Association of Longitudinal Changes in Cerebrospinal Fluid Total Tau and Phosphorylated Tau 181 and Brain Atrophy With Disease Progression in Patients With Alzheimer Disease

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Abstract

IMPORTANCE The amyloid/tau/neurodegeneration (A/T/N) framework uses cerebrospinal fluid (CSF) levels of total tau (tTau) as a marker of neurodegeneration and CSF levels of phosphorylated tau 181 (pTau181) as a marker of tau tangles. However, it is unclear whether CSF levels of tTau and pTau181 have similar or different trajectories over the course of Alzheimer disease.

OBJECTIVES To examine the rates of change in CSF levels of tTau and pTau181 across the Alzheimer disease course and how the rates of change are associated with brain atrophy as measured by magnetic resonance imaging.

DESIGN, SETTING, AND PARTICIPANTS This cohort study was set in tertiary research clinics. Each participant was a member of a pedigree with a known mutation for dominantly inherited Alzheimer disease. Participants were divided into 3 groups on the basis of the presence of a mutation and their Clinical Dementia Rating score. Data analysis was performed in June 2019.

MAIN OUTCOMES AND MEASURES Rates of change of CSF tTau and pTau181 levels and their association with the rate of change of brain volume.

RESULTS Data from 465 participants (283 mutation carriers and 182 noncarriers) were analyzed. The mean (SD) age of the cohort was 37.8 (11.3) years, and 262 (56.3%) were women. The mean (SD) follow-up duration was 2.7 (1.5) years. Two or more longitudinal CSF and magnetic resonance imaging assessments were available for 160 and 247 participants, respectively. Sixty-five percent of mutation carriers (183) did not have symptoms at baseline (Clinical Dementia Rating score, 0). For mutation carriers, the annual rates of change for CSF tTau and pTau181 became significantly different from 0 approximately 10 years before the estimated year of onset (mean [SE] rates of change, 5.5 [2.8] for tTau [P = .05] and 0.7 [0.3] for pTau181 [P = .04]) and 15 years before onset (mean [SE] rates of change, 5.4 [3.9] for tTau [P = .17] and 11 [0.5] for pTau181 [P = .03]), respectively. The rate of change of pTau181 was positive and increased at the early stages of the disease, showing a positive rate of change starting at 15 estimated years before onset until 5 estimated years before onset (mean [SE], 0.4 [0.3]), followed by a positive but decreasing rate of change at year 0 (mean [SE], 0.1 [0.3]) and then negative rates of change at 5 years (mean [SE], −0.3 [0.4]) and 10 years (mean [SE], −0.6 [0.6]) after symptom onset. In individuals without symptoms (Clinical Dementia Rating score, 0), the rates of change of CSF tTau and pTau181 were negatively associated with brain atrophy (high rates of change in CSF measures were associated with low rates of change in brain volume in asymptomatic stages). After symptom onset (Clinical Dementia Rating score, >0), an increased rate of brain atrophy was not associated with rates of change of levels of both CSF tTau and pTau181.

(continued)
CONCLUSIONS AND RELEVANCE  These findings suggest that CSF tTau and pTau181 may have different associations with brain atrophy across the disease time course. These results have implications for understanding the dynamics of disease pathobiology and interpreting neuronal injury biomarker concentrations in response to Alzheimer disease progression and disease-modifying therapies.

Introduction

The neuropathologic hallmarks of Alzheimer disease (AD) are the presence of neuritic amyloid plaques, primarily of the amyloid-β peptide, and the intraneuronal accumulation of neurofibrillary tangles (NFTs) and neuropil threads composed of hyperphosphorylated, aggregated tau protein. Our understanding of AD has evolved substantially over the past 2 decades, with neuroimaging and fluid biomarkers allowing for early detection of AD-related pathologic abnormalities. In fewer than 1% of patients, AD is caused by autosomal dominant mutations in either the presenilin 1, presenilin 2, or amyloid precursor protein genes. Dominantly inherited AD (DIAD) is considered clinically similar to sporadic AD except for a younger age at onset (AAO). Mutation carriers (MCs) have a somewhat predictable age at AD symptom onset.

Biomarker studies have contributed to hypothesized trajectories of fluid and imaging biomarker changes that occur over the course of the disease, from the preclinical phase to the end stages characterized by advanced dementia. However, models of biomarker change in AD have been based mostly on cross-sectional data. More recently, many models have classified biomarkers according to their proposed association with the biological underpinnings of the disease; the recent amyloid/tau/neurodegeneration (A/T/N) framework was developed to provide a more biological rationale to the classification of the disease. Phosphorylated tau 181 (pTau181) in the cerebrospinal fluid (CSF) has been suggested to represent NFT pathologic abnormalities, whereas total tau (tTau) in the CSF is thought to be a marker of neurodegeneration that is passively released with cell death or injury. Given these putative mechanisms, it might be expected that levels of both tTau and pTau181 would continue to become more abnormal with disease progression, as NFT pathologic abnormalities increase and neurodegeneration accelerates. However, recent longitudinal studies from the Dominantly Inherited Alzheimer Network (DIAN) and Alzheimer’s Disease Neuroimaging Initiative (ADNI) cohorts have challenged the linear model from previous cross-sectional studies, which have consistently found higher CSF levels of tTau and pTau181 as the disease progresses.

These findings highlight a need for accurate determination of the evolution of longitudinal changes in CSF levels of tTau and pTau181 and their association with disease progression. In the present study, we assessed the longitudinal pattern of changes in CSF levels of tTau and pTau181 and their association with brain atrophy as measured by magnetic resonance imaging (MRI). We hypothesized that if CSF tTau and pTau181 were passively released with neurodegeneration, they should be associated with MRI measures of neurodegeneration (e.g., rate of atrophy). To evaluate this hypothesis, we used a well-characterized cohort with DIAD from the DIAN study.

