Sex and APOE ε4 modify the effect of cardiovascular risk on tau in cognitively normal older adults

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The interaction between APOE ε4 and vascular risk factors on cognitive function is stronger in women than in men. These effects may be mediated by the amount of tau pathology in the brain. Therefore, we examined whether APOE ε4 and sex modify cross-sectional associations between cardiovascular risk and tau deposition in cognitively normal older adults from the Alzheimer’s Disease Neuroimaging Initiative. We calculated the Framingham Heart Study cardiovascular disease risk score for 141 participants (74 women, 47 APOE ε4 carriers) with complete medical history data, processed tau-PET data and a Clinical Dementia Rating global score of 0.0 at the time of the tau-PET scan, implying no significant cognitive or functional impairment. We used linear regression models to examine the effects of sex, APOE ε4, cardiovascular risk and their interactions on tau deposition in the entorhinal cortex, inferior temporal cortex and a composite meta-region of interest of temporal lobe areas. We found a significant three-way interaction among sex, APOE ε4 status and cardiovascular disease risk on tau deposition in the entorhinal cortex (β = 0.04; 95% CI, 0.01–0.07; P = 0.008), inferior temporal cortex (β = 0.02; 95% CI, 0.0–0.05; P = 0.029) and meta-region (β = 0.02; 95% CI, 0.0–0.04; P = 0.042). After stratifying by APOE ε4 status to examine interactions between sex and cardiovascular disease risk on tau in APOE ε4 carriers and non-carriers, we found a significant two-way interaction between sex and cardiovascular disease risk on tau in the entorhinal cortex (β = 0.05; 95% CI, 0.02–0.08; P = 0.001), inferior temporal cortex (β = 0.03; 95% CI, 0.01–0.05; P = 0.009) and meta-region (β = 0.02; 95% CI, 0.01–0.04; P = 0.008) only among APOE ε4 carriers. In analyses stratified by sex, higher cardiovascular risk scores were associated with higher levels of tau in the entorhinal cortex (β = 0.05; 95% CI, 0.02–0.08; P = 0.002), inferior temporal cortex (β = 0.02; 95% CI, 0.0–0.05; P = 0.023) and meta-region (β = 0.02; 95% CI, 0.01–0.04; P = 0.013) in female APOE ε4 carriers but not in male carriers. Our findings suggest that cognitively normal older women carrying at least one APOE ε4 allele, may be particularly vulnerable to the effects of cardiovascular disease risk on early tau deposition.
Introduction

While it is clear that vascular risk factors\textsuperscript{1,2} and vascular neuropathology\textsuperscript{2} are associated with the clinical diagnosis of Alzheimer’s disease, some investigators have also proposed that vascular risk factors may be aetiologically relevant to Alzheimer’s disease neuropathology.\textsuperscript{3} Risk factors such as hypertension can contribute to brain hypoperfusion and blood–brain barrier damage, potentially facilitating amyloid and tau accumulation in the brain.\textsuperscript{4} Growing evidence from animal\textsuperscript{5} and human\textsuperscript{6-9} studies indicates that vascular risk burden is associated with increases in tau pathology, while a few studies report associations between vascular risk and amyloid accumulation only,\textsuperscript{10} or both amyloid and tau pathology.\textsuperscript{11} One group found that tau deposition mediated associations between cerebrovascular disease and cognitive impairment,\textsuperscript{12} and another study showed that tau burden mediated associations of cerebral blood flow and a marker of pericyte injury with cognitive scores.\textsuperscript{13}

Studies have also shown that the impact of cardiovascular risk burden on cognitive decline and Alzheimer’s disease progression is stronger among carriers of the apolipoprotein \textit{ɛ}4 (APOE \textit{ɛ}4) allele.\textsuperscript{14-16} APOE \textit{ɛ}4, the major genetic risk factor for Alzheimer’s disease, cannot only promote Alzheimer’s disease pathogenesis directly by reducing amyloid clearance and increasing tau phosphorylation,\textsuperscript{17-20} but can also indirectly impact Alzheimer’s disease pathogenesis by increasing cardiovascular risk factors such as elevated...
plasma low-density lipoproteins, which are also risk factors for Alzheimer’s disease. A recent study of dementia-free older adults demonstrated that higher vascular risk was associated with a stronger effect of APOE ε4 on memory decline in women but not men. Although these findings are consistent with evidence of a more robust APOE ε4 effect on clinical outcomes and tau pathology in women, the mechanism for a stronger APOE ε4 effect in women remains unknown.

Sex differences in the effect of cardiovascular burden on brain health could constitute a strong candidate mechanism. Our group previously found that among cognitively normal (CN) older adults aged between 56 and 75 years, women displayed stronger associations between higher pulse pressure, a surrogate marker of arterial stiffness, and white matter microstructural abnormalities, compared with men in the same age group. We also previously demonstrated that among a cohort of community-dwelling older adults with a mean age of 67 years, higher cardiovascular risk was associated with greater executive function and memory decline among women but not men, despite higher risk scores among men. Similarly, another group found that among older adults between 60 and 100 years of age, higher systolic blood pressure was correlated with worse cognitive performance among hypertensive women but not men. Given the strong link between tau pathology and clinical presentation of Alzheimer’s disease, the exacerbated effects of cardiovascular risk on cognition in women could be related to tau pathology. Therefore, we examined the role of sex and APOE ε4 status in modifying the effect of cardiovascular disease risk on cortical tau in CN older adults.

Materials and methods

Participants

All data used in preparation of this manuscript were obtained from the Alzheimer’s Disease Neuroimaging Initiative (ADNI) database (accessed from ida.loni.usc.edu in April 2021). ADNI is a study which aims to measure the progression of MCI and early Alzheimer’s disease using multi-modal imaging methods. ADNI inclusion criteria have been described previously (refer to http://adni.loni.usc.edu/). Notably, individuals with Hachinski ischaemic scores >4 at screening are excluded from ADNI. The Hachinski Ischaemic score is a widely used questionnaire designed to distinguish individuals with vascular dementia from individuals with pure Alzheimer’s disease and mixed dementia (i.e., dementia caused by both Alzheimer’s disease and vascular pathology). Additional inclusion criteria for our study included fully processed regional Flortaucipir PET (FTP-PET) for measurement of tau deposition, clinical and medical history data required for cardiovascular disease risk calculations, as well as a Clinical Dementia Rating global score (CDR-GS) equal to 0.0 at the time of FTP-PET scan.

