Lymphocyte Circadian Clocks Control Lymph Node Trafficking and Adaptive Immune Responses

Graphical Abstract

Highlights

- Lymphocyte numbers in lymph nodes and lymph oscillate over the course of the day
- Rhythmic Ccr7 and S1pr1 expression drives rhythmic lymphocyte homing and egress
- Adaptive immune responses to immunization and pathogens are time-of-day dependent
- Loss of circadian clocks in lymphocytes ablates rhythmic adaptive immune responses

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In Brief
Lymphocyte trafficking through lymph nodes and lymph is an important immune surveillance mechanism of the body. Druzd et al. (2017) demonstrate that this trafficking occurs in a circadian manner and that adaptive immune responses are also time-of-day dependent and are ablated when circadian clock function is lost in T cells.
Lymphocyte Circadian Clocks Control Lymph Node Trafficking and Adaptive Immune Responses

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http://dx.doi.org/10.1016/j.immuni.2016.12.011

SUMMARY

Lymphocytes circulate through lymph nodes (LN) in search for antigen in what is believed to be a continuous process. Here, we show that lymphocyte migration through lymph nodes and lymph occurred in a non-continuous, circadian manner. Lymphocyte homing to lymph nodes peaked at night onset, with cells leaving the tissue during the day. This resulted in strong oscillations in lymphocyte cellularity in lymph nodes and efferent lymphatic fluid. Using lineage-specific genetic ablation of circadian clock function, we demonstrated this to be dependent on rhythmic expression of promigratory factors on lymphocytes. Dendritic cell numbers peaked in phase with lymphocytes, with diurnal oscillations being present in disease severity after immunization to induce experimental autoimmune encephalomyelitis (EAE). These rhythms were abolished by genetic disruption of T cell clocks, demonstrating a circadian regulation of lymphocyte migration through lymph nodes with time-of-day of immunization being critical for adaptive immune responses weeks later.

INTRODUCTION

Lymphocytes survey antigen by circulating through blood, lymph nodes (LNs) and lymph and shape specific immune responses in LNs. To enter LNs, lymphocytes must undergo extensive interactions with high endothelial cell venules (HEVs) (Butcher, 1991; Förster et al., 2008; Ley et al., 2007; Muller, 2011; Springer, 1994; Vestweber and Blanks, 1999; von Andrian and Mempel, 2003; Wagner and Frenette, 2008). Lymphocytes initially tether on peripheral nodal addressin (PNAd) expressed on HEVs using L-selectin (CD62L) as a ligand. Lymphocytes roll along the vascular endothelium and become activated via interactions of the chemokine receptors CCR7 and CXCR4 with their respective ligands CCL21 and CXCL12. Activated leukocytes use the integrin LFA-1 (CD11a) to bind to ICAM-1 to promote adhesion and finally emigrate into the LN parenchyma. After LN entry, lymphocytes interact with dendritic cells in order to scan presented antigen (Gasteiger et al., 2016), and finally emigrate into efferent lymphatic vessels. For this egress, expression of the sphingosine-1-phosphate-receptor 1 (S1P1, encoded by S1pr1) on lymphocytes is critical, recognizing the chemoattractant phospholipid sphingosine-1-phosphate (S1P) (Matloubian et al., 2004). S1P concentrations are high in blood and lymph but low in tissues, thus providing a gradient that guides lymphocytes out of the LN and into efferent lymph (Cyster and Schwab, 2012). This mechanism is therapeutically exploited for treating multiple sclerosis patients by antagonizing S1P function with the drug FTY720 (fingolimod) to keep autoreactive T cells from exiting LNs and entering the central nervous system (Massberg and von Andrian, 2006).

Lymphocyte trafficking into LNs is believed to occur in a continuous fashion, and not to be influenced by time-of-day variables. Moreover, it is generally unclear whether circadian rhythms regulate overall cellularity in these tissues (Arjona and Sarkar, 2005; Esquifino et al., 1996; Fortier et al., 2011; Hemmers and Rudensky, 2015).
Circadian rhythms are important drivers for most physiological processes as they align the body with rhythmically occurring daily changes in the environment (Dibner et al., 2010). They normally rely on an intricate interplay of cell-intrinsic clock genes driving circadian responses (Mohawk et al., 2012). Daily oscillations of lymphocyte counts in blood have been described (Arjona et al., 2012; Curtis et al., 2014; Haus and Smolensky, 1999; Labrecque and Cermakian, 2015; Scheiermann et al., 2013) and cells of the adaptive immune system such as T and B cells, as well as dendritic cells, possess the components of the molecular clock machinery (Bollinger et al., 2011; Hemmers and Rudensky, 2015; Silver et al., 2012a). In contrast to monocytes of the innate immune system (Nguyen et al., 2013), however, the functional relevance of these cell-intrinsic oscillations for lymphocytes is unclear (Hemmers and Rudensky, 2015). Stimulated by previous findings, which described periodic oscillations in innate immune cell function (Gibbs et al., 2014; Nguyen et al., 2013; Scheiermann et al., 2012) and T helper-17 (Th17) cell differentiation (Yu et al., 2013), we postulated that the migration of lymphocytes through murine LNs might be regulated in a circadian manner with direct relevance for the mounting of adaptive immune responses.

RESULTS

Lymphocyte Numbers Exhibit Circadian Oscillations in Lymph Nodes

In contrast to circulating blood lymphocyte numbers, which peak during the day in mice around Zeitgeber time (ZT) 5 (i.e., 5 hr after light onset) (Figure 1A), numbers for CD4+ and CD8+ T cells as well as B cells showed delayed oscillations (by ~8 hr) in inguinal lymph nodes (iLNs), with highest counts occurring at the beginning of the dark phase (ZT13, i.e., 1 hr after lights off) (Figure 1A). These rhythms were consistently observed for naive and central memory T cells, demonstrating a broad phenomenon also affecting T lymphocyte subpopulations (Figures S1A–S1C). Oscillations were not only observed in the rhythmic environment represented by 12 hr light:12 hr dark conditions (LD) but were sustained in constant darkness (dark:dark, DD), indicating their bona fide endogenous circadian nature (Figure 1B). Light exposure was an important entrainment factor, since rhythms were inverted when the light regime was reversed (DL) (Figure 1B).

Rhythms were furthermore detected across various types of LNs (Figure 1C and Figures S1D–S1F), indicating a relevant phenomenon across the LN compartment. To investigate the
underlying mechanisms driving these oscillations, we focused on the cellular LN input and output pathways by blocking lymphocyte homing or egress, both critical determinants of LN cellularity (Lo et al., 2005). Blocking homing with anti-integrin antibodies dramatically decreased LN cellularity over 24 hr while blocking lymphocyte egress with FTY720 increased LN cellularity over the same time frame, confirming the temporally highly dynamic cellular nature of this tissue (Figures 1D and 1E). Both treatments ablated rhythmicity, indicating that lymphocyte homing and egress—but not intranodal proliferation (Figures S1G and S1H)—were the central determinants of circadian oscillatory cellularity. These data demonstrate a striking circadian oscillation in lymph node cellularity, peaking at night onset.

**Lymphocyte Homing Is Dependent on Oscillations in Lymphocytes and Microenvironment**

We next used adoptive transfer techniques to determine whether lymphocyte homing to the LN was occurring in a rhythmic manner. LN infiltration of lymphocyte subpopulations peaked around night onset and remained low during the day (Figure 2A). To define whether oscillations were determined by lymphocyte-intrinsic and/or microenvironmental signals, we adoptively transferred cells harvested at ZT5 (“day”) or ZT13 (“night”) into LD-entrained recipients at either ZT5 or ZT13. While “day” (cells) into “day” (recipient) transfers exhibited the lowest homing capacity and “night” into “night” transfers the highest, a mixed contribution of both lymphocyte and microenvironment timing was observed in the “day” into “night” and “night” into “day” chimeras (Figure 2B). A screen for oscillations of promigratory factors on T and B cells revealed that expression of the chemokine receptor CCR7 exhibited rhythmicity peaking at ZT13 (Figure 2C) while the adhesion molecules CXCR4, CDT1a, and L-selectin showed either no oscillations or not for all lymphocyte subpopulations (Figures S2A and S2B). In addition, expression analyses of whole lymph node mRNA and extracellular protein on HEVs revealed oscillatory amounts of the chemokine CCL21, a ligand for CCR7—but not CXCL12 (not shown)—being high around night onset (Figures 2D and 2E). HEVs also exhibited rhythmic expression of ICAM-1 but not of PNAd (Figures S2C and S2D). Oscillations in lymphocyte chemokine receptors were critical for rhythmic homing because a titrated, short pre-treatment of adoptively transferred cells with pertussis toxin (PTX) (Lo et al., 2005), an inhibitor of chemokine receptor signaling, ablated rhythmicity (Figure 2F). To investigate the involvement of CCR7 in this process, we analyzed total lymph node cellularity of CCR7-deficient mice, as well as the rhythmic homing capacity of isolated CCR7-deficient mice. Ccr7<−/−> mice exhibited no oscillations in lymph node cell counts while also exhibiting the expected lower overall numbers (Förster et al., 1999) (Figure 2G). In addition, Ccr7<−/−> cells failed to show rhythmic lymph node homing (Figure 2H). These data demonstrated that lymphocyte recruitment to LNs is determined by rhythms in leukocytes and the microenvironment, along with in-phase expression of the CCR7-CCL21 receptor-ligand axis.

**Circadian Clocks Control Cellular Oscillations in Lymph Nodes**

LNs exhibit oscillations of clock genes (Figure 3A), prompting us to investigate the role of lymphocyte clocks in their migratory behavior. We generated mice in which the core clock gene Bmal1 (also known as Amtl) was deleted in T cells (Bmal1<fllox/flx>Cd4-cre) or B cells (Bmal1<fllox/flx>Cd19-cre) (Figure 3B and Figures S2E and S2F). Remarkably, loss of lymphocyte BMAL1 ablated the overall rhythmicity of T and B cell numbers in lymph nodes (Figure 3C and Figure S2G). In addition, rhythmic homing of Bmal1<fllox/flx>Cd4<−/−> T cells into WT recipients was ablated (Figure 3D). In agreement with these findings, rhythmic expression of CCR7 surface protein (Figure 3E) and mRNA (Figure 3F) was absent in BMAL1-deficient CD4+ T cells, indicating the regulation of the molecule at the transcriptional level by the circadian clock. Together, these data provide evidence for a functional role of cell-autonomous clocks in lymphocyte migration.

**Lymphatic Egress from Lymph Nodes Is under Circadian Control**

Because our data indicated that, in addition to a rhythmic homing component, lymphocyte egress might counterbalance LN oscillations (Figures 1D and 1E), we quantified lymphocyte numbers in lymph fluid by cannulating efferent lymphatic vessels. Prominent rhythms in cellular counts were detected, peaking at ZT9 and exhibiting a low at ZT21 (Figure 4A). These oscillations were observed for different lymphocyte populations (Figure 4A and Figures S2H and S2I) and were bona fide circadian in nature as they persisted in constant darkness (Figure 4B). Rhythms were not due to higher lymph volume or flow rates at different times of the day (Figure S2J). To verify whether oscillations in lymph cellularity were truly attributable to rhythmic egress and not secondary to rhythmic input into LNs (Figure 2A), we transferred lymphocytes at different times of the day, blocked subsequent LN entry and quantified their transit through the LN into lymph over time (Mandl et al., 2012). A higher LN retention capacity of cells injected at ZT13 was observed compared to ZT5 and a less rapid accumulation of cells in lymph (Figures 4C and 4D and Figure S3), demonstrating lymphocyte egress to be highly rhythmic. This effect resulted in longer LN half-lives of cells injected at ZT13 (CD4: 12 hr, CD8: 12 hr, B cells: 16 hr) compared to ZTS (CD4: 12 hr, CD8: 9 hr, B cells: 13.5 hr) (Figures 4C and 4D and Figure S3). T- and B cell-specific BMAL1-deletion ablated oscillations in lymph, indicating the importance of cell-autonomous clocks also for lymphocyte egress (Figure 4E and data not shown). Of importance, adoptively transferred BMAL1-deficient CD4+ T cells exhibited no time-of-day variations in their LN half-life (Figure 4F), demonstrating the relevance of T cell clocks in their rhythmic trafficking behavior.

Using a mathematical approach, we assessed whether oscillatory LN counts could be modeled with only either homing or egress to be rhythmic or whether both components needed to oscillate. Although oscillations were also observed when only one component was rhythmic, the best fit was achieved when both homing and egress were assumed to oscillate, thus supporting our experimental data (Figure 4G and Figure S4). In summary, lymphocyte clocks and the time-of-day entry of cells into LNs have functional consequences for LN transit and egress into lymph.

**Rhythmic Lymphocyte Egress Depends on Oscillatory S1pr1 Expression**

S1P-receptors are critical in regulating lymph node egress (Oyster and Schwab, 2012). We therefore investigated whether
expression of S1P-receptor family members exhibited oscillations using quantitative PCR (Q-PCR). All S1P receptors exhibited robust diurnal oscillations peaking between ZT1 and ZT9 (Figure 5A and Figure S5A), which coincided with high lymphocyte egress. In addition, FTY720, as well as the S1P1-specific functional antagonist SEW2871 strongly down-modulated lymphatic
Figure 3. Oscillations of Circadian Clock Genes in Lymph Nodes Control Cellularity

(A) Q-PCR analysis of circadian clock genes in LN over 24 hr; n = 3–5 mice, one-way ANOVA.

(B) Circadian clock gene mRNA profiles in sorted CD4+ T cells from Bmal1flox/floxCd4-cre and control animals; n = 3–10 mice, two-way ANOVA.

(C) Lymph node CD4 and CD8 T cell counts in control and T cell specific Bmal1/C0/C0 mice, n = 3–9 mice, one-way and two-way ANOVA.

(D) LN homing of lymphocytes harvested from control or T cell-specific Bmal1/C0/C0 mice at ZT5 or ZT13 into WT hosts; n = 10–34 mice, one-way ANOVA with Tukey’s multiple comparisons test.

(E) CCR7 surface expression on T lymphocyte subpopulations in LN of control and T cell-specific Bmal1/C0/C0 mice; n = 3–5 mice, one-way ANOVA.