Methods

All participants were recruited as part of the DIAN study. Participants provided written informed consent or assent with proxy consent. The institutional review boards for each of the participating DIAN sites approved all aspects of the study. This study follows the Strengthening the Reporting of Observational Studies in Epidemiology (STROBE) reporting guideline.
For our analysis, 465 participants with at least 1 CSF measurement in the DIAN cohort were available from the data freeze 12 (last data from June 30, 2017). Each participant was a member of a pedigree with a known mutation for DIAD. Genotyping was performed to determine the genetic status for each participant at risk; the presence or absence of a DIAD mutation was determined using polymerase chain reaction–based amplification of the appropriate exon followed by Sanger sequencing methods on a genetic analyzer (ABI 3130xl; ThermoFisher Scientific).21

Participants were divided into 3 groups according to the presence of a mutation and their Clinical Dementia Rating (CDR)22 score at baseline, with a CDR score of 0 indicating no symptoms and a CDR score of 0.5 or higher indicating the presence of symptoms. The participant’s estimated years before expected symptom onset (EYO) was defined as the participant’s age at baseline minus their expected AAO.6,7,23 The AAO was calculated according to either the family mutation–specific expected age at dementia onset or parental age at first progressive cognitive decline if the expected AAO for the mutation was unknown. Negative values indicate that an individual is younger than his or her expected AAO.

Clinical and Neuropsychological Assessments
Participants underwent extensive clinical evaluation, which included family history of AD, personal medical history, and physical and neurological examination. Clinical dementia status was determined with the CDR in accordance with standard protocols and criteria.22,24 Clinicians performing the assessments were blinded to mutation status of participants. Full details of participating sites, enrollment, and assessments in DIAN have been published elsewhere.20

Brain Imaging
MRI Scanning
Participants underwent volumetric T1-weighted MRI, using the magnetization-prepared rapid acquisition with gradient-echo sequence defined in the ADNI second phase.25 Sites used a 3-T scanner and were required to pass regular quality control assessments. Volumetric segmentation and cortical surface reconstruction were performed using FreeSurfer image analyzing software version 5.3 (Harvard Medical School),26 and subcortical volumes were corrected for intracranial volume using a regression approach.27 Cortical thickness and volume measures were averaged across hemispheres because there were no a priori laterality predictions. Because the focus of our study was to assess the association between longitudinal CSF tTau and pTau181 levels with atrophy rate, a limited number of regions of interest (ROIs) reflecting brain atrophy patterns across AD stages were included. We followed an ROIs approach to include posterior areas with earlier tau deposition (eg, hippocampus, entorhinal cortex, parahippocampus, posterior cingulate gyrus, precuneus gyrus, and supramarginal gyrus) and atrophy vs anterior areas with later tau deposition (eg, superior frontal gyrus, orbital frontal gyrus, and midfrontal gyrus) and atrophy. The ROIs were defined according to areas of early vs late neurodegeneration and tau deposition in the literature and our previous work.3,14,28-31

Biochemical Analysis
The CSF was collected in the morning under fasting conditions by lumbar puncture and immediately placed on ice. Samples were shipped on dry ice to the DIAN biomarker core laboratory. We have previously reported a longitudinal decrease in CSF tTau and pTau181 levels using commercially available immunoassays.14 To also test whether those findings were specific to the method of measurement, here we used a fully automated, high-performance electroluminescence immunoassay (Elecsys; Roche Diagnostics) of tTau and pTau181 in the laboratory of Leslie Shaw, PhD, in the ADNI Biomarker Core at the University of Pennsylvania according to the kit manufacturer’s instructions and as described in previous studies.32-34 Analyses were performed in a series of runs, with each sample run 1 time for each of the tau analytes, over the period of October 18, 2017, through November 9, 2017; acceptance criteria as documented according to the Roche Protocol in the University of Pennsylvania ADNI Biomarker Laboratory were followed. In each of the 14 days of
performing analyses, the manufacturer's quality control results were within stated limits to meet acceptance criteria for precision and accuracy. There were 2 runs each day of approximately 40 samples per run. Precision performance was documented according to analysis of pristine aliquots of a CSF pool in each of the 28 runs for each of the biomarkers. The percentage coefficients of variation for this CSF quality control pool were less than 2%.

**Statistical Analysis**

Data analysis was performed in June 2019. Baseline characteristics of the participants are summarized as mean (SD) for continuous variables and number (column percentage) for categorical variables. P values for comparing the difference between MCs and noncarriers (NCs) were obtained using generalized linear mixed-effects models with random intercepts for family clusters to take into account the associations between participants within the same family; the tests were 2-sided. The annualized rate of change over the longitudinal follow-up period was estimated for each participant separately using linear regression and then plotted against baseline EYO to evaluate the trajectories of the biomarker changes over EYO. To visualize the differences in the rates of change of different biomarkers as a function of EYO, locally estimated scatterplot smoothing curves of the standardized difference of individual annual rates of change for MCs for different biomarkers were plotted against baseline EYO. A linear or linear spline mixed-effects model for each biomarker (depending on the curve fit) was then used to estimate and test the rates of change at each integer EYO point. The fixed effects in the models included mutation group (MC or NC), baseline EYO, time since baseline, and all possible 2-way or 3-way interactions. Sex, years of education, and apolipoprotein ε4 allele status were considered as covariates, but only significant effects were retained in the models. Random effects included in the models are random intercepts for family clusters, individual random intercept, and random slope. An unstructured covariance matrix was used to account for the within-participant association due to repeated measures. Further details for the linear and linear spline mixed-effects models can be found in the article by McDade et al.14

Associations between the individual annual rates of change of tTau and pTau181 concentrations and the annualized rates of brain atrophy were evaluated for MCs with symptoms (CDR score, >0) and MCs without symptoms (CDR score, 0) separately, using bivariate linear mixed-effects models with random intercepts for family clusters.35,36 Statistical analyses were conducted with the PROC MIXED and PROC NLMIX procedures in SAS statistical software version 9.4 (SAS Institute Inc). A P < .05 was considered to be statistically significant. For CSF biomarkers (145 NCs and 235 MCs), the effect size that could be detected with 80% power was 0.3. For MRI biomarkers (177 NCs and 270 MCs), the effect size that could be detected with 80% power was 0.27.

