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## Stability in BMI over time is associated with a better cognitive trajectory in older adults

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### Abstract

**Objective:** Evidence on simultaneous changes in body mass index (BMI) and cognitive decline, which better reflect the natural course of both health phenomena, is limited.

**Methods:** We capitalized on longitudinal data from 15,977 initially non-demented elderly from the Alzheimer's Disease Centers followed for 5 years on average. Changes in BMI were defined as (1) last minus first BMI, (2) mean of all follow-up BMIs minus first BMI, and (3) standard deviation of BMI change from baseline and all follow-up visits (representing variability).

**Results:** Participants with significant changes in BMI (increase or decrease of 5%), or who had greater variability in BMI, had faster cognitive decline. This pattern was consistent irrespective of normal (BMI <25; N = 5747), overweight (25 ≤ BMI <30; N = 6302), or obese (BMI ≥ 30; N = 3928) BMI at baseline.

**Conclusions:** Stability in BMI predicts better cognitive trajectories suggesting clinical value in tracking BMI change, which is simple to measure, and may point to individuals whose cognition is declining.

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#### CONFLICTS OF INTEREST

All other authors have nothing to disclose.

#### SUPPORTING INFORMATION

Additional supporting information may be found in the online version of the article at the publisher's website.

## Keywords

Cognitive decline; body mass index; older adults; stability

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## 1 INTRODUCTION

Progressive loss of cognition is common in older adults and its prevention and treatment is a major public health priority. The aging research community has successfully identified diverse risk factors for late-life cognitive impairment. Consistent evidence associates body mass index (BMI), the most common measure of global adiposity, in midlife, with poor cognitive outcomes in late life.<sup>1–3</sup> Individuals who were obese in midlife have a 2- to 3-fold higher dementia risk in old age compared to individuals who were non-obese.<sup>4</sup> We and others showed more cognitive impairment and dementia with greater BMI in old age,<sup>5,6</sup> but some have found no associations, or even lower dementia risk in obese older adults.<sup>2,4,7</sup> In addition, losing weight has also been associated with higher dementia risk.<sup>8</sup>

Despite the relevance of BMI to the risk of incident cognitive impairment and the subtle but consistent decrease in BMI that accompanies old age,<sup>9</sup> there is limited evidence on the simultaneous changes in BMI and cognitive decline,<sup>10,11</sup> which better reflect the natural course of both health phenomena. There is evidence indicating that BMI stability over time is associated with less frailty, disability, and mortality in late life.<sup>12,13</sup> However, whether BMI stability is associated with better cognitive trajectories is essentially unknown. To address this gap in knowledge, we exploited longitudinal clinical data from ≈16,000 initially non-demented older adults participating in the Alzheimer's Disease (AD) Centers (ADCs) across the United States to examine associations of changes in BMI over time with changes in cognition, investigate whether both increases and decreases of BMI are related to cognitive decline, and if these associations are consistent in those who have normal, overweight or obese BMI at baseline.

## 2 | METHODS

### 2.1 | Data source and sample derivation

Data are drawn from the National Alzheimer's Coordinating Center (NACC), which maintains the Uniform Data Set (UDS) of participants enrolled in ADCs funded by the National Institute on Aging (NIA).<sup>14–17</sup> The study included all participants who were enrolled in NACC-UDS between September 2005 (UDS start date) and the December 2019 data freeze (N = 42,022). To be eligible for the current study, at baseline, participants had to be ≥60 years of age, without a diagnosis of dementia as determined by clinician consensus, have a Clinical Dementia Rating (CDR) of 0 or 0.5, and have at least one annual follow-up visit. The sample selection steps are presented in Figure 1. Recruitment methods vary between ADCs; therefore NACC-UDS should not be considered a population-based sample. Written informed consent was provided by all participants and their informants and approved by local institutional review boards (IRBs). Research using the NACC database was approved by the University of Washington IRB.

## 2.2 | Measures

**2.2.1 | Global cognition and cognitive domains**—All participants were administered a standardized neuropsychological battery at each study visit. We categorized the neuropsychological tests into the following domains<sup>18</sup>: memory (Wechsler Memory Scale - Revised [WMS]-R Logical Memory Immediate and Delayed Recall), attention/working memory (WMS-R Digit Span forward and backward, Trail Making Test [TMT] Part A), language (category fluency [mean of animals and vegetables], Short Form of the Boston Naming Test [BNT]), and executive function (TMT Part B).<sup>18</sup> For all neuropsychological tests except TMT A and B, higher scores indicate better cognition. TMT A and B scores were reversed so that higher scores also indicate better cognition. Cognitive composite scores were computed using previously published methods.<sup>19</sup> First, a baseline z-score for each test was computed. A baseline composite domain was constructed by averaging the z-scores for each test in that domain, and a baseline global cognitive z-score was calculated by averaging the z-scores from all tests. Missing values for individual tests were allowed. However, the domain score was set to missing if more than half of the tests in that domain were missing. The global score was set to missing if more than half of all tests were missing. Requiring complete data on all tests did not change study results. Mean and standard deviation (SD) from the baseline scores were used to construct corresponding z-scores for each domain and global cognition.

