Association of Early β-Amyloid Accumulation and Neuroinflammation Measured With $^{[11C]}$PBR28 in Elderly Individuals Without Dementia

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Abstract

Objective
To examine whether early β-amyloid (Aβ) accumulation and metabolic risk factors are associated with neuroinflammation in elderly individuals without dementia.

Methods
We examined 54 volunteers (mean age 70.0 years, 56% women, 51% APOE ε4 carriers) with the translocator protein (TSPO) tracer $^{[11C]}$PBR28 to assess neuroinflammation and with $^{[11C]}$ Pittsburgh compound B (PiB) to assess cerebral Aβ accumulation. $^{[11C]}$PBR28 and $^{[11C]}$PiB standardized uptake value ratios (SUVRs) were quantified in 6 regions of interests by using the cerebellar cortex as a pseudo-reference and reference region, respectively. Fasting venous glucose, insulin, and high-sensitivity C-reactive protein (hs-CRP) values were determined. Homeostatic model assessment of insulin resistance (HOMA-IR) was calculated. A subset of individuals (n = 11) underwent CSF sampling, and Aβ40, Aβ42, total tau, phospho-tau, soluble TREM2, and YKL-40 levels were measured.

Results
Among the whole study group, no significant association was found between $^{[11C]}$PiB and $^{[11C]}$PBR28 SUVR composite scores (slope 0.02, p = 0.30). However, higher $^{[11C]}$PiB binding was associated with higher $^{[11C]}$PBR28 binding among amyloid-negative ($^{[11C]}$PiB composite score ≤ 1.5) (TSPO genotype –, age- and sex-adjusted slope 0.26, p = 0.008) but not among amyloid-positive (slope −0.004, p = 0.88) participants. Higher CSF soluble TREM2 ($r_s = 0.72$, p = 0.01) and YKL-40 ($r_s = 0.63$, p = 0.04) concentrations were associated with a higher $^{[11C]}$PBR28 composite score. Higher body mass index, HOMA-IR, and hs-CRP were associated with higher $^{[11C]}$PBR28 binding in brain regions where Aβ accumulation is first detected in Alzheimer disease.

Conclusions
While there was no association between amyloid and neuroinflammation in the overall study group, neuroinflammation was associated with amyloid among the subgroup at early stages of amyloid pathology.
Neuroinflammation, that is, activation of microglia and astrocytes, seems to play an important role in the pathogenesis of Alzheimer disease (AD) and other diseases leading to dementia. Metabolic risk factors, including obesity and insulin resistance, have been linked to cognitive decline and dementia. However, little is known about the possible relationship among chronic low-grade inflammation, metabolic risk factors, neuroinflammation, and the neuropathology of AD.

Activated microglia overexpress 18-kDa translocator protein (TSPO), and thus PET imaging with tracers binding to TSPO is used to assess neuroinflammation. TREM2 is an innate immune receptor that is selectively expressed by microglia in the CNS. Its soluble variant (sTREM2) and YKL-40, a glycoprotein expressed by reactive astrocytes and microglia in the CNS, can be measured from the CSF. Studies have suggested that both of these markers of neuroinflammation are increased in the early stage of AD.

On the basis of previous literature, we hypothesized that increased TSPO expression would associate with amyloid-β (Aβ) accumulation in the cerebral cortex and that both TSPO expression and Aβ accumulation would associate with CSF biomarkers of neuroinflammation and AD. Previous reports have shown that midlife metabolic risk factors predict brain Aβ accumulation and dementia. Therefore, we hypothesized that insulin resistance, obesity, and chronic low-grade inflammation measured by high-sensitivity C-reactive protein (hs-CRP) would associate with increased neuroinflammation.

To test these hypotheses, we examined 54 elderly volunteers without dementia with brain PET imaging using a TSPO radioligand [11C]PBR28 and Aβ radioligand [11C] Pittsburgh compound B (PiB). In addition, we performed CSF analyses on a subset of the study population.

Methods

The study participants took part in a neuroimaging study carried out at the Turku PET Centre in 2014 to 2016. The study was conducted to investigate the associations among midlife insulin resistance, inflammation, and late-life brain Aβ accumulation, neuroinflammation, cerebrovascular lesions, and cognition. The results concerning Aβ accumulation, neuroinflammation, cerebrovascular lesions, and cognition have been published previously. The study participants were recruited from a nationwide Finnish health examination survey, the Health 2000 survey, according to their insulin sensitivity and APOE genotype in 2000. The recruitment process has been described in detail previously.