**Results**

We analyzed data from 465 participants, including 283 MCs (183 [65%] without symptoms and 100 with symptoms) and 182 NCs (Table 1). The mean (SD) age of the cohort was 37.8 (11.3) years, and 262 (56.3%) were women. Of the MCs, 213 (75.3%) had presenilin 1 mutations, 22 (7.8%) had presenilin 2 mutations, and 48 (16.9%) had amyloid precursor protein mutations. There were no differences between MCs and NCs in terms of age (mean [SD] age, 37.8 [10.8] vs 37.9 [11.7] years), sex (55.1% vs 58.2% female), educational level (mean [SD] years, 14.3 [3.0] vs 14.8 [2.9]), or the presence of at least 1 apolipoprotein ε4 allele (29.7% vs 30.8%). The EYO for the entire cohort ranged from 38.2 years before the parental AAO to 22.6 years after the parental AAO; however, to reduce the risk of identifying individual participants at the extremes of the EYO range, we show only the EYO interval of −25 to 10 years. The NCs had no or very little evidence of AD pathologic abnormalities and almost all (171 [94.0%]) had normal cognition. At baseline, 183 MCs (64.7%) did not have symptoms (CDR score, 0). The CDR scores of the MCs with symptoms ranged from 0.5 (very mild) to 3 (severe) (Table 1). Two or more longitudinal CSF and MRI assessments were available for 160 and 247...
participants, respectively (Table 1 and eTable 1 in the Supplement) with a mean (SD) follow-up of 2.7 (1.5) years.

**Longitudinal Change in CSF tTau and pTau181 Levels**

Previous analyses of this cohort have found that CSF tTau and pTau181 levels were increased in MCs 15 years before expected symptom onset (EYO = −15). We also examined the longitudinal change in CSF tTau and pTau181 levels in terms of annual rate of change across the EYO (Figure 1 and Table 2). Rates of change for NCs were not significantly different from 0. For MCs, the annual rates of change for CSF tTau and pTau181 became significantly different from 0 near EYO −10 (mean [SE] rates of change, 5.5 [2.8] for tTau [P = .05] and 0.7 [0.3] for pTau181 [P = .04]) and EYO −15 (mean [SE] rates of change, 5.4 [3.9] for tTau [P = .17] and 1.1 [0.5] for pTau181 [P = .03]), respectively (Table 2). Importantly, the longitudinal rates of change of CSF tTau and pTau181 levels depended on where the participant fell with respect to their EYO, and the pattern of change over the AD course was different for tTau and pTau181. Specifically, the positive rate of change of tTau gradually increased until EYO −10 and then remained constant after symptom onset (mean [SE] rate of change, 5.6 [2.3] for EYO 0, 5.6 [3.2] for EYO 5, and 5.7 [4.4] for EYO 10). In contrast, the positive rate of change of pTau181 increased in those at early stages of the disease, starting at EYO −15 until EYO −5 (mean [SE], 0.4 [0.3]), followed by a positive but decreasing rate of change at year 0 (mean [SE] 0.1 [0.3]) and then negative rates of change at EYO 5 (mean [SE], −0.3 [0.4]) and EYO 10 (mean [SE], −0.6 [0.6]) (Table 2), resulting in overall lower levels of pTau181 at later stages of disease.

### Table 1. Clinical, Cognitive, Imaging, and Biochemical Characteristics of Study Participants at Baseline*

| Clinical and Baseline Biomarker Characteristics | Mean (SD) | Noncarriers (n = 182) | Mutation Carriers (n = 283) | P Value |
|-----------------------------------------------|----------|-----------------------|-----------------------------|---------|
| Age, y                                        | 37.9 (11.7) | 37.8 (10.8) | .80 |
| Female, No. (%)                               | 106 (58.2) | 156 (55.1) | .50 |
| Education, y                                  | 14.8 (2.9) | 14.3 (3.0) | .70 |
| Mini-Mental State Examination score            | 29.0 (1.2) | 26.9 (5.1) | <.001 |
| Clinical Dementia Rating score, No. (%)       | 171 (94.0) | 183 (64.7) | <.001 |
| >0                                            | 11 (6.0) | 100 (35.3) |  
| Apolipoprotein ε4 allele carrier, No. (%)     | 56 (30.8) | 84 (29.7) | .80 |
| Estimated years before onset, y               | −9.7 (12.2) | −8.2 (11.0) | .20 |
| Length of follow-up, y                        | 3.4 (1.6) | 2.9 (1.5) | .03 |
| No. of longitudinal cerebrospinal fluid assessments, No. (%) | 55 (37.9) | 105 (42.9) |  
| 2                                            | 38 (69.1) | 69 (65.7) |  
| 3                                            | 12 (21.8) | 25 (23.8) | .24 |
| ≥4                                           | 5 (9.1) | 11 (10.5) |  
| Aβ42 level, pg/mL                             | 1385 (471) | 961 (623) | <.001 |
| Ratio of Aβ42 to Aβ40                         | 0.09 (0.01) | 0.06 (0.03) | <.001 |
| Total tau level, pg/mL                        | 171.6 (58.5) | 287.9 (159.1) | <.001 |
| Phosphorylated tau 181 level, pg/mL           | 14.2 (5.3) | 30.8 (22.7) | <.001 |
| Ratio of phosphorylated tau 181 to Aβ42      | 0.0109 (0.0057) | 0.0518 (0.0534) | <.001 |
| No. of longitudinal MRI assessments, No. (%)^ | 89 (50.3) | 158 (58.5) |  
| 2                                            | 63 (70.8) | 96 (60.8) |  
| 3                                            | 18 (20.2) | 47 (29.7) | .09 |
| ≥4                                           | 8 (9.0) | 15 (9.5) |  
| Hippocampal volume, mm³                        | 8798 (809) | 8363 (1302) | <.001 |
| Thickness, mm                                 | 3.50 (0.30) | 3.43 (0.36) | .04 |
| Entorhinal cortical                           | 2.38 (0.13) | 2.28 (0.22) | <.001 |
| Precuneus                                     | 2.66 (0.13) | 2.62 (0.17) | .02 |

