Microglia cells protect neurons by direct engulfment of invading neutrophil granulocytes: a new mechanism of CNS immune privilege

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Abstract

Microglial cells maintain the immunological integrity of the healthy brain and can exert protection from traumatic injury. During ischemic tissue damage such as stroke, peripheral immune cells acutely infiltrate the brain and may exacerbate neurodegeneration. Whether and how microglia can protect from this insult is unknown. Polymorphonuclear neutrophils (PMNs) are a prominent immunologic infiltrate of ischemic lesions in vivo. Here, we show in organotypic brain slices that externally applied invading PMNs massively enhance ischemic neurotoxicity. This, however, is counteracted by additional application of microglia. Time-lapse imaging shows that microglia exert protection by rapid engulfment of apoptotic, but, strikingly, also viable, motile PMNs in cell culture and within brain slices. PMN engulfment is mediated by integrin- and lectin-based recognition. Interference with this process using RGDS peptides and N-acetyl-glucosamine blocks engulfment of PMNs and completely abrogates the neuroprotective function of microglia. Thus, engulfment of invading PMNs by microglia may represent an entirely new mechanism of CNS immune privilege.
Microglia Cells Protect Neurons by Direct Engulfment of Invading Neutrophil Granulocytes: A New Mechanism of CNS Immune Privilege

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Microglial cells maintain the immunological integrity of the healthy brain and can exert protection from traumatic injury. During ischemic tissue damage such as stroke, peripheral immune cells acutely infiltrate the brain and may exacerbate neurodegeneration. Whether and how microglia can protect from this insult is unknown. Polymorphonuclear neutrophils (PMNs) are a prominent immunologic infiltrate of ischemic lesions in vivo. Here, we show in organotypic brain slices that externally applied invading PMNs massively enhance ischemic neurotoxicity. This, however, is counteracted by additional application of microglia. Time-lapse imaging shows that microglia exert protection by rapid engulfment of apoptotic, but, strikingly, also viable, motile PMNs in cell culture and within brain slices. PMN engulfment is mediated by integrin- and lectin-based recognition. Interference with this process using RGDS peptides and N-acetyl-glucosamine blocks engulfment of PMNs and completely abrogates the neuroprotective function of microglia. Thus, engulfment of invading PMNs by microglia may represent an entirely new mechanism of CNS immune privilege.

Key words: neuroinflammation; stroke; microglia; polymorphonuclear granulocytes; PMN; phagocytosis; time-lapse imaging

Introduction

Abundant evidence exists that an inflammatory response is mounted within the CNS after cerebral ischemia. Postischemic inflammation comprises the infiltration of polymorphonuclear granulocytes and monocytes/macrophages into the injured brain parenchyma, activation of microglia, and expression of pro-inflammatory cytokines, adhesion molecules, and other inflammatory mediators (Feuerstein et al., 1998; Dirnagl et al., 1999). Despite these well described inflammatory events after ischemia, the main impact of postischemic inflammation (beneficial or detrimental) is controversially discussed (del Zoppo et al., 2001; Feuerstein and Wang, 2001).

There is striking evidence that the infiltration of activated polymorphonuclear neutrophils (PMNs) (Kochanek and Hallenbeck, 1992; Jean et al., 1998; Prestigiacomo et al., 1999) into the injured parenchyma and the activation of microglia (Banati and Graeber, 1994; Kreutzberg, 1996; Minghetti and Levi, 1998) are playing an important role in the pathology of cerebral ischemia.

It is suggested that activated PMNs contribute to tissue damage by the release of oxygen radicals, proteases, and proinflammatory cytokines like TNFα (tumor necrosis factor α) (Barone et al., 1991; Jordan et al., 1999). As evidence, experimental strategies by avoiding PMN infiltration into the injured parenchyma are neuroprotective (Heinel et al., 1994; Beray-Berthat et al., 2003b; Miljkovic-Lolic et al., 2003). However, other studies failed to provide clear evidence of a cause–effect of PMN contribution to neuronal damage after ischemia (Heyward et al., 1996; Fassbender et al., 2002; Beray-Berthat et al., 2003a; Harris et al., 2005).

Similarly controversial findings exist regarding the role of microglia after ischemia. Several studies demonstrated the neurotoxic properties of activated microglia after ischemic or excitotoxic damage (Giulian et al., 1993; Kim and Ko, 1998; Rogove and Tisika, 1998; Yrjanheikki et al., 1999), whereas considerable evidence shows that microglia triggered by injured/dying neurons mediate a reduction of neuronal damage and induction of tissue repair (Morioka et al., 1991; Rapalino et al., 1998; Streit, 2002; Kohl et al., 2003; Kitamura et al., 2004; Shaked et al., 2005; Hayashi et al., 2006; Lalancette-Hebert et al., 2007).
Considering these controversial findings on the role of PMNs and microglia during transient ischemia, strikingly few studies have evaluated the direct effect of these immune cells on neuronal survival/damage (Dinkel et al., 2004; Mitrasinovic et al., 2005). Despite that, although the combined recruitment of both cell types to tissue sites affected by cerebral ischemia is well established, a potential direct interaction of both immune cell types has not been investigated in depth. Two recent studies demonstrated a possible microglia–PMN interaction. By histology, microglia with engulfed PMNs were observed in zones of focal cerebral ischemia in vivo (Denes et al., 2007; Weston et al., 2007).