(F) Q-PCR analysis of CD4+ T cell Ccr7 over 24 hr in control and T cell-specific Bmal1/C0/C0 mice; n = 4–8 mice, one-way and two-way ANOVA. *p < 0.05, **p < 0.01, ****p < 0.0001. All data are represented as mean ± SEM. See also Figure S2.
egress in a time- and concentration-dependent manner (Figures 5B and 5C and Figure S5B). The observation that FTY720-treated animals exhibited reduced but still rhythmic lymph cellularity indicated a daytime-sensitive role for S1P1 in mediating lymphocyte exit. Rhythmic expression of $S1pr1$ was ablated in BMAL1-deficient CD4$^+$ T cells, pointing toward a regulation of the gene by the circadian clock (Figure 5D). To investigate this in more detail, we performed an in vitro assay, in which the promoter region of $S1pr1$ was cloned in front of the luciferase ($\text{Luc}$) reporter gene. Luciferase activity in HEK293 cells transfected with the $S1pr1$-luc reporter was decreased after co-transfection of increasing amounts of $Bmal1$ and $Clock$ expression plasmids (Figure 5E). This demonstrated that expression of $S1pr1$ is regulated by BMAL1 and CLOCK.

To confirm the role of S1P1 in the time-of-day-dependent egress genetically, we generated T cell-specific mice that were heterozygous for $S1pr1$ in order not to completely block lymphocyte egress (Matloubian et al., 2004) but to titrate S1P1 amounts, as loss of one allele had been demonstrated to result in haploinsufficiency (Lo et al., 2005). $S1pr1^{-/-}\text{floxxCd4-cre}$ mice exhibited no more oscillations in LN counts and altered lymph rhythmicity, demonstrating the importance of S1P1 in the proper timing of lymphocyte egress (Figures 5F and 5G and Figure S5C). Importantly, no diurnal oscillations were observed in amounts of S1P in efferent lymph (Figure 5H) or in S1P synthesizing or degrading enzymes in lymph node (Figure S5D), suggesting that oscillatory expression of the receptor (S1P1) and not its ligand (S1P) was the driver for rhythmic lymphocyte egress. Together, these data demonstrate a critical role for S1P1 in mediating circadian lymphocyte egress from lymph nodes into efferent lymph.

Relevance of Circadian Oscillations in Lymph Node Cellularity

We hypothesized that oscillatory lymphocyte counts in LNs might have functional consequences in a potential time-of-day dependence of adaptive immune responses. We therefore tested whether the activation status of lymphocytes in LNs varied over the course of the day. More activated T cells were present in LNs at night onset as assessed by CD69 staining, coinciding with higher overall lymphocyte counts at this time (Figure 6A). Because dendritic cells (DCs) are key antigen-presenting cells critical in the activation of lymphocytes and the generation of adaptive immune responses (Girard et al., 2012), we
next investigated whether these cells also exhibited oscillatory LN counts. Migratory DC cellularity showed strong oscillations peaking in phase with lymphocytes (Figure 6B and Figure S6A). These data point to the existence of a concerted circadian migration pattern of antigen-bearing (DCs) and antigen-recognizing (T cells) cells in LNs.

Recent evidence indicates that the immune system can respond to challenges in a rhythmic fashion (Fortier et al., 2011; Gibbs et al., 2014; Nguyen et al., 2013; Scheiermann et al., 2012; Silver et al., 2012b). We therefore investigated the pathophysiological significance of circadian oscillatory LN counts in the autoimmunity model of EAE. Mice immunized during the late light phase (ZT8, when cell counts are high in LNs, Figures 1A and 1C) showed a dramatically accelerated disease progression 2 weeks later, with higher clinical scores compared to late night-immunized animals (ZT20, when LN counts trough) (Figure 6C). Differences in disease scores were associated with higher immune cell infiltration and demyelination in the spinal cord at the peak of the disease (Figures 6D and 6E). We detected elevated interleukin-2 (IL2) mRNA amounts (Figure 6F) and a higher number of IL-17 producing as well as very-late antigen (VLA)-4 integrin positive CD4+ T cells (Figures 6G and 6H).
Figure 6. T Cell Clock Function Regulates Disease Severity in EAE
(A) Oscillations of CD69+ T cell numbers in lymph node; n = 3–5 mice, one-way ANOVA.
(B) Oscillations of migratory dendritic cells (DCs) in lymph node; n = 6–12 mice, one-way ANOVA.
(C) EAE disease scores of mice immunized at ZT8 or ZT20. Disease score EC50 comparisons show accelerated symptom progression in ZT8-immunized mice; n = 5 mice, two-way ANOVA (left panel) and unpaired Student’s t test (right panel).
T cells in LNs of ZT8-immunized mice, subtypes that have been shown to be critical for the induction of EAE (Kawakami et al., 2012) (Figure 6G). This indicated that circadian regulation of immunization occurred at a very early phase of the process when T cells are activated in draining lymph nodes (Figure S6B). Two days after induction of EAE, an increase of both naive and activated CD4+ and CD8+ T cells was detected in draining lymph nodes of ZT8-immunized animals, while in ZT20-immunized animals T cell numbers remained relatively low (Figures 6H–6J and Figure S6C). Thus, oscillations in the numbers of CD4+ T cells in lymph nodes during initial encounters with antigen appear to be pivotal for the severity of EAE. To investigate whether T cell autonomous clocks regulate this response, we genetically deleted T cell circadian clock function. Although in control animals disease development depended on the time of immunization, in T cell specific Bmal1 /− /− mice it did not (Figure 6K). Two days after immunization, total and T cell counts in draining lymph nodes were different at ZT15 between day- and night-immunized control, but not in T cell specific Bmal1 /− /− mice (Figure 6L and Figure S6D). Hence, T cell clocks determine time-of-day function and, after challenge, development of autoimmune sequelae. We finally investigated whether adaptive immune responses to pathogens exhibited similar circadian rhythmicity. Mice were infected with the gastric bacterial pathogen Helicobacter pylori at three different time points during the day, and lymph node counts were quantified 3 weeks later. Also in this chronic infection model, LN counts showed strong circadian responsiveness to the initial infection with highest numbers present at ZT7, analogous to the EAE immunization experiments (Figure S6E). In addition, acute viral infection with influenza A virus led to stronger pulmonary infiltration of CD8+IFN-γ+ T cells when animals were infected at ZT8 compared to ZT20, 8 days post infection (Figure S6F). Together, these data strongly indicate that immunization reactions and the adaptive immune responses to various pathogens follow a circadian rhythm (Figure 6M and Movie S1).

**DISCUSSION**

We have described here the mechanisms that govern a circadian rhythmicity in the capacity of lymphocytes to enter and exit lymph nodes, which depend on cell-autonomous, clock-gene-controlled expression of promigratory factors. Lymphocytes entered LNs most prominently at the onset of the night phase and egressed from the tissue during the day. This resulted in oscillatory cell counts in lymph nodes and lymph and time-of-day differences in the adaptive immune response weeks after immunization. In addition, DCS were found to be present in LN in highest numbers around night onset, peaking in phase with the lymphocyte populations. Our data reveal that T cell-autonomous circadian oscillations are critical in regulating adaptive immunity.

It is surprising to note that lymph nodes exhibit circadian differences in their cellularity, given that they represent such a central tissue of the immune system and, accordingly, have thus been intensely studied. Since we observed oscillations in all investigated lymph nodes, the phenomenon appears to be broad and robust and not restricted to specific body locations. It is noteworthy that other lymphoid organs such as the thymus (data not shown) and the bone marrow do not exhibit overt circadian oscillations in absolute numbers. At least the latter, however, still displays circadian activity in cellular trafficking as hematopoietic stem and progenitor cells (HSPCs) are mobilized into blood (Lucas et al., 2008; Méndez-Ferrer et al., 2008) and recruited back into the bone marrow (Scheiermann et al., 2012) at different times. The fraction of mobilized and homed cells might be small compared with the overall numbers, though, which might explain why overt oscillations of total cells in the BM are not observed. In contrast to the BM, lymph node total cellularity is highly dynamic over 24 hr, as seen when homing or egress is blocked. Still, homing of leukocytes to lymph nodes and bone marrow occurs predominantly at night, while egress (or mobilization in the case of the bone marrow) occurs predominantly during the day. Thus, rhythmic egress of lymphocytes via efferent lymph is a major mechanism underlying the oscillatory leukocyte numbers in blood. Whether other egress routes for lymphocytes, from the thymus or the spleen, occur in a circadian manner is currently unclear.

