Neuroinflammatory Component of Gray Matter Pathology in Multiple Sclerosis

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Objective: In multiple sclerosis (MS), using simultaneous magnetic resonance–positron emission tomography (MR-PET) imaging with $^{11}$C-PBR28, we quantified expression of the 18kDa translocator protein (TSPO), a marker of activated microglia/macrophages, in cortex, cortical lesions, deep gray matter (GM), white matter (WM) lesions, and normal-appearing WM (NAWM) to investigate the in vivo pathological and clinical relevance of neuroinflammation.

Methods: Fifteen secondary-progressive MS (SPMS) patients, 12 relapsing–remitting MS (RRMS) patients, and 14 matched healthy controls underwent $^{11}$C-PBR28 MR-PET. MS subjects underwent 7T T2-weighted imaging for cortical lesion segmentation, and neurological and cognitive evaluation. $^{11}$C-PBR28 binding was measured using normalized 60- to 90-minute standardized uptake values and volume of distribution ratios.

Results: Relative to controls, MS subjects exhibited abnormally high $^{11}$C-PBR28 binding across the brain, the greatest increases being in cortex and cortical lesions, thalamus, hippocampus, and NAWM. MS WM lesions showed relatively modest TSPO increases. With the exception of cortical lesions, where TSPO expression was similar, $^{11}$C-PBR28 uptake across the brain was greater in SPMS than in RRMS. In MS, increased $^{11}$C-PBR28 binding in cortex, deep GM, and NAWM correlated with neurological disability and impaired cognitive performance; cortical thinning correlated with increased thalamic TSPO levels.

Interpretation: In MS, neuroinflammation is present in the cortex, cortical lesions, deep GM, and NAWM, is closely linked to poor clinical outcome, and is at least partly linked to neurodegeneration. Distinct inflammatory-mediated factors may underlie accumulation of cortical and WM lesions. Quantification of TSPO levels in MS could prove to be a sensitive tool for evaluating in vivo the inflammatory component of GM pathology, particularly in cortical lesions.
immunopathological characteristics of cortical demyelination differ significantly from those in white matter (WM), based on the evidence that cortical lesions lacked the inflammatory pattern typical of WM plaques.\textsuperscript{2,3} Subsequent pathological examinations, however, suggested that cortical MS lesions might develop following meningeal inflammation accompanied by cortical microglia activation.\textsuperscript{4,5} The role of activated microglia in cortical MS lesion pathogenesis is not without controversies, and is not uniformly observed.\textsuperscript{6} Nevertheless, postmortem studies demonstrate that severe cortical microglia activation and demyelination are associated with a less favorable disease outcome.\textsuperscript{4–6}

Activation of microglia/macrophages, associated with demyelination and neurodegeneration, has been pathologically observed in deep GM\textsuperscript{7} and hippocampus in MS,\textsuperscript{8,9} although their exact pathophysiologic role and clinical implications remain unclear.

The in vivo study of microglia/macrophage activation in the cortex, particularly cortical lesions, and deep GM could elucidate the role of neuroinflammation in the pathogenesis of demyelination and neurodegeneration of these structures, as they usually lack the blood–brain barrier abnormalities commonly detected in WM lesions using gadolinium magnetic resonance imaging (MRI).\textsuperscript{10,11} Additionally, activated microglia/macrophages can be present even in the absence of blood–brain barrier disruption seen on contrast-enhanced MRI.\textsuperscript{12}

Activated microglia/macrophages upregulate expression of the 18kDa translocator protein (TSPO), which can be imaged in vivo by selective positron emission tomography (PET) TSPO radioligands, of which \textsuperscript{11}C-PK11195 is the best known.\textsuperscript{13} In MS, \textsuperscript{11}C-PK11195 studies have reported increased TSPO levels in the cortex,\textsuperscript{14} thalamus,\textsuperscript{15} and central deep GM;\textsuperscript{16} however, knowledge of microglia/macrophage activation in MS cortical lesions is still lacking due to the low sensitivity of clinical imaging tools to detect cortical demyelination. Seven-tesla (7T) $T_2^*$-weighted gradient-echo protocols demonstrate increased sensitivity to in vivo visualization of focal cortical lesions in MS, confirmed pathologically to correspond to cortical areas with the greatest degree of demyelination.\textsuperscript{17,18}

\textsuperscript{11}C-PBR28 is a second-generation TSPO PET tracer\textsuperscript{19} with \textasciitilde 80 times more specific binding than \textsuperscript{11}C-PK11195\textsuperscript{20} and good reproducibility.\textsuperscript{21} In experimental studies, inflammation-induced microglial activation determined \textsuperscript{11}C-PBR28 signal increase, confirmed pathologically to mainly result from microglia binding.\textsuperscript{22}

We combined \textsuperscript{11}C-PBR28 imaging on an integrated 3T magnetic resonance (MR)-PET system with 7T $T_2^*$-weighted MRI to quantify, in a heterogeneous MS cohort, microglia/macrophage activation in the (1) cortex, particularly in cortical lesions segmented at 7T; and (2) thalamus, basal ganglia, and hippocampus.

