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Central Nervous System Stromal Cells Control Local CD8+ T Cell Responses during Virus-Induced Neuroinflammation

Graphical Abstract

Highlights
- CNS stromal cells swiftly generate CCR7 ligands during neurotropic virus infection
- CCR7-expressing antiviral CD8+ T cells prevent lethal CNS disease
- Stromal cell-derived CCR7 ligands guide CD8+ T cells to infected target cells

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In Brief
Whether CNS stromal cells regulate protective T cell immunity is unclear. Ludewig and colleagues demonstrate that local chemokine production by activated endothelial and fibroblastic stromal cells is required to transfer protective CD8+ T cell immunity from the draining lymph node to virus-infected areas of the central nervous system.
Central Nervous System Stromal Cells Control Local CD8+ T Cell Responses during Virus-Induced Neuroinflammation

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SUMMARY

Stromal cells generate a complex cellular scaffold that provides specialized microenvironments for lymphocyte activation in secondary lymphoid organs. Here, we assessed whether local activation of stromal cells in the central nervous system (CNS) is mandatory to transfer immune recognition from secondary lymphoid organs into the infected tissue. We report that neurotropic virus infection in mice triggered the establishment of such stromal cell niches in the CNS. CNS stromal cell activation was dominated by a rapid and vigorous production of CC-motif chemokine receptor 7 (CCR7) ligands CCL19 and CCL21 (Randall et al., 2008; Förster et al., 2008). Podoplanin (PDPN)- and extracellular matrix protein ER-c�7-expressing FRCs represent a major lymph node (LN) stromal cell population involved in the production of CCR7 ligands (Link et al., 2007; Scandella et al., 2008). In addition, BECs in HEVs and lymphatic endothelial cells (LECs) contribute to production of CCL21 (Gunn et al., 1998). Moreover, CCR7 ligands can be generated in peripheral, non-lymphoid organs (Christopherson et al., 2003; Lo et al., 2003; Krumbholz et al., 2007; Columbus-Cabezas et al., 2003; Kivisakk et al., 2004) where their expression further increases during inflammatory reactions (Pashenkov et al., 2003; Krumbholz et al., 2007; Rangel-Moreno et al., 2007; Burke et al., 2010). These data suggest that CCR7 ligands expressed in non-lymphoid organs might serve as inducible chemoattractants to foster immune surveillance.

Tight control of lymphocyte entry and effector function is particularly important for organs such as the CNS because immunopathological damage of this organ during active immune responses has to be kept at a minimum (Ransohoff and Engelhardt, 2012; Moseman and McGavern, 2013). The finding that T cells migrating into the inflamed CNS express CCR7 (Alt et al., 2002) suggests that T cell entry to the CNS might be controlled via this molecule. Moreover, chronic inflammatory diseases of the CNS such as multiple sclerosis (MS) are associated with elevated CCL19 levels in the cerebrospinal fluid (Pashenkov et al., 2003; Krumbholz et al., 2007) and significantly higher expression in active and inactive lesions (Krumbholz et al., 2007). In an experimental model of MS, both CCL19 and CCL21 can be detected in ECs of inflamed venules and cells that accumulate in the perivascular space (Alt et al., 2002; Columbus-Cabezas et al., 2003). However, it has remained elusive whether local activation of fibroblastic and/or endothelial stromal cells in the CNS is required to transfer immune recognition from SLOs into the infected tissue and to what extent these processes control local immune responsiveness during virus-induced neuroinflammatory CNS disease.

To address these questions, we utilized intranasal infection with a neurotropic coronavirus that leads to a transient encephalitis in immunocompetent animals but causes chronic demyelinating and lethal disease when the host is immunodeficient...
Here, we focused on the events that take place early during establishment of the infection when viral replication has caused a rapid and more than 1,000-fold upregulation of Ccl19 and Ccl21 gene expression in the olfactory bulb. We found that two distinct stromal cell populations, i.e., ECs of meningeal and sub-meningeal venules and PDPN+/ER-TR7+ FRC-like cells in the perivascular space, served as major source for CCR7 ligands. CCR7 expression on antiviral T cells was mandatory to prevent lethal neuroinflammatory disease, while provision of extra-lymphatic CCL21 ensured sufficient CD8+ T cell recruitment to and re-activation in the CNS to protect the host from severe disease. We therefore conclude that activation of CNS stromal cells is critical for optimal control of neurotropic viral infection and that this function is executed—at least partially—through the swift provision of CCR7 ligands.