**2.2.2 BMI**—BMI was calculated based on measured weight and height (weight in kilograms [kg] divided by squared height in centimeters) at baseline and at each follow-up visit. For descriptive analyses, each participant's baseline BMI ( $BMI_{bl}$ ) was categorized into three groups based on the World Health Organization (WHO) classification<sup>20</sup> of underweight or normal ( $<25 \text{ kg/m}^2$ ), overweight ( $25 \leq BMI <30 \text{ kg/m}^2$ ), and obese ( $\geq 30 \text{ kg/m}^2$ ). Few participants were underweight ( $<18.5 \text{ kg/m}^2$ ) and therefore were combined with the normal weight group. BMI outside the range of 15 to  $50 \text{ kg/m}^2$  were treated as biologically unlikely and were dropped.

**2.2.3 Change in BMI**—Change in BMI was measured in three ways: (1) difference between BMI at last visit and  $BMI_{bl}$  ( $BMI_{last-bl}$ ) as commonly used in clinical trials for weight loss,<sup>21,22</sup> (2) difference between mean BMI across all follow-up visits and  $BMI_{bl}$  ( $BMI_{mean-bl}$ ). For (1) and (2), changes in BMI were categorized as significant increase ( $\geq 5\%$  increase), stable ( $<5\%$  change), or significant decrease in BMI ( $\geq 5\%$  decrease), as a 5% change is defined as clinically significant by the U.S. Food and Drug Administration (FDA).<sup>23</sup> Finally, (3) we also constructed a variable for within-person variability, that is, SD of BMI change from baseline and all follow-up visits ( $BMI-SD$ ), categorized into quartiles<sup>1</sup>

**2.2.4 Demographics and clinical characteristics**—Demographic characteristics included age, sex, race (White, Black, vs. Other), ethnicity (Hispanic/Latino vs. Other), years of education, and years of follow-up. Cardiovascular risk factors included history of diabetes, hypertension, and hypercholesterolemia, ascertained by self-report or clinician assessment. Baseline diagnosis included cognitively normal (CN) and mild cognitive impairment (MCI). Participants' function was measured using the CDR scale,<sup>24,25</sup> based on

reports by the participant and study partner, and the Functional Assessment Questionnaire (FAQ) reported from study partners interviews.<sup>26,27</sup> Depressive symptoms were measured using the 15-item Geriatric Depression Scale (GDS-15).<sup>28,29</sup> Number of medications a participant is taking was used as a proxy for overall health status.<sup>30</sup> Apolipoprotein E (*APOE*) genotype for participants who are willing to provide samples was reported by the ADCs. We constructed an indicator variable for any *APOE*  $\epsilon$ 4 allele and an indicator for missing *APOE* information. Except for *APOE*, missing values were relatively rare (from no missing values for age, sex, education CDR, and diagnosis to a maximum of 3.3% missing values for the FAQ). Because results were unchanged when we applied mean substitution to missing values, observations with missing values were dropped in the present analyses.

**2.2.5 | Statistical analyses**—Baseline characteristics were reported as N (percent) or mean (SD) as appropriate by BMI<sub>bl</sub> groups. Our primary outcome measure was change in global cognition from baseline ( $\Delta$  zcog). The main independent variables in the analyses were time, change in the respective BMI measures ( $\Delta$  BMI<sub>last-bl</sub>,  $\Delta$  BMI<sub>mean-bl</sub>, and  $\Delta$  BMI-SD), and their interaction with time. All models were adjusted for covariates that have been associated with cognitive decline and dementia including age, sex, race/ethnicity, years of education, years of follow-up, baseline diagnosis (CN vs. MCI), FAQ, GDS, *APOE* genotype, number of medications (reflecting co-morbidities), and indicators for history of diabetes, hypertension, and hypercholesterolemia. All models were estimated using linear mixed models (LMMs) and included participant level random intercept and linear slope to allow participants to differ in their overall rate of change over time. A random effect for ADC was also included. Changes in the cognitive domain of memory, attention/working memory, language, and executive functions were examined separately. Finally, because BMI changes in old age differ in men and women,<sup>31,32</sup> we tested the interaction of each of the BMI measures with sex on cognitive decline. All analyses were performed using Stata 13.0.<sup>33</sup>