Abbreviation: MRI, magnetic resonance imaging.

*Continuous measures are presented as the mean (SD). For each type of measure, the number of individuals with at least 1 measurement is provided. The significance of the difference between the mutation carriers and noncarriers was calculated by linear mixed-effects models with random intercepts for familial clusters.

^ A total of 160 participants underwent longitudinal cerebrospinal fluid assessments.

^- A total of 247 participants underwent longitudinal MRI assessments.
Longitudinal Change in Brain Volumes

The A/T/N framework uses CSF tTau levels and/or structural MRI measurements as a marker of neurodegeneration, so we sought to determine whether CSF tTau could be used as a suitable biomarker to track MRI atrophy rate. For MRI measurements in MCs, the individual annual rate of change in precuneus thickness started to show a difference from 0 approximately 20 years before expected symptom onset (mean [SE] rate of change, −0.008 [0.003]), followed by rate of change in hippocampal volume (mean [SE] rate of change, −53.5 [17.9]), entorhinal thickness (mean [SE] rate of change, −0.015 [0.007]), and superior-frontal thickness (mean [SE] rate of change, −0.010 [0.004]) around EYO −15. The longitudinal rate of change for all measured brain ROI increased 5

Figure 1. Individual Annual Rate of Change of Cerebrospinal Fluid Levels of Total Tau (tTau) and Phosphorylated Tau 181 (pTau181) and Brain Structures by Magnetic Resonance Imaging as a Function of Baseline Estimated Years Before Onset in Mutation Carriers and Noncarriers

Values for noncarriers (blue dots), mutation carriers without symptoms (orange dots), and mutation carriers with symptoms (gray dots) are shown. Blue and orange lines are locally estimated scatterplot smoothing curves for mutation carriers and noncarriers, respectively. To maintain participant and investigator blinding to mutation status when reporting individual data points, specific estimated years before onset are not shown (vertical dashed line indicates year 0). Individual annual rate of change over the longitudinal follow-up was estimated for each participant separately using linear regressions.
years before symptoms appeared (EYO = -5) and followed a similar trend through the disease (Table 2 and Figure 1). For all ROIs included in the analysis, the rate of change followed similar trends.

To visually compare the differences in the trajectories of the rates of change of CSF tTau and pTau181 levels and structural MRI measures, locally estimated scatterplot smoothing curves were constructed for the standardized rate of change as a function of baseline EYO (Figure 2). Notably, after EYO 0, the rate of change became more negative (structures atrophied more quickly), compared with CSF tTau levels, which continued to have a stable positive rate of change.

### Association Between CSF and MRI Measures of Neurodegeneration

We found differences in nearly all posterior and limbic or paralimbic regions for associations in rates of change with tTau and pTau181 according to stage of disease when comparing MCs without symptoms vs those with symptoms (eTable 2 and eTable 3 in the Supplement). Figure 3 shows patterns of correlation coefficients between the rates of change of CSF tTau and pTau181 and brain structure stratified by the absence or presence of symptoms (CDR score of 0 vs CDR score >0) and by posterior (early atrophy) and anterior (later atrophy) cortical or subcortical regions. In individuals without symptoms (CDR score of 0), the rates of change of CSF tTau and pTau181 were inversely

### Table 2. Estimated Annual Rate of Change of Cerebrospinal Fluid Levels of tTau and pTau181 and Structural Magnetic Resonance Imaging (Regions of Interest) in Mutation Carriers by Estimated Years Before Onset

