Microglia of Prefrontal White Matter in Suicide

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Abstract

Immune functions in the brain are associated with psychiatric illness and temporary alteration of mental state. Microglia, the principal brain immunologic cells, respond to changes in the internal brain milieu through a sequence of activated states, each with characteristic function and morphology. To assess a possible association of frontal white matter pathology with suicide, we stained autopsy brain tissue samples from 11 suicide and 25 nonsuicide subjects for ionized calcium-binding adapter molecule 1, cluster of differentiation 68, and myelin. Groups were matched by age, sex, and psychiatric diagnosis. We classified ionized calcium-binding adapter molecule 1-immunoreactive cells based on shape, immunoreactivity to cluster of differentiation 68, and association with blood vessels to obtain stereologic estimates of densities of resting microglia, activated phagocytes, and perivascular cells. We found no effect of psychiatric diagnosis but 2 statistically significant effects of suicide: 1) The dorsal-ventral difference in activated microglial density was reversed such that, with suicide, the density was greater in ventral prefrontal white matter than in dorsal prefrontal white matter, whereas in the absence of suicide, the opposite was true; and 2) with suicide, there was a greater density of ionized calcium-binding adapter molecule 1-immunoreactive cells within or in contact with blood vessel walls in dorsal prefrontal white matter. These observations could reflect a mechanism for the stress/diathesis (state/trait) model of suicide, whereby an acute stress activates a reactive process in the brain, either directly or by compromising the blood-brain barrier, and creates a suicidal state in an individual at risk. They also indicate the theoretical potential of imaging studies in living vulnerable individuals for the assessment of suicide risk. Further studies are needed to investigate specific phenotypes of perivascular cells and blood-brain barrier changes associated with suicide.

Key Words: Blood-brain barrier, Human, Microglia, Perivascular cells, Suicide, White matter.

INTRODUCTION

Every year, approximately 1 million people around the world die by suicide (1). Ninety percent of people who die by suicide in Western countries are diagnosed as having a psychiatric disorder (2), but most mentally ill individuals do not die by suicide, suggesting that factors outside and in addition to the disease may play a role (3, 4).

Evidence pointing to an immunologic component in the etiology of suicide has accumulated in recent years. Distinct in vivo (5–7) and postmortem (8, 9) cytokine profiles distinguish suicide attempters and completers from nonattempters and nonsuicide sudden deaths, respectively. Microglia are the immunologic sentinel cells of the CNS and its main producers of cytokines (10, 11). Changes in microglial morphology and function have been reported in major depression, bipolar disorder, schizophrenia, anxiety disorders, autism spectrum disorders, and suicide (12). Higher microglial cell counts were also detected in the white and gray matter of autopsy brains from psychiatric patients diagnosed as having schizophrenia or affective disorders (13). Radewicz et al (14) found significantly greater densities of microglia in the temporal and frontal cortices from autopsies of individuals with chronic schizophrenia, compared with individuals with no psychiatric diagnosis. A positron emission tomography (PET) study revealed significantly greater levels of mitochondrial translocator protein, which is expressed by activated microglia (and, possibly, by reactive astrocytes) in the gray matter of subjects with recent-onset schizophrenia compared with subjects without psychiatric illness (15, 16). Microglial activation could initiate psychopathology, could be secondary to another process (central or peripheral) that initiates psychopathology, or could be a reaction to psychopathology itself. These possibilities are not mutually exclusive. On the other hand, several studies provide evidence against activation of microglia in psychiatric disease (17–22). Only one of these studies directly investigated microglial activation in brains of suicides and found greater densities of activated microglia in the dorsolateral prefrontal cortex, anterior cingulate cortex, and medio-dorsal thalamus in suicides, independent of the psychiatric diagnosis (22).

Immune reactions in the CNS are influenced by its partially privileged and protected environment; it is devoid of lymphatics, contains only small numbers of conventional
antigen-presenting cells, and is isolated by a blood-brain barrier (BBB) that impedes the influx of peripheral immune cells and antibodies (23). However, recent findings speak against the dogma of the brain’s stringent impermeability to outside molecules in psychiatric diseases (24) and suicide (25–27). A disruption in the BBB may precede or follow a peripheral or central immune response, leading to an influx of peripheral immune cells, many of which may be found within or in contact with vascular walls. Thus, altered distribution of microglia or peripheral macrophages could be a biomarker of suicidality that might be assessed by in vivo imaging. In addition, these cells may provide a useful target for pharmacologic interventions that deter their harmful effects or enhance their beneficial effects.