Clinical assessment

We used the participants’ Clinical Dementia Rating scale (CDR) score to determine clinical status at the time of FTP-PET scan. This was done to avoid potential sex-related bias from using ADNI-determined diagnoses, which rely on verbal memory tests in which women frequently outperform men despite similar levels of Alzheimer’s disease pathology. We obtained CDR-GS measured either at the same study visit as the FTP-PET scan or at the study visit closest to the FTP-PET scan (average time difference between CDR and FTP-PET measurement was 52 days). Participants with a CDR-GS equal to 0.0 were included.

Cardiovascular disease risk assessment

We quantified cardiovascular disease risk using the Framingham Heart Study cardiovascular disease (FHS-CVD) risk algorithm—a well-validated tool that has been used in both research and primary care. The FHS-CVD risk score is a sex-specific measure of cardiovascular disease risk accounting for age, systolic blood pressure, anti-hypertensive treatment, total cholesterol, high density lipoprotein cholesterol, self-reported history of diabetes and current smoking status. The risk score represents the probability of cardiovascular events (such as coronary death, myocardial infarction, etc.) occurring within 10 years of risk assessment. We chose the FHS-CVD risk measure obtained closest to the FTP-PET scan with the restriction that the period of time between FHS-CVD assessment and FTP-PET scan could not be more than 10-years. This criterion was applied to ensure this time lag did not exceed the 10-year predictive window of the cardiovascular risk assessment.

APOE genotyping

APOE genotype data were available for all participants and was obtained from the LONI database. Details regarding blood sample collection and genotyping procedures for ADNI can be found in adni.loni.usc.edu. Homozygotes and heterozygotes for the ε4 allele (including ε2/ε4, ε3/ε4 and ε4/ε4 genotypes) were combined into a single group and categorized as APOE ε4 carriers, while participants with ε2/ε3 or ε3/ε3 genotypes were categorized as APOE ε4 non-carriers. There were no ε2 homozygotes in our study sample.

Brain imaging acquisition and processing

We downloaded Aβ and tau-PET data (from ida.loni.usc.edu, accessed in April 2021) which Landau and colleagues had previously processed at the University of California, Berkeley. ADNI brain imaging acquisition and processing procedures have been described in detail elsewhere http://adni.loni.usc.edu/methods/documents. Briefly, structural
MRI scans are performed on 3T scanners using either a 3D MPRAGE or IR-SPGR T₁-weighted sequence with sagittal slices and spatial resolution of 1.1 × 1.1 × 1.2 mm³. Structural MRI scans are then skull-stripped, segmented and parcellated using the FreeSurfer (version 5.3.0; http://surfer.nmr.mgh.harvard.edu). Aβ PET images are acquired 50–70 min post Florbetapir injection in a series of four 5-min frames and tau-PET images are acquired 75–105 min after FTP injection in a series of six 5-min frames. After the raw PET data are assessed for quality, each of the acquired frames is extracted and co-registered to the first frame to account for subject motion. The motion-corrected dynamic image set is then averaged and smoothed to a uniform isotropic resolution of 8 mm full width at half maximum and then co-registered with the subject’s processed structural MRI. Standardized uptake value ratios (SUVRs) of FTP uptake are computed for each FreeSurfer-derived region by referencing the mean cerebellar grey matter deposition.41 To determine Aβ positivity we used the SUVR of a cortical summary region, which is intensity normalized by a whole-cerebellum FreeSurfer region. An SUVR cut-off of 1.11 was used to determine Aβ positivity.42,43 Lastly, regional FTP data are corrected for partial-volume effects using the Geometric Transfer Matrix approach.44,45 To examine regional tau burden, we used a bilateral volume-weighted composite of the entorhinal cortex (EC) and the inferior temporal cortex (ITC), given that these regions are cortical sites of early Alzheimer’s disease-related tau deposition.46–48 Additionally, we examined tau deposition in a larger and likely more stable composite region of interest (meta-ROI) of tau deposition, consisting of a volume-weighted average of the bilateral amygdala, fusiform gyrus, middle temporal cortex, EC and ITC.

**Statistical analyses**

All statistical analyses were computed with the R (version 4.0.3). Differences in demographic and clinical variables by sex and APOE ε4 status were assessed using Welch’s independent t-tests for continuous variables and Fisher’s exact tests for categorical variables. Next, we applied a series of linear regression models to examine the modifying role of sex and APOE ε4 on the association between FHS-CVD risk score and tau deposition covarying for the time lag between FHS-CVD risk measurement and FTP-PET scan. First, we used a linear regression model testing for a three-way interaction between sex, APOE ε4 status and FHS-CVD risk on tau. In the case of a significant three-way interaction, we stratified our sample by APOE ε4 status, and used a linear regression model testing for a two-way interaction between sex and FHS-CVD risk on tau in APOE ε4 carriers and APOE ε4 non-carriers separately. Finally, in the case of a significant two-way interaction between sex and FHS-CVD risk on tau, we further stratified by sex to examine the main effect of cardiovascular risk in men and women separately. Analyses were repeated with models additionally adjusted for age at FTP-PET scan, Aβ status and years of education to assess potential confounding effects.

**Data availability**

Data used in preparation of this manuscript were obtained from the LONI database in April 2021 (ida.loni.usc.edu).

