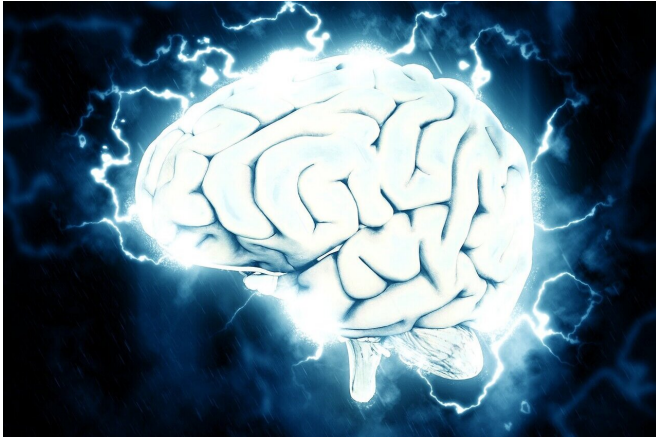


Many patients with iNPH develop Alzheimer's disease, too

15 November 2019



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Up to one in five patients treated for idiopathic normal pressure hydrocephalus, iNPH, also develop Alzheimer's disease, according to a new study from the University of Eastern Finland and Kuopio University Hospital. The researchers were able to predict the development of Alzheimer's disease by using the Disease State Index, DSI, that combines patient-specific data from various sources. The results were published in *Journal of Alzheimer's Disease*.

In iNPH, the circulation of cerebrospinal fluid (CSF) is disturbed for an unknown reason, leading to a slightly elevated brain pressure and dilation of the brain ventricles. Symptoms of NPH include gait deviations, impaired [short-term memory](#) and urinary incontinence. Patients with iNPH often have changes in brain that are related to Alzheimer's [disease](#).

The study followed patients with iNPH after they had received treatment for their disease. They were treated with shunt surgery, in which excessive CSF is drained from the brain ventricles to the [abdominal cavity](#) by using a CSF shunt.

Shunted NPH patients who had undergone brain tissue biopsy in connection with their surgery were selected for the study. The objective of the biopsy was to detect changes that are indicative of Alzheimer's disease.

During the follow-up, the researchers found that up to one in five patients with NPH were later diagnosed with Alzheimer's disease. At the end of the follow-up, patients with NPH were more frequently diagnosed with Alzheimer's disease than the general population. The researchers were able to predict the development of Alzheimer's disease, with moderate accuracy, by using patient-specific DSI. The data used in DSI included the patient's pre-surgery symptom profile, brain tissue samples and brain MRI images.

More information: Antti J. Luikku et al, Predicting Development of Alzheimer's Disease in Patients with Shunted Idiopathic Normal Pressure Hydrocephalus, *Journal of Alzheimer's Disease* (2019). [DOI: 10.3233/JAD-190334](https://doi.org/10.3233/JAD-190334)

Provided by University of Eastern Finland

APA citation: Many patients with iNPH develop Alzheimer's disease, too (2019, November 15) retrieved 13 April 2021 from <https://medicalxpress.com/news/2019-11-patients-inph-alzheimer-disease.html>

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