Migration of Perilesional Microglia after Focal Brain Injury and Modulation by CC Chemokine Receptor 5: An In Situ Time-Lapse Confocal Imaging Study

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Microglia rapidly become reactive in response to diverse stimuli and are thought to be prominent participants in the pathophysiology of both acute injury and chronic neurological diseases. However, mature microglial reactions to a focal lesion have not been characterized dynamically in adult vertebrate tissue. Here, we present a detailed analysis of long-distance perilesional microglial migration using time-lapse confocal microscopy in acutely isolated living slices from adult brain-injured mice. Extensive migration of perilesional microglia was apparent by 24 h after injury and peaked at 3 d. Average instantaneous migration speeds of ~5 μm/min and peak speeds >10 μm/min were observed. Collective, directed migration toward the lesion edge was not observed as might be expected in the presence of chemotactic gradients. Rather, migration was autonomous and could be modeled as a random walk. Pharmacological blockade of the cysteine–cysteine chemokine receptor 5 reduced migration velocity and the number of perilesional migratory microglia without affecting directional persistence, suggesting a novel role for chemokines in modulation of discrete migratory parameters. Finally, activated microglia in the denervated hippocampal stratum oriens did not migrate extensively, whereas human immunodeficiency virus-1 tat-activated microglia migrated nearly twice as fast as those at the stab lesion, indicating a nonuniform microglial response to different stimuli. Understanding the characteristics and specific molecular mechanisms underlying microglial migration after neural injury could reveal novel targets for therapeutic strategies for modulating neuroinflammation in human diseases.

Key words: brain injury; chemokines; microglia; migration; neuroinflammation; reactive gliosis

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Introduction

Microglia are thought to represent resident brain tissue macrophage precursors, which rapidly respond to diverse insults in a stereotyped manner (Kreutzberg, 1996; Becher et al., 2000). Under normal conditions, microglia exist as nonmigratory ramified cells but rapidly transform into a motile amoeboid form within hours of activation (Stence et al., 2001; Nimmerjahn et al., 2005). These activated cells are presumed to migrate toward tissue damage and transform into phagocytes. The importance of microglial function in acute neurological insults such as traumatic brain injury (TBI) is well documented (Kreutzberg, 1996; Becher et al., 2000). This is underscored by recent microarray studies, which show the largest changes in gene expression in several different animal models of TBI to be inflammation related (Kobori et al., 2002; Natale et al., 2003). Prominent roles for microglia and neuroinflammation have also been hypothesized in chronic neurological diseases in humans and animal models such as Alzheimer’s disease, human immunodeficiency virus-1 encephalopathy, prion diseases, and multiple sclerosis (Benveniste, 1997; Becher et al., 2000; Nelson et al., 2002; Streit et al., 2004; Lee et al., 2005).

Recent attempts to study microglial dynamics in mammalian tissue have been informative but restricted to acute reactions induced by tissue slicing (Nimmerjahn et al., 2005). These studies have demonstrated the transformation of ramified microglia to more motile, amoeboid forms (Stence et al., 2001) but only limited analysis of microglial migration (Stence et al., 2001; Grossmann et al., 2002; Petersen and Dailey, 2004). Studies in the leech have provided clues about migratory characteristics and mechanisms of microglia in the invertebrate nerve cord, but the relevance of these findings to mammalian systems has not been verified (McCglade-McCulloh et al., 1989; Peters et al., 1991; Duan et al., 2003).

There exist no dynamic analyses of more mature reactive microglial functions in adult mammalian tissue after focal injury. Thus, the dynamic in situ characteristics and mechanisms of microglial migration remain primarily unknown. Foremost, the...
idea that the microglial response to focal injury is characterized by directed migration toward the lesion followed by transformation into phagocytes has not been verified in vertebrate tissue. In addition, the role of cysteine–cysteine (CC) chemokines in focal lesion is unclear (Ghirnikar et al., 1996; Hausmann et al., 1998). These cytokines are thought to guide blood monocytes to the injured brain but may also play a direct role in promoting directional and collective migration of resident brain immune cells, the microglia, as predicted by studies in vitro (Hausmann et al., 1998; Marella and Chabry, 2004). It is also not known for how long activated cells retain the capacity to migrate after injury or the migration strategies used. Finally, the concept of a uniform microglial response in the face of different types of injury has not been investigated by direct observation.

