

# Association of *APOE* $\epsilon$ 4 and Plasma p-tau181 with Preclinical Alzheimer's Disease and Longitudinal Change in Hippocampus Function

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## Abstract.

**Background:** The Apolipoprotein E (*APOE*)  $\epsilon$ 4 allele has been linked to increased tau phosphorylation and tangle formation. *APOE*  $\epsilon$ 4 carriers with elevated tau might be at the higher risk for Alzheimer's disease (AD) progression. Previous studies showed that tau pathology begins early in areas of the medial temporal lobe. Similarly, *APOE*  $\epsilon$ 4 carriers showed altered hippocampal functional integrity. However, it remains unknown whether the influence of elevated tau accumulation on hippocampal functional changes would be more pronounced for *APOE*  $\epsilon$ 4 carriers.

**Objective:** We related  $\epsilon$ 4 carriage to levels of plasma phosphorylated tau (p-tau181) up to 15 years prior to AD onset. Furthermore, elevated p-tau181 was explored in relation to longitudinal changes in hippocampal function and connectivity.

**Methods:** Plasma p-tau181 was analyzed in 142 clinically defined AD cases and 126 matched controls. The longitudinal analysis involved 87 non-demented individuals (from population-based study) with two waves of plasma samples and three waves of functional magnetic resonance imaging during rest and memory encoding.

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**Results:** Increased p-tau181 was observed for both  $\epsilon$ 4 carriers and non-carriers close to AD onset, but exclusively for  $\epsilon$ 4 carriers in the early preclinical groups (7- and 13-years pre-AD). In  $\epsilon$ 4 carriers, longitudinal p-tau181 increase was paralleled by elevated local hippocampal connectivity at rest and subsequent reduction of hippocampus encoding-related activity.

**Conclusion:** Our findings support an association of APOE  $\epsilon$ 4 and p-tau181 with preclinical AD and hippocampus functioning.

Keywords: Alzheimer's disease, APOE, fMRI, hippocampus, longitudinal, magnetic resonance imaging, p-tau181, phosphorylated tau, population-based

## INTRODUCTION

The  $\epsilon$ 4 allele of the apolipoprotein E (APOE) gene is the major genetic risk factor for late-onset Alzheimer's disease (AD) [1]. A likely mechanism by which the APOE  $\epsilon$ 4 allele increases risk for AD is through influencing amyloid- $\beta$  (A $\beta$ ) deposition [2]. Independently of A $\beta$ , the elevated risk of developing AD that is conferred by the APOE  $\epsilon$ 4 genotype may also involve mechanisms associated with tau aggregation [3, 4]. Importantly, APOE  $\epsilon$ 4 is a risk factor, so not all  $\epsilon$ 4 carriers will present with AD pathology [5], and conversely 30–40% of AD cases do not possess the  $\epsilon$ 4 allele. Thus, in the preclinical phase, APOE  $\epsilon$ 4 carriers with elevated A $\beta$  and tau might be at the highest risk for AD progression. Although elevated A $\beta$  and tau levels in the clinical phases are well documented relative to controls, there is less consensus about what constitutes elevated A $\beta$  and tau levels in the preclinical phase. Longitudinal studies of population-based non-clinical samples can provide a unique opportunity to explore accumulation of A $\beta$  and tau over several years. Here we present results from the Betula longitudinal study on aging, memory, and dementia, comprising 4,425 individuals randomly sampled from the population registry and examined every five years for up to 30 years [6]. All enrolled participants were judged free of AD when included into the study. Blood was sampled at each test wave, and plasma-based measures of phosphorylated tau (p-tau181) were quantified [7]. In a case-control setting, we established p-tau181 cut-offs indicative of AD pathology in clinical as well as in early and late preclinical phases. We predicted group differences in p-tau level during both clinical and preclinical phases, such that clinically defined AD cases would exhibit elevated p-tau compared to the control group in clinical phase as well as years before clinical onset. Moreover, elevated p-tau was expected to be more pronounced among APOE  $\epsilon$ 4 carriers (c.f. [8]).

A second aim of the study was to relate longitudinal trajectories of p-tau181 levels to hippocampus functioning in APOE  $\epsilon$ 4 carriers and non-carriers. Both postmortem and *in vivo* studies using positron

emission tomography (PET) suggest that tau pathology begins early in areas of the medial temporal lobes [9, 10], indicating that tau could impact hippocampal functional integrity. Indeed, studies have linked higher tau accumulation to increased hippocampal connectivity at rest [11] and hyperactivity during memory encoding [12]. Moreover, previous genetic studies showed that APOE  $\epsilon$ 4 carriers exhibit elevated connectivity and task activation compared with non-carriers [13, 14]. Based on these and related findings, we predicted that the effect of longitudinal tau181 accumulation on hippocampal function would be more pronounced for  $\epsilon$ 4 carriers than for non-carriers.

## METHODS

### Betula study design

Data were obtained from the Swedish longitudinal Betula study [6] (<http://www.umu.se/en/betula>).

### Participants

The participants are from two longitudinal samples [6, 15]; sample S1 ( $n = 1000$ ) included at wave W1 (1988–1990), and sample S3 ( $n = 963$ ) included at W2 (1993–1995). The route from the parent cohort to the two present study samples, are illustrated in Fig. 1a and 1b.