RESULTS

Rapid Production of CCR7 Ligands during Neurotropic Virus Infection

Viral infection of the CNS triggers activation of several inflammatory cascades including the generation of chemokines (Hosking and Lane, 2010). To assess whether infection of mice with neurotropic viruses triggers a distinct program of stromal cell activation including expression of chemokines, we infected mice with the mouse hepatitis virus (MHV) strain A59 via the intranasal route. As shown in Figure 1A, infectious viral particles were first detectable in olfactory bulbs and cervical LNs (cLN) on day 2 post infection (p.i.). By day 4, viral titers had reached peak values in olfactory bulbs and the virus had spread to distal CNS areas, including the spinal cord. After 6 days, the virus had been eliminated from draining cLNs, while clearance from CNS tissues was not achieved until day 10 (Figure 1A). In order to visualize viral dissemination at the peak of the infection and to identify infected cells, we analyzed paramedian sagittal sections of infected brains (day 6 p.i.) by confocal laser scanning microscopy using antibodies against the MHV nucleoprotein (MHV-N), activated microglial cells (ionized calcium-binding adaptor molecule 1/Iba-1) and neurons (neuronal nuclei, NeuN) (Figure 1B, i). Strong viral antigen expression was detectable in rostral regions of the brain such as the olfactory bulb, anterior olfactory nucleus, and the piriform cortex (Figure 1B, i). Moreover, infected NeuN+ neurons were detected in the anterior olfactory nucleus (Figure 1B, iii) and aspartoacylase (ASPA)-expressing (data not shown) oligodendrocytes staining positive for MHV-N were found in the locus coeruleus. The finding that both neurons and glial cells stained positive for MHV-N supports the notion that the virus utilizes axonal transport (Perlman et al., 1990) and cell-to-cell spread (Gallagher and Buchmeier, 2001) as means of dissemination. To assess whether the viral infection had triggered the generation of
microenvironments that support lymphocyte activation, we performed quantitative RT-PCR analysis of olfactory bulb tissue at the peak of the infection. Interestingly, mRNA expression of inflammatory chemokines and several genes involved in lymphoid organogenesis such as Ltb, Ltbr, and II7 were substantially upregulated in the infected olfactory bulb (Figure 1C). Likewise, expression of the homeostatic chemokines Ccl19, Ccl21, and Cxcl13 increased substantially in inflamed CNS tissue compared to naive controls (Figure 1C). Since the increase in relative expression of CCR7 ligands was most prominent in olfactory bulb and cLN, we reasoned that processes driving production of these chemokines in the cLN were mirrored during local inflammatory reactions in the CNS. As shown in Figures 1D and 1E, expression of both Ccl19 and Ccl21 mRNA increased roughly 1,000-fold at their peak on day 20 in the olfactory bulb and the pattern of chemokine activation in different CNS areas was accompanied by the infiltration of inflammatory cells (Figures S1B and S1C). Elevated chemokine expression was only transient and followed the resolution of the viral infection with substantially lower expression on day 20 post infection (Figures 1D and 1E). Thus, the presence of actively replicating virus within the CNS, particularly in the olfactory bulb as virus entry point, drives a rapid increase in transcriptional activity of Ccl19/Ccl21 genes, a process that is paralleled by infiltration of immune cells and upregulation of genes associated with stromal cell activation.

Activated CNS Stromal Cells Produce CCL19 and CCL21
While CCL21 decorates extracellular matrix proteins, CCL19 is rapidly internalized and degraded following binding to CCR7 (Comerford et al., 2013). Hence, to localize CCL19-producing cells in situ, we took advantage of the recently described Ccl19-cre bacterial artificial chromosome transgenic mouse that facilitates expression of the Cre recombinase in FRCs of secondary lymphoid organs (Chai et al., 2013). We crossed Ccl19-cre mice with Rosa26-eyfp mice (referred to as Ccl19<sup>cre</sup>) to label cells exhibiting Ccl19 promoter activity with the yellow fluorescent protein. Combination of this detection method with anti-CCL21 immunostaining in confocal microscopic analysis of olfactory bulb tissue of naive Ccl19<sup>cre</sup> mice revealed only faint CCL21 staining and few cells that exhibited Ccl19 promoter activity (Figure S2A). In contrast, cells showing Ccl19-cre activity and anti-CCL21 staining were clearly detectable at day 6 post infection and were mainly confined to meningeal and sub-meningeal areas (Figures 2A and 2B, magnified in boxed areas). Nevertheless, PDPN expression, which serves as a marker for FRC activation (Chai et al., 2013), was strongly increased in regions of elevated CCR7 ligand production (Figure 2B). While mRNA expression analysis revealed rather strong induction of Cxcl13 transcript levels (Figure 1C), expression of CXCL13 protein was found only in very few cells (Figures S2B and S2C).