To better understand the complexity of cellular inflammation after experimental ischemia, we investigated the individual impact of PMNs, microglia, and macrophages on neuronal survival after ischemia and studied how these cells interact during the inflammatory response. To achieve this, we used an inflammation model that comprises oxygen–glucose deprivation (OGD) in organotypic hippocampal slice cultures (OHCs) as ischemic model and the application of immune cells onto the OHCs as simulation of the postischemic immune cell infiltration into the injured parenchyma.

Materials and Methods

Induction of focal cerebral ischemia by middle cerebral artery occlusion with endothelin-1

Focal cerebral ischemia was induced by occlusion of the left middle cerebral artery (MCA) via intracerebral microinjection of endothelin-1 (ET-1), following Sharkey and Butcher (1995) and Baldauf and Reymann (2005). Briefly, anesthesia was induced with halothane in a mixture of nitrous oxide and oxygen (50:50), and maintained with 2–3% halothane (Sigma-Aldrich) via a rat anesthetic mask (Stölting). A 29 gauge cannula was inserted through the brain close to the MCA. Ischemia was induced by injection of 376 pmol of ET-1 (Sigma-Aldrich). The animals were killed by an overdose of chloral hydrate (Sigma-Aldrich) after 1 d. Frozen brains were cut coronally and 30 μm slices were taken for additional staining.

Organotypic hippocampal slice cultures

Hippocampal interface organotypic cultures were prepared as previously described (Stoppini et al., 1991; Neumann et al., 2006) from postnatal day 7–9 Wistar rats (Harlan Winkelmann). For two-photon microscopy experiments, hippocampal slice cultures were prepared from transgenic B6.Cg-Tg(Thy1-YFP)16rs mice (The Jackson Laboratory; distributed by Charles River), which express enhanced yellow fluorescent protein (EYFP) at high levels in subsets of central neurons, including the pyramidal cells of the hippocampus (Feng et al., 2000). Hippocampi were dissected and transversely sliced at 350 μm thickness with a McIlwain tissue chopper (The Mickle Laboratory Engineering). Slices were transferred to Millicell membranes (Millipore). Cultures were maintained at 37°C in 1 ml of serum-based medium containing 50% MEM-Hanks, 25% HBSS, 17 mM HEPES, 5 mM glucose, pH 7.8 (Cell Concepts), 1 mM l-glutamine (Biochrom), 25% horse serum (Invitrogen), and 0.5% gentamicin (Biochrom). After 6 or 7 d of primary cultivation, microglia cells were separated from other cell types by shaking, placed in a 25 ml flask (TPP) at a density of 5 × 10⁶ cells/ml and maintained in 5% CO₂ at 37°C for 2 d. Cultures consisted of 95% microglia cells as determined by staining with Alexa 568-conjugated Griffonia simplicifolia isolecitin B4 (Invitrogen).

Culture of macrophage cell line RAW 264.7

The macrophage cell line RAW 264.7 was cultured in DMEM supplemented with 10% FCS (Biochrom), 1% Pen/Strep (Biochrom), and 1% l-glutamine (Biochrom). After 6 or 7 d of primary cultivation, microglia cells were separated from other cell types by shaking, placed in a 25 ml flask (TPP) at a density of 5 × 10⁶ cells/ml and maintained in 5% CO₂ at 37°C.

Application of cells onto OHCs

Isolated primary microglia or macrophages were trypsinated (trypsin/ EDTA; Biochrom), centrifuged at 500 × g for 2 min, and finally resuspended in Neurobasal medium (Invitrogen). PMNs were freshly prepared shortly before the experimental start. PMNs were also resuspended in Neurobasal medium. The cells were applied directly onto 10-d-old OHCs in a volume of 1 μl of Neurobasal medium containing 8 × 10⁴ microglia, macrophages, or 1 × 10⁵ PMNs. Viability of microglia, macrophages, and PMNs after application onto the OHC was confirmed in initial experiments by previous staining with 5-chloromethylfluorescein diacetate (CMFDA) (Invitrogen) and by time-lapse microscopy of these cells (data not shown). As indicated, OHCs were fixed with 4% paraformaldehyde (PFA) and mounted with 3:1 PBS/glycerol. OHCs were then further analyzed with the indicated microscopy approach.

Oxygen–glucose deprivation

The membrane inserts carrying up to three OHCs were placed into 1 ml of glucose-free medium in sterile six-well culture plates (TPP) that had previously been saturated with 5% CO₂/95% N₂ for 10 min. Then OHC were subjected to the OGD [40 min of OGD in a temperature-controlled hypoxic chamber (Billups-Rothenberg); no glucose medium; N₂/CO₂ atmosphere] before retransfer into normal conditions. Control cultures were kept in regular medium (plus glucose) under normoxic conditions. The cultures were analyzed as indicated by individual experiments 24 or 48 h after OGD.

