

## FEATURED ARTICLE

## Self-reported obstructive sleep apnea, amyloid and tau burden, and Alzheimer's disease time-dependent progression

Omonigho M. Bubu<sup>1,2</sup> | Ogie Q. Umasabor-Bubu<sup>3</sup> | Arlener D Turner<sup>1</sup> | Ankit Parekh<sup>4</sup> | Anna E. Mullins<sup>4</sup> | Korey Kam<sup>4</sup> | Madeline K. Birckbichler<sup>5</sup> | Fahad Mukhtar<sup>6</sup> | Alfred K Mbah<sup>6</sup> | Natasha J. Williams<sup>2</sup> | David M. Rapoport<sup>4</sup> | Mony de Leon<sup>7</sup> | Girardin Jean-Louis<sup>2</sup> | Indu Ayappa<sup>4</sup> | Andrew W. Varga<sup>4</sup> | Ricardo S. Osorio<sup>1,8</sup> | and Alzheimer's Disease Neuroimaging Initiative<sup>1</sup>

<sup>1</sup> Center for Sleep and Brain Health, Department of Psychiatry, NYU Grossman School of Medicine, New York, New York, USA

<sup>2</sup> Center for Healthful Behavior Change, Department of Population Health, New York Grossman School of Medicine, New York, USA

<sup>3</sup> Department of Epidemiology and Infection Control, State University New York Downstate Medical Center, Brooklyn, New York, USA

<sup>4</sup> Division of Pulmonary, Critical Care and Sleep Medicine at the Icahn School of Medicine at Mount Sinai, New York, New York, USA

<sup>5</sup> Department of Applied Health Sciences, Wheaton College, Wheaton, Illinois, USA

<sup>6</sup> Department of Epidemiology and Biostatistics, College of Public Health, University of South Florida, Tampa, Florida, USA

<sup>7</sup> Brain Health Imaging Institute, Department of Radiology, Weill Cornell Medicine, New York, New York, USA

<sup>8</sup> Nathan S. Kline Institute for Psychiatric Research, Orangeburg, New York, USA

## Correspondence

Omonigho M. Bubu, MD, MPH, PhD.

145 East 32nd Street, 8th floor, New York, NY 10016-6055, USA. E-mail:

[omonigho.bubu@nyulangone.org](mailto:omonigho.bubu@nyulangone.org)

Ricardo S. Osorio, M.D., 145 East 32nd Street, 5th Floor, New York, NY 10016-6055, USA.

E-mail: [ricardo.osorio@nyulangone.org](mailto:ricardo.osorio@nyulangone.org)

Alzheimer's Disease Neuroimaging Initiative: Data used in preparation of this article were obtained from the Alzheimer's Disease Neuroimaging Initiative (ADNI) database ([adni.loni.usc.edu](http://adni.loni.usc.edu)). The investigators within the ADNI contributed to the design and implementation of ADNI projects and provided data, but did not participate in the design, analysis, or writing of this report. A complete listing of ADNI investigators can be found at: [http://adni.loni.usc.edu/wp-content/uploads/how\\_to\\_apply/ADNI\\_Acknowledgement\\_List.pdf](http://adni.loni.usc.edu/wp-content/uploads/how_to_apply/ADNI_Acknowledgement_List.pdf)

Funding information: NIH/NIA/NHLBI (L30-AG064670, CIRAD P30AG059303 Pilot, T32HL129953, R01HL118624, R21AG049348, R21AG055002, R01AG056031, R01AG022374, R21AG059179, R01AG056682, R01AG056531, K07AG05268503, K23HL125939, K24HL109156).

## Abstract

**Introduction:** Obstructive sleep apnea (OSA) is associated with Alzheimer's disease (AD) biomarkers in cognitively normal (CN) and mild cognitive impaired (MCI) participants. However, independent and combined effects of OSA, amyloid beta (A $\beta$ ) and tau accumulation on AD time-dependent progression risk is unclear.

**Methods:** Study participants grouped by biomarker profile, as described by the A/T/N scheme, where "A" refers to aggregated A $\beta$ , "T" aggregated tau, and "N" to neurodegeneration, included 258 CN (OSA-positive [OSA+] [A+TN+ n = 10, A+/TN- n = 6, A-/TN+ n = 10, A-/TN- n = 6 and OSA-negative [OSA-] [A+TN+ n = 84, A+/TN- n = 11, A-/TN+ n = 96, A-/TN- n = 36]) and 785 MCI (OSA+ [A+TN+ n = 35, A+/TN- n = 15, A-/TN+ n = 25, A-/TN- n = 16] and OSA- [A+TN+ n = 388, A+/TN- n = 28, A-/TN+ n = 164, A-/TN- n = 114]) older-adults from the Alzheimer's Disease Neuroimaging Initiative cohort. Cox proportional hazards regression models estimated the relative hazard of progression from CN-to-MCI and MCI-to-AD, among baseline OSA CN and MCI patients, respectively. Multi-level logistic mixed-effects models with random intercept and slope investigated the synergistic associations of self-reported OSA, A $\beta$ , and tau burden with prospective cognitive decline.

**Results:** Independent of TN-status (CN and MCI), OSA+/A $\beta$ + participants were approximately two to four times more likely to progress to MCI/AD ( $P < .001$ ) and progressed 6 to 18 months earlier ( $P < .001$ ), compared to other participants combined (ie, OSA+/A $\beta$ -, OSA-/A $\beta$ +, and OSA-/A $\beta$ -). Notably, OSA+/A $\beta$ - versus OSA-/A $\beta$ - (CN

and MCI) and OSA+/TN– versus OSA–/TN– (CN) participants showed no difference in the risk and time-to-MCI/AD progression. Mixed effects models demonstrated OSA synergism with A $\beta$  (CN and MCI [ $\beta$  = 1.13, 95% confidence interval (CI), 0.74 to 1.52, and  $\beta$  = 1.18, 95%CI, 0.82 to 1.54]) respectively, and with tau (MCI [ $\beta$  = 1.31, 95% CI, 0.87 to 1.47]),  $P$  < .001 for all.

**Discussion:** OSA acts in synergism with A $\beta$  and with tau, and all three acting together result in synergistic neurodegenerative mechanisms especially as A $\beta$  and tau accumulation becomes increasingly abnormal, thus leading to shorter progression time to MCI/AD in CN and MCI-OSA patients, respectively.

#### KEYWORDS

Alzheimer's disease, amyloid beta<sub>42</sub>, brain amyloid-positron emission tomography, cerebrospinal fluid biomarkers, longitudinal study, obstructive sleep apnea, p-tau, t-tau

## 1 | INTRODUCTION

Obstructive sleep apnea (OSA) increases Alzheimer's disease (AD) risk<sup>1–6</sup> and at cross-section, is associated with AD biomarkers, including the presence of significant brain amyloid beta (A $\beta$ ) and tau burden, measured either by cerebrospinal (CSF) A $\beta$ <sub>42</sub> or amyloid positron emission tomography (PET), and CSF levels of tau (ie, total or hyperphosphorylated) or tau-PET, in both cognitive normal (CN) and mild cognitive impaired (MCI) participants.<sup>7–14</sup> Recently, our group found that this cross-sectional association was not found in participants with Pittsburgh compound B (PiB)-negative scans,<sup>15</sup> suggesting that the presence or absence of amyloid burden might act as a moderator in these relationships. A previous cross-sectional study suggested a similar phenomenon, with associations seen between increased amyloid deposition and higher apnea hypopnea index indices in MCI patients but not among CN controls.<sup>11</sup> More recently, we expanded the analysis of cross-sectional evaluations to 2-year follow-ups, first, in community-dwelling healthy CN elderly from New York City,<sup>15</sup> and second, from purely CN older individuals to those across the spectrum of dementia, from CN to MCI and to full AD in a large population from the Alzheimer's Disease Neuroimaging Initiative (ADNI) cohort.<sup>16</sup> In the New York University sample, we examined the association between severity of OSA and longitudinal increase in amyloid burden and found that OSA severity was associated with greater CSF A $\beta$ <sub>42</sub> changes over a 2-year follow-up in CN older adults.<sup>15</sup> In the ADNI sample, we examined the effect of self-reported clinical diagnosis of OSA on longitudinal changes in brain amyloid-PET and CSF-biomarkers (A $\beta$ <sub>42</sub>, t-tau, and p-tau) in CN, MCI, and AD older adults and observed OSA effects on longitudinal increases in amyloid burden by both CSF and PET imaging measures, in the CN and MCI groups.<sup>16</sup>

Molecular markers of AD pathology (eg, amyloid PET uptake and CSF A $\beta$ <sub>42</sub> levels), are known robust predictors of amyloid burden and of future development of AD,<sup>17,18</sup> and evidence indicates that A $\beta$  accumulation starts decades prior to the appearance of the first cognitive symptoms.<sup>19,20</sup> In previous studies, the lack of longer clinical

assessment prevented testing whether amyloid or tau deposition in OSA participants precedes subsequent cognitive decline to MCI or AD. In this study, we tried to overcome this limitation, and hypothesized that OSA's effect on MCI/AD progression risk will be synergistic with A $\beta$  and tau and this risk will significantly increase, as A $\beta$  and tau accumulation becomes increasingly abnormal leading to shorter time to MCI/AD in CN and MCI participants, respectively. Having previously demonstrated a contributory role of OSA on longitudinal increases in amyloid burden by both CSF and PET imaging measures, in CN and MCI patients, our objectives were to (1) to examine whether OSA has a direct neurotoxic and/or neurodegenerative effect that is independent of A $\beta$  or tau, sufficient to induce a prospective clinical diagnosis of cognitive decline, and (2) to examine whether OSA's direct neurotoxicity independent of A $\beta$  or tau, together with OSA's indirect effect that promotes A $\beta$  or tau accumulation, combine to act synergistically to significantly increase MCI/AD progression risk.

## 2 | METHODS

Data used in the preparation of this article were obtained from the ADNI database (adni.loni.usc.edu). Launched in 2003 as a public-private partnership, and led by Principal Investigator Michael W. Weiner, MD, ADNI's primary goal has been to measure the progression of MCI and early AD using a combination of serial magnetic resonance imaging (MRI), PET, other biological markers, and clinical and neuropsychological assessments. Currently, ADNI has recruited more than 2000 adults aged 55 to 90, consisting of CN, MCI, and early AD. PET and CSF sampling follow-up typically occurs every 1 to 2 years.

### 2.1 | Study participants

Participant selection (see Figure S1 in supporting information) using ADNI data by our group has been previously described.<sup>2,16</sup> Participants in the current study included 1043 subjects: 258 CN and 785

MCI with one or more clinical follow-up assessments. Subjects for this study must have undergone florbetapir-PET imaging while carrying a clinical diagnosis of CN or MCI. The primary outcome was time-to-progression from a clinical diagnosis of CN to a clinical diagnosis of MCI for baseline CN patients and, from MCI to a clinical diagnosis of AD, for baseline MCI patients. Neuropsychometric assessments and serial PET-MRI scans were performed at baseline and periodically on participants. Details are available at <http://www.adni-info.org>. Participants were classified as CN or MCI at the time of their baseline visit and remained as CN or MCI up to and including their 12-month visit. Patient diagnosis was recorded at 6 to 12 monthly intervals up until the download date (December 6, 2018). CN and MCI subjects were classified as converters if they converted to MCI and AD between 12 months and the download date, respectively.

## 2.2 | Standard protocol approvals, registrations, and patient consent

Written informed consent was obtained from all patients participating in ADNI, as approved by the Institutional Review Board, at each of the ADNI participating centers. ADNI inclusion and exclusion criteria are detailed elsewhere ([http://adni.loni.ucla.edu/wp-content/uploads/2010/09/ADNI\\_GeneralProceduresManual.pdf](http://adni.loni.ucla.edu/wp-content/uploads/2010/09/ADNI_GeneralProceduresManual.pdf)).

## 2.3 | OSA diagnosis

As previously described,<sup>2,16</sup> presence or absence of OSA was based on medical history of a self-reported clinical diagnosis of OSA during the clinical interview. Briefly, participants labeled OSA-positive (OSA+) reported a medical diagnosis of “sleep apnea,” “sleep disordered breathing (SDB),” “OSA,” or “SDB” and the remaining participants were considered OSA-negative (OSA-). We ensured proper group allocation from reviewed medical history clinical notes from the ADNI download.

## 2.4 | CN, MCI, and AD diagnosis

ADNI criteria for subject classification are described elsewhere.<sup>21</sup> Briefly, CN and MCI subjects scored between 24 and 30 on the Mini-Mental State Examination while AD subjects scored between 20 and 26. MCI and AD participants had global Clinical Dementia Rating scores of 0.5 and 1, respectively. The diagnosis of AD was made using established clinical criteria.<sup>22</sup>

## 2.5 | Florbetapir-PET imaging acquisition and interpretation

Information on florbetapir-PET imaging acquisition and interpretation is available at [https://adni.loni.usc.edu/wp-content/uploads/2010/05/ADNI2\\_PET\\_Tech\\_Manual\\_0142011.pdf](https://adni.loni.usc.edu/wp-content/uploads/2010/05/ADNI2_PET_Tech_Manual_0142011.pdf), <http://adni.loni.usc.edu>

### HIGHLIGHTS

- Obstructive sleep apnea (OSA) has potential for neuronal injury independent of amyloid beta ( $A\beta$ ) and tau.
- OSA appears to accelerate brain amyloid and tau deposition over time
- OSA,  $A\beta$ , and tau act synergistically leading to shorter Alzheimer's disease progression time

### RESEARCH IN CONTEXT

1. Systematic review: The authors reviewed the literature using traditional (eg, PubMed) sources. Recent research findings indicate that obstructive sleep apnea (OSA) is associated with longitudinal increases in amyloid beta ( $A\beta$ ) and tau burden in cognitive normal (CN) and mild cognitive impairment (MCI) older adults. These relevant studies are appropriately cited.
2. Interpretation: Our findings suggest that OSA acts in synergism with  $A\beta$  and with tau, with all three acting together to produce synergistic neurodegenerative mechanisms especially as  $A\beta$  and tau accumulation becomes increasingly abnormal, leading to shorter progression time to MCI/Alzheimer's disease (AD) in CN and MCI-OSA patients, respectively. These findings are consistent with recent epidemiological studies examining OSA's effect on AD biomarkers.
3. Future directions: The article proposes a framework for additional research. Examples include further understanding: (1) whether OSA- $A\beta$  synergism related to cognitive decline is independent of vascular burden and (2) OSA's effects on slow wave and rapid eye movement sleep and their mediating role in increasing  $A\beta$  accumulation

[//adni.loni.usc.edu/updated-florbetapir-av-45-pet-analysis-results/](http://adni.loni.usc.edu/updated-florbetapir-av-45-pet-analysis-results/).