**Results**

**Sex differences in demographic characteristics**

Of the sample of 141 participants who met our inclusion criteria (Fig. 1), 52% were women and 33% carried at least one APOE ε4 allele. Participants’ age at the time of cardiovascular risk assessment ranged between 55 and 90 years [mean (SD) = 72.1 (6.45)] and time lag between cardiovascular risk assessment and FTP-PET scan ranged between 2 and 10 years [mean (SD) = 4.94 (1.55)]. Women in our sample had fewer years of education (P < 0.05), lower FHS-CVD risk scores (P < 0.001) and higher total and high density lipoprotein (HDL) cholesterol (P < 0.001 and P < 0.0001, respectively) than men. Among APOE ε4 carriers, women had lower FHS-CVD risk scores (P < 0.0001), were younger at the time of FHS-CVD risk assessment (P < 0.05) and had higher total and HDL cholesterol (P < 0.05 and P < 0.01, respectively) than men. Among APOE ε4 non-carriers, women had fewer years of education (P < 0.01), higher HDL cholesterol (P < 0.001) and lower FHS-CVD risk scores (P < 0.0001) than men. Details regarding demographic and clinical variables of our sample can be found under Table 1.

**Interaction effect of sex, APOE ε4 and FHS-CVD risk on tau**

There was a significant three-way interaction among sex, APOE ε4 status and FHS-CVD risk on tau deposition in the EC (β = 0.04; 95% CI, 0.01–0.07; P = 0.008), ITC (β = 0.02; 95% CI, 0.0–0.05; P = 0.029) and meta-ROI (β = 0.02; 95% CI, 0.0–0.04; P = 0.042) (Table 2). To aid our interpretation of the three-way interaction between sex, APOE ε4 and FHS-CVD risk on tau deposition, we stratified by APOE ε4 status to examine two-way interactions between sex and FHS-CVD risk on tau in APOE ε4 carriers and non-carriers separately. We found a significant two-way interaction between sex and FHS-CVD risk on tau in the EC (β = 0.05; 95% CI, 0.02–0.08; P = 0.001) (Fig. 2A), ITC (β = 0.03; 95% CI, 0.01–0.05; P = 0.009) (Fig. 2B) and meta-ROI (β = 0.02; 95% CI, 0.01–0.04; P = 0.008) (Fig. 2C) only among APOE ε4 carriers (Table 3). Since two-way interactions between sex and FHS-CVD risk were significant only among APOE ε4 carriers, we further stratified by sex to examine main effects of FHS-CVD risk on tau in male and female APOE ε4 carriers separately. We found that higher FHS-CVD risk was associated with higher levels of tau in the EC (β = 0.05; 95% CI, 0.02–0.08; P = 0.002), ITC (β = 0.02; 95% CI, 0.0–0.05; P = 0.023) and
meta-ROI ($\beta = 0.02; 95\% \text{ CI}, 0.01–0.04; P = 0.013$) in female APOE $\varepsilon 4$ carriers but not male carriers (Table 4).

The three-way interaction between APOE $\varepsilon 4$, sex and FHS-CVD risk on EC tau deposition ($\beta = 0.04; 95\% \text{ CI}, 0.01–0.07; P = 0.012$) remained significant after additionally adjusting for age, A$\beta$ status and education, though this effect was attenuated in the ITC ($\beta = 0.02; 95\% \text{ CI}, -0.01–0.04; P = 0.058$) and meta-ROI ($\beta = 0.01; 95\% \text{ CI}, -0.01–0.02; P = 0.093$) (Table 5). Notably, A$\beta$ positive participants had higher levels of tau across all three regions, though the main effect of A$\beta$ status on tau was more robust in ITC ($\beta = 0.18; 95\% \text{ CI}, 0.10–0.26; P < 0.001$) and meta-ROI ($\beta = 0.15; 95\% \text{ CI}, 0.09–0.22; P < 0.001$) compared with the EC ($\beta = 0.13; 95\% \text{ CI}, 0.01–0.26; P = 0.030$). APOE $\varepsilon 4$ stratified analyses revealed a significant two-way interaction effect of sex and FHS-CVD risk on EC ($\beta = 0.05; 95\% \text{ CI}, 0.02–0.08; P = 0.004$), ITC ($\beta = 0.02; 95\% \text{ CI}, 0.00–0.04; P = 0.049$) and meta-ROI ($\beta = 0.02; 95\% \text{ CI}, 0.00–0.03; P = 0.045$) tau deposition among APOE $\varepsilon 4$ carriers but not non-carriers, after additionally adjusting for age, A$\beta$ status and education (Table 6). A$\beta$ status had a consistent main effect on tau deposition in the ITC and meta-ROI among both APOE $\varepsilon 4$ carriers and non-carriers. Finally, in sex-stratified analyses of APOE $\varepsilon 4$ carriers we found that higher FHS-CVD risk scores were associated with higher levels of tau in the EC ($\beta = 0.06; 95\% \text{ CI}, 0.03–0.09; P = 0.002$) and meta-ROI ($\beta = 0.02; 95\% \text{ CI}, 0.00–0.04; P = 0.028$) among female but not male carriers (Table 7). The effect of FHS-CVD risk on ITC tau in female carriers was no longer significant ($\beta = 0.02; 95\% \text{ CI}, -0.00 to 0.04; P = 0.052$) after adjusting for age, education and A$\beta$ status. Lastly, we observed no effect of age and education on tau deposition in any of the models.

**Discussion**

In this study of CN older adults, we found that sex and APOE $\varepsilon 4$ status modified the association between cardiovascular risk and tau deposition. After stratifying our sample by APOE $\varepsilon 4$ status, we found that sex modified the association between cardiovascular risk and tau burden, among APOE $\varepsilon 4$ carriers only. Finally, sex-stratified analyses showed that higher cardiovascular risk was associated with higher levels of EC, ITC and meta-ROI tau among female APOE $\varepsilon 4$ carriers but not male carriers. Adjusting for additional factors may provide further insights into the mechanisms underlying the association between cardiovascular risk and tau deposition.
Our findings are in line with other studies demonstrating stronger predictive relationships between vascular risk factors such as elevated plasma lipids and hypertension and levels of tau deposition in the ITC but not the EC. Despite the strong main effect of Aβ status on ITC and meta-ROI tau among APOE ε4 carriers and non-carriers alike, two-way interactions between female sex and cardiovascular risk on tau deposition were seen only among APOE ε4 carriers. Therefore, studies on larger samples should examine sex differences in the association between cardiovascular risk and tau deposition among groups stratified by both APOE ε4 and Aβ status.