The cross-sectional AD case-control sample consists of  $n = 142$  individuals with either manifest ( $n = 37$ ) or preclinical AD ( $n = 105$ ) and matched controls ( $n = 126$ ). The clinical AD status is defined in relation to wave W3, the timepoint from which the plasma p-tau181 analyses are based. The AD cases had a clinical onset on average 2 years (range = 0–6 years) before W3, whereas preclinical AD subsequently developed AD on average 2 (range = 1–4 years), 7 (range = 5–9 years), and 13 (range = 10–15 years) years after W3. All cases with current or forthcoming AD with available W3 plasma defines the case population. The age- and sex-matched controls, selected from the same population and age cohorts as the

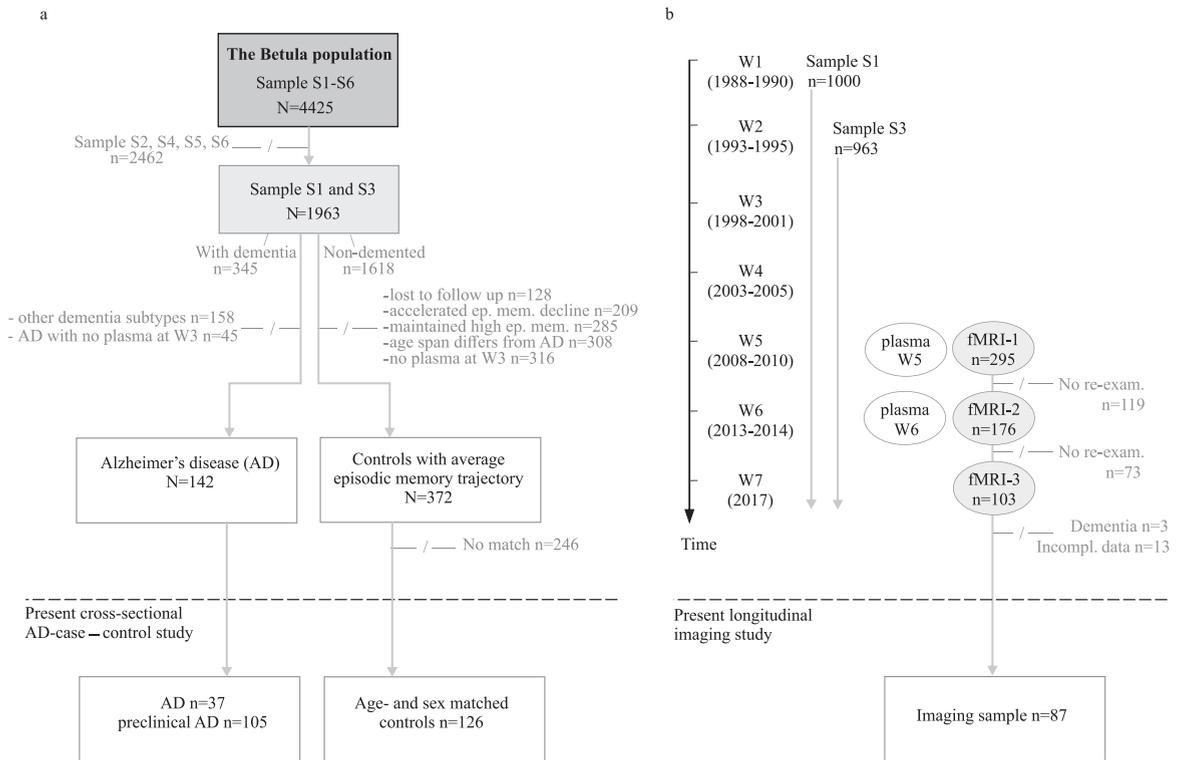


Fig. 1. a) Flowchart of cross-sectional AD-case – control study with Wave 3 (W3) as study baseline, and (b) longitudinal imaging study with Wave 5 as study baseline. –/, not included in the present study; W, wave; S, sample.

cases, were classified as having an average episodic memory decline relative to their age [16], had sufficient W3 plasma, and remained non-demented throughout the studied period (1988–2017). For 16 cases, there were no eligible controls. Characteristics of the AD case and control sample are given in Table 1.

The longitudinal imaging sample comprises 98 non-demented study participants with imaging data

obtained at W5 (2008–2010), W6 (2013–2014), and W7 (2017), with an average of 4 years between measurement points. Participants with complete p-tau181 data from waves 5–6 and fMRI observations at waves 5, 6, and 7 were included in the longitudinal analyses (n=87). Seven of these also served as controls in the case-control study. The imaging study sample, in the larger context of the Betula longitudinal study design, is displayed in Fig. 1b and presented in