Here, we address these questions in acutely isolated living slices taken from in vivo brain-injured adult mice at various postlesion intervals. These results define perilesional microglial migratory behavior and regulation in living adult mammalian tissue after focal neural injury.

Materials and Methods

Animals. Adult male C57BL/6 mice (6–8 weeks of age) were obtained commercially (Harlan, Indianapolis, IN). All animals were cared for in accordance with United States Public Health Service guidelines.

Stab lesion. Forebrain stab lesion (FSL) was performed as described previously (Carbonell and Mandell, 2003).

Intracerebroventricular injections. Recombinant HIV-1 tat protein (2.0 nM) in 100 mg/ml BSA and 0.1 mM DTT in PBS (n = 6) were stereotaxically injected into the left lateral ventricle. Tat vehicle alone (100 mg/ml BSA, 0.1 mM DTT in PBS; n = 2) or deactivated tat (n = 4) were used as volume-matched controls. Deactivated tat was prepared by exposing the 2.0 mM tat solution to light and air under a culture hood overnight (Fanales-Belasio et al., 2002).

Preparation of live brain slices. Acutely isolated coronal brain slices were collected in chilled physiological saline with a vibratome at 1, 2, 3, 5, or 7 d after FSL (n = 16, 2, 3, 2, and 2, respectively) as described previously (Murase and Horwitz, 2002) with minor modifications. We used 300 μm slices in these experiments. Migration media consisted of cell culture media-1 (HyClone, Logan, UT) with 20% heat-inactivated normal horse serum. Vital labeling of microglia was performed with 5 μg/ml Alexa 488 or 568 IB4 (Molecular Probes, Eugene, OR) in migration media as described previously (Stence et al., 2001). For vital nuclear staining of live slices, 2–6 μM Syto 16 (Molecular Probes) was added to the media for at least 2 h at 37°C followed by a brief rinse. Before recording, slices were covered with 200–500 μl of growth factor-reduced Matrigel (BD Biosciences, San Jose, CA) and gelled at 37°C for 15 min to minimize tissue settling effects/motion artifact during recording. Temporal quantitation of microglial migration was derived from measurements from three to four independent experiments per time point from slices taken from at least two different mice. For consistency between time points, quantitated subfields were all centered on the lesion.

Confocal time-lapse videomicrography and analysis. Laser-scanning confocal images were acquired on a Nikon IX-70 inverted microscope with Fluoview 300 software (Fig. 1) (Olympus Optical, Melville, NY). A 20 or 40× long working distance objective (Olympus Optical) was used in these experiments corresponding to x–y tissue fields measuring 175 × 175 and 350 × 350 μm, respectively. The microscope stage and objective were maintained at 37°C. A z-series stack covering 40 μm of slice thickness was taken every 1.5 min, creating a three-dimensional time-lapse data set. To create time-lapse movies from the data set, four to six z-plane images were collapsed as two-dimensional projections using ImageJ 1.32 (Wayne Rasband National Institutes of Health, Bethesda, MD; http://rsb.info.nih.gov/ij/) and compiled into quick-time movies with Quicktime Pro 6.3 and 7.0.1 (Apple Computers, Cupertino, CA). For quantitative analysis, we identified 47 long-distance (i.e., continuously) migratory cells from four independent 90 min experiments representing tissue from four different mice. A portion of one of these raw movies is available for viewing (movie 9, available at www.jneurosci.org as supplemental material). To estimate speed (S), the centroid of the migrating cell was traced at 90 s intervals over a 90 min period using ImageJ 1.32 (Ware et al., 1998; Knight et al., 2000; Murase and Horwitz, 2002) to derive the mean squared displacements. Instantaneous speed was derived from absolute displacements per interval (interval, 90 s) for each cell. These were corrected to reflect velocities in micrometers per minute. Persistent random walk (PRW) analysis and Chemotactic Index (CI) were performed as described previously (Maheshwari and Lauffenburger, 1998; Ware et al., 1998). All values are reported as the mean ± SEM. Rose plots, manual plots, and histograms were created in Kaleidagraph 3.0 (Abelbeck Software, Reading, PA). Automated multiple cell-track tracings were performed with the MTrack2 plug-in for ImageJ 1.32.

Inhibition of CC chemokine receptor 5 and data analysis. Recording was performed in three phases from each slice. After an initial baseline recording experiment, 0, 4, or 40 μM TAK-779, the highly selective, small-molecule CC chemokine receptor 5 (CCR5) inhibitor (Baba et al., 1999, Marella and Chabry, 2004) in solvent of water, was added to the migration media and incubated for 3 h at 37°C. Recording was resumed in the identical field for 60 min in the continued presence of a drug. For drug washout, the slice was rinsed briefly in two changes of migration media and then cultured for 24 h. For the final recording phase, the migration media were replaced, the original recording field was then identified, and acquisition continued for ≥60 min. Experiments were performed in triplicate in tissue slices from five different mice. Percentage of microglial migration represents the average number of migratory cells from three 150 × 150 μm subfields during 1 h of recording. We operationally defined a migratory cell as one demonstrating a net translocation of at least two cell lengths during the sampling period. Directional protrusive events were scored to determine effects on polarity establishment. One
point was given for each dominant protrusive event or directional orientation change that occurred ≥90° from the previous direction. PRW analysis was performed as above on continuous migrants from 30 min migration episodes in each condition. Data analysis was performed with ANOVA, and post hoc pairwise comparisons were made with a t test with the Bonferroni correction factor. For all comparisons, p < 0.05 was considered statistically significant.

Results

Focal lesion incites extensive perilesional microglial migration

To determine the optimal time period for discrete analysis of microglial migratory behavior, we recorded from slices taken 1, 2, 3, 5, and 7 after lesion and quantified microglial migration (Fig. 2A). In the perilesional thalamus 1 d after injury, the majority of IB4-positive microglia were motile and many were highly migratory (Fig. 2B, left) (movies 1 and 9, available at www.jneurosci.org as supplemental material). The numbers of migratory cells increased over subsequent time points peaking at 3 d (Fig. 2A, B, middle) (movies 2 and 3, available at www.jneurosci.org as supplemental material). However, migration of cells was greatly reduced at 5 and 7 d after lesion (Fig. 2A) (movies 4 and 5, available at www.jneurosci.org as supplemental material). Thus, widespread migration of activated perilesional microglia occurs by 1 d after stab lesion with a peak at 3 d.

To verify that the robust microglial responses we observed here were mainly a result of the FSL rather than the brain-slicing procedure, we recorded from the injured and uninjured thalamus of slices 24 h after FSL to compare the extent of cellular migration. We used Syto-16, a vital nuclear dye, as a label independent of microglial activation in these experiments. As expected, migration of Syto-16-labeled cells was observed in the injured, but not the contralateral, thalamus (Fig. 2C) (movie 6, available at www.jneurosci.org as supplemental material). In addition, when movies are captured in the cortex at midline, it is possible to examine the injured and uninjured cortices simultaneously. In doing so, we verified that microglial activation and migration were limited to the injured hemisphere with both IB4 (movie 7, available at www.jneurosci.org as supplemental material) and Syto 16 (data not shown). Finally, we collected slices from a naive, uninjured mouse and recorded from several thalamic fields from three slices after 24 h of acute culture. Microglial activation was minimal as revealed by the low-IB4-staining intensity, and only a single cell was found to be migratory (movie 8, available at www.jneurosci.org as supplemental material). Thus, these results indicate that the migration observed perilesionally after FSL was attributable to a mature microglial response established in vivo and that the slicing procedure itself does not acutely induce significant migration of microglia in the adult mouse brain.

Characterization of perilesional microglial locomotion at 1 d

To characterize perilesional microglial migration in detail, we analyzed time-lapse experiments from mice killed 1 d after injury (supplemental Table 1, available at www.jneurosci.org as supplemental material). The 3 d time point was not selected because of the vastly increased numbers of migratory cells, which complicated discrete two-dimensional analysis (e.g., overlapping paths, increased interactions). A portion of a typical experiment in 1 d tissue may be viewed on-line (movie 9, available at www.jneurosci.org as supplemental material). We found, on average, that 57 ± 3% IB4-positive cells in a 40× perilesional field were migratory. Approximately 30% were long-distance migrators (see Materials and Methods) and are the basis of the quantitative analysis below. This long-distance migration was mostly continuous and meandering (movies 1–5, 9, available at www.jneurosci.org as supplemental material). Migratory cells, which stopped for extended periods, often did so to interact with other cells, vessels, or the primary lesion (movie 9, available at www.jneurosci.org as supplemental material). Morphology of migratory cells varied, but most displayed transient forward protrusions and often long-trailing processes (Fig. 3A) (movie 10, available at www.jneurosci.org as supplemental material). Cells at the edge of the lesion were mostly nonmigratory and amoeboid in morphology and displayed behavior suggestive of phagocytosis/wound debridement (movie 9, available at www.jneurosci.org as supplemental mater-
perilesional microglia are highly locomotory 1 d after injury, and many exhibit long-distance migration. These cells represented only 5–30% of the total number of cells in each field. We conclude that perilesional microglia are highly locomotory 1 d after injury, and many exhibit long-distance migration.