### 3 | RESULTS

#### 3.1 | Baseline sample characteristics

Table 1 describes the characteristics of the sample at baseline. Participants were  $\approx$ 74 years of age, included a majority of women (59%) and White participants (81%), and had an average of  $15.6 \pm 3.6$  years of education; 15.8% were MCI, with average Mini-Mental Status Exam (MMSE) score of  $28.4 \pm 2.0$ , consistent with a sample that does not have dementia baseline. FAQ was  $1.2 \pm 2.8$ , reflecting little impairment in activities of daily living; GDS was  $1.6 \pm 2.2$ , indicating relatively little clinical depression. About 13% had diabetes, 52% had hypertension, and 53% had high cholesterol. Of those who agreed to have *APOE* genotyping (83% of the sample), 29% had at least one *APOE*  $\epsilon$ 4 allele. Between-group differences were statistically significant at  $P < .01$  for all variables except for proportion of participants with missing *APOE* status ( $P = .37$ ), years of follow-up ( $P = .40$ ), CDR score ( $P = .10$ ), and FAQ score ( $P = .11$ ). Finally, on average, participants had  $5.2 \pm 3.1$  in-person visits (initial visit + 4 annual follow-up visits) and there was no difference in attrition rate between baseline BMI groups.

Thirty-six percent of the participants had normal weight, 39.4% were overweight, and 24.6% were obese, which is consistent with the reported rates of overweight and obesity in this age group in the general US population.<sup>34</sup> As shown in Table 2, 26.7% of the sample decreased from the first to the last BMI measure by 5% or more (in initially normal, overweight, and obese group, 20.9%, 27.1%, and 34.6%, respectively), 55.8% remained stable in BMI over time (<5% BMI change; in initially normal, overweight, and obese group, 57.7%, 56.7%, and 51.8%, respectively), and 17.5% increased 5% or more in BMI (in initially normal, overweight, and obese group, 21.5%, 16.3%, and 13.6%, respectively).

### 3.2 Associations of changes in BMI with changes in global cognition

For descriptive purposes, unadjusted means (SD) of change in global cognition and in specific cognitive domain scores for each of the three BMI change measures are shown in Table S1. There were significant group differences in change in global cognition and in the specific cognitive domains (all  $P < .01$ ) for each of the BMI change measures such that the stable group/s had less cognitive decline.

LMMs were applied to test the associations of  $z_{cog}$  with the three BMI change measures and are presented in Table 3. Greater changes in  $BMI_{last-bl}$  and in  $BMI_{mean-bl}$  (both increases and decreases of 5% or more in BMI) were associated with a faster rate of cognitive decline (Table 3; Figure 2A and 2B, respectively). Similarly, the interaction of  $BMI-SD$  (categorized into quartiles) with time on global cognition was significant—greater SD (representing greater changes in BMI over time, both increase and decrease)—was associated with a faster rate of global cognitive decline (Table 3; Figure 2C).

To exemplify the differences in rate of decline according to the estimations for  $BMI_{last-bl}$ , we calculated the proportion of cognitive decline in 5% change in BMI compared to the stable BMI group. The predicted decline in the global cognition z-score for the stable group averaged 0.16 (see Table S1) over the period of the study. In contrast, in the group with 5% decrease in BMI, the predicted decline in the global cognition z-score was 0.27 and 0.26 points for those with a 5% increase in BMI. This represents a 64% and 62%, respectively, faster decline in global cognition in the groups with changes in BMI compared to the stable group.

### 3.3 | Associations of changes in BMI with changes in specific cognitive domains

We repeated analyses using the three BMI change variables for the specific cognitive domains (Table 3). Both declines and increases of 5% or more from first to last BMI, and from mean BMI to first BMI, compared to stable BMI (<5% change) were associated with faster decline in all cognitive domains; for executive functions, those whose BMI increased did not reach statistical significance. When comparing the highest quartiles of SD to the lowest SD quartile, the second quartile was associated only with a faster rate of episodic memory decline. The third and fourth quartiles, representing greater variability, were associated with a faster rate of decline in all cognitive domains (Table 3).

### 3.4 Associations of BMI changes with cognitive decline stratified by baseline BMI

To explore whether the pattern of association of greater BMI change over time with faster cognitive decline is specific to a particular baseline BMI group (for example, participants who were obese), we repeated all analyses stratifying by baseline BMI group. Overall, results remained the same, that is, in each of the BMI groups, greater changes (either increases or decreases) of BMI over time were associated with a faster rate of cognitive decline (Table S2 and Figure S1–S3).

We performed a sensitivity analysis using cut-off points of increases and decreases of 10% or more in BMI for  $BMI_{last-bl}$  and  $BMI_{mean-bl}$ . Results were essentially unchanged (Table S3). Finally, none of the interactions of the BMI measures with sex on cognitive decline were significant.

## 4 DISCUSSION

This study followed  $\approx 16,000$  well-characterized community-dwelling older adults without dementia at baseline for an average of 5 years and provides new evidence that greater increases, decreases, or variability in BMI over time are associated with an accelerated rate of cognitive decline after adjusting for a broad range of potential sociodemographic, clinical, and functional confounders. Furthermore, this pattern is consistent, irrespective of whether a person has normal, overweight, or obese BMI at baseline. As shown in Figure 2, all groups had some degree of decline including the group with stable BMI. However, the rate of decline in the non-stable groups was over 60% faster than the rate of decline in the stable BMI group. These results suggest a potential clinical value in tracking BMI in annual medical visits, which is simple, non-expensive, non-invasive, and quick to measure, as it may point to individuals whose cognition is declining and allow for early intervention. Further work is needed to identify the biological causes underlying departures from stable BMI in old age and their contribution to cognitive decline.

The current study extends prior work in several important ways. First, we assessed changes in BMI over time, measured in several different ways, and their associations with cognitive decline. Second, the large sample size and the relatively long follow-up permitted investigation of the association of both increase and decrease in BMI with cognitive decline. An increase and decrease of 5% or more in BMI was used, as it is well-accepted as clinically significant.<sup>23</sup> Third, because obese individuals have a greater range for BMI decline while normal-weight individuals have a greater range for BMI increase, we examined whether the associations of changes in BMI and cognitive decline differ by baseline BMI category (based on the World Health Organization [WHO] BMI categorization), and showed that these associations are consistent regardless of BMI category. These results indicate that in old age, the stability of BMI may confer protection against cognitive decline, irrespective of the initial BMI of the individual. Finally, analyses of the individual cognitive abilities used to construct global cognition, showed that changes in BMI over time were associated significantly with accelerated decline in episodic memory, attention/working memory, and language but not with executive function, suggesting that changes in BMI may be associated differentially with some dimensions of cognitive abilities more than others.



We and others have shown that the negative effects of a higher BMI on brain and cognition persist through late-life.<sup>32,35</sup> However, there is scarce research on changes in BMI in old age; most studies provide evidence suggesting that decreasing BMI is associated with poor health outcomes including congestive heart failure and mortality.<sup>36,37</sup> Our study showed that a decline in BMI over time was associated with accelerated cognitive decline. Consistent with our results, a 1-unit annual decline in BMI was associated with about a 35% increase in the risk of AD compared with a person experiencing no change in BMI in older adults,<sup>10</sup> suggesting that loss of BMI may reflect pathologic processes that contribute to the subsequent development of AD. Furthermore, weight loss during 1 year was associated with a faster clinical progression of MCI.<sup>11</sup> Although decline in BMI may represent a true risk factor for cognitive decline, it is also possible that it represents reverse causality, that is, that brain changes related to dementia affect BMI earlier than cognition.<sup>32</sup> Of interest, polymorphisms in *FTO*, *PPARG*, *PPARA*, and *APOE* genes,<sup>38</sup> implicated in AD,<sup>39,40</sup> are significant predictors of BMI variability,<sup>41</sup> suggesting a common cause. An alternative explanation is that both BMI variability and cognitive decline are manifestations of an underlying morbidity and are not causally related.

It is important to note that in studies examining associations of change in BMI with health outcomes, including cognitive impairment, the comparison is typically to individuals who had a stable BMI. Because in our study we showed that both decrease and increase in BMI over time were associated with accelerated cognitive decline, irrespective of the BMI at baseline, it is plausible that conceptually, in old age, BMI stability over time is associated with reduced risk of poor cognitive outcomes. The mechanisms underlying the potential health benefits of BMI stability are unknown. Aging leads to numerous physiological, endocrine, and metabolic changes, which manifest in changes in body composition, reflected in loss of bone, loss of muscle mass and strength, and increased body fat and fat redistribution.<sup>9,42,43</sup> These changes are accompanied by an increased low-grade chronic inflammation,<sup>44–46</sup> which has been consistently associated with poor health outcomes<sup>47,48</sup> and with poor cognitive outcomes and incident dementia.<sup>49,50</sup> Indeed, BMI stability over time is associated with fewer comorbidities in late life.<sup>12,13</sup> Our results suggest that BMI stability is also associated with a healthier cognitive trajectory.