Our data point to a critical role of cell-intrinsic clock-dependent mechanisms in the regulation of T and B lymphocyte trafficking. While global BMAL1 deficiency results in a diverse array of phenotypes (Bunger et al., 2000), such as altered B cell numbers (Sun et al., 2006), few studies have focused on cell-type specific deletion of BMAL1 in the immune system. Lineage-specific ablation of BMAL1 in myeloid cells results in a pro-inflammatory state (Nguyen et al., 2013), yet a similar approach targeting lymphocytes yielded no obvious phenotype (Hemmers and Rudensky, 2015). The latter finding might be due to mice with clock-deficient lymphocytes exhibiting phenotypes only at specific times, so that only when tested over multiple time points across the day can alterations be detected.
Our data showing that lack of BMAL1 in T cells ablated oscillations in EAE disease scores indicates an important role for T cell clocks in the adaptive immune response. This is in line with a previous observation that T cells harvested from Clock mutant mice exhibited altered proliferation responses at specific times (Fortier et al., 2011).

We demonstrated the critical regulators of lymphocyte trafficking, Ccr7 and S1pr1, to be under circadian control. Ccr7 and S1pr1 oscillations showed opposite phases, being high in the evening for the former and high during the day for the latter. This agrees with the additional role of CCR7 as a retention factor for T cells in lymph nodes that can antagonize S1P1 function (Pham et al., 2008). In addition, more T cells expressed the activation marker CD69 at night onset, a negative regulator of S1P1. These inverted oscillations of retention and egress factors appeared to be responsible for keeping lymphocytes retained in the node at night and promoting daytime egress. In addition to these lymphocyte intrinsic signals, the microenvironmental fluctuation of CCL21 indicated an additional rhythmic component that was non cell-autonomous. Whether direct autonomic innervation governs these oscillations—as has been shown for the bone marrow and skeletal muscle (Méndez-Ferrer et al., 2008; Scheiermann et al., 2012)—and/or humoral factors as has been shown for the lung (Gibbs et al., 2014), is outside the scope of this manuscript and should be investigated in the future.

Lymph nodes act as the body’s immunological sieve, capturing lymph-borne antigens and antigen-presenting cells (APCs) and bringing these components in close contact with lymphocytes. Our data show that not only T cells migrate to the lymph node in a circadian manner but that also dendritic cells, the major APCs, are present more prominently around night onset, peaking at the same time as lymphocytes. Both cell types need to interact to produce a functional adaptive immune response (Gasteiger et al., 2016). It therefore appears likely that having them present at the same time in the confined environment of the lymph node enhances the likelihood of antigen encounter by the very few specific T cell clones, as opposed to cells entering and exiting at random times. This process of more effective cellular interactions appears to be further helped by the fact that during the night fewer cells exit the tissue than by the very few specific T cell clones, as opposed to cells entering and exiting at random times. This process of more effective cellular interactions appears to be further helped by the fact that during the night fewer cells exit the tissue than enter at random times, thus giving APC-T cell pairs more time to interact and increase the likelihood of antigen encounter.

Circulating leukocyte counts are high during the respective resting phases, which occur during the day in mice and at night in humans. In addition, also under stimulated conditions timing matters, as administration of granulocyte colony-stimulating factor (G-CSF) to patients in order to mobilize hematopoietic stem cells from bone marrow into blood yields higher numbers at different times than in mice (Lucas et al., 2008). These data suggest that also adaptive immune responses in humans could be under circadian control. Since in our EAE model, immunization close to the onset of the activity phase produced a higher immune response in mice, in humans the early morning hours should produce a higher adaptive immune reactivity. Indeed, this appears to be the case as recent studies have unveiled that administration of hepatitis A (Phillips et al., 2008) and flu vaccines (Long et al., 2016) in the morning yielded highest antibody titers compared to other times. Together, our data provide mechanistic insights for a time-of-day difference in lymphocyte trafficking and adaptive immune responses, thus warranting further investigations into time-of-day optimization of immune therapies and vaccination programs.

**EXPERIMENTAL PROCEDURES**

**Animals**

Ccr7<sup>-/-</sup> and S1pr1<sup>YFP/ES</sup> mice were obtained from Jackson Laboratories and crossed to target T cells and B cells, respectively. Ccr7<sup>-/-</sup> mice were a gift from Reinhold Förster. 7- to 8-week-old wild-type C57BL/6 mice were purchased from Charles River and Janvier. Mice were housed under a 12 hr:12 hr light-dark cycle with food and water ad libitum. To induce changes in light regime, we placed mice in a light cycler (Park Bioservices) with a 12 hr-inverted light cycle for a minimum of 2 weeks to completely establish an inverse light cycle or kept in constant darkness. All animal experimental procedures were carried out in accordance with the German Law of Animal Welfare and approved by the Regierung of Oberbayern or the ethics committee of the Schleswig-Holstein State Ministry of Energy, Agriculture, Environment and Rural Areas.

**Flow Cytometry**

Blood was collected into EDTA-coated capillary tubes (Microvette 300). Counts were obtained using an IDEXX ProCyte DX cell counter or a Hemavet Hematology Analyzer 950FS (Drew Scientific). Erythrocytes were lysed by incubation in 0.8% NH4Cl. Lymph was collected as described below. Spleens, lymph nodes, or thymi were harvested from animals and processed through a cell strainer (40–70 µM pore size) and resuspended in PBS. For quantification of dendritic cells, lymph nodes were first chopped into small pieces and incubated in collagenase IV (1 mg/ml, C5138, Sigma) and DNase I (0.2 mg/ml, Roche) for 30 min at 37°C with gentle agitation. Following digestion, cells were passed through a 40 µM strainer and resuspended in PBS supplemented with 2% fetal bovine serum (GIBCO). A 20 min blocking step using anti-CD16/32 antibody (2.4G2; BD Biosciences) was performed on ice prior to fluorescent staining. Single-cell suspensions were stained with fluorescence-conjugated antibodies and analyzed by flow cytometry using a Galileo Flow Cytometer (Beckman Coulter) or a FACSCanto II flow cytometer (BD Biosciences). Prior to staining of intracellular cytokines, cells were fixed for 5 hr with PMA and ionomycine (cell stimulation cocktail with protein transport inhibitors, eBioscience). Data were analyzed with FlowJo software (Tree Star).

**Adaptive Transfer Studies**

Lymphocytes from spleen and peripheral lymph nodes (ratio 80:20) were labeled with 1.5 µM of carboxyfluorescein succinimidyl ester (CFSE, Immunity 46, 120–132, January 17, 2017 129
Thermo Fisher Scientific) or 2.5 µM CellTracker Red CMTPX (Thermo Fisher Scientific) in PBS containing 0.2% BSA and 2 mM EDTA for 20 min at 37°C and washed 4 times. 20 × 10^6 cells were then injected i.v. into recipient mice. In some experiments, lymph node entry was blocked two and 12 hr later by intraperitoneal (i.p.) administration of 100 µg anti-α4 (CD11a, clone M17/4; BioXCell) and anti-α7 (CD49d, clone PS2/2; BioXCell) integrin antibodies in PBS. In some experiments, cells were pre-incubated with a 200 ng/ml pertussis toxin (PTX) pulse of 10 min at 37°C and washed 4 times before injection together with control-treated cells (Lo et al., 2003).