To examine lateral (x–y) velocities of continuously migratory perilesional microglia (see Materials and Methods), we plotted instantaneous velocity versus time from individual cell data pooled from four independent 90 min experiments (Fig. 3B) (47 cells; n = 2815 observations). As a population, microglia migrated an average of 5.02 ± 0.06 μm/min (~300 μm/h) with peak bursts beyond 10 μm/min (~600 μm/h). Velocities fluctuated during migration. However, there was no dominant or consistent periodicity (data not shown). We conclude that activated perilesional microglia migrate at high, fluctuating velocities.

Autonomous, nonbiased migration of microglia in situ

In all experiments, the majority of locomotory cells appeared to migrate in a stochastic manner (Fig. 4A) (movie 11, available at www.jneurosci.org as supplemental material). Highly directed cells migrating linearly toward the lesion were rarely observed. To systematically determine whether microglia migrate collectively and directionally, we overlaid y-axis-normalized microglial cell tracks plotted from a common origin (Fig. 4B). Cells appeared to migrate as autonomous agents moving in all directions with no evidence for bulk flow. When the 47 trajectories are averaged, the resulting path is found in the positive x direction (Fig. 4B, inset). However, the magnitude (net displacement) of this mean cell path was minimal (17.72 μm). As expected, the CI for the population was only 0.09 (see Materials and Methods). We conclude that perilesional microglial migration is mostly independent and not highly directional.

Perilesional microglial migration can be modeled as a random walk

The autonomous and meandering trajectories of continuous perilesional migratory microglia is reminiscent of a random walk (Zygmourakis, 1996). Indeed, the average ratio of net to total migration distance was just 0.25. To determine whether the 47 long-distance migrating cells analyzed above could be modeled as random walkers, we determined the mean-squared displacement of the entire population (n = 2815 intervals; four independent experiments). We found that mean absolute displacement was proportional to the square root of time (Fig. 4B), consistent with a random walk process (R2 = 0.9998).

To further characterize the migration of perilesional microglia, we fitted the cell path analysis to the PRW model (Dunn, 1983; Othmer et al., 1988). All cell trajectories, individually (R2 > 0.90), reasonably fit the PRW model (data not shown). Not surprisingly, the average individual cell S was similar to the instantaneous population average at 4.98 ± 0.25 μm/min (range, 1.90–11.42 μm/min; median, 4.80 μm/min). The average P was determined to be 1.0 ± 0.01 min (range, 0.75–1.55 min; median, 0.98 min). Path length (S × P), which is the distance traveled before a change in direction, was 4.95 ± 0.23 μm (range, 1.73–10.42 μm; median, 4.86 μm).

Finally, a measure of the distribution of the cell population, or dispersion (S2 × P), was 27.02 ± 2.88 μm2/min (range, 3.41–119.02 μm2/min; median, 23.07 μm2/min). Thus, migratory perilesional microglia display high velocities but are not highly directionally persistent.