Health 2000 Survey

The Health 2000 survey is a Finnish nationwide, population-based study carried out in 2000 to 2001 and conducted by the Finnish Institute for Health and Welfare. Eight thousand twenty-eight individuals >30 years of age were randomly selected from the Finnish population register. Six thousand three hundred fifty-four volunteers (79% participation rate, mean age 55.4 years) attended the health examination proper. A thorough physical examination was performed; body mass index (BMI) was determined; and venous blood samples, including fasting insulin and glucose, and hs-CRP were drawn. APOE genotype was defined with the MassARRAY System (Sequenom, San Diego, CA) with a modified protocol. Homeostatic model assessment of insulin resistance (HOMA-IR) was calculated with the equation fasting insulin (μU/mL) times fasting glucose (mmol/L) divided by 22.5.

Study Participants and Recruitment Process in 2014

In 2014, we recruited 60 participants from those who had attended the Health 2000 survey to attend further examination at the Turku PET Center according to their HOMA-IR values in 2000 and their APOE genotype. All participants were community-dwelling individuals who did not require help with their activities of daily living. We recruited 30 participants who had insulin resistance in 2000 (IR+ group, HOMA-IR in the highest tertile [>2.17] of the Health 2000 study population) and 30 controls with normal insulin sensitivity (IR− group, HOMA-IR in the lowest tertile [<1.25] of the Health 2000 study population). Both groups were enriched for APOE E4 carriers to obtain 15 APOE E4 carriers in both groups. On the basis of a telephone interview, individuals with a history of a major stroke, a previous dementia diagnosis, any other major neurologic disorder, type 2 diabetes in 2000, and type 2 diabetes after the year 2000 for the insulin-sensitive group were excluded. Additional exclusion criteria were any contraindication for a PET or MRI scan.

The participants’ genotypes for TSPO binding were determined before [11C]PBR28 imaging because of the large interindividual variability in the binding affinity of [11C] PBR28 due to the rs6971 polymorphism of the TSPO gene. TSPO genotyping was performed in London, UK (Imperial Molecular Pathology Laboratory, Hammersmith Hospital).

Glossary

Aβ = amyloid-β; AD = Alzheimer disease; BMI = body mass index; DVR = distribution volume ratio; HAB = high-affinity binder; HOMA-IR = homeostatic model assessment of insulin resistance; hs-CRP = high-sensitivity C-reactive protein; MAB = mixed-affinity binder; MCI = mild cognitive impairment; P-tau = phosphorylated tau; PiB = Pittsburgh compound B; ROI = region of interest; sTREM2 = soluble TREM2; SUV = standardized uptake value; SUVR = SUV ratio; T-tau = total tau; TSPO = translocator protein.
DNA was extracted from peripheral blood with the Qiagen (Venlo, the Netherlands) QIAamp DNA blood mini kit. TSPO genotyping was performed with a TaqMan Allelic Discrimination assay. Among the 60 participants, there were 5 low-affinity binders. They all were excluded from the present study, leaving 55 eligible participants. In addition, 1 individual was excluded because of not being able to finish the 70 minutes of $^{[11]}$C]PBR28 scanning.

**Standard Protocol Approvals, Registrations, and Patient Consents**

The Finnish Health 2000 study was approved by the Ethics Committee for Epidemiology and Public Health in the hospital district of Helsinki and Uusimaa, Finland. The follow-up study was approved by the Ethics Committee of the Hospital District of Southwest Finland. Written informed consent was obtained from all of the participants. Separate written consent for CSF sampling was obtained from all participants who agreed to undergo a lumbar puncture. All study procedures were conducted according to Good Clinical Practice guidelines and the Declaration of Helsinki.

**Laboratory Examination at Time of PET Scans in 2014 to 2016**

The laboratory examinations were performed within 4 months of the $^{[11]}$C]PBR28 scan. Venous blood samples were drawn after an overnight fasting (minimum 10 hours) in 2014 to 2016. Serum insulin and glucose were determined as previously described. HOMA-IR was calculated. hs-CRP was determined by immunonephelometry with a BN ProSpec System (Siemens Healthcare GmbH, Marburg, Germany).