Although in vivo studies of suicide attempters and completers point to the vulnerability of white matter (28–31), most autopsy studies of microglia in psychiatric disorders have addressed only gray matter. There have been no prior studies of microglia in white matter in suicide, and there are several reasons to believe that these could be important: a) Microglia are typically much more numerous in white matter than in the corresponding regions of the neocortex (32); b) activation of white matter microglia is a sensitive indicator of subtle damage to white matter, including hypoperfusion associated with aging (33, 34) and normal-appearing white matter in multiple sclerosis (35–40); and c) in vivo imaging with PET can be used to visualize activated microglia. While this method involves exposure to radioactivity and currently is not widely available, diffusion tensor magnetic resonance imaging detects white matter disruption and, if sufficiently sensitive and specific, could be used to predict suicide risk and thus to inform decisions about hospitalization and treatment.

The goal of our study was quantitatively to evaluate microglial cell density in prefrontal white matter. We focused on the prefrontal region because of its role in planning, decision-making, and suppression of impulsiveness (41) (processes that are relevant to suicide) and because numerous studies have found prefrontal differences, largely involving the serotonin system, in autopsy brains of suicides and non-suicides. Prefrontal cortical abnormalities reported in suicide include lower neuronal density in ventral and dorsal regions (42), less binding of ligands to the serotonin transporter per unit area (43) and per neuron (42) in ventral prefrontal cortex but not in dorsal prefrontal cortex, diminished expression of serotonin 2A receptor in Brodmann area 9 (44), diminished binding of serotonin 1A antagonists in Brodmann area 47 (45), lower levels of messenger RNA for serotonin 2A receptor in dorsolateral prefrontal cortex (44), lower levels of brain-derived neurotrophic factor and its receptor trkB in Brodmann area 9 (46), and increased levels of proinflammatory cytokines in Brodmann area 10 (8).

Because the focus of this study was suicide and not a specific psychiatric illness, the suicide group included individuals with various psychiatric illnesses and without known psychiatric illnesses, and the nonsuicide comparison group included individuals with and without psychiatric disease. We hypothesized that there would be an association between suicide and a higher density of activated phagocytes in prefrontal white matter.

### MATERIALS AND METHODS

#### Human Brain Tissue

Brain tissue was selected from the Macedonian/New York State Psychiatric Institute (NYSPI) brain collection at the Department of Neuropathology and Molecular Imaging, NYSPI. Autopsies were performed at the Institute for Forensic Medicine (Skopje, Macedonia) within 24 hours of death; the median postmortem interval (PMI) for collected specimens was 15 hours. Samples of brain, blood, and urine underwent toxicologic analysis. Structured clinical and demographic evaluations were performed in Macedonia by experienced psychiatrists and psychologists who were trained in these procedures and tested for reliability at the Department of Molecular Imaging and Neuropathology, NYSPI. For all cases, clinical diagnosis was determined by applying Diagnostic and Statistical Manual of Mental Disorders IV-TR criteria (47), and the diagnosis was confirmed at a consensus conference attended by senior researchers of the Department of Molecular Imaging and Neuropathology, NYSPI. Diagnostic neuropathologic examination was performed on all fixed hemispheres.

The selection included 11 suicides (mean age, 56 years; 6 women, 5 men) and 25 individuals whose cause of death was not suicide (mean age, 55 years; 12 women, 13 men). Demographic data, clinical diagnoses, and mechanisms of death are presented in Table 1. Among suicide cases, 1 individual with schizoaffective disorder and 1 individual with bipolar disorder were included in the schizophrenic and affective disorders groups, respectively. Two nonsuicides without schizophrenia or affective disorders had a history of alcohol dependence, 1 had an adjustment disorder, 1 had fully remitted bereavement, and 1 had a history of pathologic gambling.

Cases with active infections (<1% of collected cases) or grossly visible lesions in the frontal lobe (mostly gunshot wounds or acute contusions; <5% of collected cases) were excluded in order not to have a handful of cases give rise to skewed distributions of microglial densities. On the other hand, because occult neuropathologic abnormalities are common from the sixth decade onward, other histologic lesions were not excluded so as to avoid bias.