Table 1  
Characteristics of the AD case-control sample at W3

Sample char.	clinical AD	preclinical AD			controls
		13 (10–15 y)	7 (5–9 y)	2 (1–4 y)	
N	37	35	30	40	126
Sex, %-female	87%	86%	63%	70%	74%
Age, y	83 [70–91]	71 [60–81]	72 [61–85]	79 [65–91]	76 [60–91]
Education, y	7.8 [3–18]	8.7 [4.5–17]	7.7 [6–13]	7.5 [5–17]	7.7 [3–16]
APOE ε4 carrier	43%	54%	57%	58%	17%
pτ181 (pg/ml)	4.9 [0.9–33.1]	3.0 [0.2–12.6]	3.2 [1.1–29.4]	4.9 [1.2–14.9]	2.9 [0.4–22.0]
EMC	9 [4–25]	34 [17–55]	28 [12–44]	17 [2–33]	27 [7–50]
MMSE	21 [9–27]	28 [24–30]	27 [23–30]	26 [20–30]	27 [14–30]

For age and education, values are mean [range]. For pτ181 and memory scores, values are median [range]. EMC, Episodic Memory Composite score; a composite of five tasks (max score 76). Education data, EMC, and MMSE (Mini-Mental State Examination) were available for 86–97%. Of those with clinical AD, 51% contributed complete EMC data. Age, APOE ε4 carriership, and plasma pτ181 levels were available for all.

Table 2  
Characteristics of the longitudinal imaging sample at W5–W7

Sample char.	high p $\tau$ 181 $\epsilon$ 4+	high p $\tau$ 181 $\epsilon$ 4-	low p $\tau$ 181
N	20	21	46
Sex, %-female	45%	43%	41%
Age W5, y	65 [56–77]	68 [56–81]	64 [56–76]
Education, y	14.6 [7–26]	13.1 [6–19.5]	13.6 [6–24.5]
EMC W5	43.5 [22–61]	39 [32–56]	42 [24–57]
EMC W6	39.5 [17–55]	34 [25–52]	43 [23–61] *
EM W7	12 [3–16]	10 [2–17]	10 [6–20]
MMSE W5	28 [25–30]	28 [24–30]	28 [24–30]
MMSE W6	28.5 [24–30]	28 [25–30]	28 [22–30]
MMSE W7	27 [24–30]	27 [24–30]	27 [21–30]
APOE $\epsilon$ 4 carrier	100%	0%	30%
p $\tau$ 181 W5, (pg/ml)	2.9 [1.4–6.0]	3.08 [1.0–7.1]	1.73 [0.5–4.3]
p $\tau$ 181 W6, (pg/ml)	5.5 [3.3–11.3]	5.22 [3.7–9.4]	2.19 [0.7–3.5]

EM, summed score of the two EMC-free recall tasks that were included at W7 (max score 28); units and abbreviations for all other entries are the same as in Table 1. Age, Education data, APOE  $\epsilon$ 4 carriership, plasma p $\tau$ 181 levels, EM, and MMSE were available for all. \* One subject excluded from the EMC W6-calculation due to missing data.

full details elsewhere [6]. Sample characteristics are presented in Table 2.

The Betula study was approved by the Regional Ethical Review Board and carried out in accordance with the Declaration of Helsinki. Written consent for study participation was obtained from each participant.

#### Diagnostic assessment

The diagnostic protocol was reported in full details elsewhere [6] and was based on repeated evaluations of multi-disciplinary clinical documentation, further supplemented by outcomes of the Betula study health- and memory assessments [6]. The diagnostic procedure allowed individual health trajectories, throughout the study period, to be followed at a symptom- and functional level and considered also other clinician's assessments in the diagnostic decision. Participants receiving an AD diagnosis exhibited an insidious onset and progressive cognitive decline as well as other symptoms typically attributable to clinical AD. Individuals with cardiovascular insults together with neurological signs, a fluctuating symptomatology, and stepwise progression of cognitive deficiencies received a clinical diagnosis of vascular dementia. Individuals with complex diagnostic features or insufficient clinical information were classified as dementia not otherwise specified. Less common dementia disorders such as frontotemporal dementia, Parkinson dementia, Lewy body dementia, cortico-basal syndrome, and progressive supranuclear palsy were thoroughly clinically investigated and diagnosed within the healthcare

system. Individuals with cognitive impairment close to death, which was accompanied by severe somatic conditions, were not considered as demented, nor were individuals with neurocognitive deficits of non-progressive nature after, e.g., trauma, tumor, sub-arachnoid hemorrhage. The disease onset was determined as the year at which the clinical symptoms with sufficient severity elicit interference with social functioning and instrumental activities of daily living, i.e., when the core criteria of dementia were met [17]. The diagnostic assessments were coordinated by co-author R.A. throughout the study period, applying DSM-IV classification core criteria for dementia. Additional inclusion and exclusion criteria were applied, which increased the sensitivity and specificity [18–20]. To improve the reliability, the diagnostic procedure was blindly carried out every 5<sup>th</sup> year without information on the previously determined diagnostic status at hand. Any disparity in diagnosis and/or onset age were re-examined and solved.

#### Plasma p-tau181

Plasma p-tau181 levels were measured with a fully validated in-house ultrasensitive Simoa immunoassay, previously described in detail [21].

#### Statistical analyses

A cut-off for elevated p-tau181 levels of > 6 pg/ml was determined from the distribution of values in the control group, which approximately corresponded to 1 SD (2.6 pg/ml) above the Mean (3.3 pg/ml). In a

sensitivity analysis of the relation between p-tau and APOE we also considered the stricter threshold of  $> 10$  pg/ml. Furthermore, we considered p-tau 181 as a continuous variable in a conditional logistic regression approach to predict AD.