CC chemokines acting at the CCR5 receptor, such as macrophage inflammatory protein-1α/CC chemokine ligand 3 (MIP-1α/CCL3), MIP-1β/CCL4, and regulated on activation, normal T-cell expressed and secreted (RANTES)/CCL5, are upregulated in neurological diseases and in response to brain injury (Ghirnikar et al., 1996; Hausmann et al., 1998; Bajetto et al., 2002; Babcock et al., 2003; Marella and Chabry, 2004). These factors have been thought to mediate primarily leukocyte homing to brain lesions (Bajetto et al., 2002). Recently, CCR5 function has been shown to be important for chemotaxis of resident brain macrophages, the microglia, in vitro and in vivo and has been thought to mediate similar functions in vivo (Bajetto et al., 2002; Marella and Chabry, 2004). However, this role must be reconciled with the predominantly autonomous, random walk migration of perilesional microglia that we observed. Miller et al. (2003) proposed that chemokines, in certain situations, modulate specific physical parameters of cellular migration such as speed or protrusive events. To test this hypothesis in perilesional microglia, we directly observed microglial locomotion 24 h after lesion under conditions of chemokine receptor CCR5 inhibition with the highly specific, small-molecule antagonist TAK-779 (see Materials and Methods). A dose–response study was performed, revealing that the 40 μM dose resulted in a significant 43% reduction in the number of migratory cells per subfield reflected as percentage of migratory microglia (p < 0.05) (Fig. 5A) (movie 12, middle, available at www.jneurosci.org as supplemental material). Although migration was inhibited, cell motility in general remained highly active. Many of these "arrested" cells under CCR5 blockade with 40 μM TAK-779 displayed repeated protrusive lunge in various directions as if attempting to migrate; however, little net displacement was exhibited. There was no difference in the number of directional protrusive events (see Materials and Methods) between these migration-arrested microglia and normally migrating microglia during a 30 min observation period (p = 0.67; t test; mean scores, 5.6 ± 0.5 and 5.4 ± 0.4, respectively), suggesting that directional control is independent of CCR5 activation. Microglial migration could be rescued to 86% of predrug conditions after 24 h of drug washout (Fig. 5A) (movie 12, right, available at www.jneurosci.org as supplemental material). The number of viable cells per field did not differ significantly between any of the three phases of recording, further ruling out a toxic effect of 40 μM TAK-779 treatment (Fig. 5B) (p = 0.89; ANOVA). Thus, blockade of CCR5 significantly hinders locomotion of perilesional microglia but not directional protrusive events.
To determine the effects of CCR5 blockade on the enduring migratory microglia, we performed PRW analysis at the 40 μM dose. The average cell speed decreased 27% compared with baseline conditions under CCR5 inhibition (p < 0.01) but could be rescued to 90% of baseline levels with washout (Fig. 5C). Consistent with the no-drug effect on directional protrusive events noted above, directional persistence times were unaffected (Fig. 5D). Average path length and dispersion were decreased by 28% (Fig. 5E) (p < 0.01) and 47% (Fig. 5F) (p < 0.01), respectively. Path length returned to 92% of baseline and dispersion to 83% after washout. Also supporting the PRW analysis, TAK-779 resulted in a leftward shift of the distribution histogram of instantaneous velocities (mean, 4.57 ± 0.11 μm/min) compared with pretreatment (mean, 6.65 ± 0.15 μm/min). Drug washout (mean, 6.30 ± 0.15 μm/min) rescued velocity distribution to near-baseline conditions.
modulates speed, but not directional persistence, of migrating microglia.

**Induction and quality of microglial migration is stimulus dependent**

To examine dynamic behavior of activated microglia in indirectly damaged regions, we collected slices 3 and 7 d after FSL and rendered movies from the hippocampal stratum oriens contralateral (cSO) to the lesion. This region is deafferented by the wound (Hausmann et al., 1998; Carbonell et al., 2005) (W. S. Carbonell and J. W. Mandell, unpublished observations). In contrast to the widespread, continuous migration at the primary lesion at 24 h (Fig. 6A, middle), only a small population of cells (<10%) migrated in the cSO and for brief bursts of only a few cell lengths (Fig. 6A, left). These microglia demonstrated highly dynamic processes (movie 13, available at www.jneurosci.org as supplemental material). At 7 d after injury, microglial activation in the cSO was diminished as demonstrated by very low to undetectable IB4 labeling of cells (data not shown). We conclude that activated microglia in the deafferented cSO, while possessing highly dynamic processes, do not appear to migrate extensively.

HIV-1 tat protein is a key regulator of viral replication (Rapaport et al., 1999). It has also been found to be neurotoxic in nanomolar quantities (Phillipon et al., 1994) and induces migration of microglia in vitro (Lokensgaard et al., 2001). To test the hypothesis that HIV-1 tat protein would induce migration of microglia in situ, we prepared brain slices for time-lapse confocal microscopy 24 h after an intracerebroventricular injection of 2.0 nM recombinant tat. Tat-activated microglia in situ were highly migratory (Fig. 6A, right, B) (movie 14, left, available at www.jneurosci.org as supplemental material) with an average instantaneous velocity of 9.20 ± 0.24 μm/min (~550 μm/h) and peak bursts beyond 16 μm/min (>1000 μm/h). Migratory paths were meandering, similar to cells in the perilesional thalamus (Fig. 6A, right) (movie 14, left, available at www.jneurosci.org as supplemental material). Slices from control animals injected with vehicle alone (data not shown) or deactivated HIV-1 tat protein (movie 14, right, available at www.jneurosci.org as supplemental material) demonstrated only limited microglial activation and no migration. Thus, the induction and quality of microglial migration is not uniform but highly stimulus dependent.