We investigated the relationship between \textsuperscript{11}C-PBR28 uptake in the cortex and deep GM and neurological disability, cognition, and structural MRI metrics of neurodegeneration, to better understand the prevailing effects of neuroinflammation in MS, whether mainly harmful or beneficial. Finally, we quantified \textsuperscript{11}C-PBR28 uptake in lesional and normal-appearing WM (NAWM) to investigate their MRI and clinical correlates, and to assess whether similar levels of TSPO expression could be detected in lesional tissue in the cortex and WM.

**Subjects and Methods**

**Subjects**

The institutional review board and the Radioactive Drug Research Committee approved all study procedures, and subjects gave written informed consent to participate in the study.

Twenty-seven MS subjects (15 secondary-progressive MS [SPMS], 12 relapsing–remitting MS [RRMS]) were prospectively enrolled from 33 MS subjects who were genotyped for the TSPO gene Ala147Thr polymorphism, which predicts binding affinity to \textsuperscript{11}C-PBR28.\textsuperscript{23} Only high- and mixed-affinity binders underwent subsequent study procedures. Fourteen age-matched and TSPO affinity binding–matched healthy individuals were included as controls. Inclusion criteria were: age between 18 and 65 years, a diagnosis of clinically definite MS, education $\geq$ 8 years, absence of clinical relapse within 3 months, no use of corticosteroids within 1 month of study enrollment, and being on stable disease-modifying treatment or no treatment for at least 6 months. Exclusion criteria were: treatment with benzodiazepines and blood thinners, general PET/MRI contraindications, and major medical and/or psychiatric disorders. In MS, major depression was excluded using the Beck Depression Inventory–II (cutoff score $\geq$ 28).

In MS subjects, within 1 week from imaging procedures, neurological disability was assessed using the Expanded Disability Status Scale (EDSS),\textsuperscript{24} and cognitive performance with the following tests: Symbol Digit Modalities Test (SDMT), Trail Making Test (Trails A and B), California Verbal Learning Test–II (CVLT-II), Brief Visuospatial Memory Test–Revised (BVMT-R), and Wisconsin Card Sorting Test–64 Card Version (WCST). For each patient, a $z$ score was calculated to assess information-processing speed (average SDMT and Trails A $z$ scores), and executive (average WCST and Trails B $z$ scores) and memory (average CVLT-II and BVMT-R $z$ scores) functions.

**Imaging Data Acquisition**

All subjects underwent a 90-minute \textsuperscript{11}C-PBR28 MR-PET scan on a Siemens (Erlangen, Germany) simultaneous MR-PET system, BrainPET, a brain PET scanner operating in the bore of a
3T whole-body MR system equipped with an 8-channel head coil. The spatial resolution of the BrainPET (<3mm in the center of the field of view) is superior to that of any other whole-body PET scanner, because of the smaller size of scintillator crystals and of the scanner diameter, which minimizes the noncollinearity effect. Within 1 week from MR-PET, 24 MS subjects underwent 7T MRI on a Siemens scanner using a 32-channel head coil. Three subjects were excluded due to the presence of implants not approved for 7T.

**MR-PET ACQUISITION.** All participants received an intravenous bolus injection of $^{11}$C-PBR28 produced in-house. The mean ± standard deviation (SD) administrated dose was $11.4 ± 0.7$mCi in MS and $11.7 ± 0.5$mCi in controls. PET data were acquired in list-mode format during the 90-minute scan. In 15 MS subjects (10 SPMS, 5 RRMS) and 11 controls, blood data were sampled from an arterial line during the 90-minute PET acquisition (2mL every 6 seconds during the first 3 minutes and at 5, 10, 20, 30, 60, and 90 minutes postinjection) to generate an arterial plasma input function. Plasma metabolite analysis was performed as previously outlined. In the remaining subjects, blood data were not collected either because subjects preferred not to undergo arterial line placement (n = 13) or because an anesthesiologist was not available during PET (n = 2).

MRI scans acquired simultaneously during PET included: (1) multiple gradient echo 3-dimensional (3D) magnetization-prepared rapid acquisition (ME-MPRAGE) images (1mm isotropic voxels) for cortical surface reconstruction, coregistration to PET/7T data, segmentation of deep GM, and generation of attenuation correction maps; (2) conventional 3D fluid-attenuated inversion recovery (FLAIR) images (1mm isotropic voxels) for WM lesion segmentation; and (3) diffusion images (60 diffusion-encoding directions, b value = 3,000s/mm$^2$, 8 volumes without diffusion weighting, 2.5mm isotropic voxels) for assessing microstructural integrity in the pseudoreference region used for normalizing PET data. In 9 patients, ME-MPRAGE and FLAIR images were acquired after administration of a single dose (0.1mmol/kg) of gadolinium.