The subsets of adoptively transferred cells in the recipient mice were analyzed via flow cytometry. Dyes were switched to avoid dye-specific effects.

**S1pr1 Promoter Luciferase Reporter Assay**
The S1pr1-Gaussia Luciferase (S1pr1-GLuc) GLuc-ON promoter clone containing a ~1 kb fragment upstream of the transcription start site and a ~0.3 kb fragment of the exon 1 of murine S1pr1 in pEZX-PG04 vector (also containing a constitutively expressed secreted alkaline phosphatase (SEAP) secondary reporter as an internal control) was obtained from GeneCopoeia. HEK293AAV cells (Cellbiolabs) were plated onto 96-well plates and maintained in DMEM with 2 mM stable glutamine supplemented with 10% fetal bovine serum (FBS) and 10,000 U penicillin/streptomycin at 37°C with 5% CO2. Upon 80%–90% confluence, cells were transfected with the following expression plasmids: 20 ng S1pr1-luc with various combinations of the following clock gene constructs: HA-Clock, HA-Bmal1, and pcDNA3.1 (mock transfection) as indicated per well using Lipofectamine 3000 transfection reagent (Thermo Fisher Scientific) according to the manufacturer's protocol. Medium was changed once on the next day and was collected for measurement 48 hr after transfection. The secreted GLuc and SEAP activities were measured with the Secreta-Pair Dual Luminescence Assay Kit (GeneCopoeia) according to the manufacturer’s protocol using the Berthold TriStar LB 941 plate reader (Berthold Technologies). Normalized promoter activity was calculated as the ratio of GLuc to SEAP activities.

**Lymph Cannulation**
40 min prior to cannulation, 200 µl olive oil was administered intragastrally to identify mesenteric lymph vessels that are located just upstream of the thoracic duct. Mice were anesthetized (100 mg/kg ketamine, 20 mg/kg xylazine, 1% acepromazine i.p.). Vessels were cannulated and lymph was drawn via a fine bore polyethylene tubing (Smiths Medical) that had previously been flushed with PBS containing EDTA. In some experiments, mice were injected i.p. with 1 mg/kg (or less) of FTY720 or 10 mg/kg of SEW2871 1 or 2 hours, respectively, before cannulation. Cell numbers were determined using an IDEXX ProCyte DX cell counter.

**Immunofluorescence Microscopy and Quantitative Imaging Analysis**
Harvested lymph nodes were embedded in OCT (TissueTec), frozen and sectioned with a thickness of 10 µm on a cryostat (Leica). Sections were fixed in methanol or left unfixed, incubated with fluorescently coupled antibodies and imaged on an Axio Examiner.D1 microscope (Zeiss) equipped with LEDs of 405 nm, 488 nm, 561 nm, and 642 nm excitation wavelengths with LEDs of 405 nm, 488 nm, 561 nm, and 642 nm excitation wavelengths and a 4-color laser stack and a confocal spinning-disk head (Intelligent Imaging Innovations). For investigations of protein expression in HEVs, all quantifications were performed using mask analysis (Zeiss software) based on PECAM-1 expression and quantifying expression of other fluorescent channels within the mask containing PECAM-1+ pixels delinating HEV vascular structures as previously detailed (Schliekermann et al., 2013).

**EAE Induction**
For EAE induction the MOG35-55-CFA Emulsion PTX Kit (EK-2110, Hooke Labs) was used according to the manufacturer's protocol. Mice were anesthetized (100 mg/kg ketamine and 10 mg/kg xylazine i.p.) and bilaterally s.c. injected with 200 µg myelin oligodendrocyte glycoprotein (MOG) 35-55 in 200 µl Complete Freund's Adjuvant (CFA) at the indicated time points, followed by two i.p. injections of 200 ng pertussis toxin (PTX) in phosphate buffered saline (PBS), immediately after immunization and 24 hr later. From day 8 on, paralysis was assessed using the clinical scoring system: 0, no obvious signs of disease; 0.5, distal paralysis of the tail; 1, complete tail paralysis; 1.5, mild paresis of one or both hind legs; 2, severe paresis of both hind legs; 2.5, complete paralysis of one hind leg; 3, complete paralysis of both hind legs; 3.5, complete paralysis of hind legs and paresis of one front leg. Animals with scores of 2 and up were provided access to food and water on the bottom of the cage.

**T and B Cell Isolation**
CD4+ T cells for Q-PCR analysis were purified from lymph nodes of C57BL/6 WT, × S1pr1/+xCdx4-cre, Bmal1flloxflloxxCd4-cre or littermate control mice using the EasySep Mouse CD4+ T cell enrichment kit (STEM-Cell Technologies). Purity was confirmed via flow cytometry and was generally > 96%. B220+ B cells for western blot analysis were purified from spleens of Bmal1flloxflloxCd19-flcre or littermate control mice using the EasySep Mouse B cell enrichment kit (STEMCell Technologies). The Pan T Cell Isolation Kit (Miltenyi Biotech) was used to purify total T cells from mouse splenocytes. Splenocytes were labeled with a mixture of biotin-conjugated monoclonal antibodies directed against cells other than T cells and followed by conjugate binding with anti-biotin microbeads. Magnetically labeled non-T cells were depleted with autoMACS (Miltenyi Biotech) and the negative fraction was collected, yielding an average 90% purity of (CD3+) T cells.

**Antibodies, Western Blotting, Mass Spectrometry Analyses, Quantitative Real-Time PCR, Histology, T Cell Proliferation Analyses, Helicobacter Pylori Infection and Influenza A Virus Infection**
See Supplemental Experimental Procedures.

**Statistical Analyses**
All data are represented as mean ± SEM. Comparisons between two samples were performed using the paired and unpaired Student’s t tests or Mann-Whitney test. One-way ANOVA analyses followed by Tukey’s multiple comparison test, two-way ANOVA analyses followed by Bonferroni’s post hoc test were used for multiple group comparisons. Statistical analyses were performed with GraphPad Prism 6 software. Statistical significance was assessed as *p < 0.05, **p < 0.01, ***p < 0.001, and ****p < 0.0001.

**SUPPLEMENTAL INFORMATION**
Supplemental Information includes six figures and Supplemental Experimental Procedures and can be found with this article online at http://dx.doi.org/10.1016/j.immuni.2016.12.011.

**AUTHOR CONTRIBUTIONS**
D.D. and O.M. designed and performed experiments, analyzed results, and wrote the manuscript; L.I., U.H., R.H., W.H., A.H.T., A.L., M.E., C.-S.C., K.K., A.D., S.M.H., N.K., O.U., L.Y., and L.E.S. performed experiments and provided valuable inputs on the manuscript; C. Schmal and H.H. performed modeling analyses; B.K. provided access to mass spectrometers; W.S. and H.O. designed and supervised experiments, discussed data, and wrote the manuscript; C. Scheiermann conceived and supervised the study, designed and performed experiments, discussed data, and wrote the manuscript.