| Estimated Time Before Onset, y | tTau Rate of Change, Mean (SE) | P Value | pTau181 Rate of Change, Mean (SE) | P Value | Entorhinal Thickness Rate of Change, Mean (SE) | P Value | Hippocampal Volume Rate of Change, Mean (SE) | P Value | Precuneus Thickness Rate of Change, Mean (SE) | P Value | Superior Frontal Thickness Rate of Change, Mean (SE) | P Value |
|-------------------------------|-------------------------------|--------|---------------------------------|--------|---------------------------------|--------|---------------------------------|--------|---------------------------------|--------|---------------------------------|--------|
| −25                           | 0.3 (3.8)                     | .94    | 0.1 (0.5)                       | .88    | 0.011 (0.012)                   | .33    | 5.6 (29.0)                      | .85    | −0.004 (0.004)                  | .37    | −0.006 (0.006)                  | .32    |
| −20                           | 2.9 (2.7)                     | .28    | 0.6 (0.3)                       | .09    | −0.002 (0.009)                  | .83    | −23.9 (21.7)                    | .27    | −0.008 (0.003)                 | .01    | −0.008 (0.005)                 | .09    |
| −15                           | 5.4 (3.9)                     | .17    | 1.1 (0.5)                       | .03    | −0.015 (0.007)                  | .04    | −53.5 (17.9)                    | .004   | −0.012 (0.003)                 | <.001  | −0.010 (0.004)                 | .02    |
| −10                           | 5.5 (2.8)                     | .05    | 0.7 (0.3)                       | .04    | −0.028 (0.009)                  | .001   | −83.0 (19.9)                    | <.001  | −0.016 (0.004)                 | <.001  | −0.011 (0.005)                 | .02    |
| −5                            | 5.5 (2.1)                     | .01    | 0.4 (0.3)                       | .15    | −0.041 (0.012)                  | <.001  | −112.5 (26.4)                   | <.001  | −0.020 (0.005)                 | <.001  | −0.013 (0.006)                 | .04    |
| 0                             | 5.6 (2.3)                     | .02    | 0.1 (0.3)                       | .83    | −0.066 (0.010)                  | <.001  | −232.9 (20.9)                   | <.001  | −0.048 (0.003)                 | <.001  | −0.022 (0.005)                 | <.001  |
| 5                             | 5.6 (3.2)                     | .08    | −0.3 (0.4)                      | .51    | −0.106 (0.018)                  | <.001  | −413.9 (34.4)                   | <.001  | −0.075 (0.005)                 | <.001  | −0.036 (0.008)                 | <.001  |
| 10                            | 5.7 (4.4)                     | .20    | −0.6 (0.6)                      | .28    | −0.147 (0.033)                  | <.001  | −594.9 (61.6)                   | <.001  | −0.103 (0.009)                 | <.001  | −0.050 (0.014)                 | <.001  |

Abbreviations: pTau181, phosphorylated tau 181; tTau, total tau.

*P values were for testing whether the rate of change is significantly different from 0.

Figure 2. Comparison of the Rates of Change of Cerebrospinal Fluid Levels of Total Tau (tTau) and Phosphorylated Tau 181 (pTau181) and Brain Structures by Magnetic Resonance Imaging in Mutation Carriers as a Function of Estimated Years Before Onset

Lines are locally estimated scatterplot smoothing curves for the standardized difference in individual annual rate of change. Values represent the estimated rate of change based on the longitudinal data of the mutation carriers standardized by the mean and SD of the baseline value of noncarriers.
correlated with most brain structure measures, where higher CSF tTau levels were associated with smaller cortical thickness. Generally, there were higher correlation coefficients for CSF tTau in the asymptomatic phase and mostly for the posterior neocortical and allocortical (limbic) regions. However, after symptom onset, the correlation between CSF tTau and pTau181 and brain structures changed. The intensity of neuronal damage as measured by brain atrophy continued at an increasing rate, whereas the rate of change of CSF tTau levels remained at a somewhat constant rate and the rate of change of CSF pTau181 actually switched from positive to negative. These results suggest that CSF tTau and structural brain measures have distinct patterns later in the disease course.

**Discussion**

The use of biomarkers has become an essential component of AD research and therapeutic trials. The new framework from the National Institute on Aging and Alzheimer Association capitalizes on the use of biomarkers for early identification of AD, which has substantial implications for early treatment and trial enrollment. However, little work has been done regarding comparisons of longitudinal biomarker trajectories that are currently proposed to represent similar aspects of disease (eg, CSF tTau and MRI are both proposed as markers of neurodegeneration).

In this study, on the basis of longitudinal data from the DIAN study, we evaluated the trajectories of CSF tTau and pTau181 over the course of AD. Notably, we included a bigger sample and used newer CSF tTau and pTau181 values generated with a fully automated, high-performance electroluminescence immunoassay. We compared the trajectories of CSF tTau and pTau181 with the atrophy of brain structures as measured by MRI. First, consistent with previous reports, our study found mean concentrations of CSF tTau and pTau181 to be higher in MCs from the early stage.
of AD, supporting the use of CSF tau as a marker of AD risk and progression. Second, we found that the positive rate of change of CSF tTau remained constant after EYO −10, whereas CSF pTau181 had a positive rate of change early in the disease course, which then reversed and became negative later in the disease. This indicates that our previous results were not likely to be an artifact of the measurement used because we used a different method for this study. Third, the associations identified between the rates of change of CSF tTau and pTau181 with brain atrophy do not support the assumption that CSF tTau changes follow a pattern similar to that of structural brain changes. Our findings indicate that neither CSF tTau nor pTau181 has the same pattern of change as brain measures and should be considered as associated but distinct biomarkers in AD. In addition, these findings indicate that, within the current biomarker classification, tTau is an important marker of AD but may not be the ideal marker of neurodegeneration.

Recent studies have suggested that in the presence of amyloid pathologic abnormalities, more CSF tTau and pTau181 is released. Moreover, increased tau in CSF seems to be dependent of amyloid deposition and occurs in the absence of tau brain pathologic abnormalities.

Early disease stages may also be characterized by higher cellular stress and inflammation, with higher levels of tTau and pTau181 in CSF representing a response. However, if CSF tTau directly reflected neurodegeneration, it would be expected that the rate of change of CSF tTau would increase in concert with brain atrophy during the period of maximal rate of atrophy (EYO >0). It is possible that during disease progression and neurodegeneration, the loss of neuronal cells results in less neuronal substrate to produce tau. Although this might account for some of the slowing in longitudinal changes in CSF tTau that we observed, it is unlikely that the degree of neurodegeneration is sufficient to fully explain our findings. Early elevations may also be associated with acute neuronal membrane damage, whereas apparent later reductions reflect the death of a smaller number of neurons that remain. Acute neuronal injury may be associated with a stronger inflammatory response at early stages of the disease.