Comparison of p-tau levels between cases and controls within the dementia groups were performed using paired one-sided *t*-tests as well as Wilcoxon signed-rank tests. The treatment of matched case-control data using parametric and nonparametric tests for matched samples is in accordance with previous recommendations [22]. Cases where no control was identified were discarded in the main analysis. In a further analysis we performed imputation using the control age group median as the control observation for cases without controls, and then proceeded as stated above. For investigating the association between elevated p-tau levels and APOE  $\epsilon 4$  carrier status, the Chi-square test for cross-tables was used. If there were expected cell counts below five, we performed a control analysis using Monte-Carlo simulation on the cross-tables to validate the obtained Chi-square *p*-value (the control analyses confirmed the main findings). A signed paired Wilcoxon's rank test between APOE groups for AD cases at different stages were performed.

#### Neuroimaging: Pre- and postprocessing

The acquisition of T1-weighted images, T2-weighted images, EPI-series, and experiment setup has been thoroughly described [23]. The fMRI-sequences included a 10-min blocked face-name recognition task with 6 encoding blocks, 6 retrieval blocks, and 8 blocks of an active perceptual baseline task [23, 24].

Within each encoding and retrieval block (six blocks each), four stimuli were presented for 4 s each, with a randomized interstimulus interval of 1.5, 2.5, 3, or 4.5 s. Participants were instructed before each block for 4 s. The face-name stimuli were presented during the encoding blocks, and the participants were instructed to memorize face-name associates. A button press, using the index finger, was expected to indicate that a face-name pair was seen. During the retrieval blocks, each face was presented along with three letters of which one corresponded to the first letter of the previously encoded name. Subjects were expected to indicate the letter corresponding to the name which was previously encoded with the face. The top, middle, and bottom letters corresponded to the index, the ring, and the middle fingers, respectively. Respond by guessing

was expected if participants could not remember the association. During the active baseline condition, a cross-hair was shown and this cross-hair was converted to a circle every now and then. The subjects were expected to press a button as soon as a conversion took place. This condition was used to dissociate different aspects of episodic memory from sensory motor components. The fMRI-part also included a resting-state (RS) sequence whose processing was previously described [25–28]. In brief, the T1-images were segmented, and the grey matter, white matter, and cerebrospinal fluid-likelihood-maps were used to generate, first, a subject-specific template, and then a group-specific template in Dartel-space. The fMRI-data was movement corrected, normalized to MNI (Montreal Neurological Institute) space by flow field files from the Dartel-processing, and smoothed. An adjustment for B0-inhomogeneities was performed by a B0-template calculated from W7.

For task-fMRI, a high pass filter was applied with a time constant of 200 s. A test of encoding versus baseline at W5 had revealed two peaks in left [22 –8 –16] and right [–20 –10 –16] anterior hippocampus [23], and here we extracted encoding-baseline contrast (all trials) values within two 5 mm ROIs around the peaks. We tested for between-groups differences in hippocampus activity (collapsed across the left and right hemisphere regions-of-interest) within each timepoint. At the group level, a 2-way ANOVA with the three groups as one factor and the three time points as another factor was conducted, followed by two-sample *post-hoc t*-tests when significant interactions were observed. A Pearson correlation test was performed between differences in anterior hippocampus activation and memory performance across test waves. No significant difference in time interval between T5 and T6 was observed between APOE groups ( $p = 0.44$ ).

For RS-fMRI, linear and quadratic effects of average cerebrospinal fluid and white matter-specific time courses, a global signal using matter-specific probability maps ( $p_s > 0.5$ ) and a 24-parameter model (six motion parameters, six temporal derivatives, and their squares was used) were regressed out and then a high pass filter with a time constant of 130 s were applied. For each participant and time point, average resting-state time series were extracted from 264 brain regions (3 mm radius spheres) based on a commonly used functional connectivity parcellation [29]. Here, we focused on 58 cortical ROIs that compromise the default mode network (DMN), known to include hippocampus as an important subsystem

[25, 26]. Since this functional parcellation does not include hippocampus regions, we added four anterior and posterior hippocampus regions (3 mm radius spheres; XYZ=-20 -10 -16; 22 -8 -16; -24 -30 -4; 26 -30 -2; Supplementary Figure 1) which previously showed reliable activations during a memory task [23]. To ensure that the extracted time series were derived from grey matter regions, only grey matter voxels were considered by eroding each sphere by a high probability grey matter mask ( $p < 0.001$ ).

The extracted time series were correlated to create a  $62 \times 62$  connectivity matrix (58 DMN + 4 hippocampus ROIs) for each participant and time point using Pearson's correlations, transformed to z-values using Fisher's r-to-z transformation. Then, the averaged correlation among all 4 ROIs were computed as a measure of local hippocampal connectivity. In a control analysis, given potential concerns about the validity of negative correlations [30], we set all negative correlations to zero and found that the longitudinal increase in hippocampal FC remained significant. To explore hippocampal-DMN connectivity, we computed the mean of two values: first, the

average of the left hippocampus connectivity with the left DMN ROIs and second, the average of right hippocampal connectivity with right DMN ROIs.

Preprocessing and analyses were made with the SPM12 software (<https://www.fil.ion.ucl.ac.uk/spm/>) using an inhouse program for batching and visualization (DataZ).

