

# Amyloid Positivity in the Alzheimer/Subcortical-Vascular Spectrum

Sung Hoon Kang, MD, Monica Eunseo Kim, MS, Hyemin Jang, MD, PhD, et al.

Cite as: *Neurology*® 2021;96:e2201-e2211. doi:10.1212/WNL.0000000000011833

## Correspondence

Dr. Na  
dukna@naver.com  
or Dr. Jang  
hmjang57@gmail.com

## Study Question

What is the frequency of A $\beta$  positivity in a large sample of individuals classified into 9 groups based on the combination of different levels of cognitive impairment and WMH burden? What is the association between A $\beta$  positivity and other features of cerebral small vessel disease burden, including lacunes and microbleeds, in 3 stratified cognitive levels?

## What Is Known and What This Paper Adds

Studies of A $\beta$ -positivity often dichotomize participants into those with Alzheimer disease (AD) and those with subcortical vascular cognitive impairment. This investigation's results offer a more fine-grained understanding of how A $\beta$ -positivity rates vary with severity of WMH and cognitive change and provide evidence that AD and cerebral small vessel disease lie on a continuum.

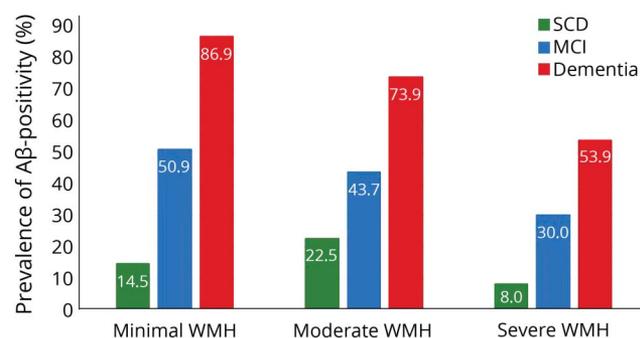
## Methods

For this cross-sectional study, the investigators analyzed data from individuals with subjective cognitive decline (SCD; n = 294), mild cognitive impairment (MCI; n = 237), or dementia (n = 516) recruited through a memory clinic at the Samsung Medical Center in Seoul. The participants underwent A $\beta$  PET scans and MRI scans. WMH burden was classified as minimal, moderate, or severe. The participants were classified into 9 groups based on the 3 cognition levels and 3 WMH burden levels. Linear trend tests were used to examine how A $\beta$ -positivity rates varied with WMH severity in participants with the same cognitive state and how they varied with cognition states in participants with the same WMH severity level.

## Results and Study Limitations

The A $\beta$ -positivity rates were higher for participants with more severe cognitive impairment (i.e., 15.7%, 43.5%, and 76.2% for the SCD, MCI, and dementia groups, respectively) but

**Figure** A $\beta$ -Positivity Rates and WMH Burdens



A $\beta$ -positivity rates for participants with 9 groups based on different cognition level and different WMH severities.

were lower for participants with greater WMH severities (i.e., 54.5%, 53.9%, and 41.0% for the minimal, moderate, and severe groups, respectively) or greater number of lacunes (i.e., 59.0%, 42.0%, and 23.4% for the 0, 1–3, and >3 groups, respectively). The present study's limitations include its cross-sectional nature, which precludes the establishment of causal relationships; the lack of pathology data; and a focus on Korean participants, which may limit generalizability to people from other ethnic groups.

## Study Funding and Competing Interests

This study was funded by the National Research Foundation of Korea (NRF) grant funded by the Korea government (grant number: NRF-2020R1A2C1009778); and by a grant of the Korea Health Technology R&D Project through the Korea Health Industry Development Institute (KHIDI) funded by the Ministry of Health & Welfare, Republic of Korea (grant number: HI19C1132). The authors report no competing interests. Go to [Neurology.org/N](https://www.neurology.org/N) for full disclosures.

*A draft of the short-form article was written by M. Dalefield, a writer with Editage, a division of Cactus Communications. The corresponding author(s) of the full-length article and the journal editors edited and approved the final version.*

# Neurology<sup>®</sup>

## Amyloid Positivity in the Alzheimer/Subcortical-Vascular Spectrum

Sung Hoon Kang, Monica Eunseo Kim, Hyemin Jang, et al.

*Neurology* 2021;96:e2201-e2211 Published Online before print March 15, 2021

DOI 10.1212/WNL.0000000000011833

**This information is current as of March 15, 2021**

<b>Updated Information &amp; Services</b>	including high resolution figures, can be found at: <a href="http://n.neurology.org/content/96/17/e2201.full">http://n.neurology.org/content/96/17/e2201.full</a>
<b>References</b>	This article cites 48 articles, 11 of which you can access for free at: <a href="http://n.neurology.org/content/96/17/e2201.full#ref-list-1">http://n.neurology.org/content/96/17/e2201.full#ref-list-1</a>
<b>Permissions &amp; Licensing</b>	Information about reproducing this article in parts (figures, tables) or in its entirety can be found online at: <a href="http://www.neurology.org/about/about_the_journal#permissions">http://www.neurology.org/about/about_the_journal#permissions</a>
<b>Reprints</b>	Information about ordering reprints can be found online: <a href="http://n.neurology.org/subscribers/advertise">http://n.neurology.org/subscribers/advertise</a>

*Neurology*® is the official journal of the American Academy of Neurology. Published continuously since 1951, it is now a weekly with 48 issues per year. Copyright © 2021 American Academy of Neurology. All rights reserved. Print ISSN: 0028-3878. Online ISSN: 1526-632X.