**ACKNOWLEDGMENTS**
We thank Reinhold Förster and Leandro Moschovakis for providing Ccr7−/− mice, Steffen Masberg, Markus Sperandio, and Martha Merrow for critical comments on the manuscript and Victor Lavilla for designing the video. This work was supported by the German Research Foundation (DFG) (Emmy-Noether grant (SCHL 1645/2-1) and SFB914 projects B09 and Z03 to C. Scheiermann, SFB914 project B05 to R.H., SFB654 project C08 to W.S. and H.O., TR128 project B10 and a Heisenberg fellowship to N.K., SFB-TR84 project C08 to L.E.S., C. Schmal and H.H. are supported by BO 3612/2-1. C. Scheiermann holds a European Research Council (ERC) starting grant (835872, CIRCODE). H.O. is a Lichtenberg fellow of the Volkswagen...
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Supplemental Information

Lymphocyte Circadian Clocks

Control Lymph Node Trafficking

and Adaptive Immune Responses

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Figure S1, related to Figure 1

Oscillatory T lymphocyte subpopulations in blood and lymph node

(A-C) Gating strategy (gated on live cells) (A) and counts of naïve, central memory (TCM) and effector memory (TEM) CD4 and CD8 T lymphocytes in (B) blood (n=3-5 mice) and (C) lymph node (n= 6-18 mice); one-way ANOVA. (D-F) Counts of CD4 and CD8 T cells and B cells in (D) mesenteric lymph node (n=7-11 mice) (E) superficial cervical lymph node (n=4-7 mice) and (F) axillary lymph node (n=7 mice); one-way ANOVA. (G) Images of lymph node sections harvested at ZT5 and ZT13 and stained with antibodies directed against Ki67, PECAM-1, and Lyve-1. Inset: isotype antibody-stained control sections. Scale bars: 200 µm. (H) Quantification of Ki67+ cells per lymph node section; n=4-5 mice. *p<0.05, **p<0.01, ***p<0.001, ****p<0.0001.
Figure S2

A. CXCR4

B. Cxcr4

C. ICAM-1

D. PNAd

E. BMAL1

F. BMAL1

G. B cell

H. Lymph counts

I. Lymph counts

J. Table:

| Time         | ZT1    | ZT7    | ZT13   | ZT21   |
|--------------|--------|--------|--------|--------|
| Lymph volume (µl) | 72 ± 5 | 70 ± 5 | 68 ± 4 | 79 ± 4 |
| Lymph flow (µl/min) | 1.2 ± 0.1 | 1.2 ± 0.1 | 1.1 ± 0.1 | 1.3 ± 0.1 |
Figure S2, related to Figures 2, 3 and 4

Oscillations of leukocyte promigratory factors and numbers

(A) Oscillations in protein surface expression on lymphocyte subpopulations in lymph nodes; CXCR4, CD11a, CD62L; n=3-5 mice, one-way ANOVA; MFI, mean fluorescence intensity. (B) Q-PCR analysis of LN Cxcr4 over 24h; n=3-5 mice, one-way ANOVA. (C-D) Quantification of expression of ICAM-1 (C) and PNAd (D) on HEV over 24h in constant darkness (CT, circadian time: the corresponding light and dark phase are indicated); n=3 mice, one-way ANOVA. (E) Amount of BMAL1 protein in isolated splenic T cells from T cell specific Bmal1−/− and control animals; n = 3 mice, each lane represents an individual animal. (F) Amount of BMAL1 protein in isolated splenic B cells from B cell specific Bmal1−/− and control animals, each lane represents a technical replicate. (G) Lymph node B cell counts in control and B-cell-specific Bmal1−/− mice; n=3-4 mice, unpaired student’s t-test. (H-I) Counts of naïve and central memory (TCM) CD4 and CD8 T lymphocytes in lymph; n=5-17 mice, one-way ANOVA. (J) Harvested lymph volume and flow rate over 24h; n=7-22 mice. *p<0.05, **p<0.01, ***p<0.001.
Figure S3

A

Lymph node: Endogenous cells

Lymph node: Adoptively transferred cells

C

Lymph: Endogenous cells

D

Lymph: Adoptively transferred cells
**Figure S3, related to Figure 4**

**Analysis of oscillations in lymph node half life**

Change in cellular numbers (in % (A-C)) or absolute numbers (D) of endogenous (A and C) and adoptively transferred (B and D) leukocytes in lymph node and lymph over 24h after block of leukocyte homing. Lymph node: n= 3-10 mice; lymph: n=3-6 mice, unpaired student’s t-test. *p<0.05, **p<0.01.
For the sake of simplicity, it was assumed that both homing and egress rhythms follow a sinusoidal oscillation with a certain amplitude and phase. The individual phases were termed $\Phi_h$ and $\Phi_e$, respectively. Let $x(t)$ be the time-dependent lymphocyte count in the lymph node:

$$\frac{dx(t)}{dt} = g(t) - l(t)$$  \hspace{1cm} (1)

Here, the time-dependent gain rate is represented by $g(t)$ and can be summarised as a non-negative sinusoidal function of the form:

$$g(t) = A(1 + \sin(\omega t + \Phi_h))$$  \hspace{1cm} (2)

where $A$ is amplitude and $\omega$ is angular frequency. The time-dependent loss rate is represented by $l(t)$ and can be replaced by a non-negative sinusoidal function of the form:

$$l(t) = d(1 + \sin(\omega t + \Phi_e)) x(t)$$  \hspace{1cm} (3)

where $d$ is decay rate and other nomenclature remains the same.

Using these formulae, a model for circadian homing with the egress replaced with a simple exponential decay rate can be summarised as:

$$\frac{dx(t)}{dt} = A(1 + \sin(\omega t + \Phi_h)) - dx(t)$$  \hspace{1cm} (4)

A model for circadian egress with a fixed gain rate can be summarised as:

$$\frac{dx(t)}{dt} = A - d(1 + \sin(\omega t + \Phi_e)) x(t)$$  \hspace{1cm} (5)

A model where both homing and egress oscillate can be summarised as:

$$\frac{dx(t)}{dt} = A(1 + \sin(\omega t + \Phi_h)) - d(1 + \sin(\omega t + \Phi_e)) x(t)$$  \hspace{1cm} (6)

Model parameters were estimated by fitting to time series data obtained under steady state and blocked homing conditions. Time series data obtained under a blocked egression was spared from the fitting and taken as a cross-validation. Comparisons of modelled and actual data in steady state conditions are shown in Figure 4G (total cell count) and below (individual subsets).

The relative quality of the models given our set of data was tested by means of the Akaike information criterion (AIC), using the formula commonly used for regression models. Its value can be given by:

$$AIC = N \ln(X^2/N) + 2K$$  \hspace{1cm} (7)

where $N$ is the number of data points, $K$ is the number of parameters plus one, and $X^2$ is the residual sum of squares. Models with smaller numbers of AIC values are assumed to outperform those with higher values.
Figure S4, related to Figure 4
Mathematical modeling of cellular oscillations in lymph nodes
Formulas and fits for optimal modeling parameter sets (curves) vs. actual data sets (points) for steady-state oscillations of CD4 and CD8 T cells and B cells using the dual oscillation mathematical model that incorporates rhythmic homing to LNs and rhythmic egress from LNs. The ZT1 time point has been double-plotted to facilitate viewing.
Figure S5

A

S1pr2

Relative mRNA expression

S1pr3

Relative mRNA expression

S1pr4

Relative mRNA expression

S1pr5

Relative mRNA expression

B

Lymph count (x10^3/µl)

ctrl FTY720

ctrl SEW2871

D

Sphk1

Relative mRNA expression

Sphk2

Relative mRNA expression

Sgpl1

Relative mRNA expression

C

S1pr1

Relative mRNA expression

ns****
Figure S5, related to Figure 5

Diurnal expression profiles of S1pr genes

(A) Q-PCR analysis of LN S1pr2, S1pr3, S1pr4 and S1pr5 over 24h; n=3-5 mice, one-way ANOVA. (B) Lymph counts in control animals or animals treated with FTY720 or SEW2871; n=3-6 mice, unpaired student’s t-test. (C) Q-PCR analysis of S1pr1 in CD4 T cells isolated from S1pr1^+/xCd4-cre mice over 24h; n=3 mice, one-way ANOVA. (D) Q-PCR analysis of LN Sphingosine kinase 1 and 2 (Sphk1 and Sphk2) and Sphingosine-1-phosphate lyase (Sgpl1) over 24h; n=3-5 mice, one-way ANOVA. *p<0.05, **p<0.01, ***p<0.001.
Figure S6, related to Figure 6
Diurnal rhythmicity in experimental autoimmune encephalomyelitis (EAE)

(A) Gating strategy (gated on live singlet cells) of migratory DCs. (B) Proliferation of MOG-specific T cells as analyzed by in vivo immunization of mice at ZT8 and ZT20, followed by dye-dilution analyses after re-stimulation with the antigen in culture; n=10 mice, unpaired student’s t-test. (C) Diurnal profiles of inguinal lymph node counts of total CD8 T cells, naïve CD8 T cells, and activated CD8 T cells on day 2 after EAE induction; n=4 mice. Black and white bars indicate the dark and light periods, respectively. (D) Naïve and activated CD4 and CD8 T cell counts in inguinal lymph nodes sampled at ZT15 on the second day after EAE induction; n=4-5 mice, unpaired student’s t-test. (E) Lymph node counts of mice that were infected at different times of the day (ZT1, ZT7 or ZT13) with Helicobacter pylori and harvested at one time 21 days post infection (dpi); n=4 mice, one-way ANOVA. (F) Pulmonary CD8+IFNγ+ T cells of mice infected with influenza A virus at different times of the day (ZT8 and ZT20). Lungs were harvested 8 dpi and infiltrating T cells were enumerated and IFNγ production was assessed by flow cytometry; n=4 mice, Mann-Whitney test. *p<0.05, **p<0.01.
| Primers | Sequences | Annealing (°C) |
|---------|-----------|---------------|
| Bmal1_F | AGA GGT GCC ACC AAC CCA TA | 62 |
| Bmal1_R | TGA GAA TTA GGT GTT TCA GTT CGT CAT | 62 |
| Bmal1_F | CCT AAT TCT CAG GGC AGC AGA T | 60 |
| Bmal1_R | TCC AGT CTT GGC ATC AAT GAG T | 60 |
| Clock_F | CAA AAT GTG AGC AGC ACT TAA TGC | 62 |
| Clock_R | ATA TCC ACT GCC CTT TGG | 62 |
| Per1_F  | TGA GAG CAG CAA GAG TAC AAA CTC A | 60 |
| Per1_R  | TCT GCA CTC AGG AGG CTG TAG | 60 |
| Per2_F  | GTC CAC CTC CCT GCA GAC AA | 60 |
| Per2_R  | TCA TTA GCC TTC ACC TGC TTC AC | 60 |
| Cry1_F  | CTC GGG TGA GGA GTT TTT CT | 62 |
| Cry1_R  | GAC TCC TCT TAC CAG TAG CT | 62 |
| Nr1d1_F | GAT AGC TCC CCT TCT GCA TCA TC | 60 |
| Nr1d1_R | TTC CAT GGC CAC TTG TAG ACT TC | 60 |
| Nr1d1_F | AGC TCA ACT CCC TGG CAC TTA C | 60 |
| Nr1d1_R | CTT TTC GGA ATG CAT GTT GTT C | 60 |
| Dbp_F   | AAT GAC CTT TGA ACC TGA TCC CGC T | 60 |
| Dbp_F   | GCT CCA GTA CTT TTC ATC CTT CTG T | 60 |
| S1pr1_F | CGG TGT AGA CCC AGA GTC CT | 64 |
| S1pr1_R | AGC AGC AGA TGA GAA TGA AC | 64 |
| S1pr2_F | ATG GGC GGC TTA TAC TCA GAG | 62 |
| S1pr2_R | GGC CAG CAC AAG ATG ATG AT | 62 |
| S1pr3_F | TTC CCG ACT GCT CTA CCA TC | 62 |
| S1pr3_R | CCA ACA GGC AAT GAC CAC AC | 62 |
| S1pr4_F | TGC GGG TGG CTG AGA GTG | 62 |
| S1pr4_R | TAG GAT CAG GGC GAA GAC C | 62 |
| S1pr5_F | CTT AGG AGC CCT GGA AAC C | 62 |
| S1pr5_R | CCC GCA CCT GAC AGT AAA TC | 62 |
| Ccr7_F  | TCA TTG CCG TGG TGG TAG TCT TCA | 62 |
| Ccr7_R  | ATG TTG AGC TGC TTG CTG GTT TCG | 62 |
| Cxcr4_F | TCA GTG GCT GAC CTC CTC TT | 60 |
| Cxcr4_R | CTT GGC CTT TGA CTG TTG GT | 60 |
| Ccl21_F | ACA GGC GCC TCC AGA AGA ACA GCG G | 64 |
| Ccl21_R | CGT GAA CCA CCC AGC TTG A | 64 |
| Gapdh_F | TGT GTC CGT GTG GGA TCT GA | 60 |
| Gapdh_R | CCT GCT TCA CCA CCT TCT TGA | 60 |
| Eef1a_F | TGC CCC AGG ACA CAG AGA CTT CA | 60 |
| Eef1a_R | AAT TCA CCA CCA CCA GCA A | 60 |
| Sphk1_F | GAAGACCTGCTCATCAACTGC | 60 |
| Sphk1_R | GTCGCCACTGTAACAGA | 60 |
| Sphk2_F | TAGATGGGGAGTTAGGGAGGATATG | 60 |
| Sphk2_R | TGCTTTTAGGCTCGTTTACGG | 60 |
| Sgpl1_F | TGATGGCCTCACAAGGCTT | 60 |
| Sgpl1_R | GCCACAATTTCTGGAGTTTTGA | 60 |
| Il2_F   | AACCTGAACACTCCCAAGGAT | 60 |
| Il2_R   | CGCAGAGGTCACAAGTTTTCATC | 60 |

Q-PCR primer sequences, related to Figure 2, 3, 5, 6, S2, S5
**Movie S1**

Movie showing the circadian migration of lymphocytes through lymph nodes. At night onset, increased homing due to higher amounts of lymphocyte CCR7 leads to enhanced lymphocyte counts in the lymph node. During the day, higher S1pr1 expression induces the eggression of lymphocytes into efferent lymph.