#### Data availability

Providing data transfer is in agreement with European Union legislation on the General Data Protection Regulation, anonymized data will be shared upon request from qualified investigators for the sole purpose of replication.

## RESULTS

### *P-tau181 levels in clinical and preclinical AD in relation to APOE $\epsilon 4$*

Figure 2a shows case-control comparisons of plasma p-tau181 levels as a function of clinical phase.

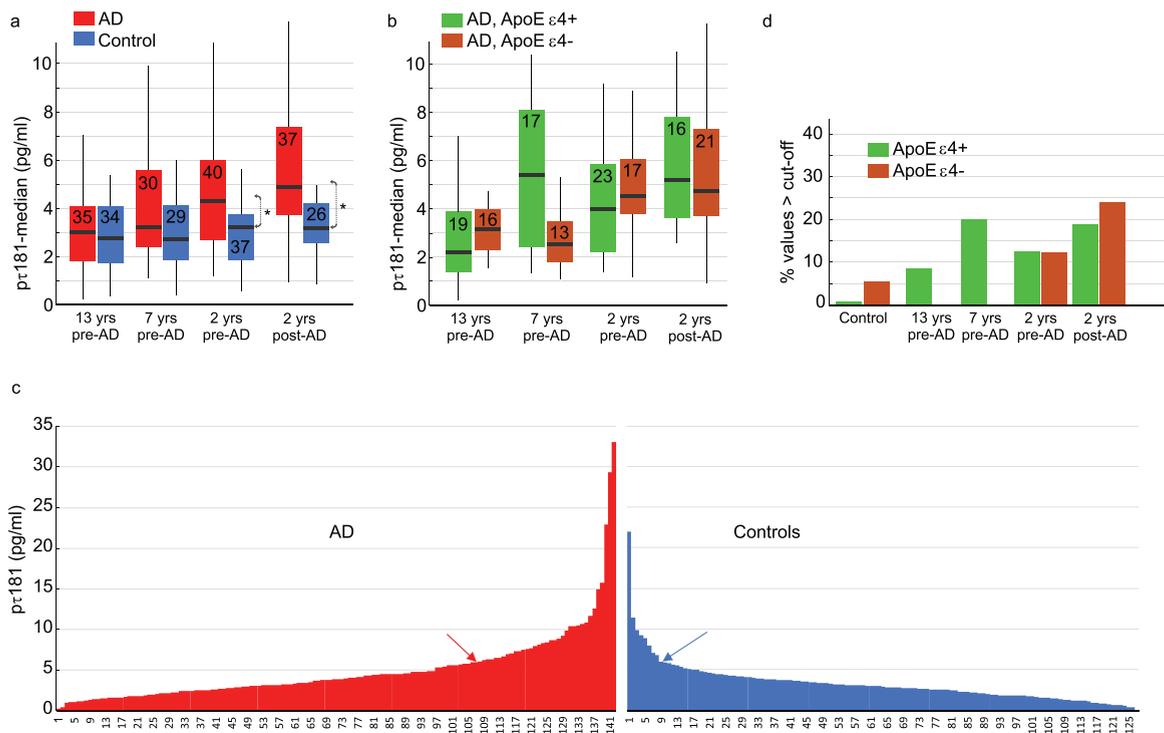


Fig. 2. P-tau181 levels for cases and controls. a) Median p-tau181 levels as a function of time to/from AD onset. Number of subjects in each group is indicated. b) Median p-tau181 levels as a function of time to/from AD onset as a function of APOE status. Number of subjects in each group is indicated. c) Distribution of individual p-tau181 levels for AD cases and controls, where arrows indicate tentative cut-off for elevated values (see text). d) Percentage with p-tau181 levels exceeding the cut-off for controls and AD cases as a function of time to/from diagnosis. Error-bars are 25/75-percentile in (a) and (b). “\*\*\*” denotes a group difference,  $p < 0.05$ .

Across phases, the cases had higher p-tau181 levels than controls, and the highest value was seen for the manifest AD-cases (ca two years post clinical AD onset). The case-control difference was significant in the clinical phase ( $p=0.0033$  for non-parametric and  $p=0.0072$  for parametric tests) and also in the late preclinical phase (ca 2 years pre-onset,  $p=0.0041$  for non-parametric and  $p=0.033$  for parametric tests). In the early preclinical phases (ca 13- and 7 years pre-onset, respectively), only a statistical trend at the group level was observed in the 7-years pre-onset group ( $p=0.058$ , parametric,  $p=0.097$  non-parametric). The significant group difference at clinical and late clinical phases survived correction for multiple comparisons ( $0.05/4=0.0125$ ). Further conditional logistic regression with p-tau181 as a continuous variable revealed that the odds ratio (OR) for AD associated with p-tau181 is 1.18 (if we transform p-tau181 to a z-score, the OR for AD=1.88). Thus, one standard deviation increase in p-tau was associated with almost a doubling of the odds for AD.

