Low body temperature governs the decline of circulating lymphocytes during hibernation through sphingosine-1-phosphate


Hibernation is an energy-conserving behavior consisting of periods of significantly inhibited metabolism (torpor) that result in a largely reduced heart and ventilation rate (1–3) and body temperature. Torpor bouts are interspersed by arousal periods with durations of 8–24 h, during which metabolism increases and body temperature rapidly returns to euthermia (2, 4). Hibernating mammals display major changes in their physiology (torpor), because administration of a specific antagonist (W146) during torpor (in a Syrian hamster at ∼8 °C) precluded restoration of lymphocyte numbers upon subsequent arousal. Furthermore, S1P release from erythrocytes via ATP-binding cassette (ABC)-transporters was significantly inhibited at low body temperature (4 °C) but was restored upon rewarming. Reversible lymphopenia also was observed during daily torpor (in a Djungarian hamster at ±25 °C), during forced hypothermia in anesthetized (summer-active) hamsters (at ±9 °C), and in a nonhibernator (rat at ±19 °C). Our results demonstrate that lymphopenia during hibernation in small mammals is driven by body temperature, via altered plasma S1P levels. S1P is recognized as an important bioactive lipid known to regulate lymphocyte egress from lymph nodes (23–25). To examine mechanisms of immunological alterations in hibernation, changes in the number of circulating leukocytes were examined in different stages of hibernation in hamster species that undergo either deep multiday torpor bouts or shallow daily torpor. Effects were compared with those found in hamsters cooled under anesthesia. Specifically, regulation of lymphocyte numbers was studied because of their importance in adaptive immunity and immunological memory. After confirmation of lymphocyte storage in secondary lymphoid tissue during torpor, we further examined the role of sphingosine-1-phosphate (S1P), a bioactive lipid known to regulate lymphocyte egress from lymph nodes (23–25).

Results

After entrance into deep torpor, the body temperature of the Syrian hamster (Mesocricetus auratus) decreases to 7.7 ± 0.8 °C during the first 24 h (Fig. 5C). During torpor, the total number...
of circulating white blood cells decreases by 95%, from 1.57 ± 0.22 to 0.07 ± 0.01 x 10^6 mL. The number of circulating lymphocytes decreases even more dramatically to about 4% of the number in summer-euthermic animals (Fig. 1 A and B). Lymphocyte numbers are restored rapidly upon arousal (Fig. 1C), when body temperature increases to reach euthermia within 2 h (Fig. S1B). The blood lymphocyte count correlates significantly with the body temperature both during entrance into torpor (Pearson R^2 = 0.41; P < 0.01) (Fig. 1B) and during arousal following torpor (Pearson R^2 = 0.71; P < 0.01) (Fig. 1C). The number of circulating erythrocytes does not change throughout the torpor–arousal cycles (Fig. S2). In an animal showing daily torpor behavior, the Djungarian hamster (Phodopus sungorus), the average body temperature during torpor is 25.2 ± 1.3 °C. During these torpor bouts, the number of circulating leukocytes was reduced by about 50%, whereas the number of circulating lymphocytes decreased by ∼30% (Fig. 1D). To examine the role of body temperature in the decrease of circulating lymphocytes, forced hypothermia was induced in anesthetized (summer-active) hamsters that were not in hibernation to reach a body temperature of 9.1 ± 0.8 °C. Forced hypothermia decreased the number of circulating leukocytes and lymphocytes (Fig. 1B), and the decrease was fully reversed upon rewarming (Fig. 1C). Furthermore, the number of circulating lymphocytes correlated significantly with body temperature during cooling (Pearson R^2 = 0.67; P < 0.01) (Fig. 1B) as well as during rewarming following forced hypothermia (Pearson R^2 = 0.29; P < 0.01) (Fig. 1C). The number of circulating lymphocytes in summer-euthermic animals that underwent forced hypothermia was, at the start of the experiment, higher than in hibernating animals before entering torpor (Fig. 1B) (P < 0.01). The number of circulating lymphocytes was significantly higher in euthermic (summer-active) animals than in aroused (winter-euthermic) animals (P < 0.05) (Fig. S3). Moreover, in rats (Rattus norvegicus), a nonhibernating animal, forced hypothermia to reach a body temperature of 19.2 ± 0.7 °C did not affect the number of circulating erythrocytes (Fig. S4A) but resulted in a significant decrease in the number of circulating lymphocytes (P < 0.01) (Fig. S4B); this decrease was restored upon rewarming (P < 0.01) (Fig. S4C).