The present findings challenge some previous assumptions about AD progression and its association with both pTau181 and tTau. Contrary to the idea that tTau and pTau181 levels continue to increase with greater neurodegeneration and the spread of NFT, we found evidence of a decrease in the rate of change, arguing against the use of these measures as a reflection of a passive release from neuronal death and NFT. These data support the relevance of CSF tTau and pTau181 as markers of amyloid deposition and accompanying changes (e.g., inflammation and neuronal membrane damage); however, the complex rates of change identified here and in our previous work suggest that using them as measures of therapeu tic response requires further investigation, because levels vary as a function of where an individual is in the neuropathological cascade. The apparent disconnect between CSF tTau and MRI measures may reflect the fact that they are measuring different stages of the neurodegenerative process, with CSF tTau accounting for the active phase of neuron injury and damage, and MRI measuring the subsequent structural sequelae of the active death process.

Our findings are consistent with previous studies of sporadic late-onset AD. Seppälä et al, in a longitudinal study in Finland, described higher levels of pTau181 level in mild cognitive impairment stages when compared with AD dementia, whereas levels of pTau181 decreased over a period of about 3 years. Similarly, Toledo et al reported a decrease in tTau in those at the dementia stage from the ADNI cohort. More recently, in a similar analysis of ADNI participants with longitudinal follow-up and using similar methods to determine CSF tTau and pTau181 levels (i.e., automated electroluminescence immunoassay), Sutphen et al reported that tTau and pTau181 showed consistent increases in the amyloid-positive participants with normal cognition and those with mild cognitive impairment, whereas pTau181 decreased substantially in those with AD dementia.

This study has potential implications for AD trials using tau-based therapies and other putative disease-modifying therapies. First, during trial design, the active group and the placebo group will have to be randomized by disease stage (according to disease severity measured using biomarkers and the severity of cognitive impairment), because minor differences in the neurodegeneration
cascade stage might translate into major differences in biomarker trajectories, and as a result, might be misinterpreted as a treatment effect. Second, one must consider how to interpret changes in biomarkers during clinical trials readout; in other words, one must consider how a successful treatment would be expected to affect CSF tau levels or rate of change. This study, along with recent work assessing neurofilament light chains,\textsuperscript{51,52} suggests that neurofilament light chains may be an advantageous marker of neurodegeneration in therapeutic trials of AD.\textsuperscript{52,53}

**Limitations**

The present study is not without limitations. First, our results are dependent on the accuracy of the DIANEYO and, like any measure, it is subject to error (i.e. symptom onset could be a few years earlier or later than expected [randomly]). Second, we recognize that DIAAD and late-onset AD are similar but not identical. Sporadic AD disease occurs later in life, and individuals often exhibit a wide range of comorbidities, including vascular disease and the presence of transactive response DNA-binding protein of 43 kDa.\textsuperscript{54-57} These differences may be associated with biomarker profiles or natural progression, so the total generalizability of the present findings to late-onset AD will have to be confirmed in future cohorts. Nevertheless, the results highlight the importance of gathering longitudinal data to refine current biomarker models.

**Conclusions**

The results of this study support the A/T/N framework, whereby increases in CSF levels of tTau and pTau181 are robust diagnostic markers of pathologic abnormalities and neuronal injury in early Alzheimer disease. However, our data suggest that levels of CSF tTau and pTau181 diverge from atrophy-based measures of neurodegeneration later in the disease. This is an important result to consider in clinical trials targeting tau brain pathologic abnormalities. Future studies should explore the longitudinal rate of change of CSF pTau181 levels and positron emission tomography of tau as markers of tau brain pathologic abnormalities.

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REFERENCES

1. Braak H, Thal DR, Ghebremedhin E, Del Tredici K. Stages of the pathologic process in Alzheimer disease: age categories from 1 to 100 years. J Neuropathol Exp Neurol. 2011;70(11):960-969. doi:10.1097/NEN.Ob013e318232a379

2. Hyman BT, Phelps CH, Beach TG, et al. National Institute on Aging: Alzheimer’s Association guidelines for the neuropathologic assessment of Alzheimer’s disease. Alzheimers Dement. 2012;8(1):1-13. doi:10.1016/j.jalz.2011.10.007.

3. Ewers M, Mattsson N, Minthon L, et al. CSF biomarkers for the differential diagnosis of Alzheimer’s disease: a large-scale international multicenter study. Alzheimers Dement. 2015;11(11):1306-1315. doi:10.1016/j.jalz.2014.12.006

4. Mattsson N, Carrillo MC, Dean RA, et al. Revolutionizing Alzheimer’s disease and clinical trials through biomarkers. Alzheimers Dement (Amst). 2015;1(4):412-419. doi:10.1016/j.dadm.2015.09.001

5. Sperling RA, Aisen PS, Beckett LA, et al. Toward defining the preclinical stages of Alzheimer’s disease: recommendations from the National Institute on Aging-Alzheimer’s Association workgroups on diagnostic guidelines for Alzheimer’s disease. Alzheimers Dement. 2011;7(3):280-292. doi:10.1016/j.jalz.2011.03.003

6. Bateman RJ, Xiong C, Benzinger TLS, et al; Dominantly Inherited Alzheimer Network. Clinical and biomarker changes in dominantly inherited Alzheimer’s disease. N Engl J Med. 2012;367(9):795-804. doi:10.1056/NEJMoa1202753

7. Ryman DC, Aisen PS, Bird T, et al. Symptom onset in autosomal dominant Alzheimer disease: a systematic review and meta-analysis. Neurology. 2014;83(3):253-260.