High resolution microscopic analysis of day 6 olfactory bulb tissue revealed considerable EYFP expression in inflamed ECs of meningeal blood vessels (Figure 2C, arrows). Moreover, PDPN<sup>ER-TR7</sup> fibroblastic cells lined the perivascular space around meningeal blood vessels and in the subarachnoid space that surrounds blood vessels protruding into the CNS parenchyma (also known as Robin-Virchow space; Ransohoff and Engelhardt, 2012) (Figure 2C, arrowheads). Likewise, CD31<sup>+</sup> ECs of inflamed blood vessels in the olfactory bulb parenchyma (Figure 2D, arrows) and fibroblastic cells lining the perivascular space (Figure 2D, arrowheads) displayed Cc19 promoter-driven EYFP activity and CCL21 expression. Stromal cell activation with expression of CCL21 was most pronounced in the vicinity of virus-infected cells (Figure S2D). In addition, ECs of meningeal blood vessels (Figure 2E, arrows) and perivascular EYP<sup>+</sup> fibroblastic cells (Figure 2F, arrowheads) had upregulated expression of the intercellular adhesion molecule 1 (ICAM1, CD54), whereas only BECs showed weak ICAM1 expression in the absence of CNS inflammation (Figure 2E, inset). The numbers of EYFP-expressing cells in infected olfactory bulbs increased almost 10-fold within the first 4 days of the infection (Figure S2E) with a substantial expansion of both PDPN<sup>+</sup> fibroblastic and CD31<sup>+</sup> endothelial stromal cells (Figure S2F). Moreover, we found that only very few of the EYP<sup>+</sup> stromal cells expressed the proliferation marker Ki67 (data not shown), indicating that the increase in EYFP<sup>+</sup> stromal cells was a result of their activation with enhanced expression of homeostatic chemokines.

Further flow cytometric analysis of CNS stromal cells (Figures 3A–3C) confirmed that both stromal cell populations responded to the infection with substantial upregulation of ICAM1, vascular cell adhesion molecule 1 (VCAM1, CD106) and MHC class I (Figures 3D and 3E). Moreover, fibroblastic stromal cells from infected CNS tissue expressed substantially less CD44 molecules, while expression of markers such as CD140a and CD140b was not affected (Figures 3D and 3E). CCL21 expression was detectable in both stromal cell populations during the infection (Figure 3F), while other CNS cells such as NeuN<sup>+</sup> neurons, CD11b<sup>+</sup> macrophages and/or microglia, ASPA<sup>+</sup> oligodendrocytes, and GFAP<sup>+</sup> astrocytes did not express CCL21 at day 6 post infection (Figure S3). These data indicate that two distinct stromal cell populations serve as the main source of CCR7 ligands at hotspots of neurotropic virus replication in the infected CNS.

Lethal Neuroinflammatory CNS Disease in the Absence of CCR7
The vigorous but locally restricted activation of stromal cells with dominant production of CCL19 and CCL21 during the response to neurotropic coronavirus infection suggested that these chemokines could contribute to the control of the virus. To test this hypothesis, we assessed whether mice lacking either CCR7 or CCR7 ligand expression in SLOs display differences in their defense against neurotropic virus infection. The latter mouse strain, known as plt/plt (paucity of lymph node T cells) (Gunn et al., 1999) shows expression of the CCL21 isoform in extra-lymphatic tissues (Nakano and Gunn, 2001). Notably, both mouse strains exhibit impaired T cell zone development in SLOs due to migration defects of DCs and lymphocytes (Gunn et al., 1999; Förster et al., 1999), hence showing a comparable loss of immunocompetence in response to systemic virus infection (Junt et al., 2002; Junt et al., 2004). While Ccr7-deficient mice successfully controlled the infection and experienced transient weight loss during the first 2 weeks, Ccr7-deficient mice presented with rapid weight loss early after infection, succumbing to neuroinflammatory disease by day 10 (Figure 4A). In sharp contrast, all plt/plt mice survived the phase of transient weight loss (Figure 4A) and successfully controlled the infection as...
shown by significantly lower viral titers in brain tissues compared to Ccr7−/− mice (Figure 4B). Impaired viral control of Ccr7-deficient and plt/plt mice was associated with a pronounced reduction in antiviral CD8+ T cell numbers (Figure 4C) and function (Figure 4D), as well as a reduction in antiviral CD4+ T cell function (Figure 4E) in the whole brain on day 10 p.i. Moreover, recruitment of CD8+ T cells into the olfactory bulb was affected already on day 4 by the Ccr7-deficiency (Figure S4A). The lack of CCR7 almost completely abolished the appearance of interferon (IFN)-γ-producing antiviral CD8+ T cells in all CNS regions (Figures S4B and S4C), while CD4+ T cell recruitment was affected to a lesser extent (Figures S4D and S4E). Importantly, Ccr7-deficient mice showed a stronger impairment of antiviral CD8+ T cell responses compared to plt/plt mice (Figures 4C and 4D). The lack of chemokine receptor CXCR3 that binds the inflammatory chemokines CXCL9 and CXCL10 and that has been implicated in the regulation of T cell migration in several models of neuroinflammation (Bagai et al., 2005), did not affect survival following intranasal MHV A59 infection (Figure S4F). Likewise, the absence of CXCR5, i.e., the receptor for the B cell zone cytokine CXCL13, did not increase the susceptibility of the animals to MHV CNS infection (Figure S4G). Therefore, it appears that the ability of T cells to sense extra-lymphatic expression of the chemokines CCL19 and CCL21 through CCR7 plays a major role in the rapid control of neurotropic coronavirus infection.

