Cerebrospinal Fluid Dendritic Cells Infiltrate the Brain Parenchyma and Target the Cervical Lymph Nodes under Neuroinflammatory Conditions

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Abstract

Background: In many neuroinflammatory diseases, dendritic cells (DCs) accumulate in several compartments of the central nervous system (CNS), including the cerebrospinal fluid (CSF). Myeloid DCs invading the inflamed CNS are thus thought to play a major role in the initiation and perpetuation of CNS-targeted autoimmune responses. We previously reported that, in normal rats, DCs injected intra-CSF migrated outside the CNS and reached the B-cell zone of cervical lymph nodes. However, there is yet no information on the migratory behavior of CSF-circulating DCs under neuroinflammatory conditions.

Methodology/Principal Findings: To address this issue, we performed in vivo transfer experiments in rats suffering from experimental autoimmune encephalomyelitis (EAE), a model of multiple sclerosis. EAE or control rats were injected intra-CSF with bone marrow-derived myeloid DCs labeled with the fluorescent marker carboxyfluorescein diacetate succinimidyl ester (CFSE). In parallel experiments, fluorescent microspheres were injected intra-CSF to EAE rats in order to track endogenous antigen-presenting cells (APCs). Animals were then sacrificed on day 1 or 8 post-injection and their brain and peripheral lymph nodes were assessed for the presence of microspheres+ APCs or CFSE+ DCs by immunohistology and/or FACS analysis. Data showed that in EAE rats, DCs injected intra-CSF substantially infiltrated several compartments of the inflamed CNS, including the periventricular demyelinating lesions. We also found that in EAE rats, as compared to controls, a larger number of intra-CSF injected DCs reached the cervical lymph nodes. This migratory behavior was accompanied by an accentuation of EAE clinical signs and an increased systemic antibody response against myelin oligodendrocyte glycoprotein, a major immunogenic myelin antigen.

Conclusions/Significance: Altogether, these results indicate that CSF-circulating DCs are able to both survey the inflamed brain and to reach the cervical lymph nodes. In EAE and maybe multiple sclerosis, CSF-circulating DCs may thus support the immune responses that develop within and outside the inflamed CNS.

Citation: Hatterer E, Touret M, Belin M-F, Honnorat J, Nataf S (2008) Cerebrospinal Fluid Dendritic Cells Infiltrate the Brain Parenchyma and Target the Cervical Lymph Nodes under Neuroinflammatory Conditions. PLoS ONE 3(10): e3321. doi:10.1371/journal.pone.0003321

Editor: Jean Kanellopoulos, University Paris Sud, France
Received August 1, 2008; Accepted September 5, 2008; Published October 2, 2008

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Funding: This work was supported by grants from INSERM and ARSEP (to S. Nataf) and by a fellowship from MRT to E. Hatterer. The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

Competing Interests: The authors have declared that no competing interests exist.

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Introduction

DCs are the most powerful antigen presenting-cells of the immune system. They sequentially capture antigens in inflamed tissues, reach the lymphatic vessels, migrate toward lymphoid organs and induce the antigen-specific proliferation of T-cells [1,2]. However, this functional scheme does not apply to CNS for the following reasons: i) in contrast with all other tissues, there is no DCs residing in the CNS parenchyma, ii) the so-called “blood-brain barrier” considerably limits the penetration of blood-circulating immune cells, including DCs and their precursors, into the CNS parenchyma; iii) the CNS is devoided of lymphatic vessels. Nevertheless, despite these limitations, DCs were shown to infiltrate several compartments of the CNS under neuroinflammatory conditions. These intra-CNS compartments communicate with each others and comprise: the CSF [3,4], the meninges [5,6], the perivascular spaces [5,6] and the CNS parenchyma [6,7]. Due to the lack of intra-CNS lymphatic vessels, the question whether and how DCs migrate from the inflamed CNS to lymphoid organs is still controversial. Previous studies performed in normal rats or mice showed that DCs are able to migrate from brain to cervical lymph nodes (CLNs) and to elicit a systemic immune response [8,9]. Also, we reported that in normal rats, DCs injected into the cerebrospinal fluid reached the CLNs while DCs injected into the brain parenchyma stayed confined to the CNS [10]. However, as these experiments were performed in normal rats, one cannot conclude on the actual behavior of CNS-infiltrating DCs under neuroinflammatory conditions. In the present paper, we specifically assessed the migratory behavior and functions of CSF-circulating DCs (CSF DCs) in a rat model of multiple sclerosis, the most common autoimmune disorder of the CNS. Two complementary experimental procedures were followed: i) in a first set of experiments, we tracked endogenous CSF-circulating antigen-presenting cells (APCs) by injecting fluorescent microspheres into the CSF of rats induced for EAE; ii) in a second set of experiments, bone marrow-derived myeloid DCs were labeled with the fluorescent marker carboxyfluorescein diacetate succinimidyl ester (CFSE) then...
injected into the CSF of EAE rats, at the clinical peak of the disease, or control rats. The brains and peripheral lymph nodes of injected animals were then examined by immunohistological methods or FACS analysis on day 1 or 8 post-injection. Data showed that in EAE rats, CSF DCs substantially infiltrated the inflamed brain. Moreover, CSF DCs reached the cervical lymph nodes and enhanced the systemic humoral response against myelin oligodendrocyte glycoprotein (MOG), a major immunogenic myelin antigen.

Results

Intra-CNS migration of CSF DCs in EAE versus control rats

An immunohistological examination of the CNS was performed in: i) EAE rats injected intra-CSF with microspheres (n = 14) and sacrificed on day 1 (n = 8) or 8 (n = 6) post-injection; ii) EAE rats injected intra-CSF with CFSE-labeled DCs (n = 18) and sacrificed on day 1 (n = 8) or 8 (n = 10) post-injection; iii) control rats injected intra-CSF with CFSE-labeled DCs and sacrificed on day 1 post-injection (n = 4) or 8 post-injection (n = 3). In EAE rats, all intra-CSF injections were performed at the clinical peak of disease, on day 12 post-immunization.