Analysis of cell death

Cell death was evaluated by cellular incorporation of PI at 24 or 48 h after OGD. Cultures were incubated with PI-containing medium (10 μM) for 2 h at 33°C. Fluorescent images were acquired in a semiautomated manner (Nikon motorized stage; LUCIA software) and analyzed by densitometry to quantify necrotic cell death (LUCIA Image analysis software). Based on transmission light images, the area of analysis was determined such that only the CA area (CA1–3) was analyzed, whereas the dentate gyrus was excluded. Background correction was performed automatically by a control square (150 × 150 μm) in the stratum moleculare outside the pyramidal cell layer. To combine data from individual experiments, the densitometric mean value of the respective insult of an indi-
vidual experiment was set to the value of 1 and given as relative fluorescence intensity of the insult damage.

Electrophysiology

Wistar rats (25 d old) (Harlan Winkelmann) were killed by a blow on the neck. After decapitation, the brain was quickly removed and placed into ice-cold artificial CSF (ACSF) having the following composition (in mM): 124 NaCl, 4.9 KCl, 1.3 MgSO4, 2.5 CaCl2, 1.2 KH2PO4, 25.6 NaHCO3, 10 d-glucose, saturated with 95% O2, and 5% CO2, pH 7.4. Transverse hip-pocampal slices (350 μm thickness) with adjacent subicular and entorhinal cortices were prepared using a vibratome (microm HM 650V). The slices were placed on Millicell membranes in a six-well culture dish (Sigma-Aldrich) with 1 ml of high K+ culture media [25% horse serum (Invitrogen); 40% Eagle’s Basal Essential Media (BME) (Sigma-Aldrich); 25% Earle’s balanced salt solution (Sigma-Aldrich); 10% 250 mM Na-glucose, pH 7.3; 0.5 mM L-glutamine (Sigma-Aldrich); 28 mM glucose, pH 7.32]. One hour after preparation, 2 μl of high K+ culture media containing 4 × 10−3 PMNs and 2 μl of cell-free media for control experiments, respectively, were applied on top of the slices and incubated overnight at 37°C. To maintain a carbonated atmosphere (95% O2/5% CO2), then, the slices were transferred into an interface-type recording chamber saturated with carbogen and carbogen at 33 ± 1°C and constantly super saturated with ACSF. Synaptic responses were elicited by stimulation of the Schaffer collateral–commisural fibers in the stratum radiatum of the CA1 region using lacquer-coated stainless-steel stimulating electrodes. Glass electrodes (filled with ACSF, 1–4 MΩ) were placed in the apical dendritic layer to record field EPSPs (fEPSPs). The initial slope of the fEPSP was used as a measure of this potential. The stimulus strength of the test pulses was adjusted to 30% of the EPSP maximum. During baseline recording, three single stimuli (10 s interval) were averaged every 5 min. After tetanization, recordings were taken as indicated in Figure 1C. Once a stable baseline had been established, long-term potentiation was induced by application of four times two-paired pulses in intervals of 200 ms (theta burst). The interval between the paired pulses was 10 ms, and the width of a single pulse was 0.2 ms.

Two-photon microscopy

For two-photon microscopy, PMNs were labeled with Cell Tracker Orange [5-(and-6)-(((4-chloromethyl)benzoyl)-amino)tetramethylrhodamine] (CMTMR) (7.5 μM in PBS, 10 min, room temperature; Invitrogen). At different time points after OGD induction, the OHCs were fixed with 4% PFA, subsequently mounted and subjected to three-dimensional two-photon microscopy. In other experiments, viable OHCs were analyzed. PMNs were labeled with CFMFA and microglia with CMTMR and both applied onto OHCs. Two hours after experimental start, the OHCs were placed in a custom-built chamber supplied with 37°C and 5% CO2 directly under the microscope. The two-photon microscopy setup was used exactly as previously described (Neumann et al., 2006). OHCs prepared from transgenic B6.Cg-TgN(Thy1-YFP)16Jrs mice (Feng et al., 2000) were imaged in a modus at 800 and 920 nm (Fig. 1A). A stable baseline had been established, first at 920 nm and then at 800 nm wavelength with no filter. The emission of EEYP at 800 nm was negligible as was the emission of Cell Tracker Orange at 920 nm. Image stacks were exported as two independent 16-bit multilayer TIFF stacks and subsequently reconstructed using the Volocity software package (Improvision).

Confocal microscopy

Cultures were examined with a confocal microscope equipped with a 40× magnification Plan Neofluar 0.75 objective, an argon laser emitting at 488 nm, and a helium/neon laser emitting at 543 nm. Multitracking was used to avoid cross talk between channels. Images were analyzed with Carl Zeiss software (Pascal; Carl Zeiss).

Time-lapse video microscopy

The cellular dynamics of PMN–microglia interaction were investigated using OHCs or an in vitro PMN–microglia coculture. Time-lapse microscopy of OHCs. Fluorescently labeled PMNs {7-amino–4-chloromethylcoumarin (CMAC)} and microglia (CMTMR) were applied onto OHCs. Two hours after start of the experiment, the OHCs were placed in a custom-made chamber (Incubator S-M; Pcon) adjusted to 37°C and 5% CO2 (CTI-Controller 3700 digital; Tempcontrol 37–2 digital; Pcon).