As described previously,<sup>16</sup> the University of California at Berkeley (UC Berkeley) uploaded ADNI florbetapir summary data to the Laboratory of Neuroimaging.<sup>23,24</sup> Calculation of florbetapir standardized uptake value ratios were done by obtaining means across four cortical regions (frontal, anterior/posterior cingulate, lateral parietal, lateral temporal) and dividing this cortical summary region of interest by one of the five reference regions (cerebellar gray matter, whole cerebellum, brainstem/pons, eroded subcortical white matter, and a composite reference region). The UC Berkeley team using procedures that involved receiver-operating-characteristic analysis,<sup>23,24</sup> defined thresholds for  $A\beta$  positive and negative status as florbetapir cutoff of  $>1.11$  and  $<1.11$ , respectively, using the whole cerebellum reference region only.

## 2.6 | Cerebrospinal fluid methods

CSF bio-specimen data collection details can be found at <http://adni.loni.usc.edu/data-samples/biospecimen-data/> and as

previously described.<sup>16</sup> A standardized protocol was implemented to quantify biomarker concentrations in each of the CSF baseline aliquots using a multiplex xMAP Luminex platform (Luminex Corp, Austin, TX) with Innogenetics (INNO-BIA AlzBio3, Ghent, Belgium; for research use only reagents) immunoassay kit-based reagents, validated in Vanderstichele et al.<sup>25</sup> and Shaw et al.<sup>26</sup> Further details can be found at (<http://www.adni-info.org/index.php>). Using the recently published 2018 National Institute on Aging-Alzheimer's Association (NIA-AA) "research framework" for the diagnosis of AD,<sup>27</sup> ADNI participants were grouped by biomarker profile, as described by the A/T/N scheme,<sup>28</sup> where "A" refers to aggregated A $\beta$ , "T" aggregated tau, and "N" to neurodegeneration. Each biomarker group is dichotomized as negative (–) or positive (+) based on specified biomarkers levels. In the present study, as noted above, florbetapir-PET cutoff of >1.11 and <1.11 defined thresholds for A $\beta$ – and A $\beta$ + status, respectively, because the correlation between ADNI CSF A $\beta$  and florbetapir biomarkers is limited to a middle range of values, is modified by the apolipoprotein E (APOE) genotype, and is absent for longitudinal changes.<sup>29</sup> Following recent studies that used ADNI data,<sup>30,31</sup> "T+" individuals had CSF p-tau181 > 21.8 pg/mL and "N+" individuals had t-tau > 245 pg/mL. We merged the aggregated tau (T) and neurodegeneration (N) groups to decrease the number of groups to be compared.<sup>30,31</sup> TN negative (TN–) was defined as having both the aggregated tau (T) and neurodegeneration (N) biomarkers in the normal range (T– and N–; that is, p-tau181  $\leq$  21.8 pg/mL and t-tau  $\leq$  245 pg/mL). Participants were classified as TN positive (TN+) if either aggregated tau (T) or neurodegeneration (N) were abnormal (T+ or N+; that is p-tau181P > 21.8 pg/mL or t-tau > 245 pg/mL). None of the individuals of the total differed between the T and N biomarkers groups.

## 2.7 | Data analyses

All analyses were conducted separately for each clinical group (ie, CN and MCI) in phases. First, survival and cumulative hazard function estimates and their 95% Hall-Wellner bands were populated for both the CN and MCI groups, comparing OSA+ versus OSA–, A $\beta$ + versus A $\beta$ –, and TN+ versus TN– patients, respectively. Second, similar survival and cumulative hazard function estimates were populated comparing OSA+ versus OSA– within dichotomized A $\beta$  and TN groups (ie, A $\beta$ +/OSA+ vs A $\beta$ +/OSA–, A $\beta$ –/OSA+ vs A $\beta$ –/OSA–, TN+/OSA+ vs TN+/OSA–, and TN–/OSA+ vs TN–/OSA–). The above analyses were also performed comparing A $\beta$ + versus A $\beta$ – and TN+ versus TN– stratifying by OSA status (ie, OSA+/A $\beta$ + vs OSA+/A $\beta$ –, OSA–/A $\beta$ + vs OSA–/A $\beta$ –, OSA+/TN+ vs OSA+/TN– and OSA–/TN+ vs OSA–/TN–), for both the CN and MCI groups. The analysis comparing CN and MCI OSA+ versus OSA– among only A $\beta$ + or TN+ participants examined the combined effects of OSA and A $\beta$ <sub>42</sub> or OSA, tau accumulation, and neurodegeneration on progression risk, respectively. The analysis comparing CN and MCI OSA+ versus OSA– among only A $\beta$ – or TN– participants examined whether OSA, independent of A $\beta$  or tau accumulation and neurodegeneration, was sufficient to induce a prospective clinical diagnosis of cognitive decline, respec-

tively. The analysis comparing CN and MCI A $\beta$ + versus A $\beta$ – or TN+ versus TN–, among only OSA+ participants, examined the combined effects of OSA and A $\beta$ <sub>42</sub> or OSA, tau accumulation, and neurodegeneration on progression risk beyond that of OSA. The analysis comparing CN and MCI A $\beta$ + versus A $\beta$ – or TN+ versus TN– among only OSA– participants examined the individual effect of A $\beta$  or tau accumulation and neurodegeneration on AD progression risk among OSA– subjects. Mean and median time-to-event for all the groups were estimated. Cox proportional hazards regression models estimated the individual and combined effects of OSA and A $\beta$  load, and OSA and TN burden on the relative hazard of progression from CN to MCI and MCI to AD, among baseline CN and MCI patients, respectively. To investigate the additive or synergistic associations of self-reported OSA, A $\beta$  and tau burden with prospective cognitive decline, multi-level logistic mixed-effects models with random intercept and slope were used. We examined interactions of self-reported OSA with time, A $\beta$  burden with time, and tau burden with time in a single model (eg, model 1: CN to MCI: OSA  $\times$  time + A $\beta$   $\times$  time + tau  $\times$  time + covariates  $\times$  time). Next, we added an interaction term between the OSA, A $\beta$  burden, and time, and OSA, tau burden, and time, to examine whether these two factors increase the likelihood of prospective cognitive decline beyond their separate effects (ie, synergistic effect; model 2: CN to MCI: OSA  $\times$  A $\beta$   $\times$  time + OSA  $\times$  tau  $\times$  time + covariates  $\times$  time). Last, we added an interaction term between the OSA, A $\beta$  burden, tau, and time, to examine whether these three factors increase the likelihood of prospective cognitive decline beyond their combined effects in model 2 (ie, synergistic effect; model 3: CN to MCI: OSA  $\times$  A $\beta$   $\times$  tau  $\times$  time + covariates  $\times$  time). We operationalized time as years from baseline for each participant. For all analyses, final models adjusted for age, sex, body mass index (BMI), education, continuous positive airway pressure (CPAP) machine use, baseline biomarker data, hypertension, diabetes, history of cardiovascular disease (eg, including ischemic heart disease, heart failure, and stroke/transient ischemic attack [TIA]), alcohol use, and history of traumatic brain injury. We also performed sensitivity analyses removing CPAP users from OSA+ participants. Statistical analyses were performed using SAS (version 9.4; SAS Institute Inc., Cary, North Carolina, USA).

## 3 | RESULTS

### 3.1 | Demographic and clinical characteristics

Tables 1 and 2 show the demographic and clinical characteristics of study participants at baseline according to their OSA and A $\beta$  status, and OSA and A/T/N status, respectively. Overall, of the 1043 participants, 506 (49%) were women. The overall mean (standard deviation [SD]) age was 74.7 (5.0) years and the overall mean (SD) follow-up time was 5.5 (1.7) years (range: 2.7–10.9 years). The mean (SD) follow-up time was 5.3 (1.4) and 5.7 (1.9) for CN and MCI groups, respectively. The mean ages of OSA+ and OSA– (CN and MCI combined) were 72.3  $\pm$  7.1 and 73.9  $\pm$  7.3 years, respectively. In the CN group, 31 participants (12% [6% A $\beta$ +, 6% A $\beta$ –] and [8% TN+, 4% TN–]) were OSA+,

**TABLE 1** Baseline descriptive characteristics of participants by obstructive sleep apnea and A $\beta$  load status

Table 1a: Characteristics of cognitively normal participants by OSA and A $\beta$ load status; and converters from cognitive normal to MCI							
Characteristics	All	Without OSA		With OSA		Stable	Converters
		A $\beta$ +	A $\beta$ –	A $\beta$ +	A $\beta$ –		
Number of participants (%)	258 (100)	95 (37)	132 (51)	16 (6)	15 (6)	187 (72)	71 (28)
Follow-up time, mean (SD), years	5.3 (1.4)	5.7 (1.1)	5.2 (1.5)	4.9 (1.7)	5.4 (1.4)	5.6 (1.1)	5.1 (1.9)
Female sex, number (%)	121 (47)	52 (43)	58 (48)	2 (1)	9 (8)	92 (76)	29 (24)
Age, years, median (interquartile range)	74 (71, 78)	75 (71, 79)	71 (68, 76)	70 (70, 86)	71 (68, 75)	71 (68, 76)	74 (70, 79)
APOE positive, number (%) <sup>a</sup>	80 (31)	37 (46)	32 (41)	10 (12)	1 (1)	38 (48)	42 (52)
Education, years, median (interquartile range)	16 (14, 18)	16 (14, 18)	17 (16, 19)	12 (12, 12)	16 (16, 18)	17 (16, 19)	16 (12, 16)
BMI (kg/m <sup>2</sup> ) <sup>a</sup>	27.2 $\pm$ 4.8	27.1 $\pm$ 4.8	27.7 $\pm$ 5.2	36.5 $\pm$ 10.5	29.7 $\pm$ 4.5	28.7 $\pm$ 7.2	31.5 $\pm$ 7.8
Hypertension, number (%) <sup>a</sup>	124 (48)	38 (31)	68 (55)	9 (7)	9 (7)	84 (68)	40 (32)
Diabetes, number (%) <sup>a</sup>	28 (11)	8 (29)	13 (46)	5 (18)	2 (7)	18 (64)	10 (36)
Thyroid disease, number (%)	59 (23)	25 (43)	28 (47)	5 (9)	1 (1)	41 (69)	18 (31)
Respiratory disease, number (%)	57 (22)	16 (28)	23 (40)	5 (9)	13 (23)	38 (67)	19 (33)
Cardiovascular disease, number (%) <sup>a</sup>	173 (67)	82 (47)	75 (44)	9 (5)	7 (4)	114 (66)	59 (33)
TBI, number (%)	10 (4)	6 (58)	4 (42)			5 (50)	5 (50)
Alcohol use, number (%)	18 (7)	7 (39)	8 (44)	1 (6)	2 (11)	12 (67)	6 (33)
CPAP use, number (%)	8 (3)			6 (70)	2 (30)	6 (70)	2 (30)
MMSE median (interquartile range)	29 (28, 29)	29 (28, 30)	29 (28, 29)	29 (27, 30)	29 (28, 29)	29 (28, 29)	29 (28, 30)
CDR median (interquartile range)	0 (0, 0)	0 (0, 0)	0 (0, 0)	0 (0, 0)	0 (0, 0)	0 (0, 0)	0 (0, 0)
CSF-A $\beta$ pg/mL median (interquartile range) <sup>a</sup>	210 (155, 241)	149 (132, 173)	221 (200, 245)	112 (93, 123)	229 (213, 268)	198 (188, 245)	132 (120, 164)
TAU pg/mL median (interquartile range) <sup>a</sup>	59 (45, 84)	71 (49, 105)	53 (43, 69)	141 (126, 156)	56 (42, 63)	61 (42, 89)	109 (86, 140)
PTAU pg/mL median (interquartile range) <sup>a</sup>	27 (20, 40)	41 (30, 55)	27 (21, 38)	91 (46, 137)	27 (21, 29)	31 (21, 48)	65 (33, 117)
A $\beta$ , median (interquartile range) <sup>a</sup>	1.1 (1.0, 1.2)	1.3 (1.2, 1.4)	1.0 (1.0, 1.1)	1.2 (1.2, 1.3)	1.0 (1.0, 1.0)	1.0 (1.0, 1.2)	1.3 (1.2, 1.4)

**Table 1b: Characteristics of MCI participants by OSA and A $\beta$  load status; and converters from MCI to Alzheimer's disease**

Characteristics	All	Without OSA		With OSA		Stable	Converters
		A $\beta$ +	A $\beta$ –	A $\beta$ +	A $\beta$ –		
Number of participants (%)	785 (100)	416 (53)	278 (35)	50 (6)	41 (5)	487 (62)	298 (38)
Follow-up time, mean (SD), years	5.7 (1.9)	5.6 (1.4)	6.1 (1.1)	5.5 (1.6)	5.6 (1.5)	5.9 (1.7)	5.5 (1.7)
Female sex, number (%)	385 (49)	236 (61)	102 (27)	27 (7)	20 (5)	250 (65)	135 (35)
Age, years, median (interquartile range)	74 (68, 79)	74 (69, 78)	71 (65, 76)	72 (69, 77)	72 (68, 75)	71 (68, 75)	74 (68, 79)
APOE positive, number (%) <sup>a</sup>	395 (50)	288 (73)	55 (14)	32 (8)	20 (5)	125 (58)	89 (42)
Education, years, median (interquartile range) <sup>a</sup>	16 (14, 18)	16 (14, 18)	16 (14, 18)	16 (12, 19)	17 (15, 19)	16 (14, 18)	16 (14, 18)
BMI (kg/m <sup>2</sup> ) <sup>a</sup>	26.9 $\pm$ 4.6	26.8 $\pm$ 4.8	27.8 $\pm$ 4.6	30.5 $\pm$ 6.1	29.6 $\pm$ 5.6	28.7 $\pm$ 4.8	30.4 $\pm$ 5.8
Hypertension, number (%) <sup>a</sup>	369 (47)	192 (52)	118 (32)	30 (8)	29 (8)	196 (53)	173 (47)
Diabetes, number (%) <sup>a</sup>	79 (10)	30 (38)	21 (27)	16 (20)	12 (15)	54 (68)	25 (32)
Thyroid disease, number (%)	165 (21)	81 (49)	56 (34)	17 (10)	11 (7)	125 (76)	40 (24)



**TABLE 1** Continued.

**Table 1b: Characteristics of MCI participants by OSA and Aβ load status; and converters from MCI to Alzheimer's disease**

Characteristics	All	Without OSA		With OSA		Stable	Converters
		Aβ +	Aβ –	Aβ +	Aβ –		
Respiratory disease, number (%)	188 (24)	83 (44)	42 (22)	24 (13)	39 (21)	137 (73)	51 (27)
Cardiovascular disease, number (%) <sup>a</sup>	502 (64)	226 (45)	201 (40)	45 (9)	30 (6)	377 (75)	125 (25)
TBI, number (%)	31 (4)	10 (33)	14 (44)	5 (17)	2 (6)	14 (44)	17 (56)
Alcohol use, number (%)	63 (8)	23 (37)	32 (51)	2 (3)	6 (9)	42 (66)	21 (34)
CPAP use, number (%)	31 (4)			10 (31)	21 (69)	23 (75)	8 (25)
MMSE median (interquartile range)	27 (24, 28)	26 (25, 28)	28 (25, 29)	26 (24, 28)	27 (24, 29)	28 (25, 29)	26 (24, 28)
CDR median (interquartile range)	0.5 (0.5, 0.5)	0.5 (0.5, 0.5)	0.5 (0.5, 0.5)	0.5 (0.5, 0.5)	0.5 (0.5, 0.5)	0.5 (0.5, 0.5)	0.5 (0.5, 0.5)
CSF-Aβ pg/mL median (interquartile range) <sup>a</sup>	153 (130, 209)	134 (120, 154)	219 (199, 243)	147 (134, 167)	212 (190, 240)	179 (159, 243)	134 (120, 154)
TAU pg/mL median (interquartile range) <sup>a</sup>	80 (54, 116)	97 (70, 140)	52 (36, 72)	93 (66, 114)	55 (42, 78)	62 (46, 82)	97 (66, 140)
PTAU pg/mL median (interquartile range) <sup>a</sup>	36 (23, 51)	50 (36, 67)	23 (17, 32)	40 (32, 59)	27 (19, 38)	25 (20, 38)	50 (33, 67)
Aβ, median (interquartile range) <sup>b</sup>	1.2 (1.0, 1.4)	1.4 (1.3, 1.5)	1.0 (1.0, 1.1)	1.3 (1.2, 1.4)	1.0 (1.0, 1.0)	1.1 (1.0, 1.2)	1.3 (1.2, 1.4)

Abbreviations: ApoE4, apolipoprotein E ε4; Aβ, amyloid beta; BMI, body mass index; CDR, Clinical Dementia Rating; CPAP, continuous pulmonary airway pressure; CSF, cerebrospinal fluid; MCI, mild cognitive impairment; MMSE, Mini-Mental State Examination; OSA, obstructive sleep apnea; PTAU, phosphorylated tau; TAU, tau protein; TBI, traumatic brain injury. All columns show column percent where indicated. OSA and Aβ columns show row percent where indicated. <sup>a</sup>Indicates significant differences between groups. Aβ+ participants had significantly lower CSF-Aβ<sub>42</sub>, and higher levels of CSF t-tau and CSF p-tau. Amyloid positive subjects in both CN and MCI groups were more likely to be APOE ε4 carriers. Participants who were OSA+/Aβ+ had significantly higher BMI in the CN group but not in the MCI group, compared to other participants with varying OSA and Aβ status. In both CN and MCI, OSA+/Aβ+ participants had higher vascular burden (ie, higher rates of hypertension, diabetes, and cardiovascular disease) compared to OSA-/Aβ+ participants (eg, 9/15 [60%] vs 38/95 [40%] for hypertension in CN participants). CN/OSA+/Aβ+ participants had significantly lower educational level. Significance level *p* ≤ .05.

and 91 participants (11% [6% Aβ+, 5% Aβ-] and [7% TN+, 4% TN-]) were OSA+ in the MCI group. In both the CN and MCI groups, and in both OSA and non-OSA participants, marked differences existed in AD pathology markers comparing Aβ+ to Aβ-, with the former having significantly lower CSF-Aβ<sub>42</sub>, and higher levels of CSF t-tau and CSF p-tau (Table 1). A+/TN+ subjects in both CN and MCI groups were more likely to be APOE ε4 carriers (Table 2). Participants who were OSA+/Aβ+ had significantly higher BMI in the CN group but not in the MCI group, compared to other participants with varying OSA and Aβ status. In both CN and MCI, OSA+/Aβ+/TN+ participants had higher vascular burden (ie, higher rates of hypertension, diabetes, and cardiovascular disease) compared to OSA-/Aβ+ participants (eg, 7/10 [70%] vs 30/84 [36%] for hypertension in CN participants; Table 2). CN/OSA+/Aβ+ participants had significantly lower educational level (Table 1).

### 3.2 | Time-dependent progression risk and cumulative hazard function estimates in CN group

Figure 1 shows survival and cumulative hazard function estimates and their 95% Hall-Wellner bands populated for the CN group. Compared to OSA- participants, OSA+ participants had a significantly shorter time-to-progression to MCI (mean ± SD [median] 4.5 ± 0.3 [4.0] years

vs 5.0 ± 0.3 [4.8] years, *P* = .03), and had a 32% increased hazard risk of developing MCI (adjusted hazard ratio [aHR]:1.32, 95% confidence interval [CI]:1.11-1.65, *P* < .01). Compared to Aβ- participants, Aβ+ participants had a significantly shorter time-to-progression to MCI (mean ± SD [median] 3.6 ± 0.5 [2.6] years vs 4.2 ± 0.5 [3.0] years, ] *P* < .001), and a significantly higher risk of developing MCI (aHR: 2.44, 95% CI: 1.99-2.89, *P* < .001). Compared to TN- participants, TN+ participants had a significantly shorter time-to-progression to MCI (mean ± SD [median] 3.4 ± 0.4 [2.5] years vs 4.7 ± 0.6 [4.3] years, *P* < .001), and a significantly higher risk of developing MCI (aHR: 3.52, 95% CI: 1.89-5.17, *P* = .01). See Table 3.

Stratifying by brain Aβ or TN burden, Aβ+/OSA+ and TN+/OSA+ participants had a significantly shorter time-to-progression to MCI (mean ± SD [median] 3.5 ± 0.4 [2.9] years vs 3.8 ± 0.3 [3.5] years, *P* = .04 and 3.3 ± 0.3 [2.8] years vs 3.8 ± 0.3 [3.0] years, *P* < .001) and a significantly higher risk of developing MCI (aHR: 2.93, 95% CI: 2.17-3.69, and aHR: 2.04, 95% CI: 1.11-2.97, *P* < .001 for both) compared to Aβ+/OSA- and TN+/OSA+ participants, respectively. Compared to Aβ-/OSA- and TN-/OSA- participants, Aβ-/OSA+ and TN-/OSA+ participants showed no significant difference in time-to-progression to MCI (mean ± SD [median] 3.9 ± 0.6 [3.3] years vs 4.1 ± 0.5 [4.0] years, and aHR: 1.01, 95% CI: 0.76-1.26, *P* = .11 and mean ± SD [median] 4.5 ± 0.4 [4.0] years vs 5.3 ± 0.6 [4.5] years, and aHR: 1.01, 95% CI: 0.76-1.26, *P* = .06, respectively). See Table 3.

**TABLE 2** Baseline descriptive characteristics of participants by obstructive sleep apnea and A/T/N status

Characteristics	Table 2a: Characteristics of cognitively normal participants by OSA and A/T/N status								
	All	Without OSA			With OSA				
		A+(TN) +	A+(TN) –	A-(TN) +	A-(TN) –	A+(TN) +	A-(TN) –		
Number of participants (%)	258 (100)	84 (33)	11(4)	96(37)	36 (14)	10(4)	6 (2)	10 (4)	5 (2)
Follow-up time, mean (SD), years	5.3 (1.4)	5.9 (1.3)	5.4 (1.9)	5.6 (1.5)	5.4 (1.5)	4.7 (1.8)	5.2 (1.6)	5.4 (1.4)	5.3 (1.7)
Female sex, number (%)	121 (47)	35 (28)	17 (14)	36 (30)	22 (18)	1 (1)	1 (1)	4 (4)	5 (4)
Age, years, median (interquartile range)	74 (71, 78)	75 (71, 79)	75 (70, 79)	73 (68, 77)	72 (68, 75)	78 (70, 86)	70 (70, 70)	72 (66, 75)	72 (71, 78)
APOE positive, number (%)*	80 (31)	28 (35)	9 (11)	24 (30)	8 (11)	8 (11)	2 (1)	8 (11)	1 (1)
Education, years, median (interquartile range)*	16 (14, 18)	16 (14, 18)	16 (14, 20)	17 (16, 19)	17 (16, 18)	12 (12, 12)	12 (12, 12)	16 (14, 18)	17 (16, 19)
BMI (kg/m2)*	27.2 ± 4.8	26.7 ± 4.5	28.5 ± 6.1	27.7 ± 5.4	27.5 ± 4.8	32.4 ± 11.2	34.5 ± 4.8	30.0 ± 5.5	29.2 ± 2.7
Hypertension, number (%)*	124 (48)	30 (24)	8 (7)	47 (38)	21 (17)	7 (6)	2 (1)	6 (5)	3 (2)
Diabetes, number (%)	28 (11)	6 (21)	2 (7)	7 (25)	6 (21)	4 (14)	1 (4)	1 (4)	1 (4)
Thyroid disease, number (%)	59 (23)	19 (32)	6 (11)	18 (30)	10 (17)	1 (2)	4 (7)	1 (1)	1 (1)
Respiratory disease, number (%)	57 (22)	12 (21)	4 (7)	13 (23)	10 (17)	4 (7)	1 (2)	7 (12)	6 (11)
Cardiovascular disease, number (%)*	173 (67)	65 (38)	17 (10)	52 (30)	23 (14)	7 (4)	2 (1)	5 (3)	2 (1)
TBI, number (%)	10 (4)	6 (58)		4 (42)					
Alcohol use, number (%)	18 (7)	5 (28)	2 (11)	7 (38)	1 (6)	1 (6)		1 (6)	2 (11)
CPAP use, number (%)	8 (3)					5 (70)	1	1	1 (30)
MMSE median (interquartile range)	29 (28, 29)	28 (27, 30)	29 (28, 30)	29 (28, 29)	29 (28, 30)	27 (27, 29)	29 (28, 29)	29 (28, 29)	29 (28, 29)
CDR median (interquartile range)	0 (0, 0)	0 (0, 0)	0 (0, 0)	0 (0, 0)	0 (0, 0)	0 (0, 0)	0 (0, 0)	0 (0, 0)	0 (0, 0)
CSF-Aβ pg/mL median (interquartile range)*	210 (155, 241)	151 (132, 166)	179 (161, 184)	224 (203, 248)	210 (188, 232)	112 (93, 132)	159 (131, 172)	246 (226, 269)	224 (201, 225)
TAU pg/mL median (interquartile range)*	59 (45, 84)	86 (60, 106)	36 (27, 46)	65 (47, 81)	42 (35, 49)	141 (126, 156)	46 (37, 53)	61 (54, 69)	44 (35, 48)
PTAU pg/mL median (interquartile range)*	27 (20, 40)	48 (34, 57)	16 (12, 19)	35 (26, 43)	17 (15, 20)	91 (46, 137)	17 (17, 19)	29 (27, 32)	19 (16, 21)
Aβ median (interquartile range)*	1.1 (1.0, 1.2)	1.1 (1.1, 1.2)	1.1 (1.1, 1.1)	1.0 (1.0, 1.0)	1.0 (1.0, 1.0)	1.2 (1.2, 1.3)	1.2 (1.1, 1.3)	1.0 (1.0, 1.0)	1.0 (1.0, 1.0)

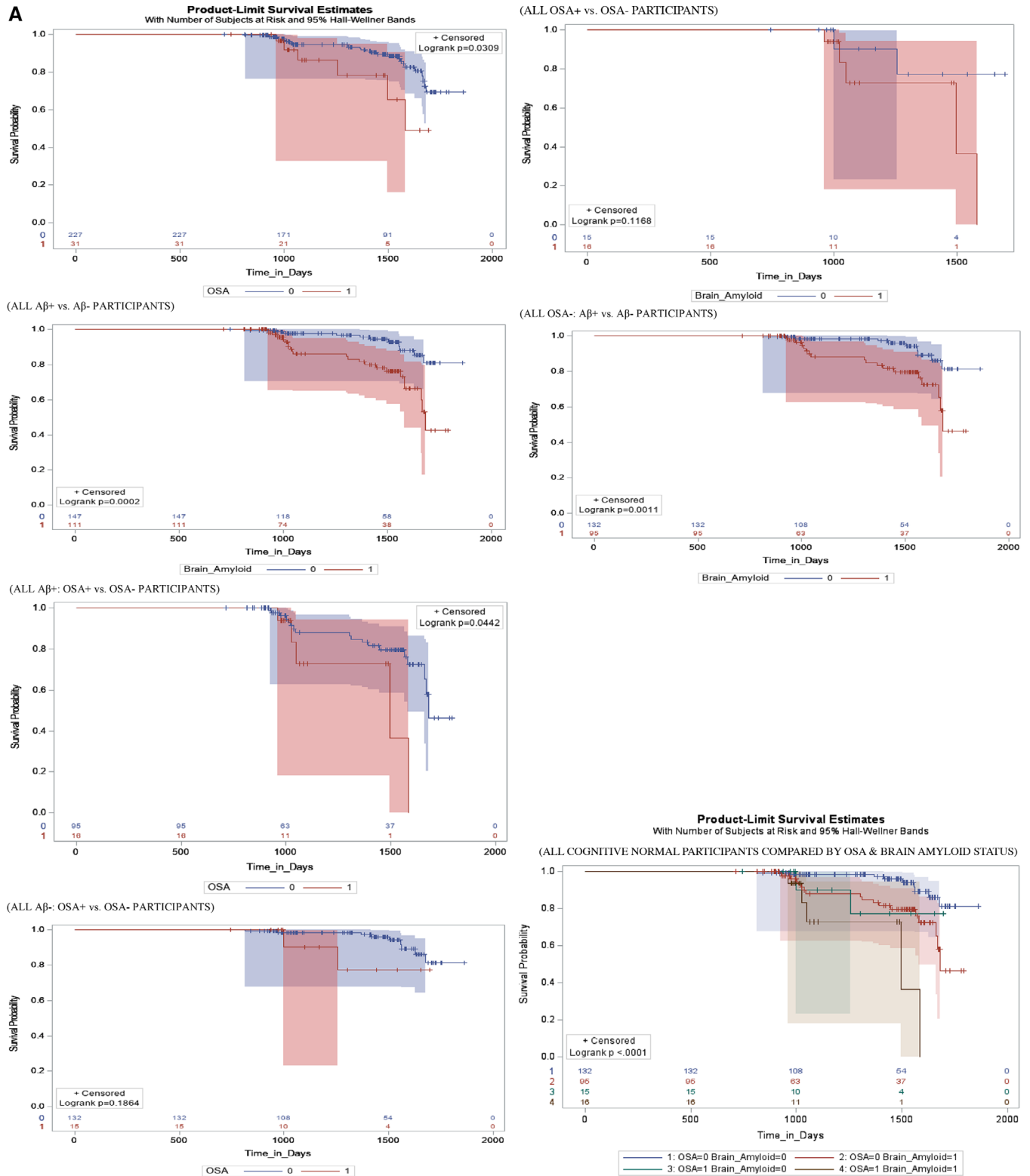
Characteristics	Table 2b: Characteristics of MCI participants by OSA and A/T/N status from MCI to Alzheimer's disease								
	All	Without OSA			With OSA				
		A+(TN) +	A+(TN) –	A-(TN) +	A-(TN) –	A+(TN) +	A-(TN) –		
Number of participants (%)	785 (100)	388 (49)	28 (4)	164 (20)	114 (15)	35 (4)	15 (2)	25 (3)	16 (2)
Follow-up time, mean (SD), years	5.7 (1.9)	5.8 (1.5)	5.6 (1.3)	6.0 (1.3)	5.9 (1.1)	5.5 (1.7)	5.7 (1.3)	5.6 (1.5)	5.7 (1.2)
Female sex, number (%)	385 (49)	210 (55)	26 (7)	63 (16)	39 (10)	20 (5)	7 (2)	11 (3)	9 (2)

**TABLE 2** Continued.

Characteristics	Table 2a: Characteristics of cognitively normal participants by OSA and A/T/N status						
	All	Without OSA			With OSA		
		A+(TN) +	A+(TN) -	A-(TN) +	A-(TN) -	A+(TN) +	A-(TN) -
Age, years, median (interquartile range)	74 (68, 79)	76 (71, 80)	71 (66, 76)	70 (65, 76)	71 (68, 77)	76 (71, 82)	70 (63, 75)
APOE positive, number (%) <sup>*</sup>	395 (50)	34 (9)	32 (8)	23 (6)	30 (6)	2 (1)	10 (3)
Education, years, median (interquartile range) <sup>*</sup>	16 (14, 18)	16 (14, 20)	16 (14, 18)	16 (14, 18)	16 (12, 19)	13 (11, 17)	17 (16, 19)
BMI (kg/m <sup>2</sup> )	26.9 ± 4.6	26.6 ± 4.7	28.0 ± 5.3	27.5 ± 4.5	30.9 ± 6.4	28.0 ± 2.4	29.0 ± 5.2
Hypertension, number (%)	369 (47)	28 (8)	61 (17)	57 (15)	26 (7)	4 (1)	18 (5)
Diabetes, number (%)	79 (10)	6 (8)	11 (14)	10 (13)	12 (15)	4 (5)	6 (7)
Thyroid Disease, number (%)	165 (21)	8 (5)	39 (24)	17 (10)	12 (7)	5 (3)	2 (1)
Respiratory Disease, number (%)	188 (24)	8 (4)	19 (10)	23 (12)	22 (12)	2 (1)	25 (14)
Cardiovascular Disease, number (%)	502 (64)	33 (7)	140 (28)	61 (12)	39 (8)	5 (1)	21 (4)
TBI, number (%)	31 (4)	10 (32)	14 (45)	8 (13)	4 (14)	1 (3)	2 (6)
Alcohol use, number (%)	63 (8)	5 (8)	24 (38)	8 (13)	2 (3)	2 (3)	6 (9)
CPAP use, number (%)	31 (4)	18 (29)	24 (38)	8 (13)	8 (26)	2 (5)	18 (58)
MMSE median (interquartile range)	27 (24, 28)	26 (25, 28)	27 (25, 29)	28 (25, 29)	26 (24, 28)	26 (24, 28)	27 (24, 29)
CDR median (interquartile range)	0.5 (0.5, 0.5)	0.5 (0.5, 0.5)	0.5 (0.5, 0.5)	0.5 (0.5, 0.5)	0.5 (0.5, 0.5)	0.5 (0.5, 0.5)	0.5 (0.5, 0.5)
CSF-ABETA pg/mL median (interquartile range) <sup>*</sup>	153 (130, 209)	178 (131, 214)	221 (202, 249)	219 (202, 239)	139 (120, 156)	145 (138, 151)	206 (189, 229)
TAU pg/mL median (interquartile range) <sup>*</sup>	80 (54, 116)	115 (77, 143)	66 (50, 79)	40 (30, 47)	103 (66, 114)	46 (37, 55)	44 (29, 51)
PTAU pg/mL median (interquartile range) <sup>*</sup>	36 (23, 51)	55 (37, 68)	32 (25, 38)	16 (13, 19)	51 (32, 58)	20 (19, 21)	18 (15, 20)
Aβ <sub>42</sub> median (interquartile range) <sup>*</sup>	1.2 (1.0, 1.4)	1.4 (1.3, 1.5)	1.3 (1.1, 1.3)	1.0 (1.0, 1.0)	1.4 (1.2, 1.4)	1.3 (1.1, 1.4)	1.0 (1.0, 1.0)

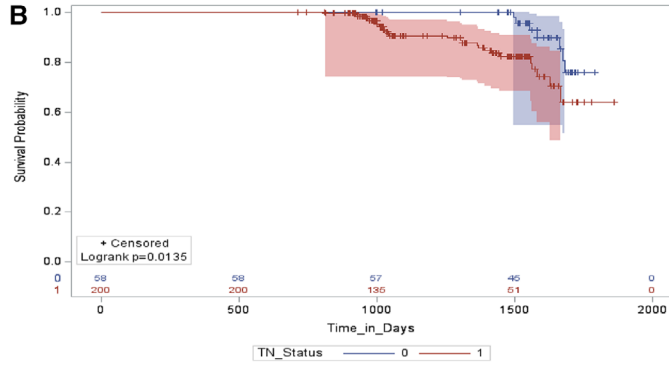
Abbreviations: ApoE4, apolipoprotein E ε4; Aβ, amyloid beta; BMI, body mass index; CDR, Clinical Dementia Rating; CPAP, continuous pulmonary airway pressure; CSF, cerebrospinal fluid; MCI, mild cognitive impairment; MMSE, Mini-Mental State Examination; OSA, obstructive sleep apnea; PTAU, phosphorylated tau; TAU, tau protein; TBI, traumatic brain injury. Notes: All columns show column percent where indicated. OSA and Aβ<sub>42</sub> columns show row percent where indicated. Superscript \* indicates significant differences between groups. Aβ<sub>42</sub> + participants had significantly lower CSF-Aβ<sub>42</sub>, and higher levels of CSF-t-tau and CSF-p-tau. Amyloid positive subjects in both CN and MCI groups were more likely to be APOE ε4 carriers. Participants who were OSA+/Aβ<sub>42</sub>+ had significantly higher BMI in the CN group but not in the MCI group, compared to other participants with varying OSA and Aβ status. In both CN and MCI, OSA+/Aβ<sub>42</sub>+ participants had higher vascular burden (ie, higher rates of hypertension, diabetes and cardiovascular disease) compared to OSA-/Aβ<sub>42</sub>+ participants (eg, 9/15 [60%] vs 38/95 [40%] for hypertension in CN participants). CN/OSA+/Aβ<sub>42</sub>+ participants had significantly lower educational level. Significance level p ≤ .05.



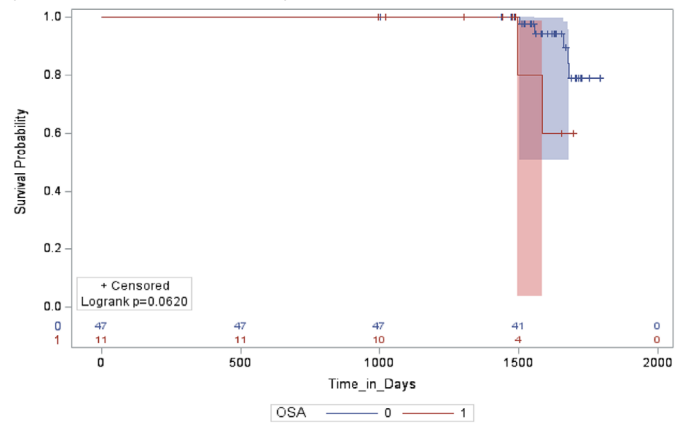


**FIGURE 1** Kaplan-Meier product limit survival estimates and time-to-progression from CN to MCI. A, (All OSA+ vs OSA- participants); (all Aβ+ vs Aβ- participants); (all OSA+: Aβ+ vs Aβ- participants); (all OSA-: Aβ+ vs Aβ- participants); (all Aβ+: OSA+ vs OSA- participants); (all Aβ-: OSA+ vs OSA- participants); (all CN participants compared by OSA & brain amyloid status). B, (all TN+ vs TN- participants); (all TN+: OSA+ vs OSA- participants); (all TN-: OSA+ vs OSA- participants); (all OSA+: TN+ vs TN- participants); (all OSA-: TN+ vs TN- participants); (all TN+ participants compared by OSA & brain amyloid status); (all TN- participants compared by OSA & brain amyloid status). Abbreviations: Aβ, amyloid beta; CN, cognitively normal; MCI, mild cognitive impairment; OSA, obstructive sleep apnea; TN, tau neurodegeneration

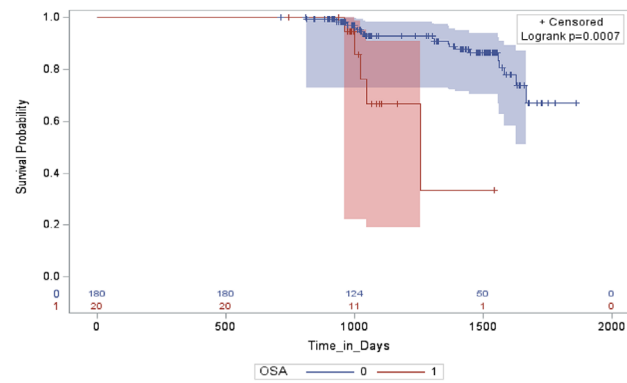
(ALL TN+ vs. TN- PARTICIPANTS)



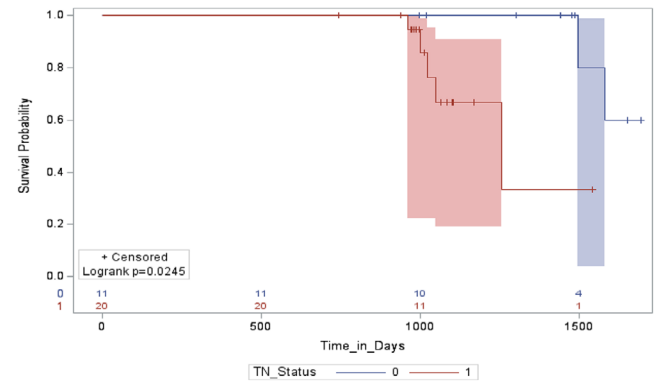
(ALL TN-: OSA+ vs. OSA- PARTICIPANTS)



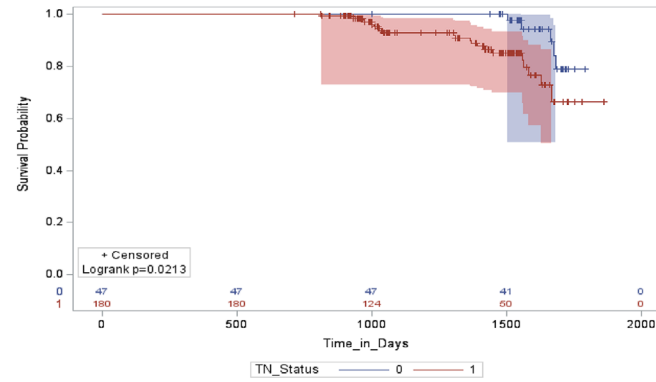
(ALL TN+: OSA+ vs. OSA- PARTICIPANTS)



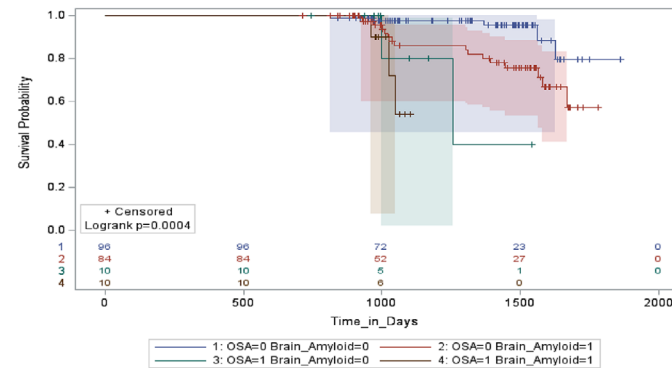
(ALL OSA+: TN+ vs. TN- PARTICIPANTS)



(ALL OSA-: TN+ vs. TN- PARTICIPANTS)



(ALL TN+ PARTICIPANTS COMPARED BY OSA & BRAIN AMYLOID STATUS)



(ALL TN- PARTICIPANTS COMPARED BY OSA & BRAIN AMYLOID STATUS)

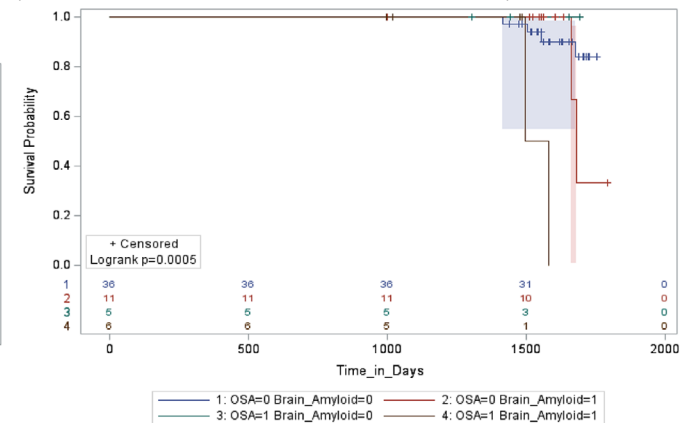
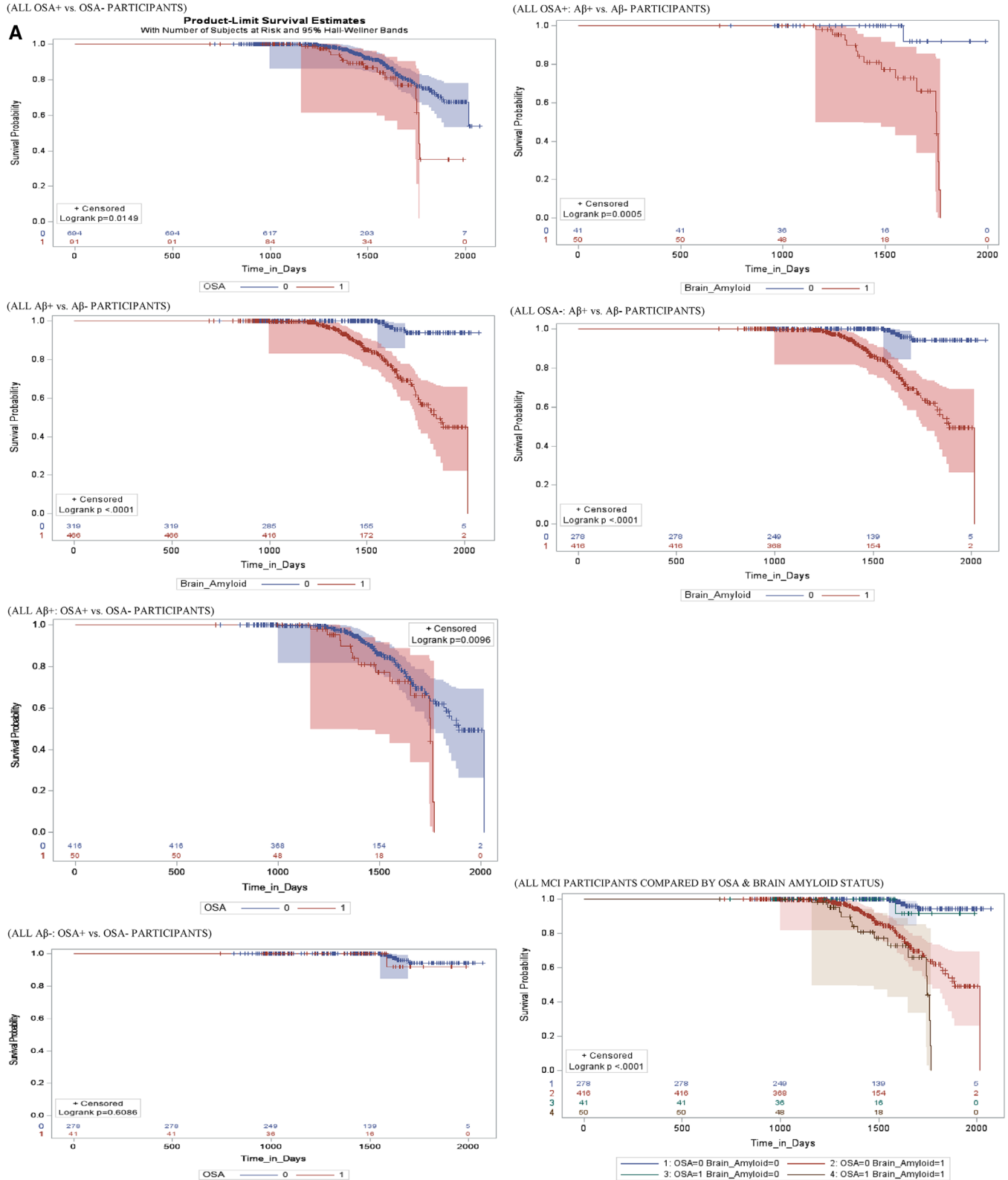