(i) Migration of CSF DCs in the periventricular parenchyma

In EAE rats injected with microspheres and sacrificed on day 1 post-injection (n = 8), phagocytic cells having engulfed fluorescent microspheres (microspheres+ cells) were detected as free-floating cells in the injected lateral ventricle (Fig 1A–D) as well as in the third ventricle (Fig 1E) and in the fourth ventricle (data not shown). These microspheres+ cells expressed CD11b/c and MHC class II molecules (Fig 1A–C). Also, microspheres+ cells harboring the same phenotype were evidenced in periventricular inflammatory lesions (Fig 1D–E). Similarly, in EAE rats injected with CFSE-labeled DCs and sacrificed on day 1 post-injection (n = 8), CFSE+/OX42+ cells were detected in the ventricular CSF compartment (Fig 1F–H) and in the periventricular parenchyma (Fig 1I). A similar distribution of CFSE+/OX42+ cells was observed in EAE rats injected with CFSE-labeled DCs and sacrificed on day 8 post-injection (n = 10) (Fig 1J). When performing an immunostaining of the myelin basic protein (MBP/Fig 2), we found that microspheres+ cells (Fig 2A–C) or CFSE+/MHC class II+ DCs (Fig 2D–I) could be evidenced within periventricular demyelinating lesions, on day 8 (Fig 2A-C) or 1 post-injection (Fig 2D-I). Such a migratory behavior was specific as the mean number of CFSE+ DCs was more than 4 times higher in periventricular demyelinating lesions than in normal-appearing periventricular white matter (9.4+/−2.5 cells/10−7 mm2 vs 1.6+/−0.5 cells/10−2 mm2) in demyelinated vs normal-appearing periventricular white matter respectively, p = 0.02, Mann and Whitney test(Fig 3A). Interestingly, some of the microspheres+ cells that localized in these periventricular demyelinating lesions displayed an intracytoplasmic MBP staining, suggesting they had ingested MBP-containing myelin debris (Data supplement S1). Finally, to further determine whether the intra-CNS migration of CSF DCs is conditioned by the neuroinflammatory environment, we performed parallel experiments in which CFSE+ DCs were injected into the CSF of normal rats (n = 4). Rats were sacrificed on day 1 post-injection and serial sections of the injected lateral ventricle were examined. CFSE+ DCs were then counted in order to establish the respective percentages of intraventricular vs periventricular CFSE+ DCs on day 1 following injection (Fig 3B). The same procedure was applied to EAE rats injected with CFSE-labeled DCs and sacrificed on day 1 post-injection (n = 7). While in normal rats, 14.5+/−0.6% cells localized in the periventricular parenchyma, we found that in EAE rats more than 40% of the injected cells localized in the periventricular parenchyma (42+/−7%, p = 0.008, Mann and Whitney test).

Altogether these data show that in EAE rats as compared to controls, CSF DCs present an increased ability to penetrate the periventricular parenchyma and, in particular, the demyelinating lesions that are adjacent to ventricles.

(ii) Migration of CSF DCs along the deep penetrating meninges

On day 1 or 8 following injection, numerous microspheres+ cells were evidenced in the superficial meninges covering the outer surface of the brain (data not shown). However, microspheres+ cells were also frequently detected in the deep penetrating meninges (Fig 4). In particular, microsphere+ cells localized along the pia matter covering the molecular layers of the cerebellum (Fig 4A–B) or lining the inner surface of the brainstem (data not shown). These microsphere+ cells expressed MHC class II molecules (Fig 4C–E) and OX42 (data not shown). Similarly, on day 1 post-injection, numerous CFSE+ DCs were evidenced in the deep penetrating meninges (Fig 4C, D). As observed for microspheres+ cells, CFSE+ DCs were frequently detected in the meninges covering the molecular layers of the cerebellum (data not shown) or in the meninges lining the inner surface of the brainstem (Fig 4F–G). These cells expressed MHC class II molecules and OX42 (Fig 4H, J and data not shown). However, on day 8 following injection, only few CFSE+ cells were still detectable in the deep penetrating meninges (data not shown). Interestingly, when CFSE+ DCs were injected into the CSF of normal rats, we could not detect CFSE+ cells in the deep penetrating meninges (Table 1).

(iii) Migration of CSF DCs in the brain parenchyma and perivascular spaces

On day 1 or 8 post-injection, microspheres+ /MHC class II+ cells as well as CFSE+ DCs localized in parenchymal areas, distant away from the site of injection (Fig 5 and 6). In some cases, these cells appeared to infiltrate a non-inflamed parenchymal area as judged by the lack of infiltrating MHC class II+ or OX42+ inflammatory cells (Fig 5A–C). In other cases, CSF-derived DCs were clearly localized within parenchymal inflammatory foci. (Fig 5D–I). Interestingly, microspheres+ /MHC class II+ cells or CFSE+ DCs were demonstrated in perivascular infiltrates (Fig 6). Indeed, as early as day 1 post-injection, microspheres+ /MHC class II+ cells or CFSE+/MHC class II+ DCs were detected in close vicinity with the vascular wall of cuffed vessels (Fig 6A–D) including small caliber venules (Fig 6B, arrow). A similar migratory pattern was observed on day 8 post-injection (Fig 6E–F). In comparison, only a few CFSE+ cells were detected in the perivascular spaces of normal rats injected with CFSE+ DCs (Table 1).