#### Processing of Tissue

At autopsy, the left hemisphere was sliced in the coronal plane at 2-cm intervals. The slices were placed in homemade cassettes to prevent warping during fixation in approximately 10 vol of phosphate-buffered 10% formalin for 5 days at room temperature. To forestall loss of immunoreactivity associated with prolonged fixation, we stored the slices at 4°C in a large volume of phosphate-buffered saline with 0.02% sodium azide. For this study, a coronal slice of the hemisphere, approximately 4 mm thick, was taken at the level of the rostral tip of the frontal horn of the lateral ventricle (Fig. 1) and subdivided into approximately 8 pieces to fit onto 75 × 25-mm slides. A record was made of the position of each piece. The blocks were embedded in paraffin. Serial 6-μm sections were cut onto charged glass slides (Fisher Super-Frost Plus; Fisher Scientific, Pittsburgh, PA).
Immunohistochemistry

Two serial sections were processed from each block. To determine microglial phenotype, we performed double bright-field immunostaining on both sections using 2 complementary markers: ionized calcium-binding adapter molecule 1 (Iba-1) and cluster of differentiation 68 (CD68). CD68 (or macrosialin) is a lysosomal membrane marker that is upregulated in phagocytosis (48). Microglia constitutively express CD68 (32). Although "CD68-immunoreactive microglia" are often interpreted as activated microglia (49, 50), this presumably refers to microglia recognizable by CD68 staining alone. Iba-1 is a well-established marker for microglia in many mammalian species, including humans (51, 52), and both cell bodies and processes of microglia are strongly immunoreactive to Iba-1. When activated, the cell body enlarges and becomes more strongly immunoreactive to CD68, and the processes become shorter and thicker (53–55). With double labeling, both of these features are apparent, thereby facilitating classification (56, 57). In our material, all Iba-1-immunoreactive cells contained some immunoreactivity to CD68. Iba-1-immunoreactive cells were classified as resting microglia if the processes were thin and CD68 staining occurred predominantly in a punctate distribution (Fig. 2A). (Similar double-labeling patterns were seen in [58] and [59].) Activated microglia had thicker processes and at least 1 focus of CD68 immunoreactivity that was greater than 2.5 μm in diameter (Fig. 2B). Macrophages had rounded cell bodies, few or no processes, and abundant immunoreactivity to CD68 (Fig. 2B).

For statistical analyses, because macrophages accounted for less than 5% of activated phagocytes and because most are probably microglia in advanced stages of activation, counts of activated microglia and macrophages were combined to estimate the density of "activated phagocytes." Because the nature of intravascular and perivascular Iba-1-immunoreactive cells is uncertain, all cells within or in contact with blood vessel walls were classified simply as perivascular. Typically, perivascular cells have ovoid bodies and no visible processes, but the vascular wall often seems to obscure their true shape (Fig. 2C). Counts of activated phagocytes, resting microglia, and perivascular cells were combined to determine total labeled cell density.

After rehydration of paraffin sections and treatment with 3% H2O2 (EMD Chemicals Inc, Gibbstown, NJ) for 30 minutes, antigen retrieval was performed by microwaving in Tris/EDTA (Sigma-Aldrich Inc, St Louis, MO) at pH 6.0 until

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**TABLE 1.** Patient Demographic Data, Psychiatric Diagnoses, and Mechanism of Death

<table>
<thead>
<tr>
<th></th>
<th>Suicide (n = 11)</th>
<th>Nonsuicide (n = 25)</th>
<th>Test Statistic</th>
<th>p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age, mean ± SD, years</td>
<td>56 ± 18</td>
<td>55 ± 16</td>
<td>t14 = −0.01</td>
<td>0.92</td>
</tr>
<tr>
<td>Sex</td>
<td>6 F/5 M</td>
<td>12 F/13 M</td>
<td>χ²1 = 0.1 (n = 36)</td>
<td>0.72</td>
</tr>
<tr>
<td>PMI, mean ± SD, hours</td>
<td>12 ± 8</td>
<td>15 ± 7</td>
<td>t14 = 1.1</td>
<td>0.28</td>
</tr>
<tr>
<td>pH, mean ± SD</td>
<td>6.5 ± 0.2</td>
<td>6.3 ± 0.3</td>
<td>t10 = −0.9</td>
<td>0.39</td>
</tr>
<tr>
<td>Psychiatric diagnosis</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Schizophrenia</td>
<td>2 F/1 M</td>
<td>4 F/3 M</td>
<td>χ²2 = 5.5 (n = 36)</td>
<td>0.05</td>
</tr>
<tr>
<td>Affective disorders</td>
<td>4 F/1 M</td>
<td>2 F/1 M</td>
<td></td>
<td></td>
</tr>
<tr>
<td>No psychiatric illness</td>
<td>3 M</td>
<td>6 F/9 M</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Mechanism of death, n</td>
<td></td>
<td></td>
<td>χ²4 = 10.5 (n = 36)</td>
<td>0.03</td>
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<tr>
<td>Respiratory</td>
<td>7</td>
<td>4</td>
<td></td>
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<tr>
<td>Traumatic</td>
<td>3</td>
<td>11</td>
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<tr>
<td>Cardiac</td>
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<tr>
<td>Other medical conditions*</td>
<td>2</td>
<td></td>
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<td></td>
</tr>
</tbody>
</table>

*Other medical conditions included iatrogenic thrombocytopenia and hemorrhagic pancreatitis.