**Rescue from Lethal Neuroinflammatory Disease by CCR7-Expressing CD8+ T Cells**

Viral replication kinetics and the fast disease course suggested that appropriate activation of antiviral CD8+ T cells and elimination of the highly cytopathic pathogen from the CNS is of utmost importance to prevent lethal disease. To assess the activation and migration pattern of MHV-specific CD8+ T cells in vivo, we utilized a transgenic mouse expressing a H2-Kb-restricted TCR that recognizes the immunodominant epitope s598–605 of the MHV spike protein (TCR-S or Spiky). Adoptive transfer of congenic carboxy-fluorescein succimidyl ester (CFSE)-labeled Spiky cells into infected recipients revealed first activation in cLNs at 72 hr p.i. followed by strong proliferation (Figure 5A). Interestingly, accumulation of activated, CFSElow Spiky cells in the infected CNS could be detected on day 4 at the earliest
indicating that the cells had to undergo at least 8 rounds of proliferation in the cLN before they gained the ability to migrate to the CNS. As expected, activation of Spiky cells in cLNs resulted in downregulation of CCR7 (Figure 5C). Activated Spiky cells circulating in blood and those that had infiltrated the CNS expressed CCR7 (Figure 5C) indicating their ability to sense CCR7 ligands produced by ECs and FRC-like cells in inflamed regions of the CNS. CCR7 expression detected on CNS-infiltrating Spiky cells was comparable to polyclonal CD8+ T cells that had infiltrated different regions of the CNS (Figure S5). Moreover, Spiky cells derived from cLNs, blood and the brain on days 4 and 8 post infection had downregulated CD62L (Figure 5D). Functional characterization of CNS-infiltrating Spiky cells revealed that these cells were fully activated effector T cells as shown by secretion of the cytokines IFN-γ and tumor necrosis factor alpha (TNF-α), and expression of the degranulation marker CD107a (lysosomal-associated membrane protein 1, LAMP1) (Figures 5E and 5F). These data support the notion that access of polyfunctional effector CD8+ T cells to the CNS is important for successful elimination of the virus from this highly vulnerable tissue.

To investigate whether CCR7 expression on virus-specific CD8+ T cells is sufficient to secure control of neurotropic virus infection, we adoptively transferred Ccr7-proficient or -deficient mice. As expected, Ccr7-deficient mice that had received Ccr7-deficient CD3+ cells were not protected from MHV CNS infection (Figures 5G and 5H, open red circles), while Ccr7-proficient CD8+ T cells provided protection (Figures 5G and 5H, gray circles). Ccr7-deficient mice receiving transfer of Ccr7-proficient Spiky CD8+ T cells showed a mild disease course and survived the infection (Figures 5G and 5H, white circles) indicating that protection against this neurotropic viral infection depends on the ability of antiviral CD8+ T cells to respond to CCR7 ligands.

Stromal Cell-Derived CCR7 Ligands Promote Recruitment of CD8+ T Cells to and Their Re-activation in the CNS

To determine whether the effects of CCR7 ligands on CD8+ T cell performance was due to the production of CCL19 and CCL21 in the draining cLN or a consequence of the substantial generation of these chemokines in the CNS, we used transfer of Ccr7-proficient and -deficient TCR Spiky cells into plt/plt mice. These mice are well-suited to address this question because they lack CCL19 and CCL21 expression in lymphoid organs (Gunn et al., 1999; Nakano and Gunn, 2001) but show production of CCL21 by stromal cells in MHV-infected olfactory bulbs (Figure 6A). To assess early CD8+ T cell activation, we transferred Ccr7-proficient or -deficient TCR Spiky cells into plt/plt mice and compared their activation status in cLNs and the brain 4 days after infection. We found that expansion and activation of antiviral CD8+ cells in cLNs was not dependent on CCR7

Controls included adoptive transfer of sorted polyclonal CD3+ cells from Ccr7-proficient or -deficient mice. As expected, Ccr7-deficient mice that had received Ccr7-deficient CD3+ T cells were not protected from MHV CNS infection (Figures 5G and 5H, open red circles), while Ccr7-proficient CD3+ T cells provided protection (Figures 5G and 5H, gray circles). Ccr7-deficient mice receiving transfer of Ccr7-proficient Spiky CD8+ T cells showed a mild disease course and survived the infection (Figures 5G and 5H, white circles) indicating that protection against this neurotropic viral infection depends on the ability of antiviral CD8+ T cells to respond to CCR7 ligands.
Figure 4. High Susceptibility to Neurotropic Viral Infection in Ccr7-Deficient Mice

(A) Weight loss of Ccr7-deficient, plt/plt, and WT mice was recorded during the indicated time period following infection with MHV A59. Values indicate mean percentage of the initial weight ± SEM from two independent experiments (n = 8 mice per group).

(B) Viral titers in CNS tissues as determined at day 10 after infection. Data indicate means of log transformed values ± SEM from two independent experiments (n = 8 mice per group).

(C) Tetramer-binding and IFN-γ production of (D) CD8+ and (E) CD4+ T cells from the whole brain was determined by flow cytometry on day 10 after infection. Representative dot plots analysis with percentage of virus-specific T cell populations indicated. Graphs indicate mean percentage ± SEM of the respective virus-specific T cell populations from two independent experiments (n = 8 mice per group). Statistical analysis was performed using the Student’s t test (**p < 0.01; ***p < 0.001; n.s. not significant). See also Figure S4.

DISCUSSION

The CNS is considered an immunologically privileged site that becomes visible to immune effector cells under particular inflammatory conditions such as infection with neurotropic viruses (Galea et al., 2007; Moseman and McGavern, 2013). Because the CNS is vital and highly vulnerable, immune responses against neurotropic infectious agents need to be well-controlled, i.e., the efficacy of pathogen elimination has to be high, while immunopathological damage has to be kept minimal. The present study shows that activated antiviral CD8+ T cells utilize CCR7 to optimize migration into the inflamed CNS and to control their functional activity during neurotropic viral infection. Moreover, our findings indicate that CCR7 ligands not only support the recruitment of virus-specific CD8+ T cells to the CNS, but is also required for the execution of their antiviral activity.

High resolution histological analysis revealed that activated CD8+ T cells were in intimate contact with endothelial cells of inflamed meningeal blood vessels that had upregulated Ccl19 expression (Figures 7A and 7B, arrowheads). Moreover, CD8+ T cells were found to interact with CCL19+PDPN+ perivascular fibroblastic stromal cells (Figures 7A and 7B, arrows). Next, we specifically ablated CCL19-expressing stromal cells using low doses of diphtheria toxin (DT) applied to Ccl19+eyfp mice that express the DT receptor in Cre recombinase-positive cells (Ccl19+emx1cre) (Buch et al., 2005; Cremasco et al., 2014). Low dose DT treatment led to a substantial loss of EYFP+ cells in the infected CNS (Figure 7C), but left the FRC network in the cLNs intact (Figure 7C, inlets). Importantly, CNS-restricted ablation of EYFP+ stromal cells resulted in reduced CD8+ T cell accumulation (Figure 7D), while CD4+ T cell recruitment was not altered (Figure 7E). As a consequence, CNS viral titers were significantly elevated in DT-treated Ccl19+emx1cre mice on day 8 p.i. (Figure 7F). CD8+ and CD4+ T cell compartments in cLNs were not changed by the DT treatment (Figures 7G and 7H). Moreover, we found that the local stimulation of antiviral CD8+ T cells via CCR7 not only improved the ability of CD8+ T cells to exit meningeal blood vessels, but substantially fostered their access to infected areas in the CNS parenchyma (Figures 7I and 7J). Taken together, these data show that activated CNS stromal cells are potent regulators of local antiviral CD8+ T cell activity and that this interaction depends to a large extent on the CCR7-CCL7 ligand axis.
Figure 5. CCR7-Dependent Activation and Recruitment of Antiviral CD8+ T Cells Protects from Virus-Induced CNS Disease

Thy1.2+ WT mice received 10^6 TCR transgenic Thy 1.1+ Spiky cells labeled with the intracellular dye carboxy-fluorescein succimidyl ester (CFSE) and were subsequently infected i.n. with MHV. CFSE dilution of transferred Thy1.1+ CD8+ T cells from the cLN (A) and brain tissue (B) was determined by flow cytometry at the indicated time points post infection. Values in histograms indicate percentage of proliferated cells (black line) compared to naive controls (gray histogram); representative data from two independent experiments (n = 5 mice).