Migration of CSF DCs in the cervical lymph nodes

A histological examination of the cervical lymph nodes and axillary lymph nodes was performed in EAE rats injected intra-CSF with microspheres (n = 14) and sacrificed on day 1 (n = 8) or 8 (n = 6) post-injection. On day 1 or 8 following injection, fluorescent microspheres were evidenced in the cervical lymph nodes (CLNs) (Fig 7A–B) but not in the axillary lymph nodes (Fig 7E–F) of EAE rats. Microspheres preferentially localized in the B-cell area of CLNs and could be demonstrated within the germinal centers of B-cell follicles (Fig 7G). In order to formally confirm that microspheres had not been passively transported in the lymph flux, an ultrastructural analysis of CLNs was performed. This allowed microspheres to be demonstrated in the cytoplasm of phagocytic cells that localized in the B-cell areas of CLNs (Fig 7H).
Figure 1. **CSF-circulating DCs infiltrate the periventricular parenchyma.** Fluorescent microspheres or DCs labeled with the cytoplasmic fluorescent marker CFSE were injected into the left lateral ventricle of EAE rats (n = 32) at the clinical peak of disease (day 12 post-immunization). EAE rats injected with fluorescent microspheres (n = 14) or CFSE-labeled DCs (n = 18) were then sacrificed on day 1 post-injection (n = 16) or 8 post-injection.
To gain quantitative insights on the migration of CSF DCs toward the CLNs, a FACS analysis was performed on the lymph nodes of EAE rats (n = 8) or control rats (n = 7) injected intra-CSF with CFSE+ DCs (Fig 8). In injected EAE rats sacrificed on day 1 post-injection (n = 4), we found that 2.26+/−2.0,11% CFSE+ cells could be detected in the CLNs as compared to 1.37+/−0.03% in the axillary lymph nodes (p = 0.02, Mann and Whitney test)(Fig 8A, C–D). However, in injected EAE rats sacrificed on day 8 post-injection (n = 4), no statistical difference was observed between the CLNs and the axillary lymph nodes regarding the

Figure 2. CSF-circulating DCs penetrate periventricular demyelinating lesion. Fluorescent microspheres or DCs labeled with the cytoplasmic fluorescent marker CFSE were injected into the left lateral ventricle of EAE rats (n = 32) at the clinical peak of disease (day 12 post-immunization). EAE rats injected with fluorescent microspheres (n = 14) or CFSE-labeled DCs (n = 18) were then sacrificed on day 1 post-injection (n = 16) or 8 post-injection (n = 16). An immunohistological analysis of brains obtained from injected EAE rats or control healthy rats (n = 4) was then performed using antibodies directed against CD11b/CD11c (OX42) or MHC class II molecules. A–C: OX42+ cells (green) harboring intracytoplasmic fluorescent microspheres (red) are observed in the lumen of the injected lateral ventricle, on day 1 post-injection. D: Microspheres+/OX42+ cells localize in the periventricular parenchyma of the injected ventricle, on day 1 post-injection. Insert (solid square) shows a high magnification view of microspheres+/OX42+ cells infiltrating a periventricular inflammatory lesion (dashed square). E: On day 1 post-injection, MHC class II+ cells (green) harboring intracytoplasmic fluorescent microspheres (red) are detectable in the lumen of the third ventricle (dashed square) and in a large periventricular area infiltrated with MHC class II+ cells. Insert (solid square) shows a high magnification view of microspheres+/MHC class II+ cells that localize in the lumen of the third ventricle (dashed square). Arrows indicate microspheres+/MHC class II+ cells in the periventricular parenchyma. F–H: CFSE+ DCs (green) that express OX42 (red) are detected in the lumen of the injected lateral ventricle, on day 1 post-injection. I: A periventricular area bordering the injected lateral ventricle is infiltrated by OX42+ cells and contains OX42+/CFSE+ DCs, on day 1 post-injection. J: On day 8 post-injection, CFSE+ DCs are observed in a periventricular inflammatory lesion, adjacent to the injected lateral ventricle. LV: lateral ventricle, V3: third ventricle. Scale bars: 50 µm (B–J), 50 µm (A–C, E–I), 10 µm (inserts in panel D and E).

doi:10.1371/journal.pone.0003321.g001

doi:10.1371/journal.pone.0003321.g002
Figure 3. CFSE+ DCs specifically target the periventricular demyelinating lesions. DCs labeled with the cytoplasmic fluorescent marker CFSE were injected into the left lateral ventricle of healthy control rats or EAE rats at the clinical peak of disease (day 12 post-immunization). Rats were then sacrificed on day 1 post-injection (EAE rats: n = 4, normal rats: n = 4) and an immunohistological analysis was performed on brain sections crossing the injected lateral ventricle. At least 3 to 5 sections per animal were examined. A: In brain sections from EAE rats, an immunostaining of the myelin basic protein (MBP) was performed and the number of CFSE+ cells/10^-2 mm^2 was counted in demyelinated vs normal-appearing periventricular white matter, as described in the Materials and Methods section. Data show that the mean number of CFSE+ cells was more than 4 times higher in periventricular demyelinating lesions than in normal-appearing periventricular white matter (9.4+/−2.5 cells/10^-2 mm^2 vs 1.6+/−0.5 cells/10^-2 mm^2 in demyelinated vs normal-appearing periventricular white matter respectively, p = 0.02, Mann and Whitney test). B: In brain sections from EAE rats, an immunostaining of the myelin basic protein (MBP) was also performed and the number of CFSE+ cells localized in the periventricular parenchyma before being phagocytized. Nevertheless, several observations which were performed on day 1 post-injection, that is, 2 periventricular parenchyma, while, in EAE rats, more than 40% of the injected cells localized in the periventricular parenchyma (42+/−7%, p = 0.008, Mann and Whitney test). Conversely, 85.5+/−0.6% CFSE+ cells localized in the intraventricular lumen of normal rats while less than 60% of the injected cells localized in the intraventricular lumen of EAE rats (58+/−7%, p = 0.008, Mann and Whitney test). *: p<0.05, **: p<0.01. doi:10.1371/journal.pone.0003321.g003

Discussion

The use of fluorescent microspheres to track endogeneous APCs has been commonly used to analyze the migratory behavior of DCs in other tissues than the CNS [1,12]. In our experimental settings, however, one has to consider that microspheres may not have been exclusively taken up by CSF-circulating APCs. Indeed, free microspheres may have penetrated into the inflamed CNS parenchyma before being phagocytized. Nevertheless, several observations which were performed on day 1 post-injection, that is, percentage of CFSE+ cells [Fig 8B]. It has to be noticed that a dilution effect may have partly biased these results as cellularity increases in the CLNs during the course of EAE (data not shown).

Overall, these results show that CSF-circulating APCs rapidly and specifically target the CLNs under neuroinflammatory conditions. This process is, at least partly, dependent on the immune processes that take place in the CNS and the CLNs.