FIGURE 1 Continued



**FIGURE 2** Kaplan–Meier product limit survival estimates and time-to-progression from MCI to AD. A, (OSA+ vs OSA– participants); (all Aβ+ vs Aβ– participants); (all OSA+: Aβ+ vs Aβ– participants); (all OSA–: Aβ+ vs Aβ– participants); (all Aβ+: OSA+ vs OSA– participants); (all Aβ–: OSA+ vs OSA– participants); (all CN participants compared by OSA & brain amyloid status). B, (all TN+ vs TN– participants); (all TN+: OSA+ vs OSA– participants); (all TN–: OSA+ vs OSA– participants); (all OSA+: TN+ vs TN– participants); (all OSA–: TN all vs TN– participants); (all TN+ participants compared by OSA & brain amyloid status); (all TN– participants compared by OSA & brain amyloid status). Abbreviations: Aβ, amyloid beta; AD, Alzheimer’s disease; CN, cognitively normal; MCI, mild cognitive impairment; OSA, obstructive sleep apnea; TN, tau neurodegeneration

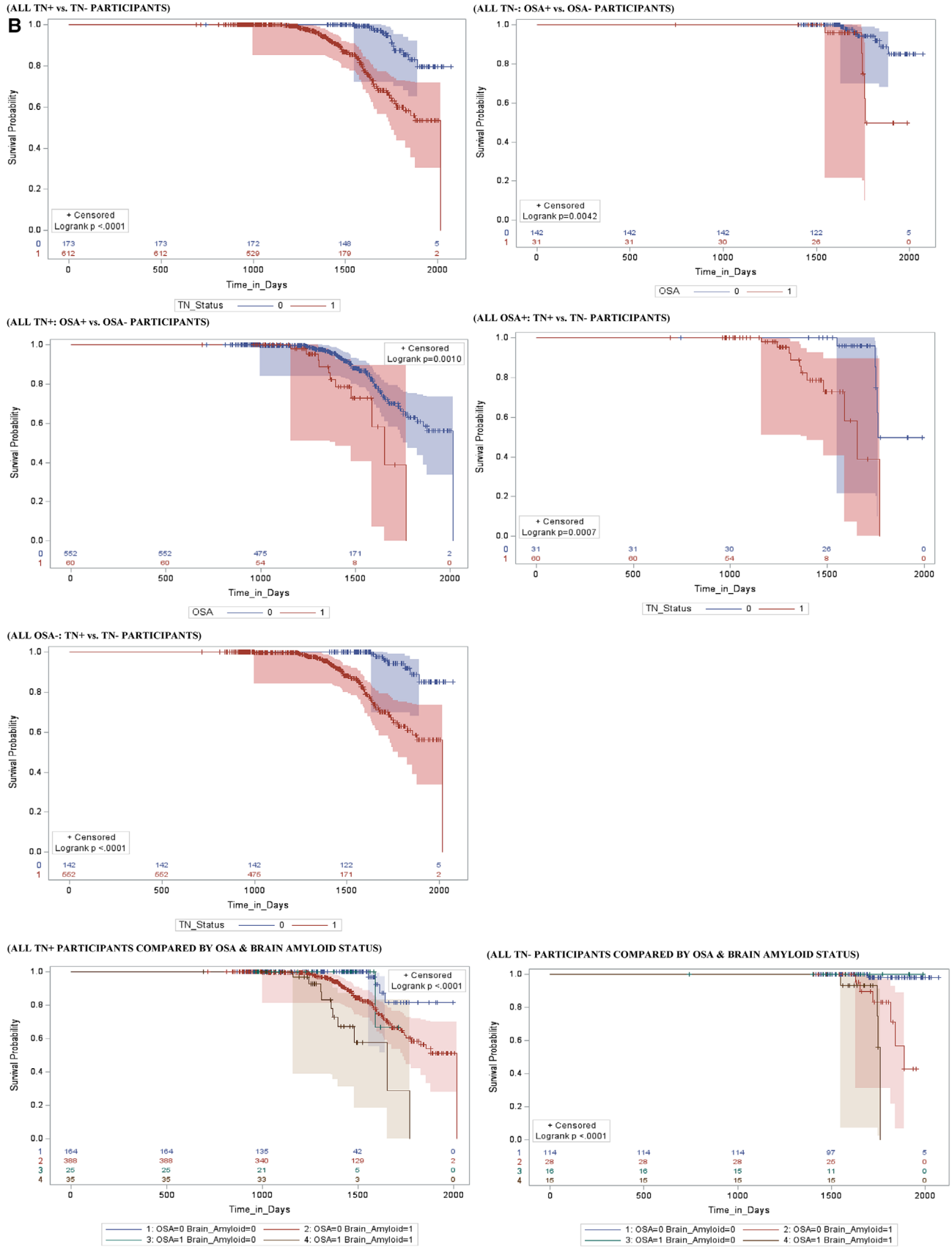


FIGURE 2 Continued

**TABLE 3** Cox proportional hazard models' estimates of the effect of OSA and A $\beta$  load on AD progression in CN and MCI participants from ADNI

Characteristics N	Mild cognitive impairment n (%)	Mean time-to-MCI Days $\pm$ SD (Years)	Median time-to-MCI Days (Years)	Model 1 HR 95% CI	P-value	Model 2 HR 95% CI	P-value
Cox proportional relative hazard of progression from cognitive normal to mild cognitive impairment.							
<b>OSA+ versus OSA- participants</b>							
OSA+ (N = 31)	11 (35)	1638 $\pm$ 105 (4.5)	1462 (4.0)	<b>1.78 (1.58, 1.98)</b>	.007	<b>1.32 (1.11, 1.65)</b>	<.01
OSA- (N = 227)	60 (26)	1833 $\pm$ 181 (5.0)	1745 (4.8)	REF		REF	
<b>A<math>\beta</math>+ versus A<math>\beta</math>- participants</b>							
A $\beta$ + (N = 111)	47 (42)	1325 $\pm$ 183 (3.6)	956 (2.6)	<b>2.96 (2.71, 3.22)</b>	<.001	<b>2.44 (1.99, 2.89)</b>	<.001
A $\beta$ - (N = 147)	24 (16)	1543 $\pm$ 170 (4.2)	1084 (3.0)	REF		REF	
<b>Amyloid positive (A<math>\beta</math>+) participants by OSA status</b>							
OSA+ (N = 16)	8 (49)	1288 $\pm$ 151 (3.5)	928 (2.5)	<b>3.61 (3.24, 3.97)</b>	<.001	<b>2.93 (2.17, 3.69)</b>	<.001
OSA- (N = 95)	39 (41)	1402 $\pm$ 125 (3.8)	1275 (3.5)	REF		REF	
<b>Amyloid negative (A<math>\beta</math>-) participants by OSA status</b>							
OSA+ (N = 15)	3 (17)	1421 $\pm$ 207 (3.9)	1206 (3.3)	1.05 (0.31, 1.79)	.11	1.01 (0.76, 1.26)	.11
OSA- (N = 132)	21 (16)	1497 $\pm$ 167 (4.1)	1475 (4.0)	REF		REF	
<b>OSA positive (OSA+) participants by A<math>\beta</math> load</b>							
A $\beta$ + (N = 16)	8 (49)	1288 $\pm$ 151 (3.5)	928 (2.5)	2.55 (0.98, 4.12)	.12	2.16 (0.87, 3.45)	.12
A $\beta$ - (N = 15)	3 (17)	1421 $\pm$ 207 (3.9)	1206 (3.3)	REF		REF	
<b>OSA negative (OSA-) participants A<math>\beta</math> load</b>							
A $\beta$ + (N = 95)	39 (41)	1402 $\pm$ 125 (3.8)	1275 (3.5)	<b>2.98 (2.66, 3.30)</b>	<.001	<b>2.47 (2.06, 2.88)</b>	<.001
A $\beta$ - (N = 132)	21 (16)	1497 $\pm$ 167 (4.1)	1475 (4.0)	REF		REF	
Cox proportional relative hazard of progression from mild cognitive impairment to Alzheimer's disease.							
<b>OSA+ versus OSA- participants</b>							
OSA+ (N = 91)	39 (43)	1678 $\pm$ 117 (4.6)	1521 (4.2)	<b>2.78 (2.25, 3.31)</b>	.01	<b>2.47 (1.79, 3.15)</b>	.01
OSA- (N = 694)	259 (37)	2196 $\pm$ 109 (6.0)	1937 (5.3)	REF		REF	
<b>A<math>\beta</math>+ versus A<math>\beta</math>- participants</b>							
A $\beta$ + (N = 466)	210 (43)	1381 $\pm$ 171 (3.8)	1096 (3.0)	<b>3.03 (2.61, 3.46)</b>	<.001	<b>2.62 (2.17, 3.07)</b>	<.001
A $\beta$ - (N = 319)	88 (28)	1526 $\pm$ 169 (4.2)	1268 (3.5)	REF		REF	
<b>Amyloid positive (A<math>\beta</math>+) participants by OSA status</b>							
OSA+ (N = 50)	27 (54)	1323 $\pm$ 141 (3.6)	1008 (2.8)	<b>3.53 (3.12, 3.94)</b>	.001	<b>2.78 (2.22, 3.34)</b>	<.01
OSA- (N = 416)	183 (44)	1720 $\pm$ 136 (4.7)	1316 (3.6)	REF		REF	
<b>Amyloid negative (A<math>\beta</math>-) participants by OSA status</b>							
OSA+ (N = 41)	12 (29)	1637 $\pm$ 102 (4.5)	1416 (3.9)	1.25 (0.67, 1.83)	.07	1.17 (0.86, 1.48)	.07
OSA- (N = 278)	76 (27)	1872 $\pm$ 117 (5.1)	1468 (4.0)	REF		REF	
<b>OSA positive (OSA+) participants by A<math>\beta</math> load</b>							
A $\beta$ + (N = 50)	27 (54)	1323 $\pm$ 141 (3.6)	1008 (2.8)	<b>2.21 (2.06, 2.36)</b>	<.001	<b>2.16 (2.04, 2.32)</b>	<.001
A $\beta$ - (N = 41)	12 (29)	1637 $\pm$ 102 (4.5)	1416 (3.9)	REF		REF	
<b>OSA negative (OSA-) participants by A<math>\beta</math> load</b>							
A $\beta$ + (N = 416)	183 (44)	1720 $\pm$ 136 (4.7)	1316 (3.6)	<b>3.05 (2.64, 3.46)</b>	<.0001	<b>2.55 (2.13, 2.98)</b>	<.001
A $\beta$ - (N = 278)	76 (27)	1872 $\pm$ 117 (5.1)	1468 (4.0)	REF		REF	

Abbreviations: ADNI, Alzheimer's Disease Neuroimaging Initiative; ApoE4, apolipoprotein E  $\epsilon$ 4; A $\beta$ , amyloid beta; BMI, body mass index; CI, confidence interval; CPAP, continuous pulmonary airway pressure; CSF, cerebrospinal fluid; MCI, mild cognitive impairment; OSA, obstructive sleep apnea; PTAU, phosphorylated tau; SD, standard deviation; TAU, tau protein; TBI, traumatic brain injury, TIA, transient ischemic attack.

Notes: Model I: Adjusted for age, sex, education, body mass index, baseline biomarker data and ApoE  $\epsilon$ 4 status.

Model II: Adjusted for age, sex, BMI, education, CPAP use, ApoE  $\epsilon$ 4 status, alcohol use, baseline biomarker data, hypertension, diabetes, history of cardiovascular disease (eg, including ischemic heart disease, heart failure, and stroke/TIA), and history of traumatic brain injury. Significance level  $p \leq .05$ .



Stratifying by OSA status, OSA+/A $\beta$ + participants showed no significant difference in time-to-progression to MCI (mean  $\pm$  SD [median] 3.5  $\pm$  0.4 [3.5] years vs 3.9  $\pm$  0.6 [3.3] years,  $P = .49$  and aHR: 2.16, 95% CI: 0.87–3.45,  $P = .12$ ), relative to OSA+/A $\beta$ – participants. However, OSA+/TN+ participants demonstrated significant difference in time-to-progression to MCI (mean  $\pm$  SD [median] 3.3  $\pm$  0.3 [2.8] years vs 4.5  $\pm$  0.4 [4.0] years, and aHR: 3.31, 95% CI: 1.36 – 5.27,  $P = .03$ ), relative to OSA+/TN– participants. Relative to OSA–/A $\beta$ – and OSA–/TN– participants, OSA–/A $\beta$ + and OSA–/TN+ participants had a significantly shorter time-to-progression to MCI (mean  $\pm$  SD [median] 3.8  $\pm$  0.3 [2.9] years vs 4.1  $\pm$  0.5 [4.0] years,  $P = .05$  and mean  $\pm$  SD [median] 3.8  $\pm$  0.3 [3.0] years vs 5.3  $\pm$  0.6 [4.5] years,  $P = .03$ , respectively), and a significantly higher risk of developing MCI (aHR: 2.47, 95% CI: 2.06–2.88,  $P < .001$  and aHR: 3.46, 95% CI: 1.78–5.14,  $P = .02$ , respectively).