**7T MRI ACQUISITION.** The 7T protocol included acquisition of 2D fast low-angle shot-T$_2$ spoiled single-echo and/or multiecho images covering the supratentorial brain (0.33 × 0.33 × 1mm$^3$ voxels, 25% gap) for cortical lesion segmentation as previously detailed. Whole brain images were registered to the 60- to 90-minute PET data. PET data were acquired, the ME-MPRAGE was coregistered to the first non–diffusion-weighted image using affine registration, eddy current correction, and fitting of the diffusion tensor model at each voxel. Mean diffusivity maps were obtained in each subject.

**Quantification of $^{11}$C-PBR28 Binding**

In all subjects, $^{11}$C-PBR28 binding was assessed across brain regions using standardized uptake values (SUV) normalized by a pseudoreference region. In 15 MS and 11 controls, we also quantified $^{11}$C-PBR28 volume of distribution ($V_T$) using a 2-tissue compartment model with a metabolite-corrected arterial plasma curve as the input function.

**$^{11}$C-PBR28 SUV.** In-house software was used to compute voxelwise, for each subject, SUV (mean radioactivity/injected dose/weight) from the 60- to 90-minute postinjection data following a 2-step process. A preliminary SUV image was created using an attenuation correction map computed from the ME-MPRAGE in its native space. To account for possible motion between the time the ME-MPRAGE and the 60- to 90-minute PET data were acquired, the ME-MPRAGE was coregistered to this preliminary image using SPM8. The resulting transformation matrix was applied to move the attenuation map into the final PET space. A final SUV map (sampled at 1.25mm isotropic voxel size) was created using the new attenuation map, now well registered to the 60- to 90-minute PET data. PET data were reconstructed using 3D ordinary Poisson ordered-subset expectation maximization reconstruction, with corrections applied for attenuation, scatter, random events, dead-time, sensitivity, and normalization.

To account for global signal differences across subjects, SUV maps were normalized by a pseudoreference region (normalized SUV [SUVR]) with mean SUV in MS around the mean of SUV in controls. Due to MS being a diffuse disease that lacks an anatomically consistent reference region, we used a cluster-based approach. Using FreeSurfer and FSL, we computed voxelwise, for each subject, normalized SUV [SUVR] with mean SUV in MS around the mean of SUV in controls. Due to MS being a diffuse disease that lacks an anatomically consistent reference region, we used a cluster-based approach. Using FreeSurfer and FSL, we...
<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Controls, n = 14</th>
<th>All MS, n = 27</th>
<th>SPMS, n = 15</th>
<th>RRMS, n = 12</th>
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<tbody>
<tr>
<td><strong>Demographics and clinical data</strong></td>
<td></td>
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<tr>
<td>Gender, F/M</td>
<td>6/8</td>
<td>21/6</td>
<td>11/4</td>
<td>10/2</td>
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<tr>
<td>Age, yr, mean (SD)</td>
<td>48 (13)</td>
<td>48 (10)</td>
<td>52 (7)</td>
<td>43 (10)</td>
</tr>
<tr>
<td>HAB/MAB</td>
<td>7/7</td>
<td>16/11</td>
<td>10/5</td>
<td>6/6</td>
</tr>
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<td>SDMT, mean (SD)</td>
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<td>−0.4 (1.8)</td>
<td>−1.4 (1.4)</td>
<td>0.9 (1.3)</td>
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<td>Information processing, mean (SD)</td>
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<td>−0.9 (2.1)</td>
<td>−1.9 (1.9)</td>
<td>0.3 (1.8)</td>
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<td>Executive function, mean (SD)</td>
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<td>−1.3 (2.3)</td>
<td>−2.0 (2.6)</td>
<td>−0.3 (1.5)</td>
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<td>Learning recall, mean (SD)</td>
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<td>−0.70 (1.30)</td>
<td>−0.05 (1.2)</td>
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<td>Disease duration, yr, median [range]</td>
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<td>7 [1–40]</td>
<td>21 [6–40]</td>
<td>2 [1–33]</td>
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<tr>
<td>Subjects with at least 1 cognitive test with z score ≤ −1.5</td>
<td>—</td>
<td>20</td>
<td>13</td>
<td>7</td>
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<td>Subjects with mild/moderate depression</td>
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<td>2/2</td>
<td>1/2</td>
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<td><strong>MS treatments</strong></td>
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<td>Dimethyl fumarate</td>
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<td>Natalizumab</td>
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<td>2</td>
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<td>Glatiramer acetate</td>
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<td>6</td>
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<tr>
<td><strong>MRI metrics</strong></td>
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<tr>
<td>WM lesion volume, cm³, mean (SD)</td>
<td>—</td>
<td>8.1 (11.5)</td>
<td>12.6 (13.8)</td>
<td>2.6 (3.3)</td>
</tr>
<tr>
<td>Cortical thickness, mm², mean (SD)</td>
<td>2.39 (0.11)</td>
<td>2.33 (0.08)*</td>
<td>2.32 (0.10)*</td>
<td>2.36 (0.06)</td>
</tr>
<tr>
<td>Thalamic fractionb</td>
<td>6.1 (0.9) × 10⁻³</td>
<td>5.5 (1.1) × 10⁻³</td>
<td>5.2 (0.9) × 10⁻³</td>
<td>5.9 (0.1) × 10⁻³</td>
</tr>
<tr>
<td>Hippocampal fractionb</td>
<td>2.8 (0.6) × 10⁻³</td>
<td>2.7 (0.7) × 10⁻³</td>
<td>2.5 (0.6) × 10⁻³</td>
<td>2.9 (0.8) × 10⁻³</td>
</tr>
</tbody>
</table>