Next, we addressed the issue of elevated p-tau181 levels in relation to APOE. Of the 142 cases, 75 were  $\epsilon 4$  carriers (53%), whereas only 21 of 126 controls were  $\epsilon 4$  carriers (17%). Figure 2b plots  $\epsilon 4$  status as a function of diagnostic stage for cases. There were no significant differences ( $p_s > 0.15$ ) between APOE  $\epsilon 4$  carriers ( $\epsilon 4+$ ) and non-carriers ( $\epsilon 4-$ ) among the AD-cases at the different clinical stages, except for a trend ( $p=0.072$ ) at 7 years pre-AD towards elevated p-tau181 levels in  $\epsilon 4$  carriers (Median = 5.41 pg/ml) compared to non-carriers (Median = 2.54 pg/ml).

Figure 2c shows the highly skewed distributions of p-tau181 levels in controls and cases. Of the controls, eight (6%) had a p-tau181  $> 6$  pg/ml ( $\cong M + 1$  SD; blue arrow in Fig. 2c). Using this control-group derived cut-off for elevated p-tau181 levels, 34 AD cases (24%) had an elevated value (red arrow in Fig. 2c).

Figure 2d plots the p-tau181 elevated controls and AD cases as a function of clinical phase and APOE status. In the early preclinical phases (13 and 7 years), all cases with elevated values were APOE  $\epsilon 4$  carriers, and there was a significant association between elevated p-tau181 and APOE genotype ( $\chi^2=8.42$ ;  $p=0.004$ ). As expected, the percentages of individuals with elevated p-tau181 values were higher in closer proximity to AD onset, but at these stages there was no APOE effect (i.e., similar rates of  $\epsilon 4$  carriers and non-carriers in the late preclinical phases;  $\chi^2=0.32$ ;  $p=0.57$ ).

A similar analysis based on a stricter cut-off for defining elevated p-tau181 levels ( $> 10$  pg/ml) yielded a similar outcome with regard to APOE. In the early preclinical AD phases (13 and 7 years), 4 individuals (6.2%) had values exceeding 10 pg/ml, and all were  $\epsilon 4$  carriers. In the later preclinical phases ( $-2$  years from diagnosis), 8 (10.5%) had values exceeding the stricter cut-off but only 2 were  $\epsilon 4$  carriers. Only 2 controls (1.6%) had a value exceeding the stricter cut-off.

#### *Hippocampus activation and connectivity in relation to p-tau181 and APOE $\epsilon 4$*

In the imaging study, analyses were based on fMRI data from waves 5–7, and plasma samples from test waves 5 and 6 (Fig. 1b). The longitudinal tau-trajectories revealed that 46 individuals (53%) had low p-tau181 levels at both waves 5 and 6 (14 of these were  $\epsilon 4$  carriers), whereas 41 (47%) had higher levels that increased from wave 5 to 6 (Fig. 3a). The latter 41 individuals could be subdivided into  $\epsilon 4$  carriers ( $N=20$ ) and non-carriers ( $N=21$ ).

First, we examined whether brain activity in the anterior hippocampus during face-name encoding was related to tau accumulation across waves, and whether this relation differed for  $\epsilon 4$  carriers and non-carriers. A repeated measure ANOVA revealed a significant group by time interaction [ $F(4,168)=4.59$   $p=0.0015$ ]. *Post-hoc t*-tests revealed no group differences at waves 5 and 6 (Fig. 3b), but a marked reduction in bilateral hippocampus activity at wave 7 was evident in high p-tau  $\epsilon 4$  carriers (green) relative to the low p-tau group (blue;  $p < 0.001$ ) and also relative to the  $\epsilon 4$  non-carriers high p-tau group (red;  $p=0.018$ ). In the low p-tau181 group, there was no difference in anterior hippocampus activity between  $\epsilon 4$  carriers and non  $\epsilon 4$  carriers at W7 [ $t(44)=1.46$ ,  $p=0.15$ ].

Next, in view of recent findings of elevated local hippocampal connectivity during resting state [11], we quantified bilateral hippocampal connectivity at each test wave for the three sub-groups. In the  $\epsilon 4$  carriers with elevated p-tau levels, a gradual increase in hippocampus connectivity was observed across waves, and this increase was significant between waves 5 and 7 [Fig. 3c;  $t(18)=2.20$ ,  $p < 0.05$ ]. No significant longitudinal change in hippocampal connectivity was seen in the other groups; thus, hippocampal hyperconnectivity was specifically seen for  $\epsilon 4$  carriers with elevated p-tau181 levels. To explore