Table 1 Demographic and clinical sample characteristics

| Variables                        | Total cohort (n = 141) | APOE ε4 carriers (n = 47) | APOE ε4 non-carriers (n = 94) |
|----------------------------------|------------------------|---------------------------|-----------------------------|
|                                  | Women (n = 74)         | Men (n = 67)              | Women (n = 24)              | Men (n = 23)                  | Women (n = 50)            | Men (n = 44)            |
| Aβ positive, n (%)               | 28 (37.8)              | 23 (34.3)                 | 14 (58.3)                   | 12 (52.2)                    | 14 (28)                   | 11 (25)                 |
| Education, years, mean (SD)      | 16.3 (2.44)            | 17.3 (2.47)               | 16.5 (2.38)                 | 16.7 (2.85)                  | 16.1 (2.48)               | 17.6 (2.21)             |
| White, n (%)                     | 70 (94.6)              | 63 (94)                   | 23 (95.8)                   | 22 (95.7)                    | 47 (94)                   | 41 (93.2)               |
| African American or Black, n (%) | 3 (4.1)                | 2 (3)                     | 1 (4.2)                     | 1 (4.3)                      | 2 (4)                     | 1 (2.3)                 |
| Asian, n (%)                     | 0                      | 2 (3)                     | 0                           | 0                            | 0                        | 2 (4.5)                 |
| More than one race, n (%)        | 1 (1.4)                | 0                         | 0                           | 0                            | 1 (2)                     | 0                       |
| FHS-CVD risk score, %, mean (SD) | 12.1 (5.76)            | 26.7 (11.3)               | 12.1 (5.89)                 | 29.6 (14.3)                  | 12.1 (5.76)               | 25.1 (6.25)             |
| Age, years, mean (SD)            | 71.7 (6.08)            | 72.6 (0.86)               | 69.0 (6.04)                 | 73.4 (7.96)                  | 73.0 (5.7)                | 72.1 (6.25)             |
| Systolic blood pressure, mm Hg, mean (SD) | 134 (14.1)      | 136 (14.9)               | 136 (14.9)                  | 136 (13.9)                   | 134 (13.8)                | 136 (15.4)              |
| Total cholesterol, mg/dL, mean (SD) | 197 (38.6)          | 181 (29.6)                | 207 (38.6)                  | 181 (35.5)                   | 192 (38)                  | 182 (26.5)              |
| HDL cholesterol, mg/dL, mean (SD) | 64.6 (15.5)           | 53.5 (11.0)               | 67.1 (15.4)                 | 53.3 (12.0)                  | 63.4 (15.6)               | 53.6 (10.6)             |
| Taking anti-hypertensive medication, n (%) | 15 (20.3)           | 11 (16.4)                | 6 (25)                      | 7 (30.4)                     | 9 (18)                    | 4 (9.1)                 |
| Diabetic, n (%)                  | 6 (8.1)                | 8 (11.9)                  | 2 (8.3)                     | 4 (17.4)                     | 4 (8)                     | 4 (9.1)                 |
| Smokers, n (%)                   | 1 (1.4)                | 3 (4.5)                   | 0                           | 1 (4.3)                      | 1 (2)                     | 2 (4.5)                 |
| Enthorinal cortex tau-PET SUVR, mean (SD) | 1.82 (0.398)        | 1.65 (0.303)             | 1.94 (0.492)                | 1.77 (0.351)                 | 1.76 (0.334)              | 1.58 (0.255)           |
| Inferior temporal cortex tau-PET SUVR, mean (SD) | 1.73 (0.282)         | 1.60 (0.172)             | 1.8 (0.309)                 | 1.67 (0.212)                 | 1.70 (0.265)              | 1.56 (0.134)           |
| Meta-ROI temporal lobe tau-PET SUVR, mean (SD) | 1.62 (0.229)        | 1.50 (0.150)             | 1.68 (0.257)                | 1.56 (0.185)                 | 1.59 (0.210)              | 1.47 (0.119)           |
| Time between FHS-CVD Assessment and tau-PET scan, years, mean (SD) | 4.77 (1.55)          | 5.13 (1.55)              | 4.79 (1.82)                 | 5.00 (1.21)                  | 4.76 (1.42)               | 5.2 (1.71)              |

Mean and SD are provided for continuous variables and n and % are provided for categorical variables. Aβ = amyloid-β; FHS-CVD = Framingham Heart Study cardiovascular disease; HDL = high density lipoprotein; SUVR = standardized uptake value ratio.

Table 2 Three-way interaction effect of APOE ε4, sex and cardiovascular risk on tau deposition

| Predictor variables | EC tau-PET SUVR | ITC tau-PET SUVR | Meta-ROI tau-PET SUVR |
|---------------------|-----------------|------------------|-----------------------|
|                     | Estimate        | CI               | P-value               | Estimate        | CI               | P-value               | Estimate        | CI               | P-value               |
| FHS-CVD risk        | 0.01            | −0.0 to 0.02     | 0.126                 | 0.00            | −0.01 to 0.01    | 0.939                 | 0.00            | −0.01 to 0.01    | 0.927                 |
| APOE ε4 carriers    | 0.28            | −0.14 to 0.70    | 0.187                 | 0.15            | −0.15 to 0.45    | 0.317                 | 0.10            | −0.15 to 0.34    | 0.425                 |
| Sex (female)        | 0.23            | −0.12 to 0.58    | 0.195                 | 0.12            | −0.13 to 0.37    | 0.358                 | 0.07            | −0.14 to 0.27    | 0.525                 |
| Time lag            | 0.03            | −0.00 to 0.07    | 0.059                 | −0.00           | −0.03 to 0.02    | 0.913                 | −0.00           | −0.02 to 0.02    | 0.920                 |
| APOE ε4 carriers × FHS-CVD risk | −0.00               | −0.02 to 0.01                   | 0.563                     | 0.00           | −0.01 to 0.01    | 0.781                 | −0.00           | −0.01 to 0.01    | 0.927                 |
| Sex (female) × FHS-CVD risk | 0.01                  | −0.01 to 0.02                   | 0.539                     | 0.00           | −0.01 to 0.02    | 0.783                 | −0.00           | −0.01 to 0.02    | 0.477                 |
| APOE ε4 carriers × sex (female) | −0.57                  | −1.12 to −0.01            | 0.048**                 | −0.33          | −0.73 to 0.06    | 0.099                 | −0.23          | −0.55 to 0.10    | 0.175                 |
| APOE ε4 carriers × sex (female) × FHS-CVD risk | 0.04                      | 0.01 to 0.07                        | 0.008**                 | 0.02           | 0.00 to 0.05     | 0.029**               | 0.02           | 0.00 to 0.04     | 0.042**               |