Time-lapse microscopy of PMN–microglia coculture. Primary microglia were cultured onto Matrigel-coated surfaces in a 12-well plate (Falcon; BD Biosciences Discovery Labware). Matrigel basement membrane matrix (BD Biosciences), containing laminin as a major component, was diluted in ice-cold DMEM (ratio, 1:20) and polymerized at 37°C for 30 min. Thereafter, 2 × 105 microglia/well were placed and recovered for 1 d. Freshly prepared human PMNs were applied to the primary microglia culture. If indicated, PMNs were CMFDA labeled prior to application. Thirty minutes after the experimental start, the coculture was placed into the chamber as described above. The time-lapse microscope was based on an Axiovert 200M (Carl Zeiss) stage equipped with a 10×, numerical aperture (NA) 0.3 lens or a 32×, NA 0.5 lens (Carl Zeiss) and a CCD camera (AxioCam MRm; Carl Zeiss). Images were recorded at defined time intervals. The data were subsequently analyzed with the Carl Zeiss software (AxioVs40 V4.5).

Statistical analysis

All data are given as mean ± SEM. Statistical analysis was performed by one-way ANOVA followed by post hoc comparison (Tukey’s test). A value of p < 0.05 was considered statistically significant.

Results

PMNs infiltrate the brain parenchyma after focal ischemia

Given the heterogeneity of the data concerning the relevance of peripheral immune cells for ischemia-induced neuronal damage, we first wanted to investigate which type of peripheral immune cell was initially recruited into the injured brain parenchyma after transient ischemia. To this end, endothelin was injected directly above the MCA. Endothelin as vasoconstrictor occluded the MCA for ~30 min. We found that, 1 d after injection, endothelin-mediated transient focal ischemia induced a cellular infiltrate that was mainly composed of PMNs as detectable from the characteristic lobulated nuclei of infiltrating cells in immunohistological sections of the damaged area (Fig. 1A). We found no marked infiltration with other peripheral immune cells such as macrophages or lymphocytes. Thus, the acute cellular infiltrate of the ischemia-injured brain in vivo was dominated by PMNs, suggesting a prominent role of this immune cell type for the additional development of neuronal damage.

Application of human PMNs onto organotypic hippocampal cultures does not influence neuronal survival

Following the results of the in vivo ischemia-induced recruitment of PMNs (Fig. 1A), one goal of this study was to clarify the role of PMNs for the development of neuronal viability. To this end, we chose a well established model of ischemic injury in situ using OHCs that maintain many characteristics of true brain–parenchyma, especially the complex three-dimensional structure of neuronal circuits, yet allow for precise control of cellular or humoral factors impacting on neuronal viability as described previously (Neumann et al., 2006). The first step was to evaluate whether PMNs as such had a detrimental effect on neuronal viability in healthy OHCs. Because of the very limited availability of primary rat PMNs, we chose to use human PMNs throughout this study and performed only key experiments with primary rat PMNs. Direct application of up to 2 × 104 human PMNs onto control OHCs had no effect on neuronal viability in the cornu ammonis (CA1–CA3) area after 24 h (Fig. 1B). Additionally, we investigated the effect of PMNs on neuronal function by electro-
physiological EPSP recording with subsequent LTP induction in 1 d in vitro (DIV) hippocampal slices. This approach provides two critical parameters of neuronal health. First, the shape and value of the synaptic signal reflects the quantity and integrity of the involved synapses. Second, the amount and persistence of the synaptic potentiation represents a highly sensitive marker for neuronal viability and functionality. According to the EPSP signal, neither the persistence nor the amount of the LTP was disturbed by application of \( \frac{2}{10^5} \) PMNs onto the 1 DIV hippocampal slices (Fig. 1C). These experiments indicated that PMNs derived from healthy human volunteers did not cause neuronal cell loss or neuronal electrophysiological disturbance in rat OHCs.

PMNs exacerbate neuronal damage after oxygen–glucose deprivation

To examine the effect of PMNs on CNS cells after ischemia, we simulated the PMN infiltration into the brain parenchyma by direct application of PMNs onto post-OGD OHCs (Fig. 2A). Application of increasing numbers of PMNs onto the OHCs after OGD resulted in a significant exacerbation of neuronal damage compared with OGD-induced neuronal damage alone (Fig. 2B). Representative fluorescence images of the densitometric quantification (Fig. 2B) are shown in Figure 2C. These data suggest that PMNs can aggravate the outcome of ischemic neurological insults after 24 h. For all the following experiments involving direct application of PMNs onto OHCs, \( \frac{1}{10^5} \) PMNs were used.