A subset of 11 participants gave permission to obtain CSF samples. The samples were centrifuged within 30 minutes and stored at -70°C. The samples were sent to the University of Gothenburg for further analysis. CSF YKL-40 concentration was measured with an YKL-40 ELISA kit (R&D Systems, Minneapolis, MN). CSF $\beta_40$ and $\beta_{12}$ concentrations were measured with electrochemiluminescence technology and the MS6000 Human Abeta 3-Plex Ultra-Sensitive Kit (Meso Scale Discovery, Rockville, MD). CSF total tau (T-tau) and phosphorylated tau (P-tau) concentrations were measured with commercially available INNOTEST sandwich ELISAs (Fujirebio Europe, Ghent, Belgium). CSF sTREM2 concentration was measured with an in-house ELISA as previously described.

All CSF biomarker measurements were performed with 1 batch of reagents in 1 round of experiments by board-certified laboratory technicians who were blinded to clinical data. Intra-assay coefficients of variation of quality control samples were <10%.

**MRI and PET Scanning With TSPO Radioligand $^{[11]}$C]PBR28 and $^{[11]}$C]PiB**

To obtain anatomic reference and to be able to exclude potential structural abnormalities, the study volunteers underwent MRI with a 3T PET-MRI scanner (Philips Ingenuity TF PET-MR device, Philips Healthcare, Best, the Netherlands).

$^{[11]}$C]PBR28 PET imaging was performed to assess microglial activation, and $^{[11]}$C]PiB PET imaging was done to assess Aβ accumulation in the brain. Both the dynamic 70-minute $^{[11]}$C]PBR28 PET and the dynamic 90-minute $^{[11]}$C]PiB PET scans were performed with a brain-dedicated high-resolution PET scanner, the ECAT HRRT (Siemens Medical Solutions, Knoxville, TN). An individually shaped thermoplastic mask was used to minimize head movement.

$^{[11]}$C]PBR28 was manufactured as previously reported. An anesthesiologist performed arterial catheterization before imaging for those who did not have any contraindication (anticoagulant therapy) for arterial cannulation (n = 44). The participants received the $^{[11]}$C]PBR28 as a rapid intravenous bolus (mean dose 496 MBq, SD 19 MBq). Arterial blood samples measuring radioactivity and metabolites were collected. The additional details of $^{[11]}$C]PBR28 PET data acquisition, reconstruction, arterial blood sampling, and radiometabolite analysis have been presented previously. The $^{[11]}$C]PiB PET scanning protocol was described earlier. PET images were realigned and coregistered to individual T1-weighted MRIs with SPM12 software (Wellcome Trust Centre for Neuroimaging, London, UK) running in MATLAB (MathWorks, Natick, MA). Automated region of interest (ROI) generation was conducted for both $^{[11]}$C]PBR28 and $^{[11]}$C]PiB data with FreeSurfer software (version 5.3.0, freesurfer.net/) and T1-weighted MRI data as input. Six ROIs were formed (parietal cortex, prefrontal cortex, anterior cingulum, posterior cingulum, precuneus, lateral temporal cortex) based on regions where Aβ accumulation typically begins in AD. 

For the analysis of regional and voxel-by-voxel $^{[11]}$C]PBR28 (n = 54) data, we used a standardized uptake value (SUV) ratio (SUVR) from 30 to 70 minutes with respect to cerebellum pseudoreference region, which has been validated previously. In addition, distribution volume (VT) was calculated for those with arterial blood data (n = 44) using Logan analysis within a 30- to 70-minute interval, after which distribution volume ratios (DVRs) were calculated by dividing the target ROI VT by the cerebellar cortex VT. We used a 2-sample t test to ensure that the SUVs and VT values of the cerebellar cortex did not differ between the amyloid-negative and amyloid-positive groups. Regional and voxel-by-voxel $^{[11]}$C]PiB SUVs were calculated using imaging data from 60 to 90 minutes after the tracer injection. SUVRs were obtained with the cerebellar cortex used as a reference region.
Composite cortical $^{[11C]}$PBR28 and $^{[11C]}$PiB SUVR scores were calculated as a volume-weighted average $^{[11C]}$PBR28 SUVR and $^{[11C]}$PiB SUVR over all 6 ROIs. The $^{[11C]}$PiB-PET scan was considered amyloid positive when the $^{[11C]}$PiB composite SUVR score was >1.5 according to previous studies on cognitively healthy elderly populations.\(^{28,29}\)