Beck Depression Inventory–II scores for mild depression = 14–19; for moderate depression = 20–28.

*p < 0.05 by Mann–Whitney U test.

bThalamic or hippocampal volume/total intracranial volume (SD).

EDSS = Expanded Disability Status Scale; F = female; HAB = high-affinity binders; M = male; MAB = mixed-affinity binders; MRI = magnetic resonance imaging; MS = multiple sclerosis; RRMS = relapsing-remitting MS; SD = standard deviation; SDMT = Symbol Digit Modalities Test; SPMS = secondary-progressive MS; WM = white matter.
identified clusters (minimum size = 9 contiguous voxels) in the WM of controls and NAWM of patients that were within ±0.5 SD of the mean $^{11}$C-PBR28 SUV for the global WM in controls. This process was conducted separately for high- and mixed-affinity binders. In MS, NAWM was defined by subtracting WM lesions from the global WM mask. To exclude patients the presence of microstructural changes due to inflammation in the pseudoreference region, mean diffusivity values in the selected NAWM clusters were compared to mean diffusivity values in WM clusters from controls using linear regression, and covarying for age, and no significant differences were found (MS = 6.1 ± 0.5 × 10^{-10} m²/s; controls = 6.3 ± 0.5 × 10^{-10} m²/s).

In each subject, masks of NAWM, WM lesions, and deep GM were registered to SUVR maps using FNIRT/FSL to extract mean SUVR. For deep GM structures, left and right SUVR were averaged together. To minimize partial volume effects due to the thin and circumvolved nature of cortex, SUVR in the cortex and/or cortical lesions (intracortical, leukocortical) were assessed using a surface-based approach. Using FreeSurfer, as previously detailed, in each subject, whole cortex and cortical lesion (intracortical, leukocortical) SUVR maps were registered onto their corresponding 3T cortical surface, thus resulting in 2D cortical maps. Mean SUVR was computed in each map at midcortical depth.

$^{11}$C-PBR28 $V_T$. PET data were binned into 28 time frames (durations: 8 × 10, 3 × 20, 2 × 30 seconds, and 1 × 1, 1 × 2, 1 × 3, 8 × 5, 4 × 10 minutes), and dynamic PET images were reconstructed following the same procedure detailed above.

Plasma activity and radioactivity were fitted to a 3-exponential model to derive an input function for the radiotracer injection. The resulting metabolite-corrected plasma input function, and time–activity curves data extracted from the same regions selected for SUVR analyses, were used to fit a 2-tissue compartment model of radiotracer binding and calculate $V_T$ for each region Pixel-wise modeling software (PMOD v3.3) from the rate constant, as previously described.

To account for (1) global signal differences across subjects and (2) intersubject variability in the input function, $V_T$ were normalized by the same pseudoreference region (normalized volume of distribution [DVR]) used for SUV. No significant differences in $V_T$ were found between MS subjects and controls in the pseudoreference region.

**Statistical Analysis**

Statistical analysis was performed using R Statistics software (v2.3). Demographics and conventional MR metrics were compared in MS versus controls using Mann–Whitney $U$ test. Linear regression models were used to: (1) compare, in MS versus controls, SUVR and DVR across GM (cortical lesions, whole cortex, deep GM) and WM (lesions, NAWM); (2) investigate in MS the relationship between SUVR and clinical (EDSS, cognitive scores) and structural MRI (cortical thickness, subcortical volume fractions) metrics; and (3) determine the relationship between SUVR and DVR. Age and TSPO affinity were included as covariates of no interest when appropriate. A correction for multiple comparisons was performed using the Holm method with a significance threshold of $p < 0.05$ for each subset of analyses including (1) SUVR and (2) DVR.
comparisons in all MS, (3) SUVR and (4) DVR comparisons in MS subgroups relative to controls, (5) correlations between SUVR and DVR, (6) correlations between SUVR and EDSS, (7) information-processing speed, (8) memory functions and, (9) structural data. The standardized effect size (Cohen $d$) was also calculated.

