' brains were evaluated both locally at the seven different hospitals and centrally at the Karolinska Institutet. The researchers also used a new method, called the centiloid method, in order to standardize the measured amyloid values on a unified scale so they can be compared. Levels of amyloid42 were measured from the samples of cerebrospinal fluid at each local hospital. All samples were also analyzed centrally at Sahlgrenska Hospital in Gothenburg, where the levels of amyloid42, as well as a second [amyloid protein](#) amyloid40, were measured.

The researchers found that the best fit was achieved when the amyloid level in the brain was compared with the ratio between the levels of amyloid42 and amyloid40 in the spinal fluid. Given this finding, the research team hypothesized that the forms of  $\beta$ -amyloid found in the brain and spinal fluid are not completely identical.

"Interestingly, there was also a difference between the values measured in the [brain](#) and spinal fluid in a smaller group of patients. This may justify that, in some unclear cases, the diagnosis should include an amyloid PET scan to complement the [cerebrospinal fluid](#) sample. These findings are also important because it is increasingly common to perform amyloid-PET studies upon the inclusion of [patients](#) in new drug studies", said the study's coordinator Agneta Nordberg.

**More information:** "Pittsburgh Compound B imaging and cerebrospinal fluid amyloid- $\beta$  in a multicentre European Memory Clinic study"; *Brain*, published online 8 July 2106, [DOI: 10.1093/brain/aww160](#)

Provided by Karolinska Institutet

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