### Statistical Analyses

The associations between explanatory variables ($^{[11C]}$PiB SUVRs/BMI/HOMA-IR/hs-CRP) and $^{[11C]}$PBR28 SUVR composite score, $^{[11C]}$PBR28 SUVRs of the 6 cortical ROIs, and $^{[11C]}$PBR28 DVRs of the same ROIs (n = 44) were studied with multivariable linear models. The explanatory variables were adjusted for age, sex, and TSPO genotype because previous studies have shown that they can affect $^{[11C]}$PBR28 binding\(^{23,25,30}\) and that SUVR levels might be paradoxically lower in high-affinity binders (HABs) than mixed-affinity binders (MABs) for TSPO genotype.\(^{27}\) To achieve a normal distribution, a logarithmic transformation was performed for HOMA-IR and hs-CRP. Normality assumption for the analyses was checked from the residuals. Bonferroni correction for multiple comparisons was performed for the analyses of individuals ROIs (uncorrected \(p\) values were multiplied by the number of ROIs studied, i.e., 6).

The association between the CSF proteins and $^{[11C]}$PiB and $^{[11C]}$PBR28 SUVR composite scores was assessed with nonparametric Spearman correlation because the parameters were not normally distributed.

Two-sided statistical significance was set at \(p < 0.05\). The statistical analyses were performed with SAS JMP Pro 14 (SAS Institute, Cary, NC). Voxel-by-voxel associations between $^{[11C]}$PBR28 SUVRs and $^{[11C]}$PiB SUVRs were assessed with SPM12 by using normalized SUVR images in Montreal Neurological Institute space. The images were smoothed with 3-dimensional gaussian 8-mm full width at half-maximum filter to ensure normality before the statistical analysis.

### Data Availability

Anonymized data are available on reasonable request for a study that has been approved by a local ethics committee.

### Results

#### Demographics

The characteristics of the study population according to amyloid status are shown in table 1. The mean age of the
participants was 70.0 years (range 65–80 years). The average BMI was 26.6 kg/m² (range 19.7–38.8 kg/m²). There were 25 MABs and 29 HABs. Twenty-five participants (46%) were amyloid positive. There were no differences between amyloid-negative and amyloid-positive participants in age, sex, BMI, fasting glucose, hs-CRP, or total cholesterol. Amyloid-positive individuals were more often APOE ε4 carriers. Amyloid-positive participants tended to have a lower Consortium to Establish a Registry for Alzheimer’s Disease total score, as expected, but this difference was not statistically significant (p = 0.06). One individual (amyloid positive, MAB) had exceptionally high [11C]PBR28 SUVR values in all ROIs ([11C]PBR28 SUVR composite score 1.40) and an exceptionally high BMI (38.8 kg/m²) and HOMA-IR value (21.4) among the study population. This outlier was excluded from all analyses.

CSF samples were obtained from 11 individuals (mean age 68.1 years), of which 6 were women. There were 5 MABs and 6 HABs. Five individuals (46%) were APOE ε4 carriers. The mean BMI was 26.1 kg/m². Only 3 of the participants who underwent CSF sampling were classified as amyloid positive (table 2).


Among all participants (amyloid negative and amyloid positive), no significant association was found between [11C]PiB and [11C]PBR28 SUVR composite scores (slope 0.02, p = 0.30) or between [11C]PiB SUVRs and [11C]PBR28 SUVRs in any of the ROIs (all p ≥ 0.21) (data not shown).

To evaluate whether the association between [11C]PiB and [11C]PBR28 SUVR composite scores would be modulated by amyloid status based on [11C]PiB PET scans, we analyzed the interaction of [11C]PiB composite score × amyloid positivity for predicting [11C]PBR28 SUVR composite score. TSPO genotype was added into the analysis as an explanatory variable. The interaction for [11C]PiB composite score × amyloid positivity was statistically significant (p = 0.02); thus, the analyses were stratified according to amyloid positivity.

A higher [11C]PiB SUVR composite score was associated with a higher [11C]PBR28 SUVR composite score among amyloid-negative participants (slope 0.26, p = 0.008). Among amyloid-positive participants, no association between [11C]PiB and [11C]PBR28 SUVR composite scores was found (slope −0.004, p = 0.88) (figure 1).


To evaluate the association between [11C]PiB and [11C]PBR28 without predefined assumptions of specific ROIs, we performed voxel-level analysis on 53 participants (outlier excluded). TSPO genotype was added as an explanatory variable in the analyses. Among amyloid-negative participants, [11C]PiB binding showed a positive association with [11C]PBR28 binding, especially in the parietal cortex, but also in the frontal cortex, in the precuneus, and in the posterior cingulum, reflecting cortical regions of early Aβ accumulation in AD (figure 1, false discovery rate–corrected p < 0.05). A positive association was found between [11C]PiB and [11C]PBR28 in the false discovery rate–corrected voxel-level analysis in the entire study population as well (data not shown). However, the association was explained by the amyloid-negative subgroup; no association was found between [11C]PiB and [11C]PBR28 for the amyloid-positive participants.
CSF sTREM2 and YKL-40 Associated With $[^{11}C]$PBR28, and CSF $A\beta_{42}$ and $A\beta_{42}/A\beta_{40}$ Ratio Associated With $[^{11}C]$PiB

We used the CSF results to support our results from the PET scans by measuring proteins related to AD and inflammation and evaluating the correlation between CSF proteins and $[^{11}C]$PiB and $[^{11}C]$PBR28 binding. The results of the CSF analysis are presented in table 2. All the participants had T-tau and P-tau levels within the normal range.

A higher CSF sTREM2 value and a higher CSF YKL-40 value was associated with a higher $[^{11}C]$PBR28 SUVR composite score (figure 2 and table 3). A lower CSF $A\beta_{42}$ level and a lower CSF $A\beta_{42}/A\beta_{40}$ ratio, but not $A\beta_{40}$, were associated with a higher $[^{11}C]$PiB SUVR composite score (table 3).

Metabolic Risk Factors Measured at Time of PET Scans in 2014 to 2016 Associated With $[^{11}C]$PBR28

A higher BMI was associated with a higher $[^{11}C]$PBR28 SUVR composite score ($p = 0.006$). A positive association between BMI and $[^{11}C]$PBR28 SUVR was found in the parietal cortex (Bonferroni corrected $p < 0.0006$) and in the precuneus (Bonferroni-corrected $p = 0.002$). In other ROIs, no significant association between BMI and $[^{11}C]$PBR28 SUVRs was found (all uncorrected $p \geq 0.10$) (table 4 and figure 3).

HOMA-IR or hs-CRP did not associate with $[^{11}C]$PBR28 composite score (HOMA-IR $p = 0.21$, hs-CRP $p = 0.29$). The associations between HOMA-IR and hs-CRP and $[^{11}C]$PBR28 SUVRs are shown in table 4. The association between
Additional analyses were performed with [11C]PBR28 DVRs (Bonferroni-corrected $p$ value) in the precuneus ($r = 0.72$, $p = 0.01$) and in the parietal cortex ($r = 0.63$, $p = 0.04$). Significant associations were found between BMI and [11C]PBR28 DVR in the parietal cortex ($p < 0.0006$) and the precuneus ($p = 0.049$) and between hs-CRP and [11C]PBR28 DVR in the parietal cortex ($p = 0.03$) after Bonferroni correction (data not shown).

**A Higher BMI in Midlife Predicted Microglial Activation Measured by [11C]PBR28 PET Imaging 15 Years Later**

To evaluate whether midlife risk factors would associate with late-life neuroinflammation, we analyzed the associations between metabolic variables measured in 2000 and [11C]PBR28 SUVRs measured in 2014 to 2016. A higher BMI in midlife associated with a higher [11C]PBR28 SUVR in the parietal cortex (uncorrected $p = 0.002$) and precuneus (uncorrected $p = 0.03$). Only the association between BMI and [11C]PBR28 SUVR in parietal cortex was significant after Bonferroni correction (Bonferroni-corrected $p = 0.01$). Neither baseline HOMA-IR nor hs-CRP associated with [11C]PBR28 SUVRs in any of the ROIs (all uncorrected $p \geq 0.07$). As expected, there was a strong correlation between midlife and late-life BMI ($r = 0.77$, $p < 0.0001$) (data not shown).