8. Trojanowski JQ, Vandeerstichele H, Korecka M, et al; Alzheimer’s Disease Neuroimaging Initiative. Update on the biomarker core of the Alzheimer’s Disease Neuroimaging Initiative subjects. Alzheimers Dement. 2010;6(3):230-238. doi:10.1016/j.jalz.2010.03.008

9. Jack CR Jr, Knopman DS, Jagust WJ, et al. Hypothetical model of dynamic biomarkers of the Alzheimer’s pathological cascade. Lancet Neurol. 2010;9(9):119-128. doi:10.1016/S1474-4422(09)70299-6

10. Petersen RC, Trojanowski JQ. Use of Alzheimer disease biomarkers: potentially yes for clinical trials but not yet for clinical practice. JAMA. 2009;302(4):436-437. doi:10.1001/jama.2009.1073

11. 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. doi:10.1016/j.jalz.2018.02.018

12. Montine TJ, Phelps CH, Beach TG, et al; National Institute on Aging. Alzheimer’s Association. National Institute on Aging-Alzheimer’s Association guidelines for the neuropathologic assessment of Alzheimer’s disease: a practical approach. Acta Neuropathol. 2012;123(1):1-11. doi:10.1007/s00401-011-0910-3

13. Fagan AM, Xiong C, Jasielec MS, et al; Dominantly Inherited Alzheimer Network. Longitudinal change in CSF biomarkers in autosomal-dominant Alzheimer’s disease. Sci Transl Med. 2014;6(226):226ra30. doi:10.1126/scitranslmed.3007901

14. McClade E, Wang G, Gordon BA, et al; Dominantly Inherited Alzheimer Network. Longitudinal cognitive and biomarker changes in dominantly inherited Alzheimer disease. Neurology. 2018;91(14):e1295-e1306. doi:10.1212/ WNL.0000000000006277

15. Sutphen CL, McCue L, Herries EM, et al; ADNI. Longitudinal decreases in multiple cerebrospinal fluid biomarkers of neuronal injury in symptomatic late onset Alzheimer’s disease. Alzheimers Dement. 2018;14(7):869-879. doi:10.1016/j.jalz.2018.01.012

16. Preische O, Schultz A, Apel A, et al. Serum neurofilament light chain levels are associated with CSF neurofilament light chain, cognitive status, and disease progression in autosomal dominant AD. Alzheimers Dement. 2018;14(7):P1170. doi:10.1016/j.jalz.2018.06.1611

17. Schindler SE, Li Y, Todd KW, et al; Dominantly Inherited Alzheimer Network. Emerging cerebrospinal fluid biomarkers in autosomal dominant Alzheimer’s disease. Alzheimers Dement. 2019;15(5):655-665. doi:10.1016/j.jalz.2018.12.019

18. Hampel H, Goernitz A, Buergler K. Advances in the development of biomarkers for Alzheimer’s disease: from CSF total tau and Abeta(1-42) proteins to phosphorylated tau protein. Brain Res Bull. 2003;61(3):243-253. doi:10.1016/S0361-9230(03)00087-X

19. Ryan NS, Nicholas JM, Weston PSJ, et al. Clinical phenotype and genetic associations in autosomal dominant familial Alzheimer’s disease: a case series. Lancet Neurol. 2016;15(13):1326-1335. doi:10.1016/S1474-4422(16)30193-4

20. Morris JC, Aisen PS, Bateman RJ, et al; Developing an international network for Alzheimer’s research: the Dominantly Inherited Alzheimer Network. Clin Investig (Lond). 2012;2(10):975-984. doi:10.4155/CLIN.12.93

JAMA Network Open. 2019;2(12):e1917126. doi:10.1001/jamanetworkopen.2019.17126

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21. Kauwe JS, Jacquart S, Chakraverty S, et al. Extreme cerebrospinal fluid amyloid β levels identify family with late-onset Alzheimer's disease presenilin 1 mutation. *Ann Neurol*. 2007;61(5):446-453. doi:10.1002/ana.21099

22. Morris JC. The Clinical Dementia Rating (CDR): current version and scoring rules. *Neurology*. 1993;43(11):2412-2414. doi:10.1212/WNL.43.11.2412-a

23. Moulder KL, Snider BJ, Mills SL, et al. Dominantly Inherited Alzheimer Network: facilitating research and clinical trials. *Alzheimers Res Ther*. 2013;5(5):48. doi:10.1186/alzrt213

24. 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. doi:10.1212/WNL.34.7.939

25. Jack CR Jr, Bernstein MA, Borowski BJ, et al; Alzheimer's Disease Neuroimaging Initiative. Update on the magnetic resonance imaging core of the Alzheimer's disease neuroimaging initiative. *Alzheimers Dement*. 2010;6(3):212-220. doi:10.1016/j.jalz.2010.03.004

26. Fischl B. FreeSurfer. *Neuroimage*. 2012;62(2):774-781. doi:10.1016/j.neuroimage.2012.01.021

27. Gordon BA, Blazey TM, Su Y, et al. Spatial patterns of neuroimaging biomarker change in individuals from families with autosomal dominant Alzheimer's disease: a longitudinal study. *Lancet Neurol*. 2018;17(3):241-250. doi:10.1016/S1474-4422(18)30028-0

28. Bittner T, Zetterberg H, Teunissen CE, et al. Technical performance of a novel, fully automated electrochemiluminescence immunoassay for the quantitation of β-amyloid (1-42) in human cerebrospinal fluid. *Alzheimers Dement*. 2016;12(5):517-526. doi:10.1016/j.jalz.2016.09.005

30. Hansson O, Seibyl J, Stromfjord E, et al; Swedish BioFINDER study group; Alzheimer's Disease Neuroimaging Initiative. CSF biomarkers of Alzheimer's disease concord with amyloid-β PET and predict clinical progression: a study of fully automated immunoassays in BioFINDER and ADNI cohorts. *Alzheimers Dement*. 2018;14(11):1470-1481. doi:10.1016/j.jalz.2018.01.010

34. Sperling R, Johnson K. Biomarkers of Alzheimer disease: current and future applications to diagnostic criteria. *Continuum (Minneap Minn)*. 2013;19(2):325-338. doi:10.1212/01.CON.0000429181.60095.99