**SURFACE-BASED ANALYSIS.** Each individual 2D SUVR cortical map, sampled at midcortical depth, was smoothed along the surface with a 10mm full-width at half-maximum Gaussian kernel and normalized to a common template in FreeSurfer. A general linear model was used to assess vertexwise across the entire cortex: (1) differences in $^{11}$C-PBR28 SUVR between MS and controls and (2) the relationship in MS between $^{11}$C-PBR28 SUVR and clinical scores (EDSS, cognitive scores). Binding affinity, age, and cortical thickness at the vertex level were used as covariates of no interest. Correction for multiple comparisons was performed using Monte Carlo simulation with 10,000 iterations and a significance level of $p < 0.05$. Localization of significant cortical clusters was performed using the Desikan–Killiany atlas in FreeSurfer.

**Results**

**Demographics, Clinical Data, and Conventional MRI Findings**

Subjects’ demographics, binding genotype, MRI, and clinical data are reported in Table 1. Age was not statistically different between patients and controls.

Mean cortical thickness was lower in all MS and SPMS subjects than in controls. There were no differences in deep GM fractions; only SPMS cases showed significant thalamic atrophy (see Table 1).

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**TABLE 2. Corrected Probability Values, ES (Cohen $d$), and SE for Comparisons by Linear Regression (Covarying for Age and Binding Affinity) of $^{11}$C-PBR28 Uptake across Different Brain Tissue Compartments in MS Subjects versus Controls**

| Compartment | MS vs Controls | | | SPMS vs Controls | | | | RRMS vs Controls | | |
|-------------|----------------|-----------------|-----------------|-----------------|-----------------|-----------------|-----------------|-----------------|-----------------|
|              | $p$ | ES | SE | $p$ | ES | SE | $p$ | ES | SE |
| SUVR        | n = 27 vs 14 | | | n = 15 vs 14 | | | n = 12 vs 14 | | |
| Cortex      | 0.007 | 0.9 | 0.06 | 0.01 | 1.3 | 0.08 | 0.1 | 0.8 | 0.07 |
| NAWM        | 0.02 | 0.9 | 0.07 | 0.01 | 1.3 | 0.08 | 0.5 | 0.5 | 0.07 |
| WML         | 0.4 | 0.3 | 0.07 | 0.2 | 0.6 | 0.08 | 0.9 | 0.1 | 0.07 |
| ICL         | 0.007 | 1.4 | 0.06 | 0.01 | 1.6 | 0.09 | 0.04 | 1.4 | 0.08 |
| LCL         | 0.02 | 1.1 | 0.07 | 0.02 | 1.2 | 0.08 | 0.1 | 1.1 | 0.1 |
| Thal        | 0.008 | 1.2 | 0.1 | 0.002 | 1.7 | 0.2 | 0.3 | 0.9 | 0.1 |
| Hipp        | 0.006 | 1.3 | 0.06 | 0.003 | 1.6 | 0.09 | 0.1 | 1.1 | 0.07 |
| BG          | 0.02 | 1.0 | 0.06 | 0.01 | 1.3 | 0.09 | 0.3 | 0.8 | 0.07 |
| DVR         | n = 15 vs 11 | | | n = 10 vs 11 | | | n = 5 vs 11 | | |
| Cortex      | 0.0007 | 1.5 | 0.07 | 0.005 | 1.7 | 0.09 | 0.03 | 1.5 | 0.07 |
| NAWM        | 0.001 | 1.5 | 0.07 | 0.004 | 1.6 | 0.09 | 0.1 | 1.3 | 0.09 |
| WML         | 0.05 | 0.9 | 0.07 | 0.05 | 1.1 | 0.08 | 1 | 0.3 | 0.06 |
| CL$^b$      | 0.02 | 1.5 | 0.1 | 0.03 | 3.1 | 0.1 | 0.03 | 1.5 | 0.07 |
| Thal        | 0.005 | 1.3 | 0.1 | 0.004 | 1.9 | 0.1 | 0.7 | 0.8 | 0.1 |
| Hipp        | 0.04 | 1.1 | 0.1 | 0.05 | 1.2 | 0.1 | 1 | 0.8 | 0.07 |
| BG          | 0.02 | 1.1 | 0.07 | 0.02 | 1.4 | 0.07 | 0.9 | 0.5 | 0.06 |

$^a$The ES was computed by adjusting the calculation of the pooled standard deviation with weights for the sample size.

$^b$SPMS, n = 6; RRMS, n = 3.