**Discussion**

Contrary to our original hypothesis, no association was seen between amyloid and neuroinflammation among the whole study group. However, we found that neuroinflammation might be associated with the early stages of Aβ accumulation in elderly individuals without dementia. The association between [11C]PiB and [11C]PBR28 SUVR composite scores was seen among amyloid-negative participants only. In addition, higher levels of metabolic risk factors (BMI, HOMA-IR, and hs-CRP) were positively associated with neuroinflammation measured with [11C]PBR28 PET in brain regions where Aβ accumulation is first detected in AD. The results are supported by the finding that CSF sTREM2 and YKL-40, 2 previously established markers for neuroinflammation, were positively associated with our main outcome, the [11C]PBR28 SUVR composite score.

To date, few studies have used second-generation TSPO ligands to assess the relationship between neuroinflammation and AD. Most of the previous studies evaluating the role of neuroinflammation in the AD pathologic process have been performed with a traditional TSPO ligand, [11C](R)PK11195. It has been suggested that [11C](R)PK11195 has limited sensitivity and specificity,$^{31}$ whereas second-generation TSPO ligands might have higher specific TSPO binding.$^{32}$ However, no head-to-head study has compared [11C]PBR28 and [11C](R)PK11195 PET in AD, and thus, there is no convincing evidence suggesting that one ligand would be more sensitive than the other.$^{3}$ A recent meta-analysis of 28 PET studies (of which 4 had been performed with [11C]PBR28) indicated that patients with AD have increased TSPO levels compared to healthy controls especially within frontotemporal regions.$^{33}$ According to one of the studies included in the meta-analysis, [11C]PBR28 binding was higher in cortical brain regions among patients with AD compared to controls, but there was no difference between patients with mild cognitive impairment (MCI) and controls.$^{34}$ However, another study showed higher [11C]PBR28 uptake in amyloid-positive patients with MCI compared to healthy controls.$^{35}$ In a longitudinal study (mean follow-up time 2.7 years), patients with AD had a greater increase in TSPO binding in temporoparietal regions than controls.$^{36}$ Studies focusing on neuroinflammation in elderly individuals at risk for AD are scarce.

### Table 3

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<tr>
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<tr>
<td></td>
<td>$r_s$ Value $p$ Value</td>
<td>$r_s$ Value $p$ Value</td>
</tr>
<tr>
<td>YKL-40</td>
<td>0.63 0.04</td>
<td>−0.31 0.36</td>
</tr>
<tr>
<td>sTREM2</td>
<td>0.72 0.01</td>
<td>−0.06 0.85</td>
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<tr>
<td>Aβ40</td>
<td>0.69 0.02</td>
<td>−0.27 0.42</td>
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<tr>
<td>Aβ42</td>
<td>0.12 0.73</td>
<td>−0.85 0.001</td>
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<tr>
<td>Aβ42/Aβ40 Ratio</td>
<td>−0.26 0.43</td>
<td>−0.87 0.0005</td>
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<tr>
<td>Total tau</td>
<td>0.47 0.14</td>
<td>−0.21 0.54</td>
</tr>
<tr>
<td>Phosphorylated tau</td>
<td>0.59 0.06</td>
<td>−0.34 0.31</td>
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</table>

Abbreviation: Aβ = β-amyloid; PiB = Pittsburgh compound B; sTREM2 = soluble TREM2; SUVR = standardized uptake value ratio.
Neuroinflammation is assumed to be both beneficial and detrimental, depending on the stage of AD pathogenesis.7 Presumably, the initial microglial activation would be protective, but it would become unfavorable with disease progression.37,38 A [11C]PBR28 study demonstrated that there could be 2 microglial activation peaks in the AD pathogenesis: an early protective peak and a later proinflammatory peak.13 In line with previous TSPO-PET studies,2,7 our results suggest that neuroinflammation occurs at early stages of the AD pathogenesis. A previous [11C]PBR28 study found a correlation between Aβ load measured with [18F]flutemetamol and [11C]PBR28 binding.12 The correlation was stronger among patients with MCI than patients with AD.12 A TSPO-PET study with [18F]DPA-714 found an increase of TSPO binding in patients with AD, especially at the prodromal stage, compared to controls.38

In addition, they found higher TSPO binding in slow decliners than in fast decliners, with no difference in Aβ accumulation, indicating that microglial activation might have a protective role.38 However, in a longitudinal study, the same study group showed that the subsequent increase in [18F]DPA-714 binding among patients with AD was associated with disease worsening.39 In the present study, we found an association between Aβ accumulation and [11C]PBR28 among amyloid-negative participants but not among amyloid-positive participants, suggesting that neuroinflammation might be associated with early stages of Aβ accumulation.