PMNs migrate rapidly into hippocampal slices independent of OGD-induced neuronal damage

We had previously shown that microglia can migrate deeply into OGD-damaged OHCs to provide neuroprotection (Neumann et al., 2006). Thus, we speculated that also PMNs might immigrate into OHCs, albeit with a neurotoxic effect after OGD. To test this assumption directly, we subjected cocultures of OGD-treated OHCs and PMNs to two-photon microscopy as previously described (Neumann et al., 2006). In this model, OHCs are made from mice that express EYFP under a specific Thy-1 promoter that leads to strong expression of the transgene in hippocampal neurons (Feng et al., 2000). We observed the rapid migration of PMNs into the brain tissue after 1 h under basal conditions. However, the PMN immigration rate into OGD-damaged slices was indistinguishable from that observed under control conditions (Fig. 3A,B). Also, at later time points, we detected a uniform distribution of PMNs within the slice and no differences between OGD conditions and basal conditions were observed (Fig. 3A–F). Additionally, we studied the morphology of EYFP-positive neurons within the OHCs after PMN application. The application of PMNs onto hippocampal slices without OGD showed morphologically intact neurons (neuronal body, dendrites, axons) at all analyzed time points (Fig. 3A,C,E). In contrast, OGD slices containing PMNs were characterized by a severe loss of axons and dendrites after 6 h (Fig. 3D) and an almost complete loss of EYFP-positive neurons associated with the appearance of subcellular granular material, presumably stemming...
from neuronal apoptosis and necrosis, in the CA1 area after 24 h (Fig. 3F). We also noted the appearance of holes in the otherwise homogenous layer of intact neuronal somata (Fig. 3F, arrowheads), which probably showed the loss of cell bodies in the respective areas. Thus, although PMNs have the ability to invade OHCs under normal conditions, they do not cause neuronal damage, unless this process is initiated by OGD.

**Exogenous microglia counteract the neurotoxicity of PMNs**

We previously showed that microglia can have a neuroprotective function in the OGD-induced neuronal damage within OHCs (Neumann et al., 2006). We next asked whether this might also hold true for the PMN-induced neurotoxicity. The direct application of PMNs (Fig. 4A), microglia, or macrophages (RAW264.7) to control OHCs had no effect on neuronal viability in the CA area after 48 h (Fig. 4B, C). However, although the presence of PMNs in OGD-treated OHCs showed a strong exacerbation of OGD-induced neuronal damage after 24 and 48 h (Fig. 4B, C), microglia application resulted in a significant reduction of OGD-induced neuronal damage after 48 h but not after 24 h (Fig. 4B, C). No significant effect on neuronal death was observed after application of macrophages (Fig. 4B, C). We next wanted to analyze whether microglia were also able to counteract the massive neurotoxicity induced by PMNs. Therefore, we applied PMNs (1 × 10^6) in combination with microglia or macrophages (0.8 × 10^5 each) directly onto the OHCs (Fig. 4D). The combined application of PMNs and microglia resulted in a significant reduction of PMN-caused exacerbation of neuronal damage after OGD (Fig. 4E). In contrast, no significant effect was detected by simultaneous application of PMNs and macrophages. These data suggested that there might be a direct interaction between microglia and PMNs, which significantly attenuated the deleterious effects of PMNs on neuronal survival.
Microglia engulf PMNs within the OHCs

Based on these findings, we investigated a possible direct microglia–PMN interaction in OGD-damaged OHCs. Therefore, we applied fluorescently labeled PMNs and microglia onto the OHCs. Confocal microscopy analyses revealed that exogenous microglia had engulfed PMNs in OGD-treated OHCs (Fig. 5A). By means of two-photon microscopy and three-dimensional reconstruction of individual microglia, we found that endogenous as well as newly applied microglia were able to fully incorporate single or multiple PMNs within the OHCs (Fig. 5B). To investigate the cellular dynamics of such an engulfing process in more detail, we applied fluorescently labeled PMNs and microglia onto OHCs and recorded their migratory behavior by time-lapse microscopy. Surprisingly, we observed that, in addition to inactive (immotile and presumably apoptotic) PMNs, microglia were able to also engulf active (motile and viable) PMNs (Fig. 5C; supplemental movie 1, available at www.jneurosci.org as supplemental material).

Microglia engulf viable, motile, nonapoptotic PMNs

The previous experiments suggested that the microglia was able to engulf fully viable PMNs. However, although the phagocytosis of apoptotic cellular material by microglia is a relatively common process (Stolzing and Grune, 2004; Takahashi et al., 2005; Chan et al., 2006), the capture of active and viable cells by microglia has not been observed before. Phototoxicity and bleaching prevented us from performing hour-long time-lapse sequences on individual OHCs. However, in several instances, we were able to document the uptake of live PMNs by parenchymal microglia within the brain slices in a manner very similar to the one observed before for isolated microglia (supplemental movie 6, available at www.jneurosci.org as supplemental material), which suggests that the behavior of the externally added microglia was similar to the parenchymal cells. The related low frequency of phagocytosis events in individual image sequences did not allow performing a thorough quantitative analysis of PMN phagocytosis in this ex vivo model system. We therefore performed cell culture experiments in vitro. PMNs (3 × 10⁵) were cocultured with primary microglia (0.75 × 10⁵) in vitro. These experiments allowed to clearly visualize the engulfing process of motile PMNs by individual microglia (Fig. 6A, B) and demonstrated that before being engulfed and phagocytosed PMNs could exhibit profound motility over long time periods (Fig. 6A, A*).