Biological homeostasis is the ability of the “normal” body to maintain equilibrium, balancing multiple factors—including blood pressure, cholesterol, as well as body weight—at stable levels despite temporary changes in life conditions.<sup>51</sup> The progressive homeostatic imbalance that occurs with aging<sup>52–54</sup> may be at least in part responsible for the change in BMI observed among older adults,<sup>55,56</sup> and facilitates deleterious processes that cause a myriad of cellular dysfunctions, including neuronal death and neurodegeneration.<sup>52,53</sup> Stability in BMI over time may represent less departure from homeostasis, possibly conferring the preservation of neuronal health. Of interest, in the stratified analysis, among those who had initially normal BMI, the rate of decline in cognition was nominally faster than among participants with baseline overweight or obesity, despite the lack of difference in numerous baseline medical and sociodemographic characteristics. We speculate that this may point to an adaptive process of the overweight and obese individuals to their higher BMI, reaching a new homeostatic balance,<sup>57–59</sup> and conferring a degree of protection against cognitive decline.

The study has limitations. The NACC data set is based on a convenience sample of participants approaching the ADCs in the United States, rather than a community-based sample. Furthermore, the sample has 80% White participants with underrepresentation of ethnic minorities. Thus caution must be applied when generalizing the results to the whole population of older adults and the results should be replicated in more diverse cohorts. The cohort includes significant majority of women, which in some studies have shown different trajectories of BMI than men. However, the interaction of BMI changes with sex was tested, and sex did not modify the associations we have found. The cardiovascular risk factors are self-reported rather than directly measured. The prevalence of diabetes and hypertension in our sample is less than that reported in national statistics (Table 1. 12.8% vs. 26.8% for diabetes<sup>60</sup> and 52.1% vs. 70% for hypertension<sup>61</sup>), whereas hyperlipidemia percentage was close to national statistics (53.4% vs. 48%<sup>62</sup>). Overall there is likelihood that some of the confounding factors were underreported in our sample. Our results remained robust after adjusting for cardiovascular risk factors and diseases, and for depression and Activities of Daily Living (ADL), but other factors such as cancer may have affected BMI. Increased adiposity has been consistently associated with higher levels of chronic inflammation markers, which in turn have been related to AD.<sup>63</sup> However, inflammatory markers are not available in the NACC data set. Because the study investigated changes in BMI over time, which may be related to increasing morbidity, we repeated the analyses replacing baseline number of medications by changes in number of medications at last follow-up compared to first follow-up, such that a greater number represented an increase in number of medications, suggesting growing morbidity. However, adjusting for this covariate did not change any of the results (data not shown). It is important to note that BMI may not be an optimal measure of adiposity in older adults due to redistribution of body fat with age,<sup>64–67</sup> indicating the need for more direct measures of adiposity in studies of older adults, such as regional fat measures, and fat-related secreted factors. Finally, overall, the rates of cognitive decline found in this study are relatively small, possibly because participants with dementia at baseline were excluded. Yet, the clinical hallmark of AD is the progressive loss of cognitive function. A long-term extrapolation of a greater decline in cognition of over 60% in the groups with BMI change compared to the group with stable BMI may represent a progressive disease process.

Confidence in our findings is enhanced by several factors. Body weight and height were directly measured (rather than self reported) at every visit. A large number of male and female participants were examined annually for up to 14 years, with structured, validated, and harmonized clinical measures of cognition permitting investigation of global cognition as well as specific cognitive domains.

In conclusion, we found that stability in BMI over time, in contrast to increases or decreases in BMI, is associated with a slower rate of cognitive decline, both in global cognition and specific cognitive domains. Disentangling the biological pathways underlying different trajectories of BMI in old age, and their contribution to brain health and disease, is necessary to develop potential therapies.



## Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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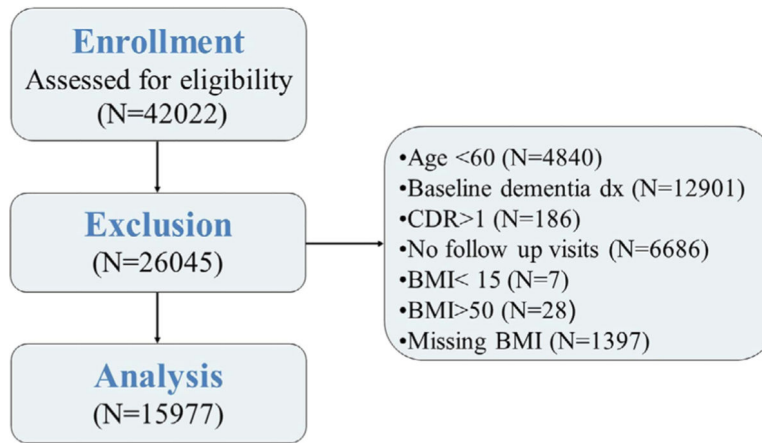
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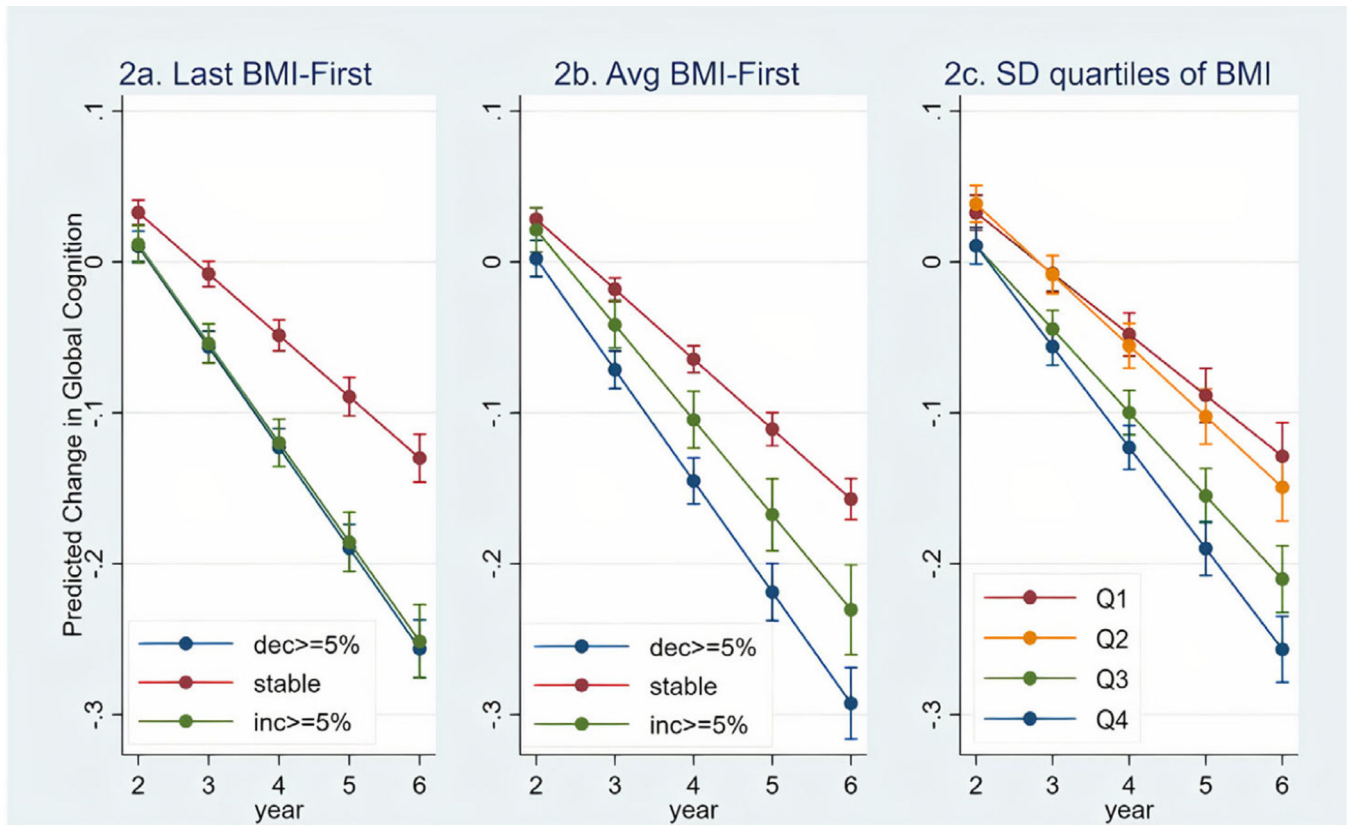
## RESEARCH IN CONTEXT

- 1. Systematic review:** Literature searches using PubMed were conducted to ascertain original and review articles, using keywords including “BMI,” “adiposity,” “trajectories,” and “cognitive decline.” These articles suggested an association of baseline BMI with cognitive decline, with some suggesting that in old age, higher BMI may be a protective factor. PubMed search did not identify studies reporting on the long-term simultaneous changes in BMI and cognitive decline.
- 2. Interpretation:** Our study provides new evidence that greater changes in BMI over time are associated with a faster rate of cognitive decline, suggesting that in initially non-demented older adults, stability in BMI predicts a better cognitive trajectory.
- 3. Future directions:** These results suggest a potential clinical value in tracking BMI change, a simple, non-invasive measure, as it may point to individuals whose cognition is declining. Further work is needed to identify mechanisms underlying departures from stable BMI in old age and their contribution to cognitive decline.



**FIGURE 1.** Sample selection steps. Abbreviation: dx, Diagnosis; CDR, Clinical Dementia Rating Scale; BMI, body mass index.





**FIGURE 2.** Associations of changes in BMI (last minus first BMI, average BMI at follow-up minus first BMI, quartiles of standard deviation) with decline in global cognition. Abbreviation: BMI, Body Mass Index; Avg, Average; SD, standard deviation; decr, decrease; inc, increase.

**TABLE 1**

Baseline characteristics by baseline BMI group

Variables	Total Sample	Normal (BMI < 25 kg/m <sup>2</sup> )	Overweight (25 BMI < 30 kg/m <sup>2</sup> )	Obese (BMI ≥ 30 kg/m <sup>2</sup> )
N (%)	15,977	5,747 (36)	6,302 (39.4)	3,928 (24.6)
Age, mean (SD)	73.5 (7.6)	74.5 (8.0)	73.8 (7.5)	71.7 (6.9)
Female (%)	59.2	66.8	50.6	62.0
Hispanic (%)	6.9	5.0	7.4	9.0
White (%)	80.9	86.1	82.7	70.5
Black (%)	14.7	7.9	13.8	26.0
Other (%)	4.4	6.0	3.5	3.5
Years of education, mean (SD)	15.6 (3.6)	16.1 (3.0)	15.7 (3.2)	14.9 (3.4)
Years of follow-up, mean (SD)	5.2 (3.1)	5.3 (3.1)	5.2 (3.1)	5.1 (3.0)
BMI, mean (SD)	27.2 (5.0)	22.5 (1.8)	27.2 (1.4)	34.1 (3.8)
MMSE (range 0 to 30), mean (SD)	28.4 (2.0)	28.4 (2.0)	28.4 (2.0)	28.3 (2.0)
FAQ (range 0 to 30), mean (SD)	1.2 (2.8)	1.3 (2.9)	1.3 (2.9)	1.2 (2.8)
Hypertension (%)	52.1	39.3	53.4	68.7
Hyperlipidemia (%)	53.4	44.8	56.4	61.3
Diabetes (%)	12.8	6.4	11.6	23.9
GDS (range 0 to 15), mean (SD)	1.6 (2.2)	1.5 (2.1)	1.6 (2.1)	1.8 (2.3)
APOE ε4 (%)	28.8	30.3	28.8	26.5
Number of medications, mean (SD)	5.7 (4.1)	5.1 (5.8)	5.7 (4.1)	6.6 (4.5)
MCI (%)	15.8	16.0	15.3	16.3
CDR at baseline (%)	0	61.7	60.9	64.5
	0.5	38.3	39.1	35.5

Abbreviation: SD, standard deviation; BMI, body mass index; MMSE, Mini Mental State Exam; FAQ, Functional Activities Questionnaire; GDS, Geriatric Depression Scale; APOE, Apolipoprotein E; MCI, mild cognitive impairment; CDR, Clinical Dementia Rating scale.

Change in BMI between baseline and last visit: total sample and by baseline BMI group

**TABLE 2**

	Total sample			Normal (BMI <25 kg/m <sup>2</sup> )			Overweight (25 BMI <30 kg/m <sup>2</sup> )			Obese(BMI ≥30 kg/m <sup>2</sup> )		
	N	%	Mean change (SD)	N	%	Mean change (SD)	N	%	Mean change (SD)	N	%	Mean change (SD)
Significant decrease, change in BMI	4,200	26.7	-3.12 (1.93)	1,179	20.8	-2.23 (1.10)	1,680	27.1	-2.95 (1.59)	1,341	34.6	-4.13 (1.93)
Stable, change in BMI <5%	8,775	55.8	-0.04 (0.72)	3,262	57.7	-0.01 (0.59)	3,510	56.6	-0.03 (0.72)	2,003	51.8	-0.11 (0.90)
Significant increase, change in BMI	2,752	17.5	2.63 (1.51)	1,215	21.5	2.38 (1.43)	1,010	16.3	2.65 (1.50)	527	13.6	3.18 (1.59)

Abbreviation: BMI, body mass index; SD, standard deviation.