F, female; M, male; PMI postmortem interval.
FIGURE 2. Representative images of microglial phenotypes in human prefrontal white matter, immunostained for Iba-1 (brown) and CD68 (black) with Verhoeff counterstain (gray blue). (A) Resting microglia were predominantly Iba-1–immunoreactive with small puncta of immunoreactivity to CD68. (B) An activated microglial cell (white arrow) and a macrophage (black arrow) showed strong immunoreactivity to CD68 (black), obscuring much of the Iba-1 staining (brown). (C) Immunoreactive cells within (red arrows) or in contact with blood vessel walls (black arrow) were counted as perivascular cells; a cell near the blood vessel, but within the parenchyma (white arrow), was counted as an activated microglial cell. Similar to activated phagocytes, perivascular cells showed strong CD68 immunoreactivity. Scale bar = 20 μm.
boiling for 3 minutes, stopping the microwaving for 5 minutes, replenishing the solution, and microwaving again until boiling for 3 minutes. The sections were then washed with phosphate buffer containing 0.1% Triton X-100 (Fisher Scientific) and covered with 10% normal goat or horse serum (Fisher Scientific) for 1 hour to block nonspecific binding. The slides were placed into individual close-fitting wells in a plastic box (ProHisto LLC, Columbia, SC), in which each was immersed in 2 mL of a 1:500 dilution of anti-CD68 mouse monoclonal IgG (DAKO Corp, Carpinteria, CA) and shaken gently for 3 days.

They were then washed, incubated with biotinylated anti-mouse IgG from horse (7.5 μg/mL; whole antibody; Vector Laboratories Inc, Burlingame, CA), washed, incubated with avidin-biotinylated peroxidase complex (Vector Laboratories Inc), washed, and incubated with 0.003% H₂O₂ (EMD Chemicals Inc), 0.02% 3,3'-diaminobenzidine-tetrahydrochloride (Acros Organics, Morris Plains, NJ), and 2% nickel sulfate (Fisher Scientific), yielding a black reaction product. The sections were washed and then incubated with agitation as before overnight at 4°C in a 1:2000 dilution of the second primary antibody, rabbit anti-Iba-1 (Wako Chemicals, Osaka, Japan). Biotinylated secondary antibody and avidin-biotinylated peroxidase complex were applied as before, except without nickel sulfate, so the reaction product was brown.

If the black and brown reaction products appear in exactly the same spot, only the black reaction product is seen because it is darker. As black staining is done first, it does not matter if the peroxidase from that reaction is still active during brown staining because the area is already black and any brown staining will be obscured. On the other hand, because CD68 and Iba-1 are distributed differently within microglial cells, double labeling of cells is easily appreciated. After immunohistochemistry, the sections were lightly counterstained for myelin by the Verhoeff method to provide a clear delineation of white matter (60). Differentiation times were optimized for each staining run to achieve the minimal staining required to produce a visible contrast between gray matter and white matter without compromising immunohistochemical evaluations. Sections were dehydrated and mounted with Permount (Fisher Scientific).

Stereologic Estimation of Microglial Density

All counts were performed, using Visiopharm Software version 3.6.5.0 (Visiopharm, Hørsholm, Denmark), by 1 observer (Tatiana P. Schnieder) who was blind to the identity of the subjects. All counts were conducted with an Olympus BX-61 microscope (Olympus, Center Valley, PA) connected to a motorized 8-slide stage (Prior, Rockland, MA) and a DP71 digital camera (Olympus). A physical disector probe (61) was used to estimate the densities of activated and resting microglia, perivascular cells, and macrophages.

The physical disector method is older and now less familiar than the optical disector (62) but is similar in concept. Both methods are for estimating the number or numeric density of objects in a volume by counting profiles of the objects in slices of the volume. In a randomly placed slice of fixed thickness, the number of profiles will be influenced by the size and orientation of the objects to be counted. Large objects and objects with their longest dimension perpendicular to the plane of section will be more likely to appear than would smaller objects or objects with their longest dimension parallel to the plane of section. This is the commonly referenced problem of “bias” (62). A related problem is that, even if one is certain that all of the objects to be counted are identical in size, shape, and orientation, if their length perpendicular to the slice is greater than the thickness of the slice of tissue in which counts are performed, it is not possible to know what that length is; therefore, there is no way to relate the number of profiles to the volume of tissue on the slide. With the optical or physical disector, however, we examine adjacent levels of the tissue to identify and count only the “first” appearance of a cell. We only count cells that are identified in a measured deeper portion of the tissue but cannot be identified more superficially. In the case of the optical disector, depth in the tissue is determined by the plane of focus in which the object appears. One designates a superficial exclusion zone, and only objects that are in focus in the deeper disector zone, but not in the exclusion zone, are counted. The physical disector works identically, but the exclusion zone is 1 of 2 serial sections, and one counts only objects that appear in the other slide, but not in the first. The counting event, with either method, can be the appearance of any aspect of the object that the observer can identify uniquely and consistently, such as the nucleus, nucleolus, or cell body. In our case, the counting event was the first recognizable appearance of the cell body because this was stained intensely and we did not counterstain for nuclei.