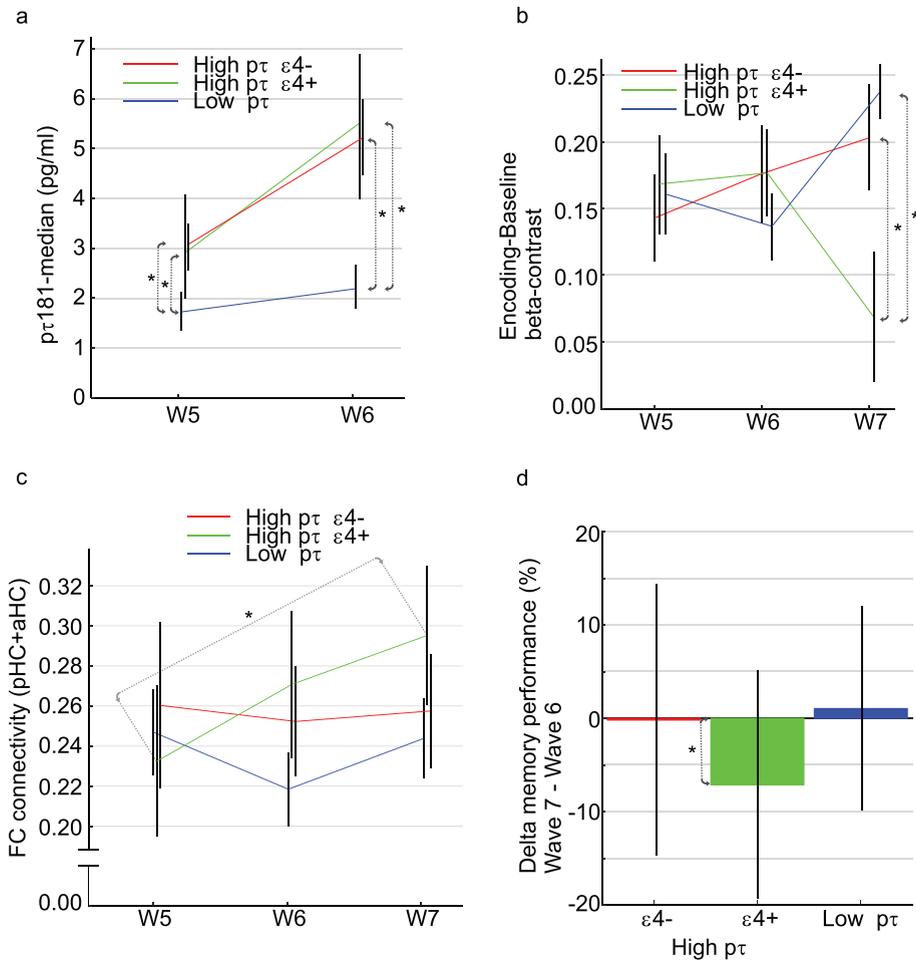


Fig. 3. Longitudinal brain-imaging study. a) Longitudinal change in p-tau181 levels in relation to APOE status. b) Longitudinal change in hippocampus activity and (c) connectivity as a function of p-tau181 levels (high/rising versus low) and APOE status ( $\epsilon 4$  carrier,  $\epsilon 4+$  versus non-carrier,  $\epsilon 4-$ ). d) Change in memory performance between waves 6–7. Error-bars are standard error of mean in (b) and (c). Error-bars are standard deviation in (d). Error-base are 25/75-percentile in (a). “\*\*\*” denotes a group difference with  $p < 0.05$  in (a) and (b), and within group change with  $p < 0.05$  in (c) and (d).

whether elevated hippocampal connectivity was accompanied by decreased hippocampus-cortical connectivity, we examined the connectivity between the hippocampus and the DMN. We found a significant decline in hippocampus-DMN connectivity from wave 5 to wave 6 ( $t = 2.62$ ,  $p = 0.0166$ ) for  $\epsilon 4$  carriers with elevated p-tau181 levels but not for the other two groups ( $ps > 0.05$ ).

The change in face-name memory performance across waves were examined for the three groups. Only for  $\epsilon 4$  carriers with elevated p-tau181 levels was a significant performance reduction from wave 6 to wave 7 observed [Fig. 3d;  $t(19) = 2.60$ ,  $p = 0.018$ ]. In further support of a relation between hippocampus activation and memory performance, across the sample, a significant change-change correlation

was observed between the wave 6–7 differences in anterior hippocampus activation and memory performance [ $r(85) = 0.28$ ,  $p = 0.0092$ ].

## DISCUSSION

### P-tau as a preclinical AD-marker

Our population-based study approach, with dementia as an exclusion criterion at initial study enrolment, provided novel information by means of early collected (W3) plasma samples from age- and sex-matched controls who were longitudinally assessed for up to 20 years to confirm that they remained non-demented. Consistent with their non-AD status,

the absolute majority of controls had low p-tau181 values. Preclinical AD as well as other pathologies or measurement error remain as possible sources of high values in a few controls, although re-assessment of those with the highest p-tau values revealed no clinical support for dementia progress during the follow-up period.

The control distribution offered a means to define a cut off for elevated p-tau181 level in preclinical and clinical AD cases. At the group level (Fig. 2a), significant case-control differences were seen close to disease onset ( $-2$  years), and at the individual level, for both control-group derived cut-offs considered, the proportion of cases with elevated p-tau181 levels exceeded that for controls. Still, with a mean for diagnosed cases of 6.4 pg/ml and less than 50% of cases having values exceeding the lenient cut-off of 6 pg/ml, it should be noted that our clinically diagnosed cases had modest p-tau181 levels relative to previous studies of p-tau181 in blood across the AD spectrum [31]. For example, p-tau181 cut-offs of 14–18 pg/ml were determined in relation to defining A $\beta$  status in the ADNI cohort [32], and p-tau181 values around 20–30 pg/ml were reported 8 years before postmortem [33]. While these differences in part might reflect different study populations, it should also be noted that the diagnostic assessment, implemented in the current study, was based on a clinical diagnostic approach [6] which together with the repeated follow-up assessments could have identified demented individuals at a quite early stage.

The present analyses converged to suggest that elevated plasma p-tau181 levels are indicative of forthcoming AD in preclinical APOE  $\epsilon 4$  carriers. Specifically, while no significant effect of APOE was seen near AD onset, in the early preclinical phases all cases with elevated values were  $\epsilon 4$  carriers. At the stage of clinically manifest AD, no influence of the genetic risk factor on tau burden is expected as all these individuals by definition have an active disease process [5]. In the preclinical phase, the APOE genotype could be regarded as a surrogate measure of time with A $\beta$  pathology, with each  $\epsilon 4$  allele being associated with  $\sim 10$  years earlier onset of A $\beta$  deposition. By this view, compared with non-carriers, similarly aged APOE  $\epsilon 4$  carriers with A $\beta$  pathology would be further along in the AD process which could explain the positive association with p-tau181 levels. The observed association between APOE and p-tau in the preclinical phase could reflect an APOE influence on tau pathology, such that the deposition of neurofibrillary tangles begins earlier and/or is accumulating

faster in  $\epsilon 4$  carriers. In support of this interpretation, APOE  $\epsilon 4$  has been linked to increased tau phosphorylation and tangle formation [34].

It should be noted that the lack of data on A $\beta$  load prevented us from assessing potential contributions to the observed p-tau181 findings of amyloid pathology [35]. Abnormal p-tau secretion may occur concomitantly with, or in response to, brain A $\beta$  pathology, including PHF-tau positive dystrophic neurites surrounding plaques, but recent studies suggest that APOE  $\epsilon 4$  may have an A $\beta$ -independent effect on tau-burden in the medial temporal cortex [36, 37]. Thus, speculatively, elevated p-tau181 levels close to disease onset in non-carriers would coincide with elevated A $\beta$  load, whereas in the preclinical phase elevated tau in APOE  $\epsilon 4$  carriers might precede overt amyloid pathology.

#### *Hippocampus functioning in relation to p-tau and APOE*

In the imaging study, the longitudinal design identified a sub-group of  $\epsilon 4$  carriers with increasing p-tau181 levels across test waves. This sub-group could be compared with an APOE  $\epsilon 4$  non-carrier group showing a similar longitudinal increase in p-tau levels, but also with a low p-tau181 group. Only for  $\epsilon 4$  carriers was increasing p-tau181 related to subsequent reduction (between waves 6–7) in hippocampus encoding-related activity. This observation supports findings that presence of APOE  $\epsilon 4$  and lower hippocampal fMRI activation is a predictor of future neurocognitive impairment [38] and extends previous reports by highlighting p-tau as a possible mechanism.

The finding that a longitudinal increase in p-tau levels translated into a reduced hippocampus activity during memory encoding is noteworthy, as several previous cross-sectional studies reported that APOE  $\epsilon 4$  [13, 39, 40] and higher tau levels [11, 12, 41, 42] are related to higher hippocampal activity and connectivity. Here, too, we observed a longitudinal increase in resting-state hippocampus connectivity for  $\epsilon 4$  carriers with high/rising p-tau levels. This constitutes novel, within-person evidence for the concept of hyperconnectivity. The trend for elevation in connectivity was seen already between waves 5–6, coinciding with tau elevation, whereas the reduction in task activity was seen at the final wave. This cascade of event is in line with previous observations that increased local hippocampal connectivity during rest translated into lower task-related hippocampus

activation during memory encoding [25], and findings that tau deposition drives elevated local hippocampal connectivity and induces hippocampal disconnection from functional networks of relevance for episodic memory [11]. Strikingly, consistent with these findings, we observed a disconnection of the hippocampus from the DMN for  $\epsilon 4$  carriers with high/rising p-tau levels with a significant connectivity decrease from wave 5 to 6 (i.e., the period during which elevated p-tau was observed). Thus, tau accumulation may induce increased local hippocampal synchrony, and eventually reduced ability to engage the hippocampus during active mnemonic processing. Although the longitudinal design is a strength of the current study, our small sample size and lack of p-tau at T7 limit the choice of analytical approach.

The apparent selectivity for  $\epsilon 4$  carriers for rising p-tau181 levels could reflect an APOE effect on rate of tau accumulation, and it is also consistent with findings that APOE  $\epsilon 4$  aggravates the negative influence of tau pathology on neuronal function [3]. Interaction of p-tau and APOE genotype on AD has been related to dysfunction of synaptic plasticity. Recent studies have shown that alterations in long-term potential plasticity are associated with cognitive decline [43] and elevated p-tau level [44]. Moreover, elevated p-tau was associated with impaired long-term potential plasticity and accelerated progress of disease in AD patients with  $\epsilon 4$  carriage [8].

### Conclusion

Our findings indicate an association of APOE  $\epsilon 4$  and tau with preclinical AD and hippocampus functioning. In APOE  $\epsilon 4$  carriers only, elevated plasma p-tau181 levels were indicative of forthcoming clinically-defined AD, and longitudinal p-tau181 increases were paralleled by increased hippocampus synchrony at rest and reduced hippocampus encoding-related activity. These findings support and extend prior observations of selective neurocognitive APOE effects in normal aging and age-related pathology [8, 45, 46].

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### SUPPLEMENTARY MATERIAL

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