(C) CCR7 and (D) CD62L expression on Thy1.1+CD8+ T cells from cLN, blood and the brain was performed at the indicated time points post infection. Values in histograms indicate mean fluorescent intensity (MFI) values of the total Thy1.1+CD8+ T population (black lines) minus MFI of isotype controls (gray shaded lines) ± SEM. Data from two independent experiments (n = 5 mice).

Expression of effector cytokines TNF-α and IFN-γ and the degranulation marker Lamp-1 by CNS infiltrating Thy1.1+CD8+ T cells (day 8 p.i.) following in vitro peptide restimulation is shown in representative dot plots (E) and summarized in a bar graph (F) with mean percentage ± SEM pooled from two independent experiments (n = 5 mice).

(Ccr7− deficient mice were adoptively transferred with a total of 10^7 of CD3+ Ccr7+ proficient, CD3+ Ccr7− deficient, or 10^6 of transgenic Spiky cells and infected with MHV.

(G) Survival and (H) weight loss of the mice were recorded during the indicated period of time. Values in (H) indicate mean percentage of the initial weight ± SEM pooled from three independent experiments (n = 9 mice per group). See also Figure S5.
Figure 6. Expression of CCR7 Ligands in the CNS Controls CD8+ T Cell Recruitment and Function

(A) Intracellular CCL21 production by PDPN+CD31−/C0 cells from cLNs and olfactory bulbs of WT (black lines) and plt/plt (red lines) mice either infected with MHV (solid lines) or left naive (dashed lines). Data are representative of two independent experiments (n = 4 mice per group).

(B and C) 10⁵ TCR Spiky Ccr7-proficient or Ccr7-deficient CD8+ T cells were adoptively transferred into plt/plt mice 12 hr before infection. Absolute numbers of Ccr7-proficient and Ccr7-deficient Spiky transgenic cells in (B) cLNs and (C) brain were recorded on day 4 post infection. Representative contour plots of T cell activation markers CD44 and CD62L expressed by Ccr7-proficient (black) and Ccr7-deficient (red) Spiky cells compared to LN-derived naive control cells (gray contour). Values in the plots represent percentage of activated cells (i.e., CD44high or CD62Llow, respectively). At day 6 post infection, absolute numbers of Ccr7-proficient and Ccr7-deficient Spiky transgenic cells in (D) brain and (E) cLNs were determined. Frequencies of brain infiltrating Spiky cells expressing markers of (F) proliferation (Ki67) and (G) apoptosis (active caspase 3) were analyzed using flow cytometry day 6 after the infection. Data represent mean value ± SEM from two independent experiments (n = 5–6 mice per group).
function as part of a multi-layered control network that secures spatiotemporal focusing of the immune response to areas of viral replication. Production of CCL19 and CCL21 by stromal cells in LNs and other SLOs is essential to recruit lymphocytes to these sites of immune activation and to induce adaptive immune responses (Forster et al., 2008). CNS infection with a neurotropic coronavirus, as shown here, triggered a moderate and transient increase in Ccl19 and Ccl21 gene expression in cLNs. It is possible that virus and viral products have been drained via lymphatic vessels that connect the dura to the cLNs (Aspelund et al., 2015). Within the CNS, both ECs and perivascular FRC-like cells showed swift and substantial production of CCR7 ligands. The expression of CCR7 ligands can be induced via the ligation of the LTβR in fibroblastic (Chai et al., 2013) and in endothelial stromal cells (Onder et al., 2013). Moreover, acute inflammatory mediators such as TNF can foster activation of the CCR7/CCL19 axis (Bose et al., 2013). It is likely that the initial upregulation of Ccl19 and Ccl21 mRNA expression in MHV infected CNS is triggered via inflammatory cytokines such as TNF. Notably, microglial cells are a prominent source of TNF during viral encephalitis (Ghoshal et al., 2007). Thus, infection of

Figure 7. Activated CNS Stromal Cells Guide Antiviral CD8+ T Cells to the CNS Parenchyma

(A and B) C57BL/6 (A) and Ccl19<sup>pyfp</sup> (B) were infected with MHV intranasally and brains were harvested for histological analysis on day 6 post infection. Meningeal and submeningeal areas were inspected for interactions of CD8<sup>+</sup> T cells with endothelial cells (arrows) and perivascular fibroblasts (arrows). Scale bar represents 10 μm in (A) and 20 μm in (B).

(C) Ccl19<sup>pyfp/cdr</sup> mice were infected with MHV intranasally, treated with PBS or diphtheria toxin (DT, 1 ng/g body weight) on days 1 and 3 post infection, and brains and cLNs (inlets) were examined by histological analysis on day 8 post infection using the indicated antibodies. Scale bar represents 30 μm.