There are practical advantages to both the physical disector and the optical disector. The physical disector requires alignment of the slides, which, although performed mostly by the software, must be fine-tuned by the operator. It allows clarity and familiarity of thin paraffin sections and eliminates concerns about penetration of antibodies through thick sections. The physical disector also avoids the optical disector’s requirement to determine the exact depth at which an object comes into focus and the reliance of that process on lenses and condensers of high numeric aperture.

To use the physical disector, the software photographed and assembled montages of low-magnification images of the 2 serial 6-μm sections. The edges of white matter from the 2 images were aligned in the software by the operator. The perimeter of white matter was outlined in one of the images (“reference”), and images of uniform random sampling fields were photographed with a 20× dry objective (Olympus UPLFLN; numeric aperture, 0.5), with a 2,000-μm step between each field. Depending on the size of the white matter region, 1 of 3 counting frame sizes was used: 150 × 100, 200 × 150, or 250 × 200 μm². The same fields were photographed in the second (“lookup”) section and carefully aligned by the operator. The counting frame was superimposed on the image by the software. Cells were counted if i) their somas were visible; ii) their somas did not touch the exclusion boundaries of the counting frame; iii) they appeared within the unbiased counting frame applied to the reference section; and iv) they did not appear in the “lookup” section (63) (Fig. 3). Except for the corpus callosum, all white matter in the sampled sections was included, regardless of its distance from the cortex.
Statistical Analysis

Statistical analyses were performed using SPSS version 21 (SPSS, Chicago, IL). The relationships between age, PMI, brain pH, sex, and cell densities were examined. Because the behavioral functions of dorsal and ventral prefrontal cortex differ (41, 64) and because the ventral cortex is more specifically implicated in suicidality (43, 65--67), we analyzed the 2 regions separately. Thus, there were 6 primary dependent variables: densities of activated phagocytes, resting microglia, and perivascular cells in 2 regions each. In addition, 3 derived variables were assessed as follows: For each cell type, the standardized difference between dorsal and ventral density and the mean of the 2 was computed as

\[
\frac{\text{ventral density} - \text{dorsal density}}{\text{ventral density} + \text{dorsal density}}
\]

Correlations among the dependent variables and between the dependent variables and age, brain pH, and PMI were determined by linear regression. Suicides and nonsuicides were compared on each dependent variable separately, including age, sex, and PMI as covariates, when significantly correlated with the dependent variable. Because the correlations and comparisons in Table 1 were determined only to inform the comparisons between groups, the significance values for these measures were not adjusted for multiple comparisons. The comparisons of suicides with nonsuicides on each of the 9 dependent variables were evaluated with and without adjustment for multiple comparisons.

On average, 256 counting frames were analyzed and 1,348 cells were counted per subject. Coefficient of error (CE) was estimated to a first approximation for each subject for every cell type in the combined coronal slice and in ventral and dorsal subregions as \( \text{CE} = \frac{1}{\sqrt{n}} \), where \( n \) is the number of cells of a particular subtype counted for that subject (68). Values of CE ranged from 0.04 to 0.26 for activated cells, from 0.04 to 0.22 for resting microglia, and from 0.05 to 0.15 for perivascular cells. Coefficient of variation (CV) was computed as \( \text{CV} = \frac{\text{standard deviation (SD)}}{\text{mean density}} \) for each cell type in dorsal, ventral, and dorsal-ventral matter for suicide and nonsuicide subjects separately (Table 2). The fractional contribution of estimation error to the variance is given by \( (\text{CE} / \text{CV})^2 \) and ranges from 0.02 to 0.15, indicating that the sampling was adequate; the accuracy of estimates of the means would be much more efficiently improved by increasing the number of subjects than by counting more cells from each subject.