Contrary to a previous study that found higher TSPO binding (measured with 18F-DPA-714) among amyloid-positive controls compared to amyloid-negative controls,38 we did

Table 4 Association Between Metabolic Variables Measured in 2014 to 2016 at Time of PET Scans and [11C]PBR28 SUVRs

<table>
<thead>
<tr>
<th>ROI</th>
<th>Prefrontal cortex</th>
<th>Parietal cortex</th>
<th>Lateral temporal cortex</th>
<th>Precuneus</th>
<th>Anterior cingulum</th>
<th>Posterior cingulum</th>
<th>Composite score</th>
</tr>
</thead>
<tbody>
<tr>
<td>BMI</td>
<td>0.0035 (0.00067 to 0.0077)</td>
<td>0.011±0.0034 (0.0066 to 0.014)</td>
<td>0.009±0.0037 (0.0044 to 0.014)</td>
<td>-0.0009 (0.0063 to 0.0044)</td>
<td>-0.0033 (0.0013 to 0.0079)</td>
<td>0.0048±0.0004 (0.00152 to 0.0082)</td>
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</tr>
<tr>
<td>HOMA-IR</td>
<td>0.002 (0.002 to 0.024)</td>
<td>0.040±0.0043 (0.017 to 0.062)</td>
<td>0.030±0.0058 (0.002 to 0.013)</td>
<td>-0.035±0.0090 (0.006 to 0.020)</td>
<td>-0.0043 (0.0028 to 0.030)</td>
<td>0.011 (0.0068 to 0.030)</td>
<td></td>
</tr>
<tr>
<td>hs-CRP</td>
<td>0.00022 (0.0015 to 0.016)</td>
<td>0.022±0.0015 (0.0054 to 0.039)</td>
<td>0.009±0.011 (0.014)</td>
<td>-0.005±0.011 (0.003)</td>
<td>-0.0045 (0.022 to 0.012)</td>
<td>0.0069 (0.0061 to 0.020)</td>
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</table>

Abbreviations: BMI = body mass index; HOMA-IR = homeostatic model assessment of insulin resistance; hs-CRP = high-sensitivity C-reactive protein; ROI = region of interest; SUVR = standardized uptake value ratio; TSPO = translocator protein. Results are shown as slope (95% confidence interval). Associations between BMI and [11C]PBR28 SUVR, HOMA-IR, and [11C]PBR28 SUVR and hs-CRP and [11C]PBR28 are analyzed in separate linear models. [11C]PBR28 SUVR composite score is calculated as the average [11C]PBR28 SUVR over all 6 ROIs. A logarithmic transformation is used for homeostatic model assessment of insulin resistance (HOMA-IR) and high-sensitivity C-reactive protein (hs-CRP). [11C]PBR28 binding is reported as standardized uptake value ratio. High-affinity binders are marked with red dots; mixed-affinity binders are marked with blue dots.
not find any group difference in microglial activation between amyloid-negative and amyloid-positive participants. However, the control groups in the previous study (6 amyloid-positive and 20 amyloid-negative controls) were smaller than our study groups, and the quantification was performed without arterial blood samples.