We frequently observed that microglia adopted a “chasing behavior” while attempting to engulf PMNs, either including cellular protrusions (supplemental movie 2, available at www.jneurosci.org as supplemental material) or the whole cell bodies (Fig. 6B; supplemental movie 3, available at www.jneurosci.org as supplemental material). Because also immotile (presumably apoptotic or preapoptotic) PMNs were engulfed by microglia...
that indeed the microglia was activated by the presence of PMNs or the supernatant of OGD-treated OHC. Thus, it was possible that the observed phagocytosis behavior was a general phenomenon of activated macrophage-like cells and not specific for microglia. To test this assumption, we compared the ability of primary microglia with peritoneal macrophages freshly isolated from rats by peritoneal lavage with PBS. Analysis showed that despite being motile and frequently touching or dragging PMNs in the culture peritoneal macrophages never engulfed PMNs, whether apoptotic or alive ( supplemental movie 7, available at www.jneurosci.org as supplemental material), whereas microglia cocultured in the same experiment phagocytosed on average 2.6 PMNs per cell ( supplemental Fig. 2 and movie 8, available at www.jneurosci.org as supplemental material). Thus, the ability to phagocytose live or dead PMNs seemed to be specific for microglia activated by the presence of PMNs or the supernatant of OGD-treated OHC and was not observed with peripheral macrophages, at least from the peritoneum.

**Blocking the engulfment process of PMNs by microglia worsens the outcome of neuronal viability after OGD**

Next, we investigated whether the engulfment of PMNs by microglia had any consequence on neuronal viability after OGD. It has been shown that apoptotic PMN cells can be internalized by binding to the \( \alpha_\beta_1 \)-integrin receptor on macrophages. In addition, lectin-like receptors have been shown to be involved in this process (Fadok et al., 1998; Meszaros et al., 1999). Thus, we evaluated the potential of the integrin-blocking tetrapeptide RGDS (Arg-Gly-Asp-Ser) and the lectin inhibitor N-acetyl glucosamine (GlcNAc) to block the engulfment process by preincubating primary microglia with the reagents before adding PMNs to the culture. We distinguished between engulfment of motile or immotile PMNs by time-lapse microscopy. RGDS and GlcNAc blocked the engulfing of both immotile and, interestingly, also motile PMNs, highly significantly (Fig. 8A). Thereby GlcNAc was more efficient than RGDS (Fig. 8A). We also noted a slight synergistic effect on the blockade of the engulfment process when both substances were combined. This synergistic effect was more pronounced in the engulfment of nonmotile cells (Fig. 8A). Additionally, we always observed altered microglia–PMN interaction patterns in the presence of RGDS and GlcNAc. Whereas untreated microglia bound and engulfed PMNs, treated microglia bound several PMNs but mostly failed to ingest them (Fig. 8B).

An important question was whether interfering with the engulfment process affected the neuronal viability after OGD. To test this, we applied PMNs and microglia simultaneously onto the OHCs after OGD. Microglia and OHCs were also preincubated with RGDS/GlcNAc before application. Indeed, the presence of

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**Figure 5.** Microglia phagocytose PMNs within the OHCs. PMNs (1 \( \times \) 10^5) were labeled with CMFDA (green) (A, B) or CMAC (blue) (C) and microglia with CMTMR (red) and then directly applied onto OHCs. A, OHC was fixed with 4% PFA and subsequently investigated using confocal microscopy. B, The images from a living slice show the three-dimensional reconstruction of one microglia that had already phagocytosed two PMNs (arrows) and was in progress to phagocytose another PMN (asterisk). The dotted line indicates the contact between microglia and PMNs. C, Time-lapse video microscopy was performed 4 h after experimental onset. Images show the engulfing of a motile PMN by the microglia. The white line indicates the migration pathway of the PMN. The microglia contacted the PMN at the time (3 min after start of time-lapse imaging) at which the PMN showed a velocity of 7.6 \( \mu \)m/min. The white dotted line shows the contact point between microglia and PMN (supplemental movie 1, available at www.jneurosci.org as supplemental material). Scale bars: A, 20 \( \mu \)m; B, 5 \( \mu \)m; C, 10 \( \mu \)m. (Fig. 6B; supplemental movie 4, available at www.jneurosci.org as supplemental material), we next addressed the question of whether engulfed PMNs always exhibited signs of proapoptosis. Therefore, we transferred PMNs to primary microglia and added FITC-labeled Annexin V to the coculture. Annexin V binds phosphatidylserine on the outer side of the membrane of cells that have already initiated the apoptotic cascade (Fadok et al., 1998; Meszaros et al., 1999). Thus, we evaluated the potential of the integrin-blocking tetrapeptide RGDS (Arg-Gly-Asp-Ser) and the lectin inhibitor N-acetyl glucosamine (GlcNAc) to block the engulfment process by preincubating primary microglia with the reagents before adding PMNs to the culture. We distinguished between engulfment of motile or immotile PMNs by time-lapse microscopy. RGDS and GlcNAc blocked the engulfing of both immotile and, interestingly, also motile PMNs, highly significantly (Fig. 8A). Thereby GlcNAc was more efficient than RGDS (Fig. 8A). We also noted a slight synergistic effect on the blockade of the engulfment process when both substances were combined. This synergistic effect was more pronounced in the engulfment of nonmotile cells (Fig. 8A). Additionally, we always observed altered microglia–PMN interaction patterns in the presence of RGDS and GlcNAc. Whereas untreated microglia bound and engulfed PMNs, treated microglia bound several PMNs but mostly failed to ingest them (Fig. 8B).