RESULTS

Group Characteristics

There were no differences in pH or PMI between groups separated by sex, manner of death (suicide or nonsuicide), cause of death, or psychiatric diagnosis. Age was different between diagnoses (\( F_{2,33} = 4.7, p = 0.02 \)) because the affective disorders group was older (mean ± SD age, 63.6 ± 11.1 years) than those without psychiatric illness (mean ± SD age, 60.7 ± 10.6 years).
TABLE 2. CV and Root Mean Square CE in Dorsal and Ventral White Matter for Suicide and Nonsuicide Deaths

<table>
<thead>
<tr>
<th>Region</th>
<th>Suicide</th>
<th>Dorsal</th>
<th>CV</th>
<th>CE</th>
<th>√CV²−CE²</th>
<th>Dorsal</th>
<th>CV</th>
<th>CE</th>
<th>√CV²−CE²</th>
<th>Dorsal</th>
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<th>CE</th>
<th>√CV²−CE²</th>
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<th>CV</th>
<th>CE</th>
<th>√CV²−CE²</th>
</tr>
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<tbody>
<tr>
<td>Activated Phagocytes</td>
<td>No</td>
<td>0.42</td>
<td>0.07</td>
<td>0.41</td>
<td>-</td>
<td>0.45</td>
<td>0.08</td>
<td>0.44</td>
<td>-</td>
<td>0.23</td>
<td>0.08</td>
<td>0.22</td>
<td>-</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Activated Phagocytes</td>
<td>Yes</td>
<td>0.46</td>
<td>0.07</td>
<td>0.45</td>
<td>-</td>
<td>0.36</td>
<td>0.07</td>
<td>0.35</td>
<td>-</td>
<td>0.19</td>
<td>0.07</td>
<td>0.18</td>
<td>-</td>
<td></td>
<td></td>
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</tr>
<tr>
<td>Resting Microglia</td>
<td>No</td>
<td>0.52</td>
<td>0.10</td>
<td>0.51</td>
<td>-</td>
<td>0.36</td>
<td>0.09</td>
<td>0.35</td>
<td>-</td>
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<tr>
<td>Resting Microglia</td>
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<td>0.34</td>
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<td>0.33</td>
<td>-</td>
<td>0.34</td>
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<td>0.23</td>
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<td>0.22</td>
<td>-</td>
<td></td>
<td></td>
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</tr>
</tbody>
</table>

The normalized variance for each measure within each group (suicide and nonsuicide), CV², contains a contribution, CE², from the error introduced into each measurement by counting systematic random samples rather than counting all of the cells of a given type, plus a contribution, CV²−CE², from the true variation among cases in the group. Thus, the minimal CV that can be obtained by counting more cells per subject is √CV²−CE². Any further reduction in the uncertainty of the mean (normalized standard error = CV/√N−1) can be achieved only by increasing N, which is the number of subjects in the group.

CV, group SD/group mean; CE, root mean square of theoretical CE for each case in the group.

Significant Correlations

Correlations among the 6 primary dependent variables and between them and age, brain pH, and PMI are listed in Table 3. The 3 measures derived from doroventral comparisons were not correlated with each other or with age, sex, or PMI, so the 3 comparisons were performed without covariates. At the level of individual slides, resting microglia were negatively correlated with activated phagocytes (r = −0.53, p < 0.0005); neither correlated significantly with perivascular cells (−0.1 < r < 0.1).

Differences Between Suicides and Nonsuicides

All density measurements are listed in Table 4. There were no significant effects of suicide on densities of resting or activated microglia in dorsal or ventral prefrontal white matter, but the density of perivascular cells in dorsal white matter was 18% higher in suicides than in nonsuicides (t = 2.2, df = 34, p = 0.033). In ventral white matter, there was no significant difference between suicides and nonsuicides (unadjusted p > 0.5).

Normalized dorsal-ventral differences for resting and perivascular cells were positive for both suicides and nonsuicides and approximately 30% greater in suicides than in nonsuicides; the differences were not statistically significant (p > 0.4). Densities of activated phagocytes were greater ventrally than dorsally among suicides, however, and greater dorsally than ventrally among nonsuicides (Fig. 4; t = 3.4, df = 34, unadjusted p = 0.0015).