Results from linear regression analysis revealed a significant three-way interaction effect of APOE ε4, sex and FHS-CVD risk on tau deposition in the EC, ITC and meta-ROI after adjusting for the time between FHS-CVD risk assessment and tau-PET scan. Significant P-values are presented in bold, where ***P < 0.005, **P < 0.01, *P < 0.05. CI = confidence interval; EC = enthorinal cortex; FHS-CVD risk = Framingham Heart Study cardiovascular disease risk; ITC = inferior temporal cortex; SUVR = standardized uptake value ratio.

covariates such as age, Aβ status and education attenuated interaction effects of sex, APOE ε4 and cardiovascular risk on tau deposition in the ITC and meta-ROI, but not the EC. The absence of a three-way interaction effect in the ITC and meta-ROI appears to be due to a more robust main effect of Aβ status on tau deposition in these regions among both APOE ε4 carriers and non-carriers. There was no such effect on tau deposition in the EC, which has been shown to be associated with APOE ε4, independently of Aβ burden. 

Furthermore, a recent study on CN older adults found that the interaction between higher cardiovascular risk and elevated Aβ burden was associated with higher
incident Alzheimer’s disease in women compared with men, and are consistent with our prior study demonstrating that among community-dwelling older adults, higher FHS-CVD risk scores were associated with greater executive function and memory decline among women but not men, despite higher risk scores among men. Tau pathology is strongly correlated with clinical symptomatology, and recent evidence suggests that associations between cerebrovascular function and cognition are mediated by tau pathology. Therefore, the exacerbated effect of cardiovascular disease risk on cognition and clinical outcomes observed among women may be, at least partly, due to elevated tau deposition.

While our study was not designed to probe mechanisms of this effect, there are potential explanations, including the impact of hormonal changes in post-menopausal women: The abrupt decrease of oestrogen levels in post-menopausal women is associated with reduced glucose tolerance, as well as increased blood pressure, endothelial dysfunction and vascular inflammation. Oestrogen-binding on endothelial and vascular smooth muscle cells has been shown to prevent neointimal responses to acute vascular injury, resulting from atherosclerosis, hypertension and other vascular diseases.

Therefore, the absence of the aforementioned vaso- and neuroprotective effects of oestrogen in post-menopausal women may contribute to a heightened inflammatory response to vascular risks, potentially leading to worse brain outcomes in women. Evidence of a more direct link between oestrogen and tau comes from animal studies showing that oestrogen depletion can lead to an increase in enzymatic activity (including protein kinase A and glycogen synthase kinase 3-β) involved in tau hyperphosphorylation.

We found that sex differences in the association between cardiovascular risk and tau deposition were restricted to APOE ε4 carriers. This finding is consistent with a recent study showing that higher vascular risk and APOE ε4 interactively predicted memory decline in women but not men. Our results extend prior evidence of a stronger APOE ε4 effect on tau pathology in women, and introduce the role of cardiovascular disease risk as a potential factor contributing to this female APOE ε4-related susceptibility to tauopathy. Notably, APOE ε4 confers greater risk of developing MCI in women specifically between 55 and 70 years of age, compared with men. In this study, we found an effect of cardiovascular disease risk on tau deposition only among female APOE ε4 carriers, who were between 55 and 78 years of age. It is

Figure 2 Sex and APOE ε4 modify associations between cardiovascular disease risk and tau deposition. Scatter plots depicting sex differences in associations between FHS-CVD risk score and tau deposition in the (A) entorhinal cortex, (B) inferior temporal cortex and (C) a composite temporal lobe meta-ROI, among APOE ε4 non-carriers (top) and APOE ε4 carriers (bottom). Shaded regions represent 95% confidence intervals. We found significant interactions between sex and FHS-CVD risk on tau deposition in the EC (β = 0.05; 95% CI, 0.02–0.08; P = 0.001), ITC (β = 0.03; 95% CI, 0.01–0.05; P = 0.009) and meta-ROI (β = 0.02; 95% CI, 0.01–0.04; P = 0.008) among APOE ε4 carriers (bottom) but not among non-carriers (top).
conceivable that the first two decades following menopause could be a critical time-window during which elevated cardiovascular vulnerability may predispose female APOE ε4 carriers to worse pathological and clinical outcomes.

Although the mechanisms of the interaction between APOE ε4 and vascular risk and its exacerbated impact on brain health among women remains unknown, the influence of sex hormones may be relevant. Oestrogen has been shown to inhibit microglial activation in response to acute inflammatory stimuli, thereby reducing the production of reactive oxygen species and proinflammatory molecules such as matrix metalloproteinase 9. However, the presence of an APOE ε4 allele inhibits the anti-inflammatory effects of oestrogen on microglial and peripheral microphage activation, suggesting that female APOE ε4 carriers are less able to benefit from the anti-inflammatory properties of circulating oestrogen, compared with non-carriers. Conversely, testosterone, which has similar anti-inflammatory properties, has also been shown to interact with APOE ε4 to impact cognitive function and Alzheimer’s disease risk. Animal studies showed that APOE ε4 can reduce androgen receptor levels in the brain, suggesting that APOE ε4 carriers may be particularly vulnerable to the effects of lower testosterone.