(D) Absolute numbers of CD8<sup>+</sup> (D) and CD4<sup>+</sup> (E) T cells, and viral titers (F) in the brains of DT-treated mice. Data are shown as mean ± SEM (n = 3–4 mice per group, pooled from two independent experiments).

(G and H) Absolute numbers of CD8<sup>+</sup> (G) and CD4<sup>+</sup> (H) T cells in cLNs of DT-treated mice. Data are shown as mean ± SEM (n = 4–6 mice per group, pooled from two independent experiments).

(I) Frequencies of Ccr7<sup>-/-</sup> and Ccr7<sup>+/+</sup> T cells in the brains of DT-treated mice. Data are shown as mean ± SEM (n = 4–6 mice per group, pooled from two independent experiments). Statistical analysis was performed using the Student’s t test (*p < 0.05; **p < 0.01; ***p < 0.001).
microglia cells, a major target cell population of MHV in the CNS, could trigger secretion of TNF that leads to the initial low-level induction of CCL19/CCL21 expression in stromal cells. With the immigration of activated, lymphotxin-expressing T cells, as demonstrated in our study, fibroblastic and endothelial stromal cells would receive additional activating signals to change into activation mode including massive production of CCR7 ligands.

In addition to their enhanced chemokine production, the phenotype of ECs and FRC-like cells in the affected regions adapted properties that are typical for SLO stromal cells with high level expression of adhesion molecules and upregulation of MHC class I (Malhotra et al., 2012). Because expression of selectins and integrins by inflamed brain vessels reduces the rolling velocity of lymphocytes and permits prolonged interactions with the endothelial wall (Engelhardt, 2006), focal activation of blood vessel ECs most likely fosters recruitment of T cells to sites of viral replication. FRC-like cells in the perivascular space in the vicinity of virus-infected areas exhibited not only elevated expression of ICAM1, and VCAM1, but also of other markers that are typical for LN FRCs including ER-TR7 and DPDPN (Cremasco et al., 2014). Importantly, ER-TR7+ reticular networks have been observed during CNS infection with the lymphocytic choriomeningitis virus (LCMV) (Kim et al., 2009) or during chronic Toxoplasma gondii infection (Wilson et al., 2009). Thus, it is likely that brain vessel ECs and the surrounding FRC-like cells are programmed by the local inflammatory milieu to generate those structures that foster and maintain lymphocyte recruitment and activation.

The presence of CCR7 was mandatory to protect mice from severe neuroinflammatory disease as Ccr7-deficient animals succumbed to infection in only 10 days. Notably, presence of CCR7 on virus-specific CD8+ T cells was sufficient to rescue Ccr7-deficient animals from lethal disease clearly indicating that CCR7 signaling can be decisive for optimal recruitment of activated CD8+ T cells into the inflamed CNS. Adoptive transfer experiments using Ccr7-proficient and -deficient antiviral CD8+ T cells suggested that the lack of CCR7 signals in SLOs can be compensated by other stimuli, but that focal, high-level production of CCR7 ligands at sites of peripheral inflammation is critical for both recruitment and performance of antiviral T cells. The finding that Ccr7-deficient spleen CD8+ T cells did not show impaired IFN-γ production during systemic lymphocytic choriomeningitis virus infection (Junt et al., 2004) supports our interpretation that the SLO environment guarantees sufficient immune stimulation to generate protective antiviral immunity. Interestingly, systemic Toxoplasma gondii infection of Ccr7-deficient mice leads to rapid replication of tachyzoites and dissemination to several peripheral organs including the brain (Noor et al., 2010). It is striking that parasite infection of the CNS provokes generation of perivascular reticular fibers that are coated with CCL21 and that anti-paraetic CDC8+ T cells migrate along these fibers (Wilson et al., 2009). Because viral CNS infection efficiently triggers activation of perivascular FRC-like cells, it is reasonable to assume that the perivascular infrastructure supporting antimicrobial immunity is built by FRC-like stromal cells. As predicted by Perry and colleagues (Galea et al., 2007) these processes therefore substantially diminish immune privilege of the CNS and permit efficient immune surveillance.