Stratifying by TN status OSA+/A $\beta$ +/TN+ participants had a significantly shorter time-to-progression to MCI (mean  $\pm$  SD [median] 2.8  $\pm$  0.4 [2.5] years vs a combined 3.9  $\pm$  0.4 [3.2] years,  $P < .01$ ) and a significantly higher risk of developing MCI (aHR: 3.17, 95% CI: 1.19 – 5.15,  $P < .01$ ) compared to other TN+ participants combined (ie, OSA+/A $\beta$ –/TN+, OSA–/A $\beta$ +/TN+, and OSA–/A $\beta$ –/TN+). Furthermore, OSA+/A $\beta$ +/TN– participants had a significantly shorter time-to-progression to MCI (mean  $\pm$  SD [median] 4.1  $\pm$  0.4 [4.0] years vs a combined 5.0  $\pm$  0.5 [4.5] years,  $P < .01$ ) and a significantly higher risk of developing MCI (aHR: 1.49, 95% CI: 1.06 – 2.84,  $P < .01$ ) compared to other TN– participants combined (ie, OSA+/A $\beta$ –/TN–, OSA–/A $\beta$ +/TN– and OSA–/A $\beta$ –/TN–).

### 3.3 | Time-dependent progression risk and cumulative hazard function estimates in MCI group

Figure 2 shows survival and cumulative hazard function estimates and their 95% Hall-Wellner bands populated for the MCI group. Compared to OSA– participants, OSA+ participants had a significantly shorter time-to-progression to AD (mean  $\pm$  SD [median] 4.6  $\pm$  0.3 [4.2] years vs 6.0  $\pm$  0.3 [5.3] years,  $P = .01$ ), and a significantly higher risk of developing AD (aHR: 2.47, 95% CI: 1.79–3.15,  $P = .01$ ). Compared to A $\beta$ – participants, A $\beta$ + participants had a significantly shorter time-to-progression to AD (mean  $\pm$  SD [median] 3.8  $\pm$  0.5 [2.8] years vs 4.2  $\pm$  0.5 [3.6] years,  $P < .001$ ), and a significantly higher risk of developing AD (aHR: 2.62, 95% CI: 2.17–3.07,  $P < .001$ ). Compared to TN– participants, TN+ participants had a significantly shorter time-to-progression to MCI (mean  $\pm$  SD [median] 4.0  $\pm$  0.3 [3.8] years vs 4.9  $\pm$  0.4 [4.6] years,  $P < .001$ ), and a significantly higher risk of developing MCI (aHR: 3.34, 95% CI: 1.79–4.89,  $P < .001$ ). See Table 4.

#### 3.3.1 | Interactive associations of OSA, A $\beta$ , and tau burden with risk of conversion in CN and MCI

In CN participants, conversion risk from CN to MCI was associated with self-reported OSA ( $\beta = 0.42$ ; 95% CI, 0.13–0.70;  $P < .01$ ), higher A $\beta$  bur-

den ( $\beta = 0.55$ ; 95% CI, 0.22–0.89;  $P < .001$ ), and higher tau burden ( $\beta = 1.2$ ; 95% CI, 0.63–1.77;  $P < .001$ ). The interactions of self-reported OSA and A $\beta$  burden, and self-reported OSA, A $\beta$ , and tau burden with time were significant ( $\beta = 1.13$ , 95% CI, 0.74–1.52;  $P < .001$  and  $\beta = 1.38$ , 95% CI, 0.99–1.76;  $P < .001$ , respectively), suggesting a synergistic effect. However, the interaction of self-reported OSA and tau burden with time was not significant ( $\beta = 0.82$ , 95% CI, –0.11–1.32;  $P = .07$ ) suggesting that the presence of OSA did not modify the relationship between tau and cognitive decline in CN participants. See Table S1 in supporting information.

In MCI participants, conversion risk from MCI to AD was associated with self-reported OSA ( $\beta = 0.84$ ; 95% CI, 0.49–1.18;  $P < .01$ ), higher A $\beta$  burden ( $\beta = 1.01$ ; 95% CI, 0.58–1.45;  $P < .001$ ), and higher tau burden ( $\beta = 1.23$ ; 95% CI, 0.57–1.68;  $P < .001$ ). The interactions of self-reported OSA and A $\beta$  burden, self-reported OSA and tau burden, and self-reported OSA, A $\beta$ , and tau burden with time were significant ( $\beta = 1.18$ , 95% CI, 0.82–1.54;  $P < .001$ ,  $\beta = 1.31$ , 95% CI, 0.87–1.47;  $P < .001$  and  $\beta = 1.39$ , 95% CI, 0.95–1.75;  $P < .001$ , respectively), suggesting a synergistic effect. See Table S1.

#### 3.3.2 | Sensitivity analysis removing CPAP users

Sensitivity analysis removing CPAP users (CN:  $n = 8$  [A $\beta$ + = 6, A $\beta$ – = 2]; MCI:  $n = 31$  [A $\beta$ + = 10, A $\beta$ – = 21]) from OSA+ participants had negligible impact on the estimates (eg, CN OSA+/A $\beta$ + vs OSA+/A $\beta$ – aHR: 2.16, 95% CI: 0.8 – 3.45,  $P = .12$  changed to aHR: 2.21, 95% CI: 0.92–3.51,  $P = .09$  and MCI A $\beta$ +/OSA+ vs A $\beta$ + / OSA– aHR: 2.78, 95% CI: 2.22–3.34,  $P < .001$  changed to aHR: 2.82, 95% CI: 2.31–3.33,  $P < .001$ ). We attribute this negligible impact on the findings to data showing low CPAP compliance rates (<50%) with majority of CPAP dropouts occurring early in treatment, and relatively fewer patients discontinuing use as time with CPAP increases.<sup>32,33</sup> Moreover, the extremely small sample size of history of CPAP users prevented stratified analyses examining the beneficial effects of CPAP especially in the CN group. Subgroup analyses as was conducted in this study precluded testing the effects of CPAP on cognitive deterioration in this ADNI group as some groups had zero participant.

### 3.4 | DISCUSSION

The major objective of this study was to evaluate the effect of OSA on AD time-dependent progression risk in older CN and MCI elderly participants from the ADNI cohort and to quantify its effect on the risk of progression as A $\beta$  and tau accumulation become increasingly abnormal. Our major findings were the following. (1) Both CN and MCI OSA+ participants had a significantly increased risk and shorter time-to-progression to MCI and AD, respectively, compared to OSA– participants. (2) Both CN and MCI A $\beta$ + and TN+ participants, respectively, had a significantly increased risk and shorter time-to-progression to MCI and AD compared to A $\beta$ – and TN– participants. (3) Among only A $\beta$ + and TN+ participants, respectively, both CN and MCI OSA+

**TABLE 4** Cox proportional hazard models' estimates of the effect of obstructive sleep apnea and tau and neurodegeneration load on Alzheimer's disease time-dependent progression risk in older CN and MCI elderly participants from the ADNI cohort

Characteristics N	Mild cognitive impairment n (%)	Mean time-to-MCI	Median time-to-MCI	Model 1		Model 2	
		Days ± SD (Years)	Days (Years)	HR 95% CI	P-value	HR 95% CI	P-value
<b>Cox proportional relative hazard of progression from cognitive normal to mild cognitive impairment.</b>							
<b>TN+ versus TN- participants</b>							
TN+ (N = 200)	64 (32)	1254 ± 128 (3.4)	914 (2.5)	<b>3.75 (2.12, 5.35)</b>	.001	<b>3.52 (1.89, 5.17)</b>	.013
TN- (N = 58)	7 (12)	1754 ± 201 (4.7)	1554 (4.3)	REF		REF	
<b>TN+ participants by OSA status</b>							
OSA+ (N = 20)	9 (45)	1194 ± 107 (3.3)	1019 (2.8)	<b>2.36 (1.15, 3.58)</b>	<.001	<b>2.04 (1.11, 2.97)</b>	<.001
OSA- (N = 180)	55 (31)	1382 ± 121 (3.8)	1084 (3.0)	REF		REF	
<b>TN- participants by OSA status</b>							
OSA+ (N = 11)	2 (18)	1652 ± 152 (4.5)	1460 (4.0)	2.61 (1.00, 4.71)	.05	2.33 (0.57, 4.09)	.06
OSA- (N = 47)	5 (11)	1917 ± 208 (5.3)	1625 (4.5)	REF		REF	
<b>OSA positive (OSA+) participants by TN status</b>							
TN+ (N = 20)	9 (45)	1194 ± 107 (3.3)	1019 (2.8)	<b>3.55 (1.71, 5.39)</b>	.02	<b>3.31 (1.36, 5.27)</b>	.03
TN- (N = 11)	2 (18)	1652 ± 152 (4.5)	1460 (4.0)	REF		REF	
<b>OSA negative (OSA-) participants by TN status</b>							
TN+ (N = 180)	55 (31)	1382 ± 121 (3.8)	1084 (3.0)	<b>3.54 (1.98, 5.15)</b>	.02	<b>3.46 (1.78, 5.14)</b>	.02
TN- (N = 47)	5 (11)	1917 ± 208 (5.3)	1625 (4.5)	REF		REF	
<b>All TN+ participants OSA &amp; Aβ load</b>							
OSA+/Aβ+ (N = 10)	5 (50)	1019 ± 151 (2.8)	904 (2.5)	<b>3.22 (1.36, 5.08)</b>	<.001	<b>3.17 (1.19, 5.15)</b>	<.001
OSA+/Aβ- (N = 10)	4 (40)	1312 ± 102 (3.6)	1051 (2.9)	REF			
OSA-/Aβ+ (N = 84)	32 (38)	1367 ± 142 (3.7)	1013 (2.8)				
OSA-/Aβ- (N = 96)	23 (24)	1597 ± 147 (4.4)	1455 (4.0)			REF	
<b>All TN- participants OSA &amp; Aβ load</b>							
OSA+/Aβ+ (N = 6)	2 (33)	1496 ± 152 (4.1)	1471 (4.0)	<b>1.58 (1.04, 3.12)</b>	<.001	<b>1.49 (1.06, 2.84)</b>	<.001
OSA+/Aβ- (N = 5)	0 (0)			REF			
OSA-/Aβ+ (N = 11)	2 (18)	1752 ± 172 (4.8)	1675 (4.6)				
OSA-/Aβ- (N = 36)	3 (8)	1847 ± 167 (5.1)	1725 (4.7)			REF	
<b>Cox proportional relative hazard of progression from mild cognitive impairment to Alzheimer's disease.</b>							
Characteristics N	Alzheimer's disease n (%)	Mean time-to-MCI	Median time-to-MCI	Model 1		Model 2	
		Days ± SD (Years)	Days (Years)	HR 95% CI	P-value	HR 95% CI	P-value
<b>TN+ versus TN- participants</b>							
TN+ (N = 612)	269 (44)	1478 ± 127 (4.0)	1376 (3.8)	<b>3.57 (2.03, 5.11)</b>	<.001	<b>3.34 (1.79, 4.89)</b>	<.001
TN- (N = 173)	29 (17)	1795 ± 129 (4.9)	1697 (4.6)	REF		REF	
<b>TN+ participants by OSA status</b>							
OSA+ (N = 60)	32 (53)	1377 ± 211 (3.8)	1356 (3.7)	<b>2.93 (1.11, 4.76)</b>	.001	<b>2.72 (1.17, 4.27)</b>	.001
OSA- (N = 552)	237 (43)	1776 ± 109 (4.9)	1652 (4.5)	REF		REF	
<b>TN- participants by OSA status</b>							
OSA+ (N = 31)	7 (23)	1753 ± 181 (4.8)	1648 (4.5)	<b>1.78 (1.14, 3.42)</b>	.001	<b>1.48 (1.09, 2.87)</b>	.004
OSA- (N = 142)	22 (15)	1879 ± 206 (5.1)	1789 (4.9)	REF		REF	

**TABLE 4** Continued.

Characteristics N	Mild cognitive impairment n (%)	Mean time-to-MCI	Median time-to-MCI	Model 1		Model 2	
		Days ± SD (Years)	Days (Years)	HR 95% CI	P-value	HR 95% CI	P-value
<b>OSA positive (OSA+) participants by OSA status</b>							
TN+ (N = 60)	32 (53)	1377 ± 211 (3.8)	1356 (3.7)	<b>3.55 (1.67, 5.43)</b>	<.001	<b>3.37 (1.86, 4.88)</b>	<.001
TN- (N = 31)	7 (23)	1753 ± 181 (4.8)	1648 (4.5)	REF		REF	
<b>OSA negative (OSA-) participants by Aβ load</b>							
TN+ (N = 552)	237 (43)	1776 ± 109 (4.9)	1652 (4.5)	<b>2.81 (1.05, 4.57)</b>	<.001	<b>2.76 (1.11, 4.41)</b>	<.001
TN- (N = 142)	22 (15)	1879 ± 206 (5.1)	1789 (4.9)	REF		REF	
<b>All TN+ participants by OSA &amp; Aβ load</b>							
OSA+/Aβ+ (N = 35)	26 (74)	1262 ± 122 (3.5)	1197 (3.3)	<b>3.98 (2.26, 5.70)</b>	<.001	<b>3.47 (1.96, 4.98)</b>	<.001
OSA+/Aβ- (N = 25)	6 (24)	1656 ± 167 (4.5)	1596 (4.4)	REF			
OSA-/Aβ+ (N = 388)	174 (45)	1581 ± 206 (4.3)	1524 (4.2)				
OSA-/Aβ- (N = 164)	63 (38)	1697 ± 154 (4.7)	1583 (4.3)			REF	
<b>All TN- participants by OSA &amp; Aβ load</b>							
OSA+/Aβ+ (N = 15)	7 (47)	1596 ± 155 (4.4)	1483 (4.1)	<b>1.56 (1.07, 3.05)</b>	<.001	<b>1.23 (1.03, 2.43)</b>	<.001
OSA+/Aβ- (N = 16)	0 (0)			REF			
OSA-/Aβ+ (N = 28)	13 (46)	1801 ± 156 (4.9)	1717 (4.7)				
OSA-/Aβ- (N = 114)	9 (8)	2190 ± 115 (6.0)	1979 (5.4)			REF	

Abbreviations: ADNI, Alzheimer's Disease Neuroimaging Initiative; ApoE4, apolipoprotein E ε4; Aβ, amyloid beta; BMI, body mass index; CI, confidence interval; CPAP, continuous pulmonary airway pressure; CSF, cerebrospinal fluid; MCI, mild cognitive impairment; OSA, obstructive sleep apnea; PTAU, phosphorylated tau; SD, standard deviation; TAU, tau protein; TBI, traumatic brain injury, TIA, transient ischemic attack.

Notes: Model I: Adjusted for age, sex, education, body mass index and ApoE ε4 status.