35. Toledo JB, Xie SX, Trojanowski JQ, Shaw LM. Longitudinal change in CSF Tau and Aβ biomarkers for up to 48 months in ADNI. *Acta Neuropathol*. 2013;126(5):659-670. doi:10.1007/s00401-013-1151-4

36. Schelle J, Häsler LM, Göpfert JC, et al. Prevention of tau increase in cerebrospinal fluid of APP transgenic mice suggests downstream effect of BACE1 inhibition. *Alzheimers Dement*. 2017;13(6):701-709. doi:10.1016/j.jalz.2016.09.005

37. Sato C, Barthélemy MR, Mawuenyega KG, et al. Tau kinetics in neurons and the human central nervous system. *Neuron*. 2018;97(6):1284-1298. doi:10.1016/j.neuron.2018.02.015

38. Sato C, Barthélemy MR, Mawuenyega KG, et al. Changes in amyloid-β and Tau in the cerebrospinal fluid of transgenic mice overexpressing amyloid precursor protein. *Sci Transl Med*. 2013;5(194):194re2. doi:10.1126/scitranslmed.3006446

42. Sperling R, Johnson K. Biomarkers of Alzheimer disease: current and future applications to diagnostic criteria. *Continuum (Minneap Minn)*. 2013;19(2):325-338. doi:10.1212/01.CON.0000429181.60095.99

43. Sato C, Barthélemy MR, Mawuenyega KG, et al. Tau kinetics in neurons and the human central nervous system. *Neuron*. 2018;97(6):1284-1298. doi:10.1016/j.neuron.2018.02.015
44. Jones DT, Knopman DS, Graff-Radford J, et al. In vivo 18F-AV-1451 tau PET signal in MAPT mutation carriers varies by expected tau isoforms. *Neurology*. 2018;90(11):e947-e954. doi:10.1212/WNL.0000000000005117

45. Suárez-Calvet M, Araque Caballero MA, Kleinberger G, et al; Dominantly Inherited Alzheimer Network. Early changes in CSF sTREM2 in dominantly inherited Alzheimer’s disease occur after amyloid deposition and neuronal injury. *Sci Transl Med*. 2016;8(369):369ra178. doi:10.1126/scitranslmed.aag1767

46. Venegas C, Kumar S, Franklin BS, et al. Microglia-derived ASC specks cross-seed amyloid-β in Alzheimer’s disease. *Nature*. 2017;552(7685):355-361. doi:10.1038/nature25158

47. Quiroz YT, Sperling RA, Norton DJ, et al. Association between amyloid and tau accumulation in young adults with autosomal dominant Alzheimer disease. *JAMA Neurol*. 2018;75(5):548-556. doi:10.1001/jamaneurol.2017.4907

48. Seppälä TT, Koivistoinen AM, Hartikainen P, Helisalmi S, Soininen H, Herukka S-K. Longitudinal changes of CSF biomarkers in Alzheimer’s disease. *J Alzheimers Dis*. 2011;25(4):583-594. doi:10.3233/JAD-2011-101911

49. Mollenhauer B, Bibi M, Trenkwalder C, et al. Follow-up investigations in cerebrospinal fluid of patients with dementia with Lewy bodies and Alzheimer’s disease. *J Neural Transm (Vienna)*. 2005;112(7):933-948. doi:10.1007/s00702-004-0235-7

50. Kinnunen KM, Cash DM, Poole T, et al; Dominantly Inherited Alzheimer Network (DIAN). Presymptomatic atrophy in autosomal dominant Alzheimer’s disease: a serial magnetic resonance imaging study. *Alzheimers Dement*. 2018;14(1):43-53. doi:10.1016/j.jalz.2017.06.2268

51. Weston PSJ, Poole T, Ryan NS, et al. Serum neurofilament light in familial Alzheimer disease: a marker of early neurodegeneration. *Neurology*. 2017;89(21):2167-2175. doi:10.1212/WNL.0000000000004667

52. Preische O, Schultz SA, Apel A, et al; Dominantly Inherited Alzheimer Network. Serum neurofilament dynamics predicts neurodegeneration and clinical progression in presymptomatic Alzheimer’s disease. *Nat Med*. 2019;25(2):277-283. doi:10.1038/s41591-018-0304-3

53. Mattsson N, Cullen NC, Andreasson U, Zetterberg H, Blennow K. Association between longitudinal plasma neurofilament light and neurodegeneration in patients with Alzheimer disease. *JAMA Neurol*. 2019;76(7):791-799. doi:10.1001/jamaneurol.2019.0765

54. Wilson AC, Dugger BN, Dickson DW, Wang D-S. TDP-43 in aging and Alzheimer’s disease: a review. *Int J Clin Exp Pathol*. 2011;4(2):147-155.

55. Hamilton RL. Lewy bodies in Alzheimer’s disease: a neuropathological review of 145 cases using alpha-synuclein immunohistochemistry. *Brain Pathol*. 2000;10(3):378-384.

56. Nelson PT, Smith CD, Abner EL, et al. Hippocampal sclerosis of aging, a prevalent and high-morbidity brain disease. *Acta Neuropathol*. 2012;126(2):161-177. doi:10.1007/s00401-013-1554-1

57. Gefen T, Gasho K, Rademaker A, et al. Clinically concordant variations of Alzheimer pathology in aphasic versus amnestic dementia. *Brain*. 2012;135(S5):i554-i565. doi:10.1093/brain/aws076

**SUPPLEMENT.**

eTable 1. Number of Participants With Longitudinal CSF Biomarkers and MRI Assessments

eTable 2. Associations Between CSF tTau and MRI Regions by Disease Stage (Asymptomatic vs Symptomatic)

eTable 3. Associations Between CSF pTau181 and MRI Regions by Disease Stage (Asymptomatic vs Symptomatic)