Effects of intra-CSF injection of DCs on EAE outcome

The clinical course of EAE was compared between control EAE rats, EAE rats injected intra-CSF with microspheres and EAE rats injected intra-CSF with CFSE-labelled DCs. As shown in Figure 9, we observed that in the time period between day 12 (intra-CSF injections) and day 20 (sacrifice), EAE rats injected with DCs (n = 10) presented higher clinical scores (cumulative clinical score: 22.25+/−1.4) than EAE control rats (n = 11, cumulative clinical score: 17.45+/−7.09, p = 0.04 as compared to EAE rats injected with DCs, Student’s t test) or EAE rats injected with microspheres (n = 6, cumulative clinical score: 14.83+/−1.6; p = 0.002 as compared to EAE rats injected with DCs, Student’s t test). We then attempted to determine whether intra-CSF injections of DCs had induced an increased peripheral immune response against CNS antigens. As microspheres’ CSF-derived APCs targeted the B-cell follicles of CLNs, we focused our analysis on the B-cell response against CNS antigens. On day 20 post-immunization, blood samples were withdrawn from control EAE rats (n = 4) or EAE rats that had been injected intra-CSF with DCs on day 12 post-immunization (n = 4). Sera were then assessed by Western blot or ELISA for the presence of antibodies directed against CNS antigens. We first probed the serum antibody repertoire against whole spinal cord antigens, using Western blot analysis (Fig 10A). Data showed that, as compared to control EAE rats, sera from injected EAE rats contained higher concentrations of antibodies directed against a 28 Kda protein, distinct from myelin basic protein (MBP). As myelin oligodendrocyte glycoprotein, a major immunogenic myelin antigen, is a 28 Kda molecule [11], we then performed ELISA experiments to determine the serum concentrations of anti-MOG antibodies (Fig 10B). The concentration of antibodies directed against the MOG peptide 35–55 was greatly increased in injected EAE rats as compared to control EAE rats (84.23+/−22.8 ng/ml vs 20.53+/−4.1 ng/ml in EAE and control rats respectively, p = 0.02, Mann and Whitney test) (Fig 10B). Such an increased antibody response was not observed when considering myelin basic protein (MBP) as a target antigen (Fig 10C).

Overall, these data demonstrate that injection of DCs into the CSF of EAE rats induces an exacerbation of clinical signs along with an increased systemic antibody response against MOG.

Traffic of CSF Dendritic Cells

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Overall, these data demonstrate that injection of DCs into the CSF of EAE rats induces an exacerbation of clinical signs along with an increased systemic antibody response against MOG.
Figure 4. CSF-circulating DCs migrate along the deep penetrating meninges. Fluorescent microspheres or DCs labeled with the cytoplasmic fluorescent marker CFSE were injected into the left lateral ventricle of EAE rats (n = 32) at the clinical peak of disease (day 12 post-immunization). EAE rats injected with fluorescent microspheres (n = 14) or CFSE-labeled DCs (n = 18) were then sacrificed on day 1 post-injection (n = 16) and an immunohistological analysis of brains was performed using antibodies directed against CD11b/CD11c (OX42) or MHC class II molecules. Nuclei were counterstained with the fluorescent nuclear dye DAPI. A, B: Counterstaining of nuclei with DAPI coloration (A) shows that microspheres (visualized as white spots in B) localize in the deep penetrating meninges lining the inner parts of the brainstem (Bs) and cerebellum (Cb). C–E: Microspheres+ cells (white in D, red in E) expressing MHC class II molecules (green in C and E) are observed in the deep penetrating meninges covering the cerebellar convolutions. Insert in E shows a high magnification view of a ramified microsphere+/MHC class II+ cell.
observed in the penetrating pia matter. F, G: Counterstaining of nuclei with DAPI coloration (F) shows that CFSE<sup>+</sup> cells (G) localize along the pia matter lining the inner parts of the brainstem (Bs) and cerebellum (Cb). H-J: An OX42<sup>+</sup> infiltrate is observed in the deep penetrating meninges covering the inner parts of the brainstem (Bs) and cerebellum (Cb) (H). These infiltrating cells comprise CFSE<sup>+</sup> cells (I) that express OX42<sup>+</sup> (J). Scale bars: 100 µm (F–G), 50 µm (A–E, H–J), 10 µm (insert in panel E).

doi:10.1371/journal.pone.0003321.g004

Another interesting observation is the particular ability of CSF-circulating DCs to infiltrate the deep penetrating meninges of the inflamed brain. The pia-matter not only covers the hemispheric and cerebellar cortical convolutions, but it lines deep cerebral structures such as the hippocampus or the inner part the brainstem. The pre- and post-capillary vessels are also covered by the pia matter, which thus connects the CSF compartment with the perivascular spaces (Virchow-Robin spaces) [14]. Our data indicate that the pia matter is a major route of traffic for CSF-circulating DCs. In this view, it has to be noticed that meninges have been described as an important site of antigen-presentation and/or B-cell proliferation in EAE or MS [15,16,17]. The molecular mechanisms supporting the migration of CSF-DCs along the pia matter have to be clarified. One may hypothesize that chemokines accumulate specifically in the CSF compartment delineated by the penetrating pia-matter. Alternatively, leptomeningeal cells themselves may secrete chemokines [18]. One may speculate that such a peculiar routage via the pia-matter may allow CSF-DCs: i) to sample soluble antigens that are drained from the interstitial fluid of CNS parenchyma [19,20,21], ii) to eventually reach intra-parenchymal locations that are distant away from the ventricles, iii) to target the perivascular spaces of post-capillary venules, where T-cells preferentially infiltrate the CNS [14].