Sundermann et al. found that the association between lower testosterone and higher levels of cerebrospinal fluid phosphorylated tau were strongest among female APOE ε4 carriers. Consequently, the heightened vascular inflammation seen in post-menopausal women along with the heightened cerebrovascular impact of APOE ε4 in the absence of the neuroprotective effects of sex hormones may interact to exacerbate the effects of cardiovascular risk factors on Alzheimer’s disease tauopathy in female APOE ε4 carriers. Future experimental research is needed to probe this possible explanation for our findings.

**Limitations**

The comparatively higher mortality rates at younger ages due to cardiovascular disease in men may have resulted in a survival bias in our study. This survival bias along with

### Table 3 Two-way interaction effect of sex and cardiovascular disease risk on cortical tau deposition among APOE ε4 carriers but not non-carriers

| Predictor variables | EC tau-PET SUVR | ITC tau-PET SUVR | Meta-ROI tau-PET SUVR |
|---------------------|-----------------|------------------|-----------------------|
|                     | Estimate        | CI               | P-value               | Estimate        | CI               | P-value               |
| APOE ε4 carriers    |                 |                  |                       |                 |                  |                       |
| FHS-CVD risk        | 0.00            | −0.01 to 0.01    | 0.553                 | −0.00           | −0.01 to 0.01   | 0.753                 | −0.00           | −0.01 to 0.01   | 0.968                 |
| Sex (female)        | −0.36           | −0.88 to 0.16    | 0.169                 | −0.22           | −0.57 to 0.13   | 0.21                  | −0.16           | −0.45 to 0.13   | 0.275                 |
| Time lag            | 0.06            | −0.02 to 0.13    | 0.132                 | 0.00            | −0.05 to 0.05   | 0.965                 | 0.00            | −0.04 to 0.04   | 0.995                 |
| Sex (female) × FHS-CVD risk | 0.05         | 0.02–0.08        | **0.001***          | 0.03            | 0.01–0.05       | **0.009**            | 0.02            | 0.01–0.04       | **0.008**            |
| APOE ε4 non-carriers|                 |                  |                       |                 |                  |                       |
| FHS-CVD risk        | 0.01            | −0.00 to 0.02    | 0.105                 | 0.00            | −0.01 to 0.01   | 0.942                 | 0.00            | −0.01 to 0.01   | 0.925                 |
| Sex (female)        | 0.22            | −0.10 to 0.54    | 0.176                 | 0.12            | −0.12 to 0.36   | 0.341                 | 0.07            | −0.13 to 0.26   | 0.503                 |
| Time lag            | 0.02            | −0.02 to 0.06    | 0.236                 | −0.00           | −0.03 to 0.03   | 0.863                 | −0.00           | −0.02 to 0.02   | 0.894                 |
| Sex (female) × FHS-CVD risk | 0.01         | −0.01 to 0.02 | 0.490 | 0.00            | −0.01 to 0.01   | 0.771                 | 0.00            | −0.01 to 0.01   | 0.418                 |

Results from linear regression analysis on APOE ε4 stratified groups revealed a significant interaction effect of sex and FHS-CVD risk on tau deposition in the EC, ITC and meta-ROI only among APOE ε4 carriers, after adjusting for the time between FHS-CVD risk assessment and tau-PET scan. Significant P-values are presented in bold, where *P < 0.05, **P < 0.01, ***P < 0.005. CI = confidence interval; EC = entorhinal cortex; FHS-CVD risk = Framingham Heart Study cardiovascular disease risk; ITC = inferior temporal cortex; SUVR = standardized uptake value ratio.

### Table 4 Main effect of cardiovascular disease risk on cortical tau deposition in female APOE ε4 carriers but not male carriers

| Predictor variables | EC tau-PET SUVR | ITC tau-PET SUVR | Meta-ROI tau-PET SUVR |
|---------------------|-----------------|------------------|-----------------------|
|                     | Estimate        | CI               | P-value               | Estimate        | CI               | P-value               |
| Female APOE ε4 carriers |                 |                  |                       |                 |                  |                       |
| FHS-CVD risk        | 0.05            | 0.02–0.08        | **0.002***           | 0.02            | 0.00–0.05       | **0.023**           | 0.02            | 0.01–0.04       | **0.013**           |
| Time lag            | 0.03            | −0.07 to 0.13    | 0.511                 | −0.02           | −0.09 to 0.04   | 0.475                 | −0.02           | −0.08 to 0.03   | 0.349                 |
| Male APOE ε4 carriers |                 |                  |                       |                 |                  |                       |
| FHS-CVD risk        | 0.00            | −0.01 to 0.01    | 0.768                 | −0.00           | −0.01 to 0.00   | 0.371                 | −0.00           | −0.01 to 0.00   | 0.494                 |
| Time lag            | 0.12            | −0.01 to 0.25    | 0.064                 | 0.07            | −0.02 to 0.15   | 0.107                 | 0.07            | −0.00 to 0.13   | 0.062                 |

Results from linear regression analysis on sex-stratified groups of APOE ε4 carriers revealed a significant main effect of FHS-CVD risk on tau deposition in the EC, ITC and meta-ROI only among female APOE ε4 carriers, after adjusting for the time between FHS-CVD risk assessment and tau-PET scan. Significant P-values are presented in bold, where *P < 0.05, **P < 0.01, ***P < 0.005. CI = confidence interval; EC = entorhinal cortex; FHS-CVD risk = Framingham Heart Study cardiovascular disease risk; ITC = inferior temporal cortex; SUVR = standardized uptake value ratio.
women’s resilience against cardiovascular disease up until mid-life and heightened vascular vulnerability after menopause may lead to greater vascular and Alzheimer’s disease comorbidity among women. Given that participants with a Hachinski ischaemic score $>4$ are excluded from the ADNI study, our sample was composed of participants in relatively good cardiovascular health. Larger studies in samples that are more representative of cardiovascular health in the aging population are needed to investigate potential sex-disparities further. ADNI also does not provide details on women’s health such as age at menopause and exposure to female-specific cardiovascular risk factors such as pre-eclampsia, which could be critical to the interpretation of our findings. Furthermore, since our sample was

Table 5 Three-way interaction effect of APOE ε4, sex and cardiovascular risk on tau deposition, after additionally adjusting for age, Aβ status and education

| Predictor variables | EC tau-PET SUVR | ITC tau-PET SUVR | Meta-ROI tau-PET SUVR |
|---------------------|-----------------|------------------|-----------------------|
|                     | Estimate | CI | P-value | Estimate | CI | P-value | Estimate | CI | P-value |
| FHS-CVD risk        | 0.01     | -0.01 to 0.02 | 0.239 | 0.00     | -0.01 to 0.01 | 0.793 | 0.00     | -0.00 to 0.01 | 0.652 |
| APOE ε4 carriers    | 0.19     | -0.24 to 0.61 | 0.390 | 0.01     | -0.28 to 0.30 | 0.945 | -0.03    | -0.27 to 0.20 | 0.773 |
| Sex (female)        | 0.25     | -0.11 to 0.61 | 0.166 | 0.11     | -0.14 to 0.35 | 0.392 | 0.05     | -0.15 to 0.24 | 0.641 |
| Age                 | 0.00     | -0.01 to 0.01 | 0.532 | 0.00     | -0.01 to 0.01 | 0.643 | 0.00     | -0.00 to 0.01 | 0.210 |
| Time lag            | 0.02     | -0.01 to 0.06 | 0.224 | -0.01    | -0.04 to 0.01 | 0.389 |-0.01    | -0.03 to 0.01 | 0.223 |
| Aβ status (Aβ+)     | 0.13     | 0.01–0.26 | **0.030** | 0.18     | 0.10–0.26 | < 0.001**** | 0.15     | 0.09–0.22 | < 0.001**** |
| Education           | 0.01     | -0.01 to 0.04 | 0.226 | 0.00     | -0.01 to 0.02 | 0.600 | 0.00     | -0.01 to 0.02 | 0.588 |
| APOE ε4 carriers ×  | -0.00    | -0.02 to 0.01 | 0.809 | 0.00     | -0.01 to 0.02 | 0.687 | 0.00     | -0.00 to 0.01 | 0.455 |
| FHS-CVD risk        | 0.00     | -0.01 to 0.02 | 0.681 | 0.00     | -0.01 to 0.01 | 0.911 | 0.00     | -0.01 to 0.01 | 0.534 |
| Sex (female) ×      | -0.50    | -1.06 to 0.07 | 0.084 | -0.23    | -0.61 to 0.16 | 0.242 | -0.11    | -0.42 to 0.20 | 0.475 |
| FHS-CVD risk        | 0.04     | 0.01–0.07 | **0.012** | 0.02     | -0.00 to 0.04 | 0.058 | 0.01     | -0.00 to 0.04 | 0.093 |

Results from linear regression analysis revealed a significant three-way interaction effect of APOE ε4, sex and FHS-CVD risk on tau deposition in the EC, but not the ITC or meta-ROI after adjusting the time between FHS-CVD risk assessment and tau-PET scan, age, Aβ status and education. Significant P-values are presented in bold, where *P < 0.05, **P < 0.01, ***P < 0.005, ****P < 0.001. Aβ = amyloid-β, CI = confidence interval; EC = entorhinal cortex; FHS-CVD risk = Framingham Heart Study cardiovascular disease risk; ITC = inferior temporal cortex; SUVR = standardized uptake value ratio.

Table 6 Two-way interaction effect of sex and cardiovascular disease risk on cortical tau deposition among APOE ε4 carriers but not non-carriers, after additionally adjusting for age, Aβ status and education

| Predictor variables | EC tau-PET SUVR | ITC tau-PET SUVR | Meta-ROI tau-PET SUVR |
|---------------------|-----------------|------------------|-----------------------|
|                     | Estimate | CI | P-value | Estimate | CI | P-value | Estimate | CI | P-value |
| APOE ε4 carriers    | 0.01     | -0.01 to 0.02 | 0.239 | 0.00     | -0.01 to 0.01 | 0.793 | 0.00     | -0.00 to 0.01 | 0.652 |
| FHS-CVD risk        | -0.28    | -0.81 to 0.25 | 0.294 | -0.10    | -0.42 to 0.23 | 0.541 | -0.05    | -0.32 to 0.22 | 0.714 |
| Sex (female)        | 0.05     | -0.03 to 0.13 | 0.200 | -0.02    | -0.06 to 0.03 | 0.490 | -0.02    | -0.06 to 0.02 | 0.402 |
| Age                 | -0.00    | -0.02 to 0.01 | 0.612 | 0.00     | -0.00 to 0.01 | 0.845 | 0.00     | -0.01 to 0.01 | 0.575 |
| Aβ status (Aβ+)     | 0.19     | -0.05 to 0.43 | 0.116 | 0.25     | 0.10–0.39 | 0.002**** | 0.21     | 0.08–0.33 | **0.001**** |
| Education           | 0.02     | -0.02 to 0.06 | 0.318 | -0.01    | -0.04 to 0.01 | 0.297 | -0.01    | -0.03 to 0.01 | 0.358 |
| Sex (female) ×      | 0.05     | 0.02–0.08 | **0.004**** | 0.02     | 0.00–0.04 | 0.049* | 0.02     | 0.00–0.03 | **0.045** |
| FHS-CVD risk        | 0.00     | -0.01 to 0.01 | 0.499 | 0.00     | -0.01 to 0.01 | 0.992 | 0.00     | -0.00 to 0.01 | 0.729 |
| APOE ε4 non-carriers| 0.18     | -0.16 to 0.52 | 0.304 | 0.15     | -0.10 to 0.39 | 0.242 | 0.07     | -0.12 to 0.27 | 0.452 |
| FHS-CVD risk        | 0.00     | -0.04 to 0.05 | 0.887 | -0.01    | -0.04 to 0.02 | 0.561 | -0.01    | -0.04 to 0.01 | 0.351 |
| Sex (female)        | 0.01     | -0.00 to 0.02 | 0.103 | 0.00     | -0.01 to 0.01 | 0.821 | 0.00     | -0.00 to 0.01 | 0.339 |
| Aβ status (Aβ+)     | 0.08     | -0.06 to 0.23 | 0.268 | 0.16     | 0.05–0.26 | 0.004**** | 0.14     | 0.05–0.22 | **0.002**** |
| Education           | 0.00     | -0.02 to 0.03 | 0.739 | 0.01     | -0.01 to 0.03 | 0.170 | 0.01     | -0.01 to 0.03 | 0.185 |
| Sex (female) ×      | 0.00     | -0.01 to 0.02 | 0.617 | 0.00     | -0.01 to 0.01 | 0.998 | 0.00     | -0.01 to 0.01 | 0.615 |