**SUPPLEMENTAL EXPERIMENTAL PROCEDURES**

**Antibodies**

Antibodies targeting the following antigens were used: CD3e (clone 145-2C11; Biolegend), CD3 (17A2; Biolegend), CD4 (GK1.5; Biolegend), CD8a (53-6.7; Biolegend), CD16/32 (2.4G2; BD Biosciences), CD44 (IM7; Biolegend), CD45 (30-F11; Biolegend), CD62L (MEL-14; Biolegend), B220 (RA3-6B2; Biolegend), NK1.1 (PK136; Biolegend), Gr-1 (RB6-8C5, Biolegend), CD11b (M1/70; Biolegend), CD11c (N418; Biolegend), MHCI I-A/I-E (M5/114.15.2, Biolegend); CCR7 (4B12; Biolegend), CXCR4 (L276F12; Biolegend), CD11a (M17/4; Biolegend), IL-17A (TC11-18H10, BD Horizon), CD9d (R1-2, Biolegend), Ki67 (SolA15; eBioscience), PECAM-1 (Mec13.3; Biolegend), Lyve-1 (ALY7; eBioscience), CCL21 (goat anti-mouse polyclonal; R&D Systems), ICAM-1 (YN1/1.7.4; Biolegend), PNA (MECA-79, Biolegend), blocking antibodies against CD49d (PS/2) and CD11a (M17/4; both BioXCell), BMAL1 (NB 100-2288; Novus Biologicals), beta-tubulin (ab6046; Abcam).

**Mass spectrometry analyses**

Proteins of 20 µL sample matrix were precipitated by adding 200 µL methanol including D-erythro-sphingosine-1-phosphate C17 base (860641P, Avanti Polar Lipids) as internal standard. After centrifugation the supernatant was used for the analysis of S1P by liquid chromatography mass spectrometry (LC-MS/MS) analyses with a 1200 SL HPLC system (Agilent) coupled to a 4000QTRAP tandem mass spectrometer (AB Sciex). Separation was achieved with a Zorbax SB-C18, particle size 3.5 µm, 150 mm × 2.1 mm HPLC column (Agilent) and water with 0.1% formic acid as mobile phase A and methanol/ isopropanol (1:1) with 0.1% formic acid as mobile phase B. A gradient elution was used for optimal separation from 60% B to 80% B within 7 minutes. After column cleaning at 100% B for 1 minute, the gradient was equilibrated for 2.5 min at 60% B before the next injection. The LC-MS/MS system operated in positive electrospray ionization with electron voltage of 5500V at 400 °C. Auxiliary gas was set to 50 psi and 60 psi was used for nebulizer gas. Curtain gas was set to 30 psi and collision gas was set to 8 psi. Mass transitions and individual energies (decluster potential, collision energy, collision cell exit potential) were optimized by direct infusion and set as follows: m/z 380.2 → 264.2 (66, 23, 18) and 380.2 → 82.1 (66, 49, 6) for S1P and 366.2 → 250.2 (61, 23, 18) and 366.2 → 82.0 (61, 45, 6) for internal standard. Mass transitions were detected in multiple reaction monitoring mode with a dwell time of 100 ms. Quantification was achieved by standard calibrating curves prepared from D-erythro-sphingosine-1-phosphate C18:1 base (860492P, Avanti Polar Lipids) at different concentration points in the range from 0.005 to 0.5 µmol/L.
**Western blotting**

To obtain total cell lysates, MACS-purified T cells were incubated with RIPA buffer (150 mM NaCl; 50 mM Tris-HCl, 0.1 % SDS, 0.5 % Na-Dox, 0.1 % SDS, 1 % NP40, 1 mM EDTA, and proteinase inhibitor). Nuclei were pelleted in a microfuge at 14,000 x g for 15 min, and the supernatant was used as the cytoplasm fraction. Cell lysates were normalized by measuring total protein concentrations using the Pierce BCA Protein Assay Kit (Thermo Scientific) according to the manufacturer's protocol. Proteins were separated on a 10 % SDS-PAGE and transferred to PVDF membranes (Roche). Membranes were blocked in 5 % milk powder for 1 h and then probed with anti-BMAL1 antibody (1:2000 in 5 % milk-TBS; NB 100-2288, Novus Biologicals, Littleton, USA) overnight at 4 °C. Anti-β-Tubulin antibody (Abcam) was used for normalisation. HRP-conjugated anti-rabbit or anti-mouse IgGs were used as secondary antibodies. Proteins were visualized by enhanced chemiluminescence (Thermo, No. 34080) and a Fusion SL Image Acquisition System (Vilber Lourmat).

**Histology**

Mice were perfused intracardially with ice-cold PBS followed by fixation with 4% paraformaldehyde (PFA) for 10 minutes. Spinal cords were post-fixed overnight and then dissected from the spinal canal. Paraffin embedded 10-µm transverse sections of the lumbar spinal cord were stained for total myelin with Luxol Fast Blue Stain/ Cresyl Echt Violet (Abcam) according to the manufacturer’s instructions, dehydrated, cleared with xylene, and mounted for microscopy. Images of lumbar spinal cord were taken at levels L1/2, L3/4 and L5/6 for each mouse and demyelinated areas measured using ImageJ software (National Institutes of Health).

**Quantitative real-time PCR**

RNA was extracted using RNeasy Plus Mini Kit (Qiagen) or with TRIzol reagent (Life Technologies), in accordance with the manufacturers’ instructions. Reverse transcription was carried out using the High Capacity cDNA Reverse Transcription Kit (Applied Biosystems), in accordance with the manufacturer’s instructions. Q-PCR was performed with SYBR GREEN on a StepOnePlus Real-Time PCR System (Applied Biosystems). A primer concentration of 0.5µM was found to be optimal in all cases. The sequences of the oligonucleotides used are included in the Supplemental Experimental Procedures. The PCR protocol consisted of one cycle at 95 °C (10 min) followed by 40 cycles of 95 °C (15 s) and 60-64 °C (1 min). Expression of glyceraldehyde-3-phosphate dehydrogenase (GAPDH) was generally used as a standard. For EAE experiments Q-PCR was conducted with the GoTaq Q-PCR Master Mix (Promega) and a CFX96 Real-Time PCR Detection System (Bio-Rad) according to the manufacturer’s instructions, along with gene-specific primers. Eef1a was used as reference gene. Relative quantification was performed using the ΔΔCT method.

**T-cell proliferation analyses**

Spleens were harvested from mice 10 days after EAE immunization. CD3+ T cells were negatively selected using the Pan T Cell Isolation Kit II (Miltenyi Biotec) and labelled with a proliferation dye (CellTrace™ Violet Cell Proliferation Kit, Life Technologies). T-cell depleted splenocytes (1 × 10^5/well) isolated from EAE mice were
cultured in the presence or absence of the MOG peptide (10 µg/ml) or irrelevant peptide (ovalbumin) (10 µg/ml) or no antigen at 37°C in 5% CO2 for 1 h. CD3+ T cells were co-cultured with antigen-pulsed splenocytes at 37°C in 5% CO2 for 3 days RPMI medium 1640 supplemented with 10% FCS, 10mM HEPES and 55µM beta-mercaptoethanol in 96-well flat-bottom plates, at 2×10^5 cells per well.

**Helicobacter pylori infection**
Mice were infected orogastrically at ZT1, ZT7 and ZT13 with the H. pylori strain PMSS1wt with a dose of 10^9 bacteria. Animals were sacrificed after 3 weeks and LN cellularity was analyzed by flow cytometry.

**Influenza A virus infection**
Female 8-10 week old C57B/6 (CD45.1+) mice were intranasally infected with 100 plaque forming units (pfu) of Influenza A virus (strain A/Puerto Rico/8/1934 H1N1) at ZT8 and ZT20. Animals were sacrificed on day 8 post infection and lymphocyte infiltration and activation was assessed in lung tissue homogenates by flow cytometry. Animal experiments were approved by the local animal ethics committee (LAGeSo).