The intra-CSF injection of DCs in EAE rats induced an exacerbation of clinical signs and an increased systemic antibody response against MOG. Interestingly, in different EAE models developed in rats, mice or monkeys, anti-MOG antibodies are considered as highly pathogenic [22,23,24,25]. Similarly, in MS patients, anti-MOG antibodies were evidenced in the serum, CSF and, most importantly, within CNS inflammatory lesions [22,26]. Based on these results, one may assume that in EAE rats injected with DCs, increased concentrations of serum anti-MOG antibodies may be partly responsible for the aggravation of clinical signs. Moreover, that intra-CSF injected DCs enhanced the systemic antibody response further demonstrates that DCs are able to migrate from the CSF to the CLNs. Interestingly, preliminary data indicate that, in EAE rats, a diversification of the antibody repertoire occurs in the CLNs but not the axillary lymph nodes (data supplement S2). It is thus thinkable that, when targeting the B-cell area of CLNs, DCs injected intra-CSF induce an amplification of the anti-MOG antibody response that develops in the CLNs. Supporting this view, a sub-population of DCs was previously evidenced in the B-cell zone of lymphoid organs [27,28,29] and DCs were shown to support the proliferation/differentiation of B-cells [29,30]. Besides the effects of DCs on B-

| Table 1. Semi-quantitative distribution of CSF-injected CFSE<sup>+</sup> DCs in the CNS of normal or EAE rats. |
| --- |
| CNS compartment | Normal rat | EAE rat |
| Perivascular | ++/− | + |
| Intraventricular | +++ | +++ |
| Free-floating | +++ | +++ |
| Adherent to choroid plexuses | +++ | +++ |
| Periventricular | LV | + (SVZ) | +++ |
| | V3 | ++/− | ++ |
| | V4 | ++/− | ++ |
| Brain meninges | Superficial | +++ | +++ |
| Penetrating | − | ++ |
| Cerebellum meninges | Superficial | +++ | +++ |
| Penetrating | − | +++ |
| Cerebellum parenchyma | − | + |
| Brain stem parenchyma | − | + |

In parallel experiments, DCs labeled with the cytoplasmic fluorescent marker CFSE were injected intra-CSF to normal rats (n = 7) or EAE rats (n = 8). Serial brain sections crossing different brain areas (anterior brain, midbrain, brain stem and cerebellum) were then assessed for the presence of CFSE<sup>+</sup> cells, on day 1 post-injection (EAE rats: n = 4, normal rats: n = 4). A semi-quantitative analysis was performed as follows: −: none; +: 1–5 cells per section; ++: 5 to 10 cells per section; +++: >10 cells per section. LV: lateral ventricle, V3: third ventricle, V4: fourth ventricle, SVZ: subventricular zone. doi:10.1371/journal.pone.0003321.t001
lymphocytes, it is likely that T-cell driven mechanisms are also triggered by the infiltration of CSF DCs in multiple CNS compartments, as previously shown in a murine model of EAE [6].

In many neuroinflammatory disorders, the intra-CSF accumulation of immune cells is a major diagnosis criterion. Paradoxically, little is known about how and where these CSF-circulating cells migrate, under neuroinflammatory conditions. The present work unravels the complex migratory behavior of CSF-circulating DCs in EAE rats. From the ventricular CSF compartment, DCs have the ability to target the CLNs but also inflammatory lesions that localize in periventricular areas, in the deep penetrating meninges, and to a lesser extend, around parenchymal blood vessels. Our work shows that CSF DCs are thus able to support CNS-targeted immune responses inside as well as outside the inflamed CNS.

### Materials and Methods

#### Animals

Animal care and procedures have been conducted according to the guidelines approved by the French regional ethical committee in animal experimentation (CREEA - agreement number 0106) and meet the Neuroscience Society guidelines. Eight to ten weeks old female Dark Agouti rats were obtained from Harlan (Gannat, France).

#### Reagents

Murine GM-CSF, human Flt3-L and murine IL-4 were obtained from Peprotech (Tebu). Mouse monoclonal antibodies recognizing rat MHC class II molecules (OX6 antibody), rat
CD11b/CD11c (OX42 antibody) or human myelin basic protein (MBP), were purchased from Becton Dickinson Biosciences (Pharmingen). For immuncytochemistry, a fluorescein-conjugated goat anti-mouse antibody (Alexa Fluor 488 or 546, Molecular Probes) was used as a secondary antibody.

**Generation and characterization of rat bone marrow-derived DCs**

Rat myeloid DCs were generated from whole bone marrow cultures as previously described [10]. Briefly, rats were sacrificed, and bone marrow was flushed from femurs and tibias using 10 ml of DMEM in a 10 ml syringe with a 26-gauge needle. Bone marrow cells were then re-suspended and passed through a cell strainer (70 μm pore). Cells were washed once in DMEM (Gibco) then cultured in 25 cm² culture plates at a density of 10⁶ cells/ml in IMDM (Gibco) supplemented with 15% FCS (Fetal Clone II, Perbio Science) and antibiotics (penicillin/streptomycin, Invitrogen). Bone marrow cultures were then grown for 7 days at 37°C, 5% CO₂ in the presence of murine GM-CSF (10 ng/ml) and human Flt-3 ligand (10 ng/ml). By the end of this period, clusters of non-adherent cells had formed that were removed, dispersed and re-plated in fresh media consisting of DMEM supplemented with 10% FCS (BioWest), penicillin/streptomycin (Invitrogen), murine GM-CSF (10 ng/ml) and murine IL-4 (10 ng/ml). After 3 days, large numbers of free-floating cells could be harvested, washed once in PBS and used for labeling and in vivo experiments. The phenotype and morphology of injected cells was assessed by FACS or immunocytochemistry, as previously described [10] (data

**Figure 6. CSF-circulating DCs migrate toward intraparenchymal perivascular infiltrates.** Fluorescent microspheres or DCs labeled with the cytoplasmic fluorescent marker CFSE were injected into the left lateral ventricle of EAE rats (n = 32) at the clinical peak of disease (day 12 post-immunization). EAE rats injected with fluorescent microspheres (n = 14) or CFSE-labeled DCs (n = 18) were then sacrificed on day 1 post-injection (n = 16) or 8 post-injection (n = 16). Their brains were then examined by immunohistology using antibodies directed against MHC class II molecules. Nuclei were counterstained with the fluorescent nuclear dye DAPI. A, B: Two cuffed vessels (dashed square in A) are observed in the brain parenchyma, distant away from the injected lateral ventricle, on day 1 post-injection. These perivascular infiltrates contain microspheres+/MHC class II⁺ cells (arrow heads in B). One of the cuffed vessels presents a venule-like morphology (arrow). Inserts in B show high magnification views of these microspheres+/MHC class II⁺ cells. C, D: In the brainstem, a perivascular infiltrate is formed by MHC class II⁺ cells (C) and contains CFSE+/MHC class II⁺ cells (arrow heads in C and D) on day 1 post-injection. Insert in C shows a DAPI coloration of the brainstem area where this cuffed vessel localizes (arrow in dashed square). E, F: In the hippocampus, on day 8 post-injection, three cuffed vessels harboring a venule-like morphology (white stars in E) are surrounded by CFSE⁺ cells (F). Scale bars: 100 μm (A), 50 μm (B, E, F), 20 μm (C, D).