If one corrects conservatively for multiple comparisons, the association of suicide with an altered dorsal-ventral gradient in the density of activated phagocytes remains significant. On Holm-Bonferroni correction (69), for the most significant of 9 members in a family of tests, the stated significance level is divided by 9. Thus, for an α of 0.05, the criterion becomes p < 0.0056, which is met. Once the first null hypothesis is rejected, the criterion for the second most significant result (higher perivascular cell density in dorsal white matter of suicide cases than of nonsuicide cases) becomes α/8 or p < 0.0063, which is not met by p = 0.033. The probability of type I error for at least 1 of 8 independent tests at p = 0.033 is 1 − (1 − 0.033)³ = 0.24. However, the tests are not independent; in fact, both the increased number of dorsal perivascular cells and the increased ratio of ventral to dorsal activated cell density could result from a preferential localization of activated phagocytes in dorsal prefrontal white matter to blood vessels and perivascular space in suicidal

TABLE 3. Correlation Coefficients Among Primary Variables

<table>
<thead>
<tr>
<th></th>
<th>Activated Phagocytes</th>
<th>Activated Phagocytes</th>
<th>Resting Microglia, Dorsal</th>
<th>Resting Microglia, Ventral</th>
<th>Perivascular Cells, Dorsal</th>
<th>Perivascular Cells, Ventral</th>
<th>PMI</th>
<th>pH</th>
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</thead>
<tbody>
<tr>
<td>Activated phagocytes, ventral</td>
<td>0.787*</td>
<td>-0.596*</td>
<td>-0.444*</td>
<td>0.738*</td>
<td>-0.006</td>
<td>-0.035</td>
<td>-0.404</td>
<td>0.084</td>
</tr>
<tr>
<td>Resting microglia, dorsal</td>
<td>-0.449*</td>
<td>-0.463*</td>
<td>0.36*</td>
<td>-0.011</td>
<td>-0.015</td>
<td>-0.171</td>
<td>-0.237</td>
<td>-0.113</td>
</tr>
<tr>
<td>Resting microglia, ventral</td>
<td>-0.06</td>
<td>0.147</td>
<td>0.238</td>
<td>-0.14</td>
<td>0.188</td>
<td>0.033</td>
<td>0.314</td>
<td>0.108</td>
</tr>
<tr>
<td>Perivascular cells, dorsal</td>
<td>0.029</td>
<td>0.029</td>
<td>0.053</td>
<td>-0.110</td>
<td>0.029</td>
<td>0.053</td>
<td>0.057</td>
<td>0.204</td>
</tr>
<tr>
<td>Perivascular cells, ventral</td>
<td>0.12</td>
<td>0.12</td>
<td>0.033</td>
<td>0.45*</td>
<td>0.033</td>
<td>0.461*</td>
<td>0.068</td>
<td>-0.131</td>
</tr>
<tr>
<td>PMI</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>0.204</td>
<td>-0.112</td>
</tr>
<tr>
<td>pH</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>-0.421†</td>
<td></td>
</tr>
</tbody>
</table>

Correlation coefficients are expressed as r values.
*Correlation is significant at the 0.01 level (2-tailed).
†Correlation is significant at the 0.05 level (2-tailed).
The enhanced activation produces behavioral changes and activates D-aspartate receptor, enhancing its activation by glutamate.

Potential Role of Microglia in Suicide

Suicide is often viewed as a combination of 2 components: a long-standing trait or diathesis of susceptibility and an acute stress or trigger. Autoimmunity can have a parallel structure. For example, in a mouse model of systemic lupus erythematosus, there is a state of production of antibodies to double-stranded DNA; this state does not affect the brain until an acute stress or trigger. Autoimmunity can have a parallel structure. For example, in a mouse model of systemic lupus erythematosus, there is a state of production of antibodies to double-stranded DNA; this state does not affect the brain until at an acute stress or trigger. Autoimmunity can have a parallel structure. For example, in a mouse model of systemic lupus erythematosus, there is a state of production of antibodies to double-stranded DNA; this state does not affect the brain until stress compromises the BBB. This allows the entry of the tumor necrosis factor (TNF)-a receptor, enhancing its activation by kynurenine pathway.

Speculation on serotonin

The possibility of an autoimmune basis for some cases of suicide is supported by the finding that antibodies to serotonin were present in more than half of cases of major depression (compared with fewer than 10% of individuals without depression) and were found in all cases of depression with elevated proinflammatory cytokines. Presumably, an autoimmune process directed against serotonin could remove serotonin axons or their terminals and account for deficits in serotonin, its transporter, and its presynaptic receptors. This speculation is also consistent with the dorsal-ventral specificity of our findings because the transporter deficiency associated with suicide is predominantly ventral.

Low serotonin transporter binding could be caused by reduced intrasynaptic serotonin. It is possible that serotonin levels are depleted by activated microglia that have enhanced metabolism of tryptophan to kynurenine via the kynureinic acid pathway (76). Cytokines are known to enhance the kynureinic acid pathway in microglia, and quinolinic acid can activate N-methyl-D-aspartate receptors and possibly disrupt the BBB (77). As a result, quinolinic acid could increase stimulation of neurons in ventral prefrontal cortex and lead to a loss of normal connectivity with the limbic system and impaired impulse control. Because our study was confined to white matter, correlations with cortical abnormalities are speculative.