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RGDS or GlcNAc alone, and, more efficiently, in combination strongly reduced the neuroprotective function of coapplied microglia in this model (Fig. 8C). In addition, we obtained similar results by using primary rat PMNs, confirming that the observed behavior was specific for PMNs (Fig. 8C).

Thus, blocking the engulfment of PMNs by microglia severely compromises the neuroprotective function that microglia exerted on neurons on OGD exposure pointing to a physiological role of this cellular function of microglia.

**Discussion**

This study was designed to determine the role of individual cell types of the innate immune system that contribute to the postischemic inflammation after cerebral ischemia. Taking advantage of our neuroinflammation model, we were able to simulate the migration and infiltration of these cells to the site of damage in the neuronal tissue. It is generally accepted that the main immune cells involved in the inflammation-induced secondary neuronal damage are PMNs, microglia, and macrophages, which all are recruited as early as the postischemic inflammation is initiated. However, PMNs and local microglia are the first cells present on site, followed by peripheral microglia and monocytes/macrophages. Although all three cell types potentially exhibit cytotoxicity by releasing noxious substances such as cytokines, oxygen radicals, and proteases (Hallenbeck et al., 1986; Barone et al., 1991; Minghetti and Levi, 1998), their individual contribution for the overall damage remains unclear.

There is still an ongoing debate as to the importance of the individual cell types in brain ischemia. In an attempt to overcome this limitation, we chose to use a well established ex vivo model of neuroinflammation (Ullrich et al., 2001; Mitrasinovic et al., 2005; Neumann et al., 2006). Using this model, we were now able to demonstrate that indeed only PMNs sharply increased neuronal damage associated with transient ischemia. This was not a general phenomenon of infiltrating phagocytes, because neither local nor externally added microglia nor macrophages appeared to be neurotoxic after cerebral ischemia. In fact, we and others previously showed that local (Kohl et al., 2003) or externally added (Neumann et al., 2006) microglia may even be neuroprotective in this model (Fig. 8C). In addition, we obtained similar results by using primary rat PMNs, confirming that the observed behavior was specific for PMNs (Fig. 8C).

Interestingly, the inhibition of reactive oxygen species (ROS) production by NAD(P)H-oxidases, one of the key actions of inflammatory PMNs (Segal, 2005), results in strong neuroprotective effects in rodent models of ischemic stroke (Wang et al., 2006; L. L. Tang et al., 2007). Moreover, interfering with the homing of immune cells by an antiadhesive therapy has been shown to be neuroprotective (Connolly et al., 1996; Yanaka et al., 1996). However, because this approach affects indiscriminately all immune cell types, it did not provide any suggestions as to whether PMNs might play a specific role in this context. In addition, it is assumed that also cells of the adaptive immune system can invade areas of ischemia-induced neuroinflammation (Arumugam et al., 2005). Because their contribution is still unclear, our model is well suited to examine this matter in the future. In summary, in the line of these previous data and our study, it seems likely that PMNs constitute an important neurotoxic cell type under ischemic conditions. Because we did not find evidence that microglia and macrophages were neurotoxic, at least 48 h after ischemia, we assume that interfering with the function of PMNs might be a particularly promising option to limit the extent of neuronal damage after stroke.

A major conclusion from our study is that there seems to be a natural mechanism aiming precisely at this goal. After coappling microglia and PMNs onto ischemically damaged OHCs, we found a strong decrease of neuronal damage compared with the application of PMNs alone. This was a rather unexpected finding, because numerous studies demonstrated that microglia preacti-
activated by inflammatory mediators are strongly neurotoxic (Arai et al., 2004; Minghetti et al., 2004; Qin et al., 2004; Fordyce et al., 2005; Huang et al., 2005). However, a close inspection of these studies shows that microglial activation was achieved by exposure to microbial products (Arai et al., 2004; Qin et al., 2004; Huang et al., 2005) or viral particles (Lim et al., 2003; Minghetti et al., 2004). Lipopolysaccharide (LPS) or viral particles are well known to trigger a massive immune cell reaction via Toll-like receptors (TLRs) (Kawai and Akira, 2006). Recent evidence suggests that, even in the situation of stroke, which constitutes a sterile inflammation, Toll-like mediated pathways can be triggered on microglia (Lehnardt et al., 2007) or even directly on neurons (S. C. Tang et al., 2007) and that this response increases neuronal damage after stroke. However, the signaling induced by bacterial TLR triggers might well be different from the one induced by endogenous inflammatory mediators because neurons, which express both TLR-2 and -4 do not respond to classical triggers of these receptors such as peptidoglycan or LPS (S. C. Tang et al., 2007). It might, therefore, be possible that also microglia are able to distinguish between microbial and other inflammatory stimuli of the TLR system. In this concept, microglia activated by nonmicrobial inflammation might exert neuroprotective effects, whereas microbial activation of microglia preferentially triggers neurotoxicity.