In conclusion, our study shows that the rapid generation of CCR7 ligands by stromal cells of the CNS is critical for the containment of neurotropic virus infection. The chemokines can be generated rapidly and at high levels by meningial and sub-weengeal blood vessel ECs and perivascular FRC-like cells. Once appropriately activated, CNS stromal cells generate a confined micro-environment that permits optimal T cell recruitment and activation. It remains to be determined whether some areas in the CNS provide such T cell activation hubs in a constitutive fashion and thereby foster activation of self-reactive T cells by perivascular dendritic cells (Greter et al., 2005). Moreover, it will be important to assess whether and how immune cells that have entered the CNS via the stromal cell-determined entry points leave the organ after the pathogen has been cleared. It is likely that the recently discovered lymphatic connection to the cLNs (Aspelund et al., 2015; Louveau et al., 2015) serve as exit points for immune cells that had accumulated in the inflamed tissue. Accordingly, dysregulation of immune cell influx versus efflux might foster self-perpetuation of focal immune cell aggregations with development of ectopic lymphoid-like structures (Pitzalis et al., 2014). Thus, stromal cells function as critical gate-keepers for activated immune cells and neuroinflammation might be exacerbated if activation of stromal cells following resolution of CNS infection is not restrained. Further dissection of cellular and molecular mechanisms that control stromal cell activation in the CNS is warranted to delineate potential avenues for attenuation of inflammatory processes in this vulnerable organ.

**EXPERIMENTAL PROCEDURES**

**Mice**

The TCR transgenic mouse strain C57BL/6N-Tg(Tcra,Tcrb)577Biat (Spiky) was generated as described previously (Nindi et al., 2012), Ccr7-/-, Cxcr3-/-, Cxcr5-/-, iDTR and plt/plt mice were obtained from the Institute for Laboratory Animal Sciences at the University of Zürich. BAC-transgenic C57BL/6N-Tg(Ccl19-Cre)489Biat (Ccr7-/-Cre) mice have been previously described (Chai et al., 2013). C57BL/6N (B6) mice were purchased from Charles River Laboratories. All mice were maintained in individually ventilated cages and were used in the age of 6 to 9 weeks. Experiments were performed in accordance with federal and cantonal guidelines (Tierschutzgesetz) under the permission numbers SG09/87 and SG13/04 following review and approval by the Veterinary Office of the Canton of St. Gallen.

**Virus Infection and Determination of Virus-Induced Pathology**

Mice were infected intranasally using 5 × 10³ pfu MHV A99. At indicated time points mice were sacrificed, organs were collected, weighed and stored at −80 °C until further analysis. Virus titers were determined from homogenized organs using standard plaque assay on L929 cells.

**Adaptive T Cell Transfer**

Single-cell suspensions from spleens were prepared by mechanical disruption of the organ and subjected to hypotonic red blood cell lysis. For in vivo proliferation, splenocytes were labeled using CFSE (Molecular Probes) according to the manufacturer's protocol, and 10⁴ cells (corresponding to 1 × 10⁶ CDB TCR transgenic T cells) were transferred intravenously (i.v.) into recipient mice. For in vivo protection and co-transfer study, C68 T cells or CD3+ cells from spleens were obtained using anti-CD8 or anti-CD3 MACS beads (Miltenyi Biotec). For assessment of in vivo protection, recipients received 10⁶ cells, while 10² and 10⁴ cells were transferred to plt/plt mice in separate or competitive transfer assays (respectively). At twelve hours post adoptive transfer, recipient mice were infected with MHV A99.

**Cell Isolation from Brain Tissues**

Mice were sacrificed at the indicated time points and immediately perfused with PBS. CNS-infiltrating lymphocytes were isolated using mechanical disruption of the organ followed by enrichment based on 70–30% Percoll
gradients (GE Healthcare) and centrifugation for 25 min at 800 x g. For isolation of stromal cells from olfactory bulbs, the tissue was cut into small pieces, transferred into a 24-well dish filled with RPMI 1640 medium containing 2% FCS, 20 mM HEPES (all from Lonza), 1 mg/ml Collagenase Type P (Sigma-Aldrich), and 25 µg/ml DNaseI (AppliChem), and incubated at 37°C for 30 min. After enzymatic digestion, cell suspensions were washed with PBS containing 0.5% FCS and 10 mM EDTA (MACS buffer). To enrich the stromal cell fraction, we removed myelin-producing cells using 30% Percoll and 15 min centrifugation at 700 x g followed by hematopoietic cell depletion using MACS anti-CD45 and anti-Ter119 Microbeads (Miltenyi).

**Statistical Analyses**

All statistical analyses were performed with Prism 4.0 (GraphPad). Data were analyzed with the unpaired Student’s t test. A p value of < 0.05 was considered significant.

**SUPPLEMENTAL INFORMATION**

Supplemental Information includes five figures and Supplemental Experimental Procedures can be found with this article online at http://dx.doi.org/10.1016/j.immuni.2015.12.022.

**AUTHOR CONTRIBUTIONS**

J.C. and L.O. designed and conducted the experiments and wrote the paper; C.G.—C., E.W., S.C.—F., and C.P.—S. conducted experiments and analyzed data; T.R. generated transgenic mice; I.B. analyzed and discussed data; B.L. designed experiments, supervised the study, and wrote the paper.

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