doi:10.1371/journal.pone.0003321.g006

![Image](https://example.com/image1.png)

![Image](https://example.com/image2.png)
supplement S3). Data showed that injected cells expressed a dendritic cell phenotype as demonstrated by the detection of OX42 (CD11b/c), CD11c and MHC class II molecules. Only 25–35% of cells expressed the co-stimulatory molecule CD86, indicating that a great majority of cells displayed an immature DC phenotype. Accordingly, on cytosin preparations, immunocytochemical staining of MHC molecules showed that most cells harbored intracytoplasmic MHC class II vesicles, a hallmark of immature DCs (data supplement S3).

Cell labelling
DCs were labeled with CFSE (carboxyfluorescein diacetate succinimidyl ester, Molecular Probes) as previously described [10]. Briefly, cells were washed once in PBS and incubated for 5 min at

Figure 7. CSF-derived microspheres’ cells target the B-cell zone of CLNs. Fluorescent microspheres were injected into the left lateral ventricle of EAE rats (n = 14) at the clinical peak of disease (day 12 post-immunization). EAE rats were then sacrificed on day 1 post-injection (n = 8) or 8 post-injection (n = 6) and their cervical lymph nodes and axillary lymph nodes were examined by histological methods. Nuclei were counterstained with the fluorescent nuclear dye DAPI. A–D: In the cervical lymph nodes, on day 1 (d1)(A, B) or 8 (d8)(C, D) post-injection, microspheres’ cells (visualized as white spots) localize preferentially in the cortical, B-cell rich zone. E, F: On day 8 post-injection (d8), there is no detectable microspheres’ cell in the axillary lymph nodes. G: On day 8 post-injection (d8), in the cervical lymph nodes, microspheres’ cells (arrow heads)(red spots) are observed in a B-cell follicle (arrow heads). H: On day 8 post-injection, analysis of cervical lymph nodes by transmission electron microscopy (TEM) shows, in the cortical zone, a phagocytic cell containing a latex bead (Lb, dashed square) and localizing in close contact with lymphocytes (Ly). A high magnification view of this engulfed latex bead (Lb) is shown in the insert (solid square). Scale bars: 100 μm (A–F), 50 μm (G), 1 μm (H).
doi:10.1371/journal.pone.0003321.g007
Figure 8. CSF-circulating DCs target the CLNs. In parallel experiments, DCs labeled with the cytoplasmic fluorescent marker CFSE were injected into the left lateral ventricle of control rats (n = 7) or EAE rats (n = 8) at the clinical peak of disease (day 12 post-immunization). Rats were then sacrificed on day 1 (d1) or 8 (d8) following injections. The cervical lymph nodes (CLNs) and axillary lymph nodes (ALN) were assessed by FACS analysis.
for the presence of CFSE<sup>+</sup> cells. The level of autofluorescence was established on cells obtained from the CLNs or ALNs of non-injected EAE rats (n = 4) or control rats (n = 3). A–D: In injected EAE rats sacrificed on day 1 post-injection (n = 4), we found that 2.26 ± 0.11% CFSE<sup>+</sup> cells could be detected in the CLNs as compared to 1.37 ± 0.03% in the axillary lymph nodes (p = 0.0202, Mann and Whitney test)(A). However, in injected EAE rats sacrificed on day 8 post-injection (n = 4), the percentage of CFSE<sup>+</sup> cells was not statistically different between the CLNs and the ALNs (B), Pannels C and D show representative dot plots obtained from the analysis of injected EAE rats sacrificed on day 1 post-injection. E–F: In the CLNs of injected EAE rats, a great majority of CFSE<sup>+</sup> cells express MHC class II molecules, on day 1 post-injection (94.25 ± 3.3%) or 8 post-injection (87.65 ± 5.2%). A representative dot plot is shown in F. G–H: When comparing injected EAE rats to injected control rats, data showed that on day 1 post-injection, a greater percentage of CFSE<sup>+</sup> cells was detectable in the CLNs of injected EAE rats as compared to injected control rats (2.26 ± 0.11% vs 1.49 ± 0.13% in injected EAE and injected control rats respectively, p = 0.0339, Mann and Whitney test)(G). This difference did not reach significance on day 8 post-injection (1.71 ± 0.03% vs 1.65 ± 0.13% in injected EAE and injected control rats respectively)(H). *: p < 0.05, NS: not significant. doi:10.1371/journal.pone.0003321.g008

Experimental allergic encephalomyelitis

Chronic relapsing EAE was induced in a total of 47 female Dark Agouti rats, as previously described [31]. Briefly, 0.5 gram guinea pig spinal cord obtained from Dunkin-Hartley guinea pigs was homogenized with 3 ml saline and the homogenate was then emulsified with 3 ml of Difco’s bacto complete Freund’s adjuvant (CFA) H37RA supplemented with 15 mg of Mycobacterium tuberculosis H37RA (Difco Laboratories, Detroit, MI). Rats were injected intradermally into each footpad, under Isoflurane anesthesia, with 0.1 ml of this emulsion. In this EAE model, clinical signs peak on day 12 post-immunization then animals partially recover and experience a second relapse. Rats were吱血而 boxed of clinical signs; 0 = no clinical signs, 1 = tail weakness, 2 = tail paralysis, 3 = incomplete paralysis of one or two hind legs, 4 = complete hind limb paralysis, 5 = moribund.

Stereotaxic injections of fluorescent microspheres

On day 12 post-immunization, EAE rats (n = 14) received an intracerebroventricular (intra-CSF) injection of sterile latex fluorescent microspheres diluted 1/1000. All injections were performed using red-phenol free DMEM as a vehicle. Briefly, each rat was deeply anesthetized by pentobarbital injection, placed in a stereotaxic frame and its head tilted slightly by raising the tooth bar at 5 mm. Ten µl of the solution was then slowly injected in the left lateral ventricle (stereotaxic coordinates: 1.4 mm lateral to the bregma and 4.5 mm down from the surface of the skull) over a period of 3 min, using a Hamilton syringe. The animal remained in the stereotaxic frame with the needle in place thereafter for 1 min and the needle was then slowly removed over a period of 2 min. Following intra-CSF injections, animals were kept in our animal facility before being sacrificed on day 1 post-injection (n = 8), or on day 8 post-injection (n = 6).

Stereotoxic injections of labeled DCs

Following the same protocol than described above, EAE rats (n = 18) were injected intra-CSF with vehicle (10 µl) containing 3.10<sup>5</sup> CFSE-labeled DCs. Animals were then sacrificed on day 1 (n = 8) or on day 8 (n = 10) post-injection. In parallel experiments, normal rats (n = 7) were injected intra-CSF with the same preparation of cells and sacrificed on day 1 (n = 4) or 8 (n = 3) post-injection.

Histological analysis

On day 1 or 8 after intra-CSF injections of DCs or microspheres, animals were anesthetized by halothan inhalation and killed by intraperitoneal injection of an overdose of pentobarbital sodium [100 mg/kg]. Brains, cervical lymph nodes and axillary lymph nodes were dissected out and frozen in dry-ice. When needed, frozen tissues were embedded in polyethylene glycol and cut in 14 µm thick sections with a cryostat. After fixation in ethanol, sections were rinsed 3 times in PBS then incubated for 30 min at room temperature with a blocking solution containing 4% bovine serum albumin (Sigma-Aldrich) and 10% normal goat serum. Sections were then incubated overnight at 4°C with mouse monoclonal antibodies directed against CD11b/CD11c (OX42), MHC class II molecules (OX6) or myelin basic protein. OX42 (anti-CD11b/CD11c). All primary antibodies were diluted 1:100 in blocking solution. After several washes in PBS, sections were incubated for 50 min in blocking solution containing a fluorescein-conjugated goat anti-mouse antibody (dilution 1:100) then rinsed in PBS and mounted using an aqueous preparation (Fluoroprep, BioMérieux). Images were recorded and analyzed using a computer-assisted system consisting of specific image analysis software (analysis auto, Soft Imaging System GmbH, Münster, Germany), coupled to a light and fluorescent microscope (Zeiss Axioplan II, Oberkochen, Germany) and in some cases, confocal laser scanning microscopy (LSM META Zeiss) was performed to discriminated between phagocytic cells internalization of microspheres and attachment to the cell membrane. For analysis of fluorescence, two views of the same sections were recorded, using appropriate filters, as black and white image, which were then color-coded by software as blue (DAPI, cell nuclei), green (CFSE or cell type markers) or red

37 °C in 1 µM CFSE. Then, 250 µl of FCS was added and cells were further incubated for 5 min at 37°C before being washed in PBS and re-suspended at a dilution of 3.10<sup>4</sup> cells/µl in red phenol-free DMEM.
(fluorescent microspheres or cell type markers). When needed, in order to better visualize red fluorescent microspheres on a black background, red spots were represented as white spots. The number and localization of CFSE\(^+\) cells was assessed on serial sections of different brain regions (anterior brain, midbrain, brainstem and cerebellum) in EAE rats (n = 8) or in control rats (n = 7). On sections obtained from animals sacrificed on day 1 post-injection (EAE rats: n = 4, normal rats: n = 4), a semi-quantitative analysis was performed to evaluate the presence of CFSE\(^+\) cells in the following CNS compartments: intraventricular CSF (free-floating or adherent to the choroid plexuses), periventricular parenchyma, superficial meninges, deep-penetrating meninges, perivascular spaces, CNS parenchyma. The following semi-quantitative scale was used: −: none, +/−: occasional cells, +: 2 to 3 cells per section, ++: 5 to 10 cells per section, +++: more than 10 cells per section. Also, the injected lateral ventricle was specifically examined and the numbers of intraventricular vs periventricular CFSE\(^+\) cells were counted in EAE or control rats. Finally, in EAE rats, sections crossing the injected lateral ventricle (3 to 5 sections per animal, n = 4 animals) were immunostained with MBP and the number of CFSE\(^+\) cells was counted in normal-appearing or demyelinated periventricular white matter. Using a computer-assisted system consisting (analysis auto, Soft Imaging System GmbH, Munster, Germany), we then determined the number of CFSE\(^+\) cells per mm\(^2\) surface in demyelinated vs non-demyelinated periventricular white matter.

Electron microscopy

In some experiments, cervical lymph nodes obtained from control EAE rats (n = 3), EAE rats injected intra-CSF with microspheres (n = 4) or healthy control rats (n = 3) were processed for electron microscopy examination. Briefly, tissues were fixed for 30 min in 2% glutaraldehyde-0.1 M NaCacodylate pH 7.4. They were then washed three times in 0.1 M NaCacodylate/sucrose, pH 7.4 for 15 min and post-fixed with 1% OsO\(_4\)-0.15 M NaCacodylate pH 7.4 for 30 min. After dehydration in a growing gradient of ethanol, 5 min for each step: 30, 50, 70 and 95%, impregnation steps and inclusion were performed in Epon.

Figure 10. DCS injected intra-CSF stimulates the antibody response against myelin oligodendrocyte glycoprotein. On day 20 post-immunization, blood samples were withdrawn from control EAE rats (n = 4) or EAE rats that had been injected intra-CSF with DCs on day 12 post-immunization (n = 4). Sera were then assessed by Western blot analysis or ELISA for the presence of antibodies directed against CNS antigens. A. The serum antibody repertoire against whole spinal cord homogenate (obtained from healthy rats) was profiled by Western blot analysis (left panel) followed by measures of optical densities (right panel). Data showed that, as compared to control EAE rats, sera from injected EAE rats contained higher concentrations of antibodies directed against a 28 Kda protein, distinct from myelin basic protein (MBP, a major immunogenic myelin antigen) (optic density: 5583.9+/−1161.5 vs 2343+/−677 in EAE+DC and EAE rats respectively; p = 0.0433, non parametric Mann and Whitney test). B. As myelin oligodendrocyte glycoprotein (MOG, another major immunogenic myelin antigen) is a 28 Kda protein, we performed ELISA experiments allowing serum antibodies against MOG peptide 35–55 to be measured. Data showed that higher concentrations of anti-MOG antibodies were detectable in injected EAE rats (EAE+DCs) as compared to control EAE rats (EAE) (84.23+/−22.8 ng/ml vs 20.53+/−4.1 ng/ml in EAE+DCs vs EAE rats respectively, p = 0.0273, non parametric Mann and Whitney test). C. Western blot experiments (left panel) followed by quantitative analysis of optical densities (right panel) showed that the serum antibody response against purified myelin basic protein (MBP) was not significantly different in EAE injected rats (EAE+DCs) as compared to control EAE rats (EAE) (optic density: 20572+/−1683.93 vs 18209+/−2932.34 in EAE+DCs vs EAE rats respectively, p: not significant). Membranes were stained with Red ponceau (lower left panel) to ensure that similar amounts of purified MBP had been loaded in the different lanes. *: p<0.05.

doi:10.1371/journal.pone.0003321.g010
finally polymerized at 60°C for 48 h. 60–80 nm sections were obtained using an ultramicrotome RMC-MTX (Tucson), and contrasted with uranyl acetate and lead citrate. Observations were performed on a JEOL 1200EX transmission electron microscope (Jeol LTD) equipped with a MegaView II high resolution TEM camera and Analysis Soft Imaging system (Eloıse SARL).

### Flow cytometry

Following intra-CSF injection of CFSE-labeled DCs in normal rats (n = 7) or EAE rats (n = 8), CLNs and axillary lymph nodes were mechanically dissociated, washed once in PBS 1x then filtered through a 70 μm cell strainer (PharMingen, San Diego, USA). Cells were then fixed in PFA 1% before analysis of CFSE fluorescence. The level of autofluorescence was established on cells obtained from the CLNs or ALNs of non-injected EAE rats or control rats. Alternatively, cells were stained with OX6 antibody (recognizing MHC Class II molecules) as previously described [10]. Briefly, 2 to 5.10^5 cells were sequentially incubated on ice with a blocking CD16/CD32 monoclonal antibody (PharMingen, San Diego, USA) and a mouse PE-conjugated OX6 antibody (PharMingen, San Diego, USA) diluted 1:50 in PBS containing 2% FCS. After two washes in PBS 1x, cells were then fixed in PFA 1% before being analyzed on a Galaxy flow cytometer (Partec, Germany) with FlowMax software (Partec, Germany).

### Western blot

Western blot experiments were performed to assess the presence of serum antibodies directed against purified myelin basic protein (MBP) (obtained from guinea pig brain, Sigma-Aldrich) or whole spinal cord protein extracts (obtained from normal healthy rats). Purified MBP or pooled spinal cord samples that had been boiled in SDS-PAGE reducing buffer for 5 min at 95°C, were separated on a SDS-PAGE gel and transferred onto a nitrocellulose membrane. The membrane was incubated in PBS containing 0.1% Tween 20 and 5% milk, 1 h at room temperature and incubated with serum samples (dilution 1/10) obtained from control EAE rats (n = 4) or EAE rats injected intra-CSF with DCs (n = 4). Membranes were then washed in PBS containing 0.1% Tween 20 and 1% Milk and incubated with peroxidase-conjugated goat anti-rat IgG immunoglobulin (Jackson, USA) for 1 hour at room temperature. After washing, membranes were incubated with Western Blot Chemoluminescence Reagent (Covalab, France) for 1 min at room temperature and blots were exposed to X-ray films. Semiquantitative analyses of signals were performed by measuring optical densities with the software Image Quant.

### ELISA

Serum samples were tested by ELISA for the presence of anti-MOG 35–55 antibodies, using a commercially available kit (Anaspec). Assays were performed in duplicate and according to the manufacturer’s recommendations. Briefly, 100 μl of undiluted serum samples were incubated in wells of the 96-well microtitration plate pre-coated with MOG 35–35 peptide and a blocking solution. After washing, peroxidase-conjugated antibodies to rat/mouse IgG were added and further incubated for 1 h at room temperature. The reaction was developed after another series of washes by addition of 50 μl of TMB (Tetramethylbenzidine) color substrate solution. Optical densities were then measured on a Multiskan EX ELISA spectrophotometer (Thermo Electron Corporation, Courtaboeuf, France). A calibration curve was established using standard dilutions of mouse anti-MOG IgG and individual values of serum anti-MOG IgGs were then calculated and expressed as ng/ml.

### Statistical analyses

Statistical analyses were performed with the non-parametric Mann and Whitney test. Alternatively, the Student’s t test was performed when the n reached at least 10 in each group and only if values had a Gaussian distribution as assessed by the Shapiro and Wilk test.

### Supporting Information

Data Supplement S1  Microspheres+/MBP+ cells in a periventricular demyelinating lesion. EAE rats were injected intra-CSF with fluorescent microspheres (red in A and D, white in C) and were then sacrificed on day 8 post-injection. Brain sections were immunostained with an anti-MBP antibody in order to visualize myelin (green in A and D, white in B). Microphotograph in A shows that a large demyelinated area adjacent to the third ventricle, is filled with microspheres+ cells. Within this large demyelinated area, a partially demyelinated area (solid square) contains microspheres+/MBP+ cells. Higher magnification views of this area are shown in B (MBP staining), C (microspheres) and D (merge). V3: third ventricle. Scale bars: 100 mm (A), 10 mm (B–D) Found at: doi:10.1371/journal.pone.0003321.s001 (0.92 MB TIF)

### Acknowledgments

We thank Dr Simone Peyrol and the CeCIL (Centre Commun d’Imagerie Laennec) for the use of electron microscopic facilities. We are also grateful to Ivan Natal (in memoriam) for its past continuous support. Thank you also to JF Gherzi-Egea for its helping discussions.

### Author Contributions

Conceived and designed the experiments: EH SN. Performed the experiments: EH MT SN. Analyzed the data: EH SN. Contributed reagents/materials/analysis tools: MFB JH. Wrote the paper: SN.

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Traffic of CSF Dendritic Cells

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