**TABLE 3**  
Associations of changes in BMI with changes in global cognition and cognitive domains

Independent variables	Global cognition		Episodic memory		Attention/working memory		Language		Executive functions	
	Coef. (SE)	P-value	Coef. (SE)	P-value	Coef. (SE)	P-value	Coef. (SE)	P-value	Coef. (SE)	P-value
Last BMI – Baseline BMI ( BMI <sub>last-hi</sub> , reference group: BMI change <5%)	Time	-0.041 (0.002)	<.001	-0.001 (0.003)	.739	-0.046 (0.002)	<.001	-0.059 (0.002)	-0.073 (0.003)	<.001
	Decrease 5%	0.030 (0.009)	.001	0.041 (0.017)	.017	-0.012 (0.011)	.292	0.039 (0.012)	-0.017 (0.017)	.323
	Increase 5%	0.029 (0.010)	.005	-0.012 (0.020)	.541	0.016 (0.013)	.205	0.025 (0.014)	-0.032 (0.020)	.109
Average BMI – Baseline BMI ( BMI <sub>mean-hi</sub> , reference group: BMI change <5%)	Decrease 5%•time	-0.026 (0.0030)	<.001	-0.033 (0.004)	<.001	-0.014 (0.003)	<.001	-0.027 (0.004)	-0.163 (0.004)	<.001
	Increase 5%•time	-0.025 (0.003)	<.001	-0.024 (0.005)	<.001	-0.017 (0.003)	<.001	-0.036 (0.004)	-0.010 (0.005)	.051
	Time	-0.046 (0.002)	<.001	-0.006 (0.003)	.013	-0.049 (0.002)	<.001	-0.066 (0.002)	-0.076 (0.003)	<.001
Quartiles of SD of BMI ( BMI-SD, reference group: lowest quartile of SD)	Decrease 5%	0.029 (0.010)	.003	0.074 (0.019)	<.001	-0.016 (0.012)	.177	0.034 (0.013)	-0.016 (0.019)	.402
	Increase 5%	0.026 (0.012)	.025	0.018 (0.022)	.405	0.028 (0.014)	.047	0.010 (0.016)	-0.003 (0.023)	.881
	Time	-0.273 (0.003)	<.001	-0.040 (0.005)	<.001	-0.015 (0.003)	<.001	-0.028 (0.004)	-0.018 (0.004)	<.001
Quartiles of SD of BMI ( BMI-SD, reference group: lowest quartile of SD)	Increase 5%•time	-0.017 (0.004)	<.001	-0.017 (0.006)	.003	-0.015 (0.004)	<.001	-0.013 (0.005)	-0.008 (0.006)	.147
	Time	-0.040 (0.003)	<.001	0.006 (0.004)	.166	-0.044 (0.003)	<.001	-0.059 (0.003)	-0.067 (0.004)	<.001
	Quartile 2	0.019 (0.012)	.099	0.032 (0.023)	.163	0.022 (0.015)	.121	0.005 (0.016)	0.013 (0.022)	.548
Quartiles of SD of BMI ( BMI-SD, reference group: lowest quartile of SD)	Quartile 3	0.008 (0.012)	.496	0.017 (0.023)	.447	0.009 (0.015)	.531	0.003 (0.016)	0.029 (0.022)	.193
	Quartile 4	0.031 (0.012)	.008	0.011 (0.023)	.622	0.023 (0.015)	.121	0.018 (0.016)	0.039 (0.023)	.086
	Time	-0.007 (0.004)	.069	-0.016 (0.006)	.003	-0.005 (0.004)	.144	-0.007 (0.005)	-0.003 (0.005)	.508
Quartiles of SD of BMI ( BMI-SD, reference group: lowest quartile of SD)	Quartile 3•time	-0.015 (0.004)	<.001	0.028 (0.006)	<.001	-0.011 (0.004)	.002	-0.016 (0.005)	-0.017 (0.005)	.001
	Quartile 4•time	-0.027 (0.004)	<.001	-0.030 (0.006)	<.001	-0.022 (0.004)	<.001	-0.026 (0.005)	-0.031 (0.005)	<.001
	Time	-0.041 (0.002)	<.001	-0.001 (0.003)	.739	-0.046 (0.002)	<.001	-0.059 (0.002)	-0.073 (0.003)	<.001

*Note:* Covariates included age, sex, race/ethnicity, years of education, years of follow-up, baseline diagnosis (CN vs. MCI), FAQ, GDS, *APOE* genotype, number of medications (reflecting comorbidities), and indicators for history of diabetes, hypertension, hypercholesterolemia.

Abbreviation: BMI, body mass index; Coef, Coefficient; SE, standard error.

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