BG = basal ganglia; CL = cortical lesions; DVR = normalized volume of distributions; ES = effect size; Hipp = hippocampus; ICL = intracortical lesions; LCL = leukocortical lesions; MS = multiple sclerosis; NAWM = normal-appearing white matter; RRMS = relapsing–remitting MS; SE = standard error; SPMS = secondary-progressive MS; SUVR = normalized standardized uptake values; Thal = thalamus; WML = white matter lesions.
11C-PBR28 Binding

TSPO LEVELS IN GM. Figure 1A shows 11C-PBR28 SUVR distribution across brain regions in controls and MS subjects, according to their binding genotype. DVR comparisons between MS and controls are illustrated in Figure 1B. Table 2 reports significant differences between groups, and their effect size. Relative to controls’ cortex, 11C-PBR28 SUVR were increased by 28% in intracortical lesions and by 24% in leukocortical lesions. Increased 11C-PBR28 uptake in cortical lesions did not differ significantly from that in normal-appearing cortex (paired t test).

The 11C-PBR28 DVR were significantly increased in MS (n = 15) relative to controls (n = 11) in the whole cortex (39%), thalamus (45%), hippocampus (25%), and basal ganglia (19%; see Fig 1B, Table 2). Cortical lesion DVR were also significantly increased (29%) relative to controls’ cortex in MS patients (n = 9). Due to the small number of cases with both cortical lesions and blood data, cortical lesion DVR were assessed by grouping intracortical and leukocortical lesions.

TSPO LEVELS IN WM. Higher TSPO levels were found in MS NAWM relative to control WM (see Fig 1, Table 2), with a 20% increase in SUVR, and a 29% increase in DVR. For WM lesions, we found a modest increase (7%) in 11C-PBR28 SUVR, and higher TSPO levels could be detected in 11C-PBR28 DVR (15%).

TSPO LEVELS IN RRMS AND SPMS. Corrected probability values and effect sizes for comparisons of 11C-PBR28 SUVR and DVR in each MS subgroup relative to controls are reported in Table 2. RRMS showed significant 11C-PBR28 increases in intracortical lesions and whole cortex. SUVR were also increased in thalamus, and hippocampus by ~29 to 22%, although at an uncorrected significance level. SPMS showed significantly increased SUVR and DVR in the whole cortex (28%, 32%), cortical lesions (ranging from 24% to 36%), thalamus (64%, 58%), hippocampus (33%, 30%), and basal ganglia (34%, 25%). SUVR and DVR were increased by 27% and 32% in NAWM. Increased 11C-PBR28 uptake in WM lesions was significant only for DVR (21%).

ASSOCIATION BETWEEN SUVR AND DVR THROUGHOUT THE BRAIN. In all 15 MS subjects (7 with high-affinity and 8 with mixed-affinity binding) and in 11 controls (5 with high-affinity and 6 with mixed-affinity binding) with blood data, there was a positive correlation between SUVR and DVR across all brain tissue compartments (Fig 3).
FIGURE 3: Plots illustrating, in the whole study cohort, the correlation between $^{11}$C-PBR28 normalized volume of distributions (DVR) and standardized uptake values (SUVR) across different brain tissue compartments by linear regression, adjusting for binding genotype and correcting for multiple comparisons. HC = healthy controls; MS = multiple sclerosis; NAWM = normal-appearing white matter; WM = white matter. [Color figure can be viewed at wileyonlinelibrary.com]
**Correlation with Clinical and Structural MRI Data**

In the MS cohort, EDSS correlated positively with increased $^{11}$C-PBR28 SUVR in the whole cortex, thalamus, hippocampus, and basal ganglia. A positive association was also found with $^{11}$C-PBR28 uptake in the NAWM and WM lesions (Fig 4).

Impaired memory function scores were associated with increased TSPO levels in the thalamus, whereas we did not find any association with information-processing speed function obtained by averaging SDMT and Trails A scores. Looking at individual tests, we found that SDMT $z$ scores negatively correlated with $^{11}$C-PBR28 SUVR in thalamus, hippocampus, and NAWM (see Fig 4).
FIGURE 5: (A–C) Overlay of the general linear model significance maps ($p<0.05$, corrected for multiple comparisons, adjusting for age, binding genotype, and cortical thickness) on the average pial surface showing, relative to healthy controls (HC; $n=14$), regions of increased $^{11}$C-PBR28 normalized standardized uptake values (A) in the whole multiple sclerosis (MS) cohort ($n=27$), as well as (B) in the secondary-progressive MS (SPMS; $n=15$) and (C) in the relapsing–remitting MS (RRMS; $n=12$) subgroups. (D, E) Regions in all MS cases where $^{11}$C-PBR28 uptake correlated negatively ($p<0.05$, corrected for multiple comparisons) with (D) memory and (E) information-processing speed function.
4). There was no association between $^{11}$C-PBR28 uptake and executive function.

In all patients and in each MS subgroup, cortical thinning was associated with increased TSPO levels in the thalamus ($p < 0.05$ corrected, covarying for age and binding affinity). In SPMS only, cortical thinning also correlated with neuroinflammation in the cortex and NAWM, although at an uncorrected significance level.