Results from linear regression analysis on APOE ε4 stratified groups revealed a significant interaction effect of sex and FHS-CVD risk on tau deposition in the EC, ITC and meta-ROI only among APOE ε4 carriers, after adjusting for the time between FHS-CVD risk assessment and tau-PET scan, age, Aβ status and education. Significant P-values are presented in bold, where *P < 0.05, **P < 0.01, ***P < 0.005, ****P < 0.001. Aβ = amyloid-β, CI = confidence interval; EC = entorhinal cortex; FHS-CVD risk = Framingham Heart Study cardiovascular disease risk; ITC = inferior temporal cortex; SUVR = standardized uptake value ratio.
predominantly composed of non-Hispanic white men and women, our results may not generalize to other racial and ethnic groups. Studies have shown that risk factors and cardiovascular disease manifestations can differ significantly between racial groups, highlighting the importance of studying racially and ethnically diverse samples. Another limitation of our study was its small sample size, which may have particularly affected the performance of tau deposition models in male and female APOE ε4 carriers (n = 23 and n = 24, respectively). Nevertheless, given the similar numbers of APOE ε4 carriers among men and women in the study, it is probably unlikely that the effect seen in women was driven by greater statistical power. Finally, the cross-sectional nature of the study along with the time lag between cardiovascular risk and tau-PET assessments limits our ability to draw conclusions about the temporality of associations between cardiovascular disease and tau deposition. Therefore, future longitudinal studies are needed to examine the extent to which sex and APOE ε4 impact the association of cardiovascular risk and accumulation of tau over the course of Alzheimer’s disease.

### Conclusion

Our results suggest that female APOE ε4 carriers may be particularly vulnerable to the impact of cardiovascular risk on cortical tau deposition, despite having lower cardiovascular risk scores than men. Our findings are clinically relevant, given the modifiable nature of many cardiovascular risk factors such as smoking and high blood pressure, and may contribute to our understanding of the observed heightened susceptibility to tauopathy among female APOE ε4 carriers. If replicated in larger, population-based samples, these findings could have important implications for the treatment of even low levels of vascular risk in seemingly healthy CN older women who have a genetic risk for Alzheimer’s disease. Future studies exploring associations between cardiovascular risk and Alzheimer’s disease-related outcomes should consider stratifying by sex and APOE ε4.

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| Predictor variables | EC tau-PET SUVR | ITC tau-PET SUVR | Meta-ROI tau-PET SUVR |
|---------------------|----------------|----------------|----------------------|
|                     | Estimate | CI    | P-value | Estimate | CI    | P-value | Estimate | CI    | P-value |
| **Female APOE ε4 carriers** | | | | | | | | |
| FHS-CVD risk        | 0.06    | 0.03−0.09 | **0.002*** | 0.02    | −0.00−0.04 | 0.052 | 0.02    | 0.00−0.03 | **0.028*** |
| Time lag            | 0.03    | −0.07−0.12 | 0.549 | −0.03    | −0.09−0.03 | 0.293 | −0.03    | −0.08−0.02 | 0.180 |
| Age                 | −0.02   | −0.06−0.02 | 0.279 | −0.00    | −0.03−0.02 | 0.811 | −0.00    | −0.02−0.02 | 0.734 |
| Aβ status (Aβ+)     | 0.27    | −0.10−0.63 | 0.138 | 0.33     | 0.11−0.56 | **0.005*** | 0.28     | 0.11−0.46 | **0.003*** |
| Education           | 0.03    | −0.04−0.11 | 0.364 | 0.01     | −0.04−0.05 | 0.809 | 0.00     | −0.04−0.04 | 0.952 |
| **Male APOE ε4 carriers** | | | | | | | | |
| FHS-CVD risk        | 0.00    | −0.01−0.02 | 0.675 | −0.00    | −0.01−0.00 | 0.434 | −0.00    | −0.01−0.00 | 0.568 |
| Time lag            | 0.11    | −0.08−0.30 | 0.227 | 0.04     | −0.06−0.15 | 0.409 | 0.04     | −0.05−0.13 | 0.389 |
| Age                 | −0.00   | −0.03−0.03 | 0.964 | 0.00     | −0.01−0.02 | 0.797 | 0.00     | −0.01−0.02 | 0.625 |
| Aβ status (Aβ+)     | 0.07    | −0.30−0.43 | 0.703 | 0.10     | −0.11−0.30 | 0.340 | 0.08     | −0.10−0.26 | 0.357 |
| Education           | 0.00    | −0.06−0.06 | 0.893 | −0.03    | −0.06−0.00 | 0.067 | −0.02    | −0.05−0.01 | 0.143 |

Results from linear regression analysis on sex-stratified groups of APOE ε4 carriers revealed a significant main effect of FHS-CVD risk on tau deposition in the EC and meta-ROI only among female APOE ε4 carriers, after adjusting for the time between FHS-CVD risk assessment and tau-PET scan, age, Aβ status and education. Significant P-values are presented in bold, where *P < 0.05, **P < 0.01, ***P < 0.005, ****P < 0.001. Aβ = amyloid-beta; CI = confidence interval; EC = entorhinal cortex; FHS-CVD risk = Framingham Heart Study cardiovascular disease risk; ITC = inferior temporal cortex; SUVR = standardized uptake value ratio.
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Competing interests
The authors report no competing interests.

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