Model II: Adjusted for age, sex, BMI, education, CPAP use, ApoE ε4 status, alcohol use, baseline biomarker data, hypertension, diabetes, history of cardiovascular disease (eg, including ischemic heart disease, heart failure, and stroke/TIA), and history of traumatic brain injury

participants had a significantly increased risk and shorter time-to-progression to MCI and AD compared to OSA- participants. (4) Among only Aβ- and TN- participants, respectively, there was no difference in the risk and time-to-progression to MCI between CN OSA+ versus OSA- participants. This association varied in MCI OSA+ versus OSA- participants, with significantly increased risk and shorter time-to-progression to AD seen only in TN- participants. (5) Among only OSA+ participants, only MCI Aβ+ participants had a significantly increased risk and shorter time-to-progression to AD, respectively compared to Aβ- participants. Both CN and MCI TN+ participants had a significantly increased risk and shorter time-to-progression to AD, respectively, compared to TN- participants. (6) Among only OSA- participants, both CN and MCI Aβ+ and TN+ participants had a significantly increased risk and shorter time-to-progression to MCI and AD, respectively, compared to Aβ- and TN- participants, respectively. (7) In both CN and MCI participants, the interactions of self-reported OSA and Aβ burden, and self-reported OSA, Aβ, and tau burden with time were significant, suggesting a synergistic effect. However, in CN participants, the interaction of self-reported OSA and tau burden with time was not significant.

This is the first study showing a shorter progression time to MCI/AD in both CN and MCI OSA+ participants, respectively. This finding is consistent with our previous study showing that OSA patients had

an earlier onset age to MCI or AD.<sup>2</sup> In this study, both CN and MCI OSA+ participants progressed to MCI and AD, respectively, 6 to 8 months earlier than did OSA- participants. In addition, our recent study showed both CN and MCI OSA+ subjects experiencing faster annual increase in florbetapir uptake and decrease in CSF Aβ<sub>42</sub> levels, as well as increases in CSF t-tau and p-tau compared to OSA- participants,<sup>16</sup> thereby suggesting that OSA appears to accelerate increases in amyloid deposition, CSF t-tau, and p-tau levels over time, both in CN and MCI individuals, thus possibly significantly reducing the time to MCI or AD progression.<sup>2</sup> Our results on OSA increasing AD progression risk are consistent with previous prospective studies indicating that individuals with OSA have an elevated risk of developing MCI or AD.<sup>1,3,4,34,35</sup>

Our results showing Aβ load increasing MCI or AD progression risk in both CN and MCI participants are consistent with well-established findings from previous studies showing that CN and MCI elderly participants with positive PiB-PET and low CSF Aβ<sub>42</sub> show associations with cognitive decline,<sup>36,37</sup> and have an elevated risk of AD progression, respectively.<sup>38</sup> Our results showing Aβ load predicting a shorter progression time to MCI/AD in both CN and MCI participants are consistent with well-established findings from previous studies showing Aβ+ CN and MCI subjects more likely to progress to MCI<sup>39,40</sup> and AD,<sup>41-43</sup> in short-term follow-up than Aβ- CN and MCI individuals, respectively.

Our tau findings showing tau accumulation and neurodegeneration increasing MCI or AD progression risk and predicting a shorter progression time to MCI/AD are in line with established evidence of strong associations between cortical neurofibrillary tangle load and cognitive impairment.<sup>44</sup>

In both A $\beta$ + CN and MCI participants (ie, with evidence of AD pathologic change), the risk of progression to MCI and AD was approximately two and three times that of A $\beta$ - participants, respectively. In both TN+ CN and MCI participants (ie, evidence of tau accumulation and neurodegeneration), the risk of progression to MCI and AD was approximately three and four times that of TN- participants, respectively (regardless of OSA status in TN+ CN participants). The combined risk of progressing to MCI and AD in the CN and MCI OSA+/A $\beta$ + and MCI OSA+/TN+ participants was approximately three times that of CN and MCI OSA-/A $\beta$ + participants, and four to five times that of MCI OSA-/TN+ participants, respectively. In addition, the combined risk of progression to MCI and AD among CN and MCI OSA+/A $\beta$ + and OSA+/TN+ participants was approximately three and four times that of OSA-/A $\beta$ - and OSA-/TN- participants, respectively ( $P < .01$ ). The interactions of self-reported OSA and A $\beta$  burden and self-reported OSA and TN burden with time were significant, thereby suggesting that OSA's effect on MCI/AD progression risk is: (1) independently synergistic with A $\beta$  and tau, and (2) significantly increases as A $\beta$  and tau accumulation becomes increasingly abnormal. The literature suggests that intermittent hypoxia<sup>45,46</sup> and sleep fragmentation<sup>47,48</sup> that are causes for excessive daytime sleepiness (EDS), are two main processes by which OSA may induce neurodegenerative changes and promote the accumulation of A $\beta_{42}$ . However, we also know that chronic intermittent hypoxia, hypercapnia, and hypertension in OSA can also induce neuronal damage, including axons,<sup>49</sup> white matter,<sup>50</sup> and reduced diffusion tensor imaging based mean diffusivity in multiple brain regions.<sup>51</sup> This suggests that OSA could promote neurotoxicity that is independent of hypoxia-induced A $\beta_{42}$  accumulation, thereby resulting in possible synergistic neurodegenerative mechanisms with A $\beta_{42}$  accumulation on AD progression risk. Notably, the synergistic effect of being OSA+/TN+ was absent in CN group. However the effect of being either OSA+/A $\beta$ + or OSA+/TN+ was more marked in the MCI group (eg, median time-to-AD progression of 3 years in A $\beta$ + vs 2.8 years in MCI OSA+/A $\beta$ + and median time-to-MCI progression of 2.6 years in A $\beta$ + vs 2.5 years in OSA+/A $\beta$ + in the CN group,  $P < .01$  for all), thereby lending credence to our hypothesis that OSA's effect on MCI/AD progression risk significantly increases as A $\beta$  and tau accumulation becomes increasingly abnormal.

We did not detect significant differences in progression risk when we compared CN and MCI OSA+ versus OSA- among only A $\beta$ - participants and CN OSA+ versus OSA- among only TN- participants. These analyses examined whether established effects of OSA's neurodegenerative effect that is independent of hypoxia-induced A $\beta_{42}$  accumulation was sufficient to induce cognitive decline. We favor a model of AD that implies that one of the contributory roles of OSA is the potential for neuronal injury independent of A $\beta$  in which chronic intermittent hypoxia, and hypercapnia, may induce axonal, glial, or white matter damage, in multiple brain regions.<sup>49,50,52</sup> Plausible expla-

nations for these findings are discussed below. First, this analysis used self-report for clinical diagnosis of OSA. Possible misclassification of OSA status occurring more frequently in one of the A $\beta$  or TN groups could lead to fewer OSA patients considered to have progressed to MCI/AD therefore biasing the risk toward the null. Second, our inability to determine OSA severity could also be another reason. In our previous study,<sup>15</sup> it was OSA severity that was associated with increases in brain amyloid burden. Third, our previous study using ADNI data<sup>16</sup> showed largely in CN and MCI (early stage subjects) groups that self-reported OSA+ subjects experienced faster annual increase in florbetapir uptake and decrease in CSF-A $\beta_{42}$  levels as well as increases in CSF t-tau and p-tau compared to self-reported OSA- participants, thereby suggesting that OSA+ subjects are more likely to convert to A $\beta$ + and TN+ statuses. Therefore, OSA effects may not be specific to the presence of A $\beta$  or tau; rather it may affect different risk profiles based on the disease stage examined. As such it is not that A $\beta$ - subjects are protected from progression to MCI or AD, but rather that there is a stepwise progression whereby, first, OSA increases risk for transition from A $\beta$ - to A $\beta$ +, and, once A $\beta$ +, increases risk for developing cognitive decline. Fourth, independent of A $\beta$  and tau pathology, progression to MCI/AD may be dependent on additional factors such as comorbid hypertension and microvascular changes,<sup>53,54</sup> all of which were adjusted for in our analyses. Fifth, it could be that OSA's effect on MCI/AD progression risk is only apparent at certain A $\beta$  or TN threshold levels or that longer follow-up times are needed. As such, CN A $\beta$ - and TN- participants each may be too early in the process and power may have been an issue for this time scale and sample size. Last, MCI A $\beta$ - participants may represent a less pure population of subjects on an AD trajectory and are generally less likely to progress to clinical AD diagnosis. Because pathological definition of AD requires substantial presence of A $\beta$ , histopathological examination may actually reveal these individuals may have other diagnoses (eg, depression) or might be at risk for other forms of dementia, such as a diagnosis of primary age related tauopathy.

Our result showing increased risk and a shorter time-to-progression to AD in MCI A $\beta$ + participants compared to A $\beta$ - participants among only OSA+ participants is consistent with well-established findings of MCI A $\beta$ + elderly participants having elevated risk of AD<sup>38,55</sup> and in short-term follow-up, being more likely to progress to AD than MCI A $\beta$ - participants.<sup>41-43</sup> The median time to AD was 2.8 years in OSA+/A $\beta$ + participants compared to 3.9 years in OSA+/A $\beta$ - participants. This clearly suggests the effects of OSA on AD progression risk significantly increases as A $\beta$  accumulation becomes increasingly abnormal. However, these results should be interpreted with caution as A $\beta$ - MCI ADNI patients have shown a variety of clinical and biomarker features that differ from their A $\beta$ + counterparts, suggesting that one or more non-AD etiologies (which may include vascular disease and depression) account for their AD-like phenotype.<sup>56</sup> There was no significant difference in the risk and time-to-progression to MCI in CN A $\beta$ + participants compared to A $\beta$ - participants mainly due to lack of statistical power in the subgroup analyses.

The OSA- subset analysis, comparing A $\beta$ + versus A $\beta$ - participants' risk and shorter time-to-progression to MCI and AD, respec-

tively, in CN and MCI participants revealed findings that are consistent with the OSA+ subset analysis comparing A $\beta$ + versus A $\beta$ - participants. Both eliminate OSA's effect. Our findings from examining A $\beta$ 's effect independent of OSA positivity (ie, OSA-/A $\beta$ + vs OSA-/A $\beta$ -), and comparing that effect to A $\beta$ 's effect contingent on OSA positivity (ie, OSA+/A $\beta$ + vs OSA+/A $\beta$ -), helped to further highlight OSA's synergism with amyloid burden with respect to MCI/AD progression risk. For example, the median time-to-MCI/AD progression in MCI OSA-/A $\beta$ + participants was 3.6 years versus 4 years in MCI OSA-/A $\beta$ - participants, while progression time in MCI OSA+/A $\beta$ + participants was 2.8 years versus 3.9 years in MCI OSA+/A $\beta$ - participants.

The analysis showing statistical comparison of progression times of each category (OSA+/A $\beta$ +, OSA+/A $\beta$ -, OSA-/A $\beta$ +, OSA-/A $\beta$ -) within the two subpopulations (CN and MCI), as well as the respective TN+ and TN- subset analyses, each comparing OSA+/A $\beta$ + participants to other TN+ and TN- participants combined (ie, OSA+/A $\beta$ -, OSA-/A $\beta$ +, and OSA-/A $\beta$ -) further helped to highlight OSA and A $\beta$  synergism. The latter analysis eliminated the effects of tau. OSA+/A $\beta$ + participants were approximately two to four times more likely to progress to MCI or AD and had significantly shorter time-to-progression to MCI and AD in CN and MCI participants, respectively, depending on whether they were TN+ or TN-. In addition, results from the mixed effects models examining the interactive associations of OSA, A $\beta$ , and tau burden with risk of conversion to MCI/AD demonstrated that OSA had a synergistic effect with A $\beta$  in both CN and MCI participants, and with tau only in MCI participants, thereby suggesting that OSA and tau's synergistic effect is observed and markedly increased with growing susceptibility to tau accumulation.

Strengths of our study include a well-characterized cohort, longitudinal design, relatively long follow-up, objective assessment of amyloid and tau burden, and robust statistical analytic methods and large enough sample that allowed subgroup examinations for the most part. As we previously described,<sup>2,16</sup> measurement of OSA by self-report is an important limitation. Self-reported sleep measures can be impacted by diminished cognition<sup>57</sup> and in certain situations might not be correlated with objective methods.<sup>58</sup> The significantly lower than expected prevalence of reported OSA in this elderly cohort is possibly due to underdiagnosis, as epidemiological and sleep laboratory studies document much higher OSA prevalence in elderly populations. The prevalence of OSA (with or without symptoms) is estimated at 30% to 50% in older subjects;<sup>59</sup> therefore, misclassification of some OSA+ subjects into the OSA- group could have occurred; however, this would have driven our findings toward the null, with lower estimates than the true ones, therefore attenuating OSA's true effect. OSA classification by self-report suggests that those with self-reported OSA were more likely to be symptomatic (ie, with EDS), prompting these subjects to seek diagnosis. Therefore, further research differentiating the risk of OSA for AD with and without associated daytime symptoms is necessary. Notably, a recent study demonstrated that all-cause EDS defined by Epworth sleepiness scores  $\geq 10$  was associated with longitudinal brain A $\beta$  accumulation in elderly subjects.<sup>60</sup>

## CONCLUSION

Our findings in a cohort of self-reported OSA patients support the hypothesis of an overarching model of late-onset AD with brain amyloid deposition and tau aggregates proceeding at different rates,<sup>61</sup> influenced by a combination of protective/risk factors, of which OSA is part. This model of AD implies a contributory role of OSA, first with the direct potential for neuronal injury independent of A $\beta$  by inducing intermittent hypoxia, sleep fragmentation, arousal-induced hypertensive surges, systemic inflammation, and impaired glucose handling<sup>62</sup> irrespective of MCI/AD progression risk. Second, there may be an indirect contributory role of OSA in which acute and intermittent hypoxia, sleep fragmentation, and EDS may accelerate A $\beta$  accumulation in the presence of A $\beta$  plaques via a feedback loop<sup>8,46,60</sup> and tau accumulation possibly influenced by impaired clearance through the glymphatic pathway.<sup>63,64</sup> This role of OSA significantly increases MCI/AD progression risk as suggested by our findings. Third, there may be a contributory role of OSA in which the direct neurotoxicity effect that is independent of A $\beta$  accumulation together with OSA's indirect effect that promotes A $\beta$  accumulation, combine to act synergistically to significantly increase MCI/AD progression risk, possibly affecting and/or accelerating AD biomarker change and leading to shorter time to MCI/AD in OSA CN and MCI participants, respectively. This OSA-A $\beta$  synergism related to cognitive decline can be independent of tau as well as synergistic with tau deposition. Future research using objective measures of OSA is needed to replicate these findings and examine OSA's effects on slow wave and rapid eye movement sleep and their mediating role in increasing A $\beta$  and tau accumulation.

## ACKNOWLEDGMENTS

The authors would like to gratefully acknowledge the dedication, time, and commitment of the participants and staff of the Alzheimer's Disease Neuroimaging Initiative. The National Institute on Health funded this study: NIA: L30-AGO64670 (Bubu), Columbia Center for Interdisciplinary Research on Alzheimer's disease Disparities (CIRAD) P30AG059303 Pilot (Bubu), R21AG049348 (Osorio), R21AG055002 (Osorio), R01AG056031 (Osorio), R21AG059179 (Varga), R01AG056682 (Varga), R01AG056531 (Osorio, Jean-Louis), NHLBI: T32HL129953 (Jean-Louis, Ogedegbe), K07AG05268503 (Jean-Louis), R01HL118624 (Osorio), K23HL125939 (Williams), K24HL109156 (Ayappa).