We found that higher CSF sTREM2 and YKL-40 levels were associated with higher $[^{11}C]$PBR28 binding in the cerebral cortex. According to a meta-analysis, CSF sTREM2 and YKL-40 levels are higher in both patients with MCI and patients with AD compared to healthy controls. Another meta-analysis showed that CSF YKL-40 is associated with clinically diagnosed AD, but the relationship seems to be notably weaker than that of the core biomarkers $\beta$-amyloid and $\tau$-tubulin. This is supported by epidemiologic studies showing that chronic low-grade inflammation, obesity, and insulin resistance are associated with cognitive decline. It has been suggested that environmentally modifiable risk factors of cognitive decline such as chronic inflammation and obesity would affect the risk through a sustained neuroinflammatory drive. Epidemiologic studies have suggested a link between nonsteroidal anti-inflammatory drugs and reduced risk of clinically diagnosed AD. However, most of the randomized controlled trials assessing the use of nonsteroidal anti-inflammatory drugs in AD treatment have yielded negative results; thus, they are not recommended for the prevention of AD. We have previously shown that midlife insulin resistance predicts $\beta$-amyloid accumulation in the brain, and midlife obesity has been shown to independently predict late-life brain $\beta$-amyloid accumulation. In the present study, no association was found between midlife insulin resistance or low-grade inflammation and neuroinflammation, possibly reflecting that neuroinflammation is a dynamic process whereas brain $\beta$-amyloid accumulation presumably begins decades before the onset of the first symptoms of AD. Our study suggests that the association between metabolic risk factors and neuroinflammation is seen in brain regions where $\beta$-amyloid is first accumulated in AD. However, our results concerning metabolic risk factors are to be considered preliminary and warrant further investigation. Contrary to our findings regarding the association between BMI and $[^{11}C]$PBR28 binding, a multicenter study on healthy controls from our center and 2 other centers found a negative correlation between BMI and $[^{11}C]$PBR28 $V_T$ values in gray matter, thalamus, and hippocampus by using arterial plasma curves as an input function in kinetic modeling. Most likely, these contrasting results are emerging because, in the present study, a cerebellar gray matter pseudoreference region was used as a model input instead of plasma input function. The previous study focused on $[^{11}C]$PBR28 binding in healthy controls across different age groups, while the present study focused on neuroinflammation that would be related to AD pathology. Because AD pathologic changes are rarely seen in the cerebellar cortex, we used the cerebellar cortex as a pseudoreference region in the analyses, consistent with previous reports on AD.

Our study had limitations. TSPO quantification by PET is challenging partly because of the large interindividual variability in the binding affinity of $[^{11}C]$PBR28 due to the rs6971 polymorphism. In addition, there is no true reference region in the brain for $[^{11}C]$PBR28 imaging. TSPO expression is seen throughout the brain and blood vessels; there is no brain region totally devoid of specific TSPO binding sites. Traditionally, analysis of $[^{11}C]$PBR28 PET binding requires arterial blood sampling. However, a study group has previously validated a SUVR method in $[^{11}C]$PBR28-PET imaging in patients with AD. The cerebellum is used as a pseudoreference region in the method on the basis of the findings that the cerebellar cortex is typically devoid of neuropathologic changes typical for AD, including inflammation. We did not perform arterial blood sampling data on all participants, which can be considered a limitation. However, we performed our analyses both with the SUVR method (n = 54) using cerebellum as a pseudoreference region and with the DVR method using the arterial data (n = 44). Both analysis methods yielded similar results, confirming previous reports that SUVR can be used to detect cortical $[^{11}C]$PBR28 binding to detect AD-related neuroinflammation. We did not perform partial volume corrections, but the partial volume effect was minimized by the use of a high-resolution PET scanner. Moreover, the participants in our study did not have dementia or significant brain atrophy. Nevertheless, lack of partial volume correction could be regarded as a limitation. According to a recent postmortem study, TSPO is expressed not only in microglia but also in astrocytes, endothelial cells, and vascular smooth muscle cells. Thus, PBR28 binding can be influenced by other cell types than microglia. The main advantage of this study is the possibility of combining $[^{11}C]$PBR28 and $[^{11}C]$PiB PET results in elderly individuals without dementia. A strength is that we used the CSF results to support our results from the PET scans, even if the CSF samples were obtained from 11 participants only, which limits firm conclusions based on the CSF results. Additional benefits are the arterial blood sampling in $[^{11}C]$PBR28 PET imaging from the majority of the study.
population and the long follow-up time (15 years) in terms of the metabolic risk factors.

While neuroinflammation was not associated with amyloid among all participants, this study suggests that neuroinflammation may be associated with the early stages of brain Aβ pathology. We also demonstrate that metabolic risk factors (higher BMI, insulin resistance, and chronic low-grade inflammation) are associated with neuroinflammation in brain regions where Aβ accumulation is first detected in AD. We propose that the association between metabolic risk factors and clinically diagnosed AD seen in epidemiologic studies could be mediated at least partly through neuroinflammation. It should be noted, however, that the early stages of Aβ accumulation in elderly individuals with normal cognition might not lead to symptomatic AD. Larger studies with longitudinal protocols are needed to understand the complex relationship between metabolic risk factors, neuroinflammation, and AD and particularly to detect the time course of neuroinflammation in the pathogenesis of AD. Our ongoing follow-up study on the same study population will provide insights into the longitudinal associations among neuroinflammation, amyloid accumulation, and cognitive performance. Future studies combining CSF analyses with TSPO imaging and amyloid and tau-PET imaging could provide new targets to develop novel therapies for AD.

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