**Surface-Based Analysis of Cortical $^{11}$C-PBR28 Binding**

The cortical surface-based analysis revealed several clusters of increased $^{11}$C-PBR28 SUVR in both hemispheres in the entire MS cohort versus controls (Fig 5A–C, Table 3). The increase in $^{11}$C-PBR28 cortical uptake was diffuse in SPMS, but more localized in RRMS.

Negative correlations were detected between information-processing speed and memory function and $^{11}$C-PBR28 uptake in widespread frontoparietal, temporal, and occipital regions, and right cingulate cortex (see Fig 5D, E, Table 3).

No vertexwise associations were found with executive function or EDSS.

**Discussion**

Using $^{11}$C-PBR28, a second-generation TSPO radiotracer, we demonstrated diffuse microglia/macrophage activation in the cortex, including cortical lesions, and deep GM of MS cases.

Cortical demyelination is a well-established neuropathological feature of MS, and a main substrate of disease progression.\(^4\) In histopathological examinations, cortical lesions, particularly in progressive MS, usually
lacked the typical inflammatory features of WM plaques, and the prominent cellular population was activated microglia. We used a surface-based analysis to assess, in MS relative to controls, either in cortical lesions imaged with ultra–high-resolution 7T MRI or vertexwise across the whole cortex, differences in TSPO expression sampled at midcortical depth. Compared to volumetric approaches, this method has been shown to improve detectability of PET cortical signal changes. Previous 11C-PK11195 PET data, using a volume of interest approach, showed increased TSPO levels in MS in frontal, parietal, and occipital cortices. Here, we found that all MS disease stages exhibited abnormally increased TSPO levels, indicative of microglia/macrophage activation, in cortical lesions and several areas of the cortical ribbon. The highest concentration of cortical microglia/macrophage activation was localized in the occipital and temporal cortex in RRMS, the latter often reported as the preferential location for cortical MS demyelination, whereas it extended to widespread frontal and parietal regions in SPMS.

In some neuropathological studies of progressive MS, cortical lesions appeared to be topographically associated with meningeal inflammatory infiltrates, leading to the hypothesis that cortical demyelination may be induced by soluble factors, which diffuse from the meninges into the cortex and trigger demyelination directly or indirectly through microglia activation. The extent of meningeal inflammation correlated with microglia activation and the severity of demyelination in the underlying cortex. Meningeal inflammation, by means of perivascular lymphocytic infiltration, in association with cortical lesions, has also been described in biopsies at MS onset. The possibility of quantifying in vivo TSPO levels in cortical lesions could provide a tool for assessing the innate immune system inflammatory component of cortical lesions, its severity, and its association with structural cortical integrity. Additionally, it could offer relevant information on the role of microglia/macrophage activation in cortical lesions development at any disease stage. Interestingly, in contrast to WM plaques, RRMS and SPMS showed similarly increased TSPO levels in cortical lesions.

Histopathological findings on the role of inflammation in subcortical GM demyelination and neurodegeneration in MS are sparse and heterogeneous, and variable degrees of inflammation by means of activated microglia/macrophages have been reported in deep GM and hippocampus. In this study, we showed that microglia/macrophage activation extensively involved the thalamus, hippocampus, and basal ganglia. Previous 11C-PK11195 PET studies reported increased thalamic TSPO levels in MS. Abnormally high 11C-PK11195 binding has also been observed in clinically isolated syndrome in the brain central deep GM, although no significant differences were found in any single deep GM structure relative to controls. Here, neuroinflammation in thalamus and deep GM was seen at both disease stages, although more prominently in SPMS. Our MS cohort showed, relative to controls, significant global cortical thinning, whereas there were no differences in deep GM volumes with the exception of thalamic atrophy in SPMS. Cortical thinning in MS correlated with increased thalamic 11C-PBR28 binding. Increased thalamic 11C-PK11195 uptake secondary to distant cortical pathology has been described in stroke, implying the existence of a “projected neuroinflammatory” response from distant lesions along shared anatomical pathways. The thalami are major hubs in the brain, which integrate and process the signal originating from cortical projections. Spreading of cortical pathology to the thalami, or vice versa, could occur through thalamocortical connections. The direction of such changes could be tested in MS by assessing, longitudinally, microglia/macrophage activation changes along corticothalamic pathways.

The exact relationship between cortical neurodegeneration and locally activated microglia needs to be elucidated. Pathological evidence suggested that, as for cortical lesion development, neurodegeneration may be triggered by meningeal inflammation with cortical microglia activation. In a series of RRMS and SPMS cases, leptomeningeal inflammation, as seen on postcontrast FLAIR MRI, was found to be associated with more severe cortical atrophy. According to some experimental studies, microglial activation may also constitute a specialized danger signal following neuronal death or injury. In our cohort, however, especially in RRMS, neuroinflammation was not strictly associated with neurodegeneration, suggesting that it might not simply represent an indirect marker or consequence of neuroaxonal loss.