## CONFLICTS OF INTEREST

The authors have no conflicts of interest to declare.

## REFERENCES

1. Chang WP, Liu ME, Chang WC, et al. Sleep apnea and the risk of dementia: a population-based 5-year follow-up study in taiwan. *PLoS One*. 2013;8(10):e78655.
2. Osorio RS, Gumb T, Pirraglia E, et al. Sleep-disordered breathing advances cognitive decline in the elderly. *Neurology*. 2015;84(19):1964-1971.
3. Yaffe K, Laffan AM, Harrison SL, et al. Sleep-disordered breathing, hypoxia, and risk of mild cognitive impairment and dementia in older women. *JAMA*. 2011;306(6):613-619.



4. Yaffe K, Nettiksimmons J, Yesavage J, Byers A. Sleep quality and risk of dementia among older male veterans. *Am J Geriatr Psychiatry*. 2015;23(6):651-654.
5. Bubu OM, Brannick M, Mortimer J, et al. Sleep, cognitive impairment, and Alzheimer's disease: a systematic review and meta-analysis. *Sleep*. 2017 Jan 1;40(1). PMID: 28364458 DOI: <https://doi.org/10.1093/sleep/zsw032>.
6. Shi L, Chen SJ, Ma MY, et al. Sleep disturbances increase the risk of dementia: A systematic review and meta-analysis. *Sleep Med Rev*. 2018;40:4-16.
7. Elias A, Cummins T, Tyrrell R, et al. Risk of Alzheimer's disease in obstructive sleep apnea syndrome: amyloid-beta and tau imaging. *J Alzheimers Dis*. 2018;66(2):733-741.
8. Ju YE, Finn MB, Sutphen CL, et al. Obstructive sleep apnea decreases central nervous system-derived proteins in the cerebrospinal fluid. *Ann Neurol*. 2016;80(1):154-159.
9. Liguori C, Mercuri NB, Izzi F, et al. Obstructive sleep apnea is associated with early but possibly modifiable Alzheimer's disease biomarkers changes. *Sleep*. 2017 May 1;40(5). doi: <https://doi.org/10.1093/sleep/zsx011>.
10. Osorio RS, Ayappa I, Mantua J, et al. The interaction between sleep-disordered breathing and apolipoprotein E genotype on cerebrospinal fluid biomarkers for Alzheimer's disease in cognitively normal elderly individuals. *Neurobiol Aging*. 2014;35(6):1318-1324.
11. Spira AP, Yager C, Brandt J, et al. Objectively Measured sleep and beta-amyloid burden in older adults: a pilot study. 2014 Aug 12;2:2050312114546520. doi: <https://doi.org/10.1177/2050312114546520>.
12. Yun CH, Lee HY, Lee SK, et al. Amyloid burden in obstructive sleep apnea. *J Alzheimers Dis*. 2017;59(1):21-29.
13. André C, Rehel S, Kuhn E, et al. Association of sleep-disordered breathing with Alzheimer disease biomarkers in community-dwelling older adults: a secondary analysis of a randomized clinical trial. *JAMA Neurol*. 2020;77(6):1-10.
14. Carvalho DZ, St Louis EK, Schwarz CG, et al. Witnessed apneas are associated with elevated tau-PET levels in cognitively unimpaired elderly. *Neurology*. 2020;94(17):e1793-e1802.
15. Sharma RA, Varga AW, Bubu OM, et al. Obstructive sleep apnea severity affects amyloid burden in cognitively normal elderly: a longitudinal study. *Am J Respir Crit Care Med*. 2018;197(7):933-943.
16. Bubu OM, Pirraglia E, Andrade AG, et al. Obstructive sleep apnea and longitudinal Alzheimer's disease biomarker changes. *Sleep*. 2019;42(6):zsz048.
17. Brys M, Pirraglia E, Rich K, et al. Prediction and longitudinal study of CSF biomarkers in mild cognitive impairment. *Neurobiol Aging*. 2009;30(5):682-690.
18. Mattsson N, Zetterberg H, Hansson O, et al. CSF biomarkers and incipient Alzheimer disease in patients with mild cognitive impairment. *JAMA*. 2009;302(4):385-393.
19. Aizenstein HJ, Nebes RD, Saxton JA, et al. Frequent amyloid deposition without significant cognitive impairment among the elderly. *Arch Neurol*. 2008;65(11):1509-1517.
20. Sperling RA, Laviolette PS, O'Keefe K, et al. Amyloid deposition is associated with impaired default network function in older persons without dementia. *Neuron*. 2009;63(2):178-188.
21. Petersen RC, Aisen PS, Beckett LA, et al. Alzheimer's Disease Neuroimaging Initiative (ADNI): clinical characterization. *Neurology*. 2010;74(3):201-209.
22. McKhann G, Drachman D, Folstein M, Katzman R, Price D, Stadlan EM. Clinical diagnosis of Alzheimer's disease: report of the NINCDS-ADRDA Work Group under the auspices of Department of Health and Human Services Task Force on Alzheimer's Disease. *Neurology*. 1984;34(7):939-944.
23. Landau SM, Breault C, Joshi AD, et al. Amyloid-beta imaging with Pittsburgh compound B and florbetapir: comparing radiotracers and quantification methods. *J Nucl Med*. 2013;54(1):70-77.
24. Landau SM, Lu M, Joshi AD, et al. Comparing positron emission tomography imaging and cerebrospinal fluid measurements of beta-amyloid. *Ann Neurol*. 2013;74(6):826-836.
25. Vanderstichele H, DMG SF, et al. *Biomarkers for Early Diagnosis of Alzheimer's Disease*. Hauppauge, NY: Nova Science Publishers; 2008.
26. Shaw LM, Vanderstichele H, Knapiak-Czajka M, et al. Cerebrospinal fluid biomarker signature in Alzheimer's disease neuroimaging initiative subjects. *Ann Neurol*. 2009;65(4):403-413.
27. Jack CR Jr, Bennett DA, Blennow K, et al. NIA-AA Research Framework: Toward a biological definition of Alzheimer's disease. *Alzheimers Dement*. 2018;14(4):535-562.
28. Jack CR Jr, Bennett DA, Blennow K, et al. A/T/N: An unbiased descriptive classification scheme for Alzheimer disease biomarkers. *Neurology*. 2016;87(5):539-547.
29. Toledo JB, Bjerke M, Da X, et al. Nonlinear association between cerebrospinal fluid and florbetapir F-18  $\beta$ -amyloid measures across the spectrum of Alzheimer disease. *JAMA Neurol*. 2015;72(5):571-581.
30. Suárez-Calvet M, Morenas-Rodríguez E, Kleinberger G, et al. Early increase of CSF sTREM2 in Alzheimer's disease is associated with tau related-neurodegeneration but not with amyloid- $\beta$  pathology. *Mol Neurodegener*. 2019;14(1).
31. Tan MS, Ji X, Li JQ, et al. Longitudinal trajectories of Alzheimer's ATN biomarkers in elderly persons without dementia. *Alzheimers Res Ther*. 2020;12(1):55.
32. Collard P, Pieters T, Aubert G, Delguste P, Rodenstein DO. Compliance with nasal CPAP in obstructive sleep apnea patients. *Sleep Med Rev*. 1997;1(1):33-44.
33. Loubé DI, Gay PC, Strohl KP, Pack AI, White DP, Collop NA. Indications for positive airway pressure treatment of adult obstructive sleep apnea patients: a consensus statement. *Chest*. 1999;115(3):863-866.
34. Lee JE, Yang SW, Ju YJ, Ki SK, Chun KH. Sleep-disordered breathing and Alzheimer's disease: A nationwide cohort study. *Psychiatry Res*. 2019;273:624-630.
35. Tsai MS, Li HY, Huang CG, et al. Risk of Alzheimer's disease in obstructive sleep apnea patients with or without treatment: Real-world evidence. *Laryngoscope*. 2020;130(9):2292-2298.
36. Fagan AM, Head D, Shah AR, et al. Decreased cerebrospinal fluid Abeta(42) correlates with brain atrophy in cognitively normal elderly. *Ann Neurol*. 2009;65(2):176-183.
37. Morris JC, Roe CM, Grant EA, et al. Pittsburgh compound B imaging and prediction of progression from cognitive normality to symptomatic Alzheimer disease. *Arch Neurol*. 2009;66(12):1469-1475.
38. Hansson O, Zetterberg H, Buchhave P, Londos E, Blennow K, Minthon L. Association between CSF biomarkers and incipient Alzheimer's disease in patients with mild cognitive impairment: a follow-up study. *Lancet Neurol*. 2006;5(3):228-234.
39. Fagan AM, Roe CM, Xiong C, Mintun MA, Morris JC, Holtzman DM. Cerebrospinal fluid tau/beta-amyloid(42) ratio as a prediction of cognitive decline in nondemented older adults. *Arch Neurol*. 2007;64(3):343-349.
40. Gustafson DR, Skoog I, Rosengren L, Zetterberg H, Blennow K. Cerebrospinal fluid beta-amyloid 1-42 concentration may predict cognitive decline in older women. *J Neurol Neurosurg Psychiatry*. 2007;78(5):461-464.
41. Forsberg A, Engler H, Almkvist O, et al. PET imaging of amyloid deposition in patients with mild cognitive impairment. *Neurobiol Aging*. 2008;29(10):1456-1465.
42. Okello A, Koivunen J, Edison P, et al. Conversion of amyloid positive and negative MCI to AD over 3 years: an 11C-PIB PET study. *Neurology*. 2009;73(10):754-760.

43. Wolk DA, Price JC, Saxton JA, et al. Amyloid imaging in mild cognitive impairment subtypes. *Ann Neurol*. 2009;65(5):557-568.
44. Nelson PT, Alafuzoff I, Bigio EH, et al. Correlation of Alzheimer disease neuropathologic changes with cognitive status: a review of the literature. *J Neuropathol Exp Neurol*. 2012;71(5):362-381.
45. Shiota S, Takekawa H, Matsumoto SE, et al. Chronic intermittent hypoxia/reoxygenation facilitate amyloid-beta generation in mice. *J Alzheimers Dis*. 2013;37(2):325-333.
46. Zhang X, Zhou K, Wang R, et al. Hypoxia-inducible factor 1alpha (HIF-1alpha)-mediated hypoxia increases BACE1 expression and beta-amyloid generation. *J Biol Chem*. 2007;282(15):10873-10880.
47. Lim AS, Kowgier M, Yu L, Buchman AS, Bennett DA. Sleep Fragmentation and the Risk of Incident Alzheimer's Disease and Cognitive Decline in Older Persons. *Sleep*. 2013;36(7):1027-1032.
48. Schlosser Covell GE, Dhawan PS, Lee Iannotti JK, et al. Disrupted daytime activity and altered sleep-wake patterns may predict transition to mild cognitive impairment or dementia: a critically appraised topic. *Neurologist*. 2012;18(6):426-429.
49. Ludemann P, Dziewas R, Soros P, Happe S, Frese A. Axonal polyneuropathy in obstructive sleep apnoea. *J Neurol Neurosurg Psychiatry*. 2001;70(5):685-687.
50. Kiernan TE, Capampangan DJ, Hickey MG, Pearce LA, Aguilar MI. Sleep apnea and white matter disease in hypertensive patients: a case series. *Neurologist*. 2011;17(5):289-291.
51. Kumar R, Chavez AS, Macey PM, Woo MA, Yan-Go FL, Harper RM. Altered global and regional brain mean diffusivity in patients with obstructive sleep apnea. *J Neurosci Res*. 2012;90(10):2043-2052.
52. Macey PM, Kumar R, Woo MA, Valladares EM, Yan-Go FL, Harper RM. Brain structural changes in obstructive sleep apnea. *Sleep*. 2008;31(7):967-977.
53. Esiri MM, Nagy Z, Smith MZ, Barnettson L, Smith AD. Cerebrovascular disease and threshold for dementia in the early stages of Alzheimer's disease. *Lancet*. 1999;354(9182):919-920.
54. Snowdon DA, Greiner LH, Mortimer JA, Riley KP, Greiner PA, Markesbery WR. Brain infarction and the clinical expression of Alzheimer disease. The Nun Study. *JAMA*. 1997;277(10):813-817.
55. Jack CR Jr, Wiste HJ, Vemuri P, et al. Brain beta-amyloid measures and magnetic resonance imaging atrophy both predict time-to-progression from mild cognitive impairment to Alzheimer's disease. *Brain*. 2010;133(11):3336-3348.
56. Landau SM, Horng A, Fero A, Jagust WJ. Amyloid negativity in patients with clinically diagnosed Alzheimer disease and MCI. *Neurology*. 2016;86(15):1377-1385.
57. Van Den Berg JF, Van Rooij FJ, Vos H, et al. Disagreement between subjective and actigraphic measures of sleep duration in a population-based study of elderly persons. *J Sleep Res*. 2008;17(3):295-302.
58. Unruh ML, Redline S, An MW, et al. Subjective and objective sleep quality and aging in the sleep heart health study. *J Am Geriatr Soc*. 2008;56(7):1218-1227.
59. Heinzer R, Marti-Soler H, Haba-Rubio J. Prevalence of sleep apnoea syndrome in the middle to old age general population. *Lancet Respir Med*. 2016;4(2):e5-6.
60. Carvalho DZ, St Louis EK, Knopman DS, et al. Association of Excessive Daytime Sleepiness With Longitudinal beta-Amyloid Accumulation in Elderly Persons Without Dementia. *JAMA Neurol*. 2018;75(6):672-680.
61. Jack CR Jr, Lowe VJ, Weigand SD, et al. Serial PIB and MRI in normal, mild cognitive impairment and Alzheimer's disease: implications for sequence of pathological events in Alzheimer's disease. *Brain*. 2009;132(Pt 5):1355-1365.
62. Lim DC, Veasey SC. Neural injury in sleep apnea. *Curr Neurol Neurosci Rep*. 2010;10(1):47-52.
63. Iliff JJ, Chen MJ, Plog BA, et al. Impairment of glymphatic pathway function promotes tau pathology after traumatic brain injury. *J Neurosci*. 2014;34(49):16180-16193.
64. Xie L, Kang H, Xu Q, et al. Sleep drives metabolite clearance from the adult brain. *Science*. 2013;342(6156):373-377.

#### SUPPORTING INFORMATION

Additional supporting information may be found online in the Supporting Information section at the end of the article.

**How to cite this article:** Bubu OM, Umasabor-Bubu OQ, Turner AD, et al. Self-reported obstructive sleep apnea, amyloid and tau burden, and Alzheimer's disease time-dependent progression. *Alzheimer's Dement*. 2021;17:226-245. <https://doi.org/10.1002/alz.12184>