**Discussion**

In neurological diseases and diverse brain insults, microglial cells incite and promote the innate neuroinflammatory response responsible for brain repair (Kreutzberg, 1996; Becher et al., 2000; Streit et al., 2004). However, this inflammatory cascade, like that of the peripheral tissues in the body, can become pathogenic and contribute to the progression of disease processes. Therefore, understanding microglial function is essential for designing rational approaches for therapeutic modulation of this response. In this study, we combined an in vivo focal lesion model with laser-scanning confocal time-lapse microscopy to investigate the migratory behavior of activated microglia in acutely isolated living brain slices from adult mice. These results define dynamic parameters of the migratory response of activated perilesional microglia to focal brain injury in living tissue from adult mice for the first time.

Dailey and Waite (1999) demonstrated that the FITC-tagged microglial-specific lectin IB4 from *Griffonia simplicifolia* can be used to vitally label microglia in hippocampal slices from early postnatal rats. They exploited these properties to study microglial dynamics after tissue injury with confocal microscopy using the trauma of the slicing procedure as the activation stimulus. Their important investigations shed new light on the dynamics of the early microglial reaction (Stence et al., 2001; Grossmann et al., 2002). However, likely because of the acute nature of their activation stimulus (i.e., the brain slicing procedure), they observed only a small degree of cell migration (Stence et al., 2001). Indeed, Nimmerjahn et al. (2005) found no migration of microglia for up to 5.5 h after a laser lesion in mice in vivo. In contrast, our methods allow the neuroinflammatory response to develop unimpeded in vivo before slice collection and, therefore, are particularly suited for the study of mature reactive phenotypes such as long-distance microglial migration. These phenotypes are not displayed in brain slices from naïve adult mice after 24 h of culture, demonstrating that the slice models are not equivalent (movie 8, available at www.jneurosci.org as supplemental material). These results validate the use of acutely isolated brain slices after in vivo lesion in adult mice for the dynamic study of mature microglial reactions.

Here, we focused our investigation on resident microglial migration. We selected the 1 d time point after stab lesion to perform discrete analysis of migratory parameters. Although greater numbers of migratory cells were observed at the 3 d time point, the excessive numbers of overlapping and interacting cells hindered discrete two-dimensional analysis. We were also concerned about a larger contribution of blood macrophages to the IB4-positive cell population because of the nature of our injury model (Peters et al., 1991). Indeed, it has been classically thought that the number of macrophages of vascular origin in brain injury models with blood-brain barrier disruption was extensive.
ever, recent studies have demonstrated that resident microglia considerably outnumber invading macrophages at all time points after various injuries (Fujita et al., 1998; Babcock et al., 2003; Schilling et al., 2003; Vallières and Sawchenko, 2003). For instance, Fujita et al. (1998) estimated 102 ± 10 macrophages per cubic millimeter of tissue 24 h after stab lesion based on immunohistochemical and morphological criteria. Using this estimate, we would expect less than one cell per 350 × 350 × 40 μm field to be of blood origin in our analyzed movies at 1 d. Therefore, the great majority of migratory cells observed at 1 d after stab lesion are likely resident microglia.

Our finding of autonomous, random walk locomotion in the perilesional thalamus is not consistent with the classical chemotactic gradient hypothesis of microglial migration after injury. The suggestion that chemokines may exert effects via modulation of motile behavior of individual cells rather than promotion of directional bulk flow of cell populations seems particularly salient in light of these data (Miller et al., 2003; Wei et al., 2003). A prominent chemokine receptor implicated in resident microglial migration in the literature is CCR5. It has been shown to be constitutively expressed in resting microglia (Simpson et al., 2000; Otto et al., 2001) and upregulated in diverse pathological states (Mennicken et al., 2002; Glabinski et al., 2003; Natale et al., 2003). Additionally, many CCR5 ligands such as MIP-1α/CCL3, MIP-1β/CCL4, and RANTES/CCL5 are released after neural injury (Ghirnikar et al., 1996; Babcock et al., 2003; Marella and Chabry, 2004). Consistent with this idea, we found that in situ inhibition of the CC chemokine receptor CCR5 with the potent and highly specific small-molecule inhibitor TAK-779 (Baba et al., 1999) could reduce the extent of perilesional microglial migration by >40% and decrease migration speed of enduring migratory cells by nearly 30%. In contrast, directional persistence was not affected. CCR5 is also known to be variably expressed in astroglia in normal and pathological states (Mennicken et al., 2002; Lee et al., 2005). However, the acute effects of TAK-779 on microglial migration as well as the rescue experiments with drug washout strongly support a microglial receptor-based effect rather than an indirect effect through inhibition of astrocytic CCR5 or effects at the level of transcription. Therefore, we conclude that activation of CCR5 is an important regulator of biophysical parameters of microglial migration.