In line with previous observations, we found increased TSPO levels in WM lesions and NAWM. We found that TSPO levels in WM lesions, measured with both SUVr and DVR, were generally lower than those in cortical lesions. Additionally, in RRMS, the highest 11C-PBR28 uptake, relative to controls, was measured in intracortical lesions and focal cortical areas, whereas it was relatively modest in WM. Overall, these observations suggest that the innate immune system–mediated inflammatory events underlying the development of MS lesions might be dissociated.
in cortex and WM, especially at earlier disease stages, or there may be more heterogeneity in the inflammatory response associated with WM plaques than with cortical lesions. It is possible, however, that even if 7T MRI was used for cortical lesion detection, some cortical lesions were missed at visual inspection of scans. Nevertheless, cortical lesions identified on 7T T₂-weighted images likely represent the areas with greatest cortical demyelination, especially in RRMS.  

In our MS cohort, neurological disability correlated with microglial/macrophage activation in deep GM, in addition to neuroinflammation in the cortex and WM as previously reported. Here, we also demonstrated that microglia/macrophage activation correlated with reduced cognitive performance in MS. Increased TSPO levels in thalamus, hippocampus, and NAWM were associated with impaired SDMT function, a measure of information-processing speed, which is frequently affected in MS and, thus, commonly evaluated in the clinical setting. Diffusion imaging data highlighted that hippocampal–thalamic–prefrontal disruption affects SDMT performance in RRMS. An association between depression and hippocampal neuroinflammation, as measured by ¹⁸F-PBR11, has been recently reported in RRMS. Our study excluded cases with major depression or other psychiatric conditions, and moderate depression was present in only 3 MS patients. Interestingly, in our cohort, information-processing speed dysfunction also correlated with increased ¹¹C-PBR28 binding in frontoparietal, temporal, cingulate, and occipital cortices, in line with previous functional MRI experiments that described large areas of activation in the same cortical regions during SDMT execution. Finally, neuroinflammation in the thalamus, and in widespread temporal and occipital regions, and extending to the cingulate and prefrontal cortex, was associated with decreased memory function as measured by the BVMT-R and CVLT-II, likely reflecting the different neuroanatomical correlates for memory processing of verbal and visual material.

It has been hypothesized that microglia may play a dual role in MS pathogenesis: either detrimental, by causing neuronal dysfunction and promoting oxidative burst, or neuroprotective. Our findings suggest that the prevailing effects, in both GM and NAWM, are detrimental. The observation that SPMS cases showed overall higher brain TSPO levels than RRMS cases further supports this notion. However, it cannot be excluded that, at least at early disease stages, microglia might exert protective functions. Recent data, however, demonstrated that in clinically isolated syndrome persistently increased TSPO levels in NAWM predicted conversion to MS.

We assessed TSPO levels using ¹¹C-PBR28 SUVR and quantitative blood analysis, with both measures showing similar changes in MS. As previously suggested, we used DVR to account for ¹¹C-PBR28 intersubject variability in the input function. Our and previous ¹¹C-PBR28 data seem to suggest that quantitative blood data may be more sensitive than SUVR to microglia/macrophage activation in MS WM lesions, although this needs to be further confirmed. However, the strong positive correlation between SUVR and DVR in all tissue compartments assessed indicates that SUVR estimation is a reliable method for characterizing microglial/macrophage pathology in MS. In humans, ¹¹C-PBR28 SUVR has also been used to reliably distinguish healthy individuals from cases with neurological disorders. This method is expected to improve subject tolerability by allowing shorter scan time and not requiring arterial catheterization, making it less invasive and more feasible for clinical studies. The use of an integrated MR-PET system further increases patients’ compliance, in addition to improving spatial fidelity and registration accuracy.

In this study, we could assess blood–brain barrier integrity with contrast-enhanced MRI only in one-third of MS subjects; thus, we cannot definitely exclude that some of the TSPO changes in the remaining patients were secondary to acute systemic inflammation. However, blood–brain barrier disruption in cortex and deep GM is uncommonly observed in MS using contrast-enhanced MRI, and all patients were stable in terms of clinical relapses and MS-related treatments, suggesting that microglia/macrophage activation likely largely reflected chronic inflammation, especially in SPMS. Microglia could constitute a potential target for monitoring disease course, especially in the compartmentalized inflammation in progressive MS, and the effects of therapies. Our findings could guide longitudinal evaluations in larger MS cohorts to definitely establish the role of neuroinflammation in disease pathogenesis and progression.

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**Author Contributions**

E.H. and C.G. contributed equally to this work. Concept and design of the study: C.M.; acquisition and analysis of data: all authors; drafting manuscript and/or figures: E.H., C.G., T.G., C.M.

**Potential Conflicts of Interest**

Nothing to report.

**References**


33. Fujita M, Imaizumi M, Zoghbi SS, et al. Kinetic analysis in healthy humans of a novel positron emission tomography radioligand to...