Potential Clinical Importance

A potentially important implication of the different dorsoventral distribution of activated phagocytes in suicide is that it could be detected in vivo PET with ligands for the mitochondrial translocator protein (formerly the peripheral benzodiazepine receptor) (30). These receptors are expressed by reactive microglia and possibly astrocytes. Although

![Figure 4](http://jnen.oxfordjournals.org/)

### Table 4: Cell Densities, by Manner of Death, in Dorsal and Ventral Prefrontal White Matter

<table>
<thead>
<tr>
<th></th>
<th>Suicide (n = 11)</th>
<th>Non-suicide (n = 25)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Activated phagocytes</td>
<td>5,592 ± 2,578</td>
<td>6,901 ± 2,887</td>
</tr>
<tr>
<td>Resting microglia</td>
<td>6,292 ± 2,267</td>
<td>6,309 ± 3,120</td>
</tr>
<tr>
<td>Perivascular cells</td>
<td>4,684* ± 884</td>
<td>3,967 ± 893</td>
</tr>
<tr>
<td>Total labeled cells</td>
<td>16,567 ± 1,666</td>
<td>16,736 ± 2,888</td>
</tr>
</tbody>
</table>

Data are expressed in cells per cubic millimeter and presented as mean ± SD.

*p < 0.05 (significantly different from nonsuicides).
the affinity of the most specific radioligand currently available is also influenced by a genetic polymorphism (79), this can be determined by genotyping. In any event, genetic and other factors that globally influence PET signal would affect dorsal and ventral prefrontal cortex binding comparably, without obscuring a dorsal-ventral difference. Similarly, the dorsal-ventral comparison in our study should not be influenced by shrinkage, extent of fixation, PMI, or other confounds that could globally influence measurements of cellular density. With regard to the implications of our findings for measurable biomarkers of suicide, it should also be noted that methods are undergoing development to track the influx of peripheral phagocytes to the CNS by transport of radioactive or paramagnetic tracers (80). Less specifically, the microglial changes might be associated with changes in diffusion of water in white matter, which could in theory be detected by diffusion tensor imaging.

Implications of Perivascular Cells

Between-subjects analyses failed to reveal any significant difference in the densities of resting or activated cells between suicide and nonsuicide subjects. Moreover, contrary to our prediction, in suicides, there was a trend toward a lower density of activated cells in dorsal white matter. However, in the same region, suicide was associated with an 18% greater density of perivascular cells. Although we do not yet know their source, the finding of perivascular accumulations of phagocytes in suicide strengthens the idea of changes in the properties of the BBB. If both peripheral and CNS immune cells increase cytokine secretion and increase exchange across the BBB, a feed-forward inflammatory prelude to suicide could result (81).

At first glance, our findings of increased density of perivascular cells, but not of activated phagocytes, in dorsal white matter contrast with the findings of increased density of activated microglia in dorsolateral frontal white matter in suicide (22). However, Steiner et al (22) classified perivascular human leukocyte antigen D-related (HLA-DR) -immunoreactive cells as amoeboid reactive microglia, whereas we chose to place perivascular cells in a separate category because their nature is uncertain. These cells could be juxtavascular microglia (82, 83), perivascular macrophages (84–86), pericytes (87, 88), or some mixture of these.

Limitations

Our study includes certain limitations. The sample size was small, and only white matter was analyzed. The main drawback of any postmortem brain study is the inability to draw causal conclusions from correlational data at a single—albeit in this case critical—point in time. Another concern relates to the evaluation of microglial activation. Microglia with a classical “resting” morphology perform sentinel and reparative functions in the brain. Nonetheless, there is little doubt that the microglia that we classified as activated, with short and thick or absent processes and increased immunoreactivity to CD68, represent phagocytically active cells.

Another limitation is that we did not use a specific stain for blood vessels and hence may have underestimated the density of perivascular cells. However, such a bias would presumably be independent of suicide. Finally, although we used a physical disector probe to obtain unbiased cell density measurements within the processed tissue, we cannot know how much these were affected by shrinkage of tissue during fixation and processing for paraffin embedding. Although this means that the absolute densities that we have obtained may be different from those obtained with tissue processed differently, we obtain essentially identical levels of statistical significance, or lack thereof, if we eliminate effects of shrinkage by analyzing densities of individual cell types as a fraction of the total density of Iba-1-immunoreactive cells.

In conclusion, this study, though small and requiring replication, provides evidence for the presence of abnormal densities of phagocytic cells in prefrontal white matter in individuals who committed suicide. These may prove useful for future prediction and prevention of suicide.

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