Our study also provides insights into the mechanism whereby this protection is accomplished. We demonstrate that microglia were very effective in phagocytosing invading PMNs. Macrophage-like cells are well known to clear the inflamed tissue from cell debris, such as apoptotic PMNs. Two recent histological studies have also demonstrated brain-resident microglia associated with PMN-related debris in areas of ischemia-induced neuronal damage (Denes et al., 2007; Weston et al., 2007). However, our data show that, in addition to proapoptotic PMNs, also fully viable, Annexin V-negative, PMNs were effectively engulfed by microglia.

In contrast, we have never observed microglia engulfing other types of immune cells such as freshly prepared lymphocytes and monocytes, and in addition we did not find engulfing of PMNs by macrophage lines or freshly prepared peritoneal macrophages. These findings underscore the specificity of the microglia–PMN engagement. By time-lapse microscopy, we observed microglia exerting a PMN chasing behavior, either including cellular protrusions or the whole cell bodies. To the best of our knowledge, the phagocytosis of active and viable immune cells by other immune cells has not been described so far and therefore might represent an entirely novel way of immune control. This is probably attributable to the fact that such a mechanism can only be revealed by life cell imaging, which is technically challenging, especially with the cell types investigated in the current study.

Given the fact that phagocytosis was inhibitable by RGDS peptides and GlcNAc, we assumed that integrins such as vitronectin receptors (Ruoslhlahti, 1996) or lectins mediated the engulfment process. Although this is well known for the uptake of apoptotic cells (Fadok et al., 1998; Meszaros et al., 1999), this surprisingly also held true for the uptake of presumably live cells. We cannot formally exclude that even Annexin V-negative cells were already proapoptotic thereby exposing vitronectin or lectin ligands. However, the fact that the cells were very motile argued against this hypothesis, because one of the earliest events of apoptosis is loss of migration (Savill et al., 2002).
Our study also indicates that interference with the phagocytosis process by RGDS and GlcNAc abolished the neuroprotective function of microglia. It is well conceivable that dying PMNs mediate neurotoxicity by the release of toxic intracellular compounds (Denes et al., 2007; Weston et al., 2007), and that, consequently, prompt phagocytosis of apoptotic PMNs by microglia might prevent the secretion of toxic compounds. In the light of these findings, we propose that clearance of PMNs from the nervous tissue might be an effective strategy to protect neurons from PMN neurotoxicity. Whether the engulfment of viable PMNs contributes to this protection remains to be determined and requires to be examined in additional studies. In our system, phagocytosis of motile (presumably viable) PMNs was even more frequent than the uptake of nonmotile PMNs was added to the microglia. Time-lapse microscopy was performed to distinguish the engulfment of immotile and motile PMNs by the microglia. Column scatterplots show the number of PMNs engulfed by microglia under the indicated conditions. Significance values are shown above the columns. The images show representative microglia that were exposed to PMNs. The arrows on the top image display the engulfed PMN within the microglia. The bottom image shows PMNs that adhere to the microglia without being internalized. Microglia were preincubated with RGDS (1 mM) or GlcNAc (20 mM) or both for 20 min before they were applied together with either human-derived (hPMN) or rat-derived (rPMN) PMNs onto OGD-treated OHCs. Quantification of neuronal death in CA1–3 was performed by PI incorporation after 24 h (***p < 0.001, RPMN plus MIC vs RPMN plus MIC plus RGDS/GlcNAc (OGD); **p < 0.001, RPMN plus MIC (OGC) vs RPMN plus MIC plus RGDS/GlcNAc (OGD); n = 5–7/bar]. Error bars indicate SEM. CTRL, Control; MIC, microglia.

**Figure 8.** Blockage of the engulfment process of PMNs by microglia worsens the outcome of neuronal viability after OGD. A, Microglia were preincubated with RGDS (1 mM) or GlcNAc (20 mM) or both for 20 min before PMNs were added to the microglia. Time-lapse microscopy was performed to distinguish the engulfment of immotile and motile PMNs by the microglia. Column scatterplots show the number of PMNs engulfed by microglia under the indicated conditions. Significance values are shown above the columns. The images show representative microglia that were exposed to PMNs. The arrows on the top image display the engulfed PMN within the microglia. The bottom image shows PMNs that adhere to the microglia without being internalized. B, Microglia were preincubated with RGDS (1 mM) and GlcNAc (20 mM) or both for 20 min before they were applied together with either human-derived (hPMN) or rat-derived (rPMN) PMNs onto OGD-treated OHCs. Quantification of neuronal death in CA1–3 was determined by PI incorporation after 24 h (***p < 0.001, hPMN plus MIC (OGD) vs hPMN plus MIC plus RGDS/GlcNAc (OGD); **p < 0.001, rPMN plus MIC (OGC) vs rPMN plus MIC plus RGDS/GlcNAc (OGD); n = 5–7/bar]. Error bars indicate SEM. CTRL, Control; MIC, microglia.

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**References**