The incomplete inhibition of microglial migration with TAK-779, which we observed in the present study, is consistent with the findings of Marella and Chabry (2004) in an in vitro prion model. Together, our results suggest that other chemokine and nonchemokine receptors (Nolte et al., 1997; Walter et al., 2003) likely contribute to microglial migration after brain insult. Other possible chemokine receptor candidates include CCR1, CCR3, CX3CR1, and CXCR3 (Maciejewski-Lenoir et al., 1999; Bajetto et al., 2002; Flynn et al., 2003); however, relative contributions likely depend on injury stimulus. The intrinsic chemokines for CCR5 have been studied previously after stab lesion (Ghirnikar et al., 1996) and in other injury models (Baker et al., 1999; Otto et al., 2001; Babcock et al., 2003; Marella and Chabry, 2004). A candidate common to all models is RANTES/CCL5. However, Marella and Chabry (2004) demonstrated that maximal inhibition of microglial migration in vitro in response to prion with anti-RAN-

tes/CCL5 antibodies was only ~30% (compared with ~50% with 40 μM TAK-779), supporting a contribution of other CCR5 ligands to microglial migration. This topic deserves additional investigation.

The role of microglial migration in the neuroinflammatory response has not been studied extensively but is thought to be related to cytokine receptor profiles as well as release and expression of ligands (Bajetto et al., 2002; Flynn et al., 2003). These parameters, in turn, highly depend on the injury stimulus and extent of damage. Indeed, microglial responses are thought to manifest in a graded stereotypic pattern (Kreutzberg, 1996). Thus, the degree of microglial migration in tissue may represent a barometer of neuropathologic severity. However, the dynamic responses of microglia in the face of varying injury types has not been studied previously. In contrast to the robust locomotory response in the perilesional thalamus, we found that microglia in the contralateral denervated stratum oriens of the hippocampus were primarily nonmigratory, multipolar cells with highly active cellular processes. This may be similar to the graded response of microglia in the lethal and sublethal facial nerve lesion models (Streit and Kreutzberg, 1988). Additionally, microglia activated by an intracerebroventricular injection of HIV-1 tat protein migrated nearly twice as fast as those in the perilesional thalamus. This may be explained by both the direct neurotoxic effects of lentiviral tat (Philippon et al., 1994) and the known mimicry of CC chemokines by tat (Albini et al., 1998). HIV-1 tat effects on CC chemokine and chemokine receptor expression, including CCR5 and its ligands (Weiss et al., 1999; McManus et al., 2000), are also noted. Indeed, these data indirectly support our findings (see above) that CCR5 is involved in modulation of microglial migration velocity. Thus, our results are consistent with the idea of graded, context-dependent microglial activation and regulation.

In summary, the in situ confocal time-lapse approach used in the current study revealed the remarkable extent and diversity of the microglial migratory response to focal brain injury. These results put forth a new model of microglial migration after neural injury, whereby a stochastic, rather than directed, migration strategy is used to patrol certain areas of damage. This is similar to what has been proposed for lymphocyte trafficking and detection of antigen in lymph nodes (Wei et al., 2003). This strategy is advantageous over linear trajectories, because it allows a wider and more uniform coverage of tissue (Fig. 2B, middle), assuring proper allocation of resources to all areas in need of attention, not just the primary lesion. However, this does not exclude directed microglial migration in other injury contexts (McGlade-McCulloh et al., 1989; Carbonell et al., 2005) or during development. The role of chemokines in certain situations may be to enhance the probability of microglia locating damage by modulating physical parameters of migration such as velocity. Finally, recruitment and quality of microglial migration depends on injury context. These results further our understanding of the complex microglial response to neural injury and have important implications for the development of treatment strategies for modulating neuroinflammation in human disease and injury.

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