To establish the role of the spleen in the regulation of lymphocyte numbers during hibernation, we measured the expression of markers for lymphocytes and performed splenectomies before and during torpor. mRNA expression of the T-lymphocyte marker CD3ε and the B-lymphocyte marker CD20 is unaffected by hypothermia by torpor (Tables S1 and S2). Although this measurement does not directly demonstrate the number of cells present in the spleen, we have no indication that mRNA expression of these specific markers changes during torpor. Hence, the similar expression levels of these lymphocyte markers during torpor and arousal strongly argue against massive cellular migration into the spleen during torpor or substantial apoptosis of these cells. Surgical removal of the spleen (splenectomy) preceding the hibernation season does not influence the induction of lymphopenia during torpor (Fig. 1E). Conversely, splenectomy during torpor does not preclude restoration of lymphocyte numbers during arousal (Fig. 1F). To detect retention sites of lymphocytes other

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**Fig. 1.** Depletion of circulating lymphocytes during torpor is temperature dependent and unaffected by splenectomy. (A) Normal blood lymphocyte count of summer-euthermic Syrian hamster. (B) Body temperature-dependent decrease in blood lymphocytes upon entrance into torpor and to a similar extent in forced hypothermia in nonhibernating animals. (C) Circulating lymphocyte counts are restored rapidly during rewarming from torpor and from forced hypothermia. (D) Circulating lymphocytes are restored rapidly upon arousal (Fig. 1C). The blood lymphocyte count correlates significantly with the body temperature during cooling (Pearson R^2 = 0.67; P < 0.01) (Fig. 1B) as well as during rewarming following forced hypothermia (Pearson R^2 = 0.29; P < 0.01) (Fig. 1C). The number of circulating lymphocytes in summer-euthermic animals that underwent forced hypothermia was, at the start of the experiment, higher than in hibernating animals before entering torpor (Fig. 1B) (P < 0.01). The number of circulating lymphocytes was significantly higher in euthermic (summer-active) animals than in aroused (winter-euthermic) animals (P < 0.05) (Fig. S3). Moreover, in rats (Rattus norvegicus), a nonhibernating animal, forced hypothermia to reach a body temperature of 19.2 ± 0.7 °C did not affect the number of circulating erythrocytes (Fig. S4A) but resulted in a significant decrease in the number of circulating lymphocytes (P < 0.01) (Fig. S4B); this decrease was restored upon rewarming (P < 0.01) (Fig. S4C). To establish the role of the spleen in the regulation of lymphocyte numbers during hibernation, we measured the expression of markers for lymphocytes and performed splenectomies before and during torpor. mRNA expression of the T-lymphocyte marker CD3ε and the B-lymphocyte marker CD20 is unaffected by hypothermia by torpor (Tables S1 and S2). Although this measurement does not directly demonstrate the number of cells present in the spleen, we have no indication that mRNA expression of these specific markers changes during torpor. Hence, the similar expression levels of these lymphocyte markers during torpor and arousal strongly argue against massive cellular migration into the spleen during torpor or substantial apoptosis of these cells. Surgical removal of the spleen (splenectomy) preceding the hibernation season does not influence the induction of lymphopenia during torpor (Fig. 1E). Conversely, splenectomy during torpor does not preclude restoration of lymphocyte numbers during arousal (Fig. 1F). To detect retention sites of lymphocytes other
Plasma levels of S1P decreased by ∼50-60% during deep torpor in Syrian hamsters and during daily torpor in Djungarian hamsters (Fig. 3 A and B). During arousal, plasma S1P levels rose rapidly to normal (euthermic) values in both species. Involvement of the S1P system in hibernation-associated lymphopenia was examined further by intracardial injection of a specific type 1 S1P receptor (S1P1) antagonist (W146; 5 mg/kg). No significant difference in the number of lymphocytes was found in torpid animals and aroused animals that were treated with W146, but the number of circulating lymphocytes in the subsequent arousal period was significantly lower following intracardial injection of W146 than in vehicle-treated animals (P < 0.01) (Fig. 3C). In contrast, W146 does not affect the restoration of the number of circulating neutrophils (Fig. 3D).

Upon arousal, the intracellular level of S1P in erythrocytes decreased (Fig. S5A) without a significant change in the intracellular level of sphingosine (Fig. S5B). Plasma S1P and intracellular S1P levels of erythrocytes correlate negatively in torpor and arousal periods (Pearson $R^2 = -0.77$, $P < 0.001$; Fig. 3E). To establish the influence of body temperature on the release of S1P from erythrocytes, washed erythrocytes derived from torpid animals were rewarmed to 37 °C. Although erythrocytes maintained at 4 °C do not release S1P into the medium, ex vivo rewarmed inducing substantial S1P release. Release of S1P from rewarmed erythrocytes is reduced significantly by inhibitors of ATP-binding cassette (ABC)-A1 (glyburide) and ABC-C1 (MK571) transporters and is reduced even further when both inhibitors are combined (Fig. 3F).

Discussion

Our data imply that body temperature is the driving force of lymphopenia during hibernation as shown by (i) the strong correlation between body temperature and blood lymphocyte count, (ii) the occurrence of lymphopenia during both deep and daily torpor, and (iii) the effect of forced cooling of (summer-active) hamsters that were not in hibernation on the number of circulating lymphocytes. At the start of the experiment, however, the number of circulating lymphocytes was significantly higher in animals that underwent forced hypothermia than in hibernating animals that were about to enter torpor. Because no differences were found between summer-euthermic animals that did and did not undergo forced hypothermia, we speculate that this phenomenon is caused by temperature-independent, seasonal changes affecting the number of circulating lymphocytes during euthermia. During daily torpor in the Djungarian hamster, the number of circulating lymphocytes also decreased, in spite of the shorter duration of a torpor bout with a substantially higher body temperature than during deep torpor (2). Although the decrease in the number of circulating lymphocytes during daily torpor was smaller than observed during deep torpor, the fitted curve (Fig. 1B) shows the blood lymphocyte count is ∼40% lower at an average body temperature of ∼25 °C during either natural torpor or forced hypothermia in the Syrian hamster and is about the same as observed during daily torpor in the Djungarian hamster. The temperature dependency of this process is supported further by the fact that hypothermia in rats (nonhibernators) leads to the induction of lymphopenia as well. Because rewarming of severely hypothermic rats affected their breathing, we cooled and rewarmed a second group of ventilated rats to demonstrate the reversibility of lymphopenia induced by hypothermia in rats. In addition, forced hypothermia of anesthetized rats demonstrates that hypothermia-induced lymphopenia is not specific to hibernators but also occurs in nonhibernating animals.

Lymphopenia during hibernation is caused by storage of cells in peripheral lymphoid organs but not in spleen, liver, lung, and kidney. The observation that the number of splenic lymphocytes is stable throughout hibernation and that the lymphopenia during torpor bouts is rapidly reversed upon rewarming strongly favors a storage-and-release mechanism over an apoptosis-and-replenishment mechanism to explain the dynamics of lymphocytes during hibernation. Further, splenectomies preceding the hibernation season demonstrated that the spleen is not necessary for the induction of lymphopenia. Conversely, splenectomy during torpor established that recirculating lymphocytes can be obtained from a source other than spleen. Although we cannot rule out the possibility that (some) lymphocytes are retained in the spleen during torpor, the spleen certainly is not the main organ involved. Indeed, injected fluorescently labeled lymphocytes are found in cervical lymph nodes during the subsequent torpor bout and, to a lesser extent, also in GALT.

Inkovaara et al. (26) previously showed that number of leukocytes increases in lung (mainly neutrophils) and gut (mainly lymphocytes) during torpor compared with summer euthermia or arousal in hibernating hedgehogs (Erinaceus europaeus). The number of intraepithelial lymphocytes and lamina propria lymphocytes increases about threefold, whereas the number of lymphocytes in Peyer’s patches increases only slightly compared with summer euthermia in the thirteen-lined ground squirrel (Ictidomys tridecemlineatus) (27). Because T lymphocytes in the blood are mainly T-cell receptor (TCR)αβ-positive, influx and retention of circulating T lymphocytes into GALT would result in an altered TCRαβ⁺:TCRγδ⁺ ratio. Logically, the authors speculate that the unaffected TCRαβ⁺:TCRγδ⁺ ratio in GALT reflects absence of influx, and thus that the increased lymphocyte count in GALT is caused by local expansion of cells (27). However, our results suggest that there is some influx of circu-
lating lymphocytes into GALT during torpor. Influx of circulating lymphocytes might not alter the TCRαβ:TRCγδ ratio as long as the number of cells that migrate into GALT remains relatively small compared with the number of cells already present in GALT. Therefore, the increased number of lymphocytes in GALT might well result from a combination of influx and local expansion of cells. Taken together, these observations show that secondary lymphoid tissue is the main site for the retention of lymphocytes during torpor.

In our study, we demonstrate direct effects of S1P on the restoration of the number of circulating lymphocytes. Because lymphocytes continuously recirculate among various lymphoid organs via the blood, the number of circulating lymphocytes depends on the balance between influx from the blood into peripheral lymphoid organs (homing) and egress out of these organs (as mediated by S1P). Logically, a decreased S1P plasma level, such as occurs during torpor, leads to a reduced egress from peripheral lymphoid organs and consequently to a lymphopenic state. Our data demonstrate that the retention of lymphocytes in peripheral lymphoid organs is regulated by plasma S1P levels acting via S1P1. Although the effect of S1P on lymphocyte egress from peripheral lymphoid organs has been demonstrated previously in knockout models or by using synthetic agonists and antagonists (23–25, 28), our study demonstrates the importance of S1P in regulating lymphocyte numbers in the absence of pharmacological interventions or genetic changes. Lymphocyte egress from peripheral lymphoid organs normally is favored when the S1P concentration is low in lymphoid tissue interstitium and is high at exit sites, i.e., in blood (23).

Although the extent of lymphopenia is larger during deep torpor, the relative decrease in S1P is larger during daily torpor. Other factors, such as time, blood flow velocity, and the level of expression of S1P receptors, which might differ among species, influence the egress of lymphocytes. Therefore, levels cannot be compared directly between species. However, during both deep and daily torpor the number of circulating lymphocytes correlated significantly with the S1P plasma level. Involvement of the S1P system is substantiated by (i) the rapid change in plasma S1P levels, (ii) the correlation between plasma S1P level and blood lymphocyte count, and (iii) the observation that blockade of S1P1 during arousal completely blocks the restoration of circulating lymphocytes. Thus, the blockade of lymphocyte egress by antagonism of the S1P1 receptor is a causative mechanism in the restoration of circulating lymphocyte numbers during arousal. Because blockade of S1P1 did not affect the restoration of the number of circulating neutrophils and monocytes or change erythrocyte counts, a nonspecific action of S1P1 blockade seems highly un-

![Fig. 3. S1P governs lymphocyte dynamics during hibernation through the S1P1 receptor. (A and B) Plasma S1P levels are reduced significantly in (A) deep torpor (Syrian hamster) and (B) daily torpor (Djungarian hamster). (C and D) Upon arousal, the specific S1P1 receptor antagonist W146 precludes the restoration of circulating lymphocytes but not of neutrophils compared with vehicle-treated animals. AR, arousal; TO, torpor. (E) Significant correlation between the level of S1P in erythrocytes and plasma of animals in torpor (closed circles) and arousal (open circles). (F) Temperature governs S1P release from erythrocytes isolated from torpid animals. Erythrocytes were obtained from torpid animals and washed with PBS (pre); continued incubation for 30 min at 4°C (gray bars) did not induce release of S1P from erythrocytes; ex vivo rewarming for 30 min at 37°C (hatched bars) induces release of S1P that is blocked by the ABC-transporter inhibitors MK571 (50 μM), glyburide (1 mM), and by MK571 plus glyburide. Bars represent mean ± SEM of four to eight animals per group. Groups were compared using student’s t-test or one-way ANOVA and post hoc least significant difference. Different letters above bars represent significant differences at P < 0.05.](image-url)
likely. Together, these data suggest that S1P acting via S1P₁ has a major role in regulating lymphocyte number in peripheral blood during hibernation.

Our data suggest that S1P release from erythrocytes constitutes an important regulatory mechanism in S1P plasma levels. This proposal is consistent with previous observations in sphingosine kinase-deficient mice that the plasma level of S1P is derived mainly from erythrocytes (23). Indeed, erythrocytes derived from torpid animals release S1P upon rewarming ex vivo, thus demonstrating that body temperature is the primary factor that can stimulate release. Furthermore, the strong negative correlation between plasma S1P levels and intracellular S1P content of erythrocytes suggests a prominent role for erythrocytes in regulating the plasma level of S1P. The fact that release of S1P can be reduced by inhibitors of ABC-A1 (glyburide) and ABC-C1 (MK571) transporters and is reduced even further when both inhibitors are combined demonstrates a role for ABC transporters in regulating S1P release from erythrocytes upon rewarming. The combination of both inhibitors demonstrates a role for both ABC-A1 and ABC-C1 transporters in the release of S1P from erythrocytes. Incomplete inhibition, however, might suggest the involvement of other (yet unknown) transporters in the release of S1P from erythrocytes as well. Taken together, the data show that, in the absence of potential regulating factors from plasma, an increase in body temperature is sufficient to induce a rapid and substantial release of S1P from erythrocytes of torpid animals.

Our study identifies the reduction of body temperature resulting from metabolic suppression in torpor as the major driving force in modulating lymphocyte egress from lymphoid tissue that constitutes a versatile system of a rapidly reversible reduction of immune function. The induction of lymphopenia by low body temperature is widely conserved, because it seems to be a common response of mammals (shared by hibernating hamsters, hypothermic hibernating hamsters, and nonhibernating rats). Although this response seems to be common among mammals rather than an adaptation specific for hibernators, it may be beneficial during hibernation. The induction of an immune response during arousal increases the time before animals go back into torpor (7). Because periodic arousals account for 80–90% of the energy used throughout hibernation, an immune response might increase the energy costs of hibernation (29). Because, in general, no massive death of hibernating animals occurs, and most microbes do not proliferate well at low temperatures, we speculate that the energy benefits of a reduced immune function outweigh the risk of infection. However, some pathogens, such as the psychrophilic fungus Geomyces destructans that causes white-nose syndrome in bats, grow well at low temperatures (14, 17–19, 22). Hence, immune suppression during torpor may be detrimental. Unfortunately, reports on leukocyte dynamics during hibernation in bats are lacking. Thus, whether immune suppression might be involved in the etiology of white-nose syndrome remains to be elucidated.

In this study we show that release of S1P from erythrocytes is reduced significantly by low body temperature. S1P may serve a key function during the induction of torpor by regulating additional aspects of the immune system as well as other physiological changes that are not related primarily to immune function (30). Not only does S1P regulate the function of other types of leukocytes (31); it also is involved in other biological processes (32) and is implicated in governing protection against ischemia/reperfusion-induced injury in brain, heart, liver, and kidney (33–36). Therapeutic hypothermia is used in patients with clinical conditions such as cardiac arrest and brain trauma and in patients undergoing cardiac or brain surgery because reducing cerebral metabolism has neuroprotective properties in periods of low oxygen supply (9). However, hypothermia is associated with increased renal injury postoperatively (10). Given that the plasma level of S1P in humans also is regulated mainly by transport of S1P from erythrocytes (37, 38) and the S1P-system exerts protective effects following ischemia/reperfusion in different organs (33–36), our results may be of direct consequence in understanding the benefits and detriments of therapeutic hypothermia.

**Conclusion**

Our study identifies the reduction of body temperature resulting from metabolic suppression in torpor as the major driving force in modulation of lymphocyte egress from lymphoid tissue because of decreased S1P plasma levels caused by inhibition of S1P release from erythrocytes (Fig. S6). Understanding the mechanisms of specific physiological alterations involved in the induction of torpor increases our knowledge of natural hibernation and has relevance for therapeutic hypothermia as well as for pharmacologically induced suspended animation.

**Materials and Methods**

**Hibernation.** To induce hibernation in Syrian hamsters (Mesocricetus auratus), the light-dark (L:D) cycle was shortened to 8 h:16 h for ∼10 wk followed by continuous dim light (∼5 lux) at an ambient temperature of 5 °C. Movement detectors connected to a computer allowed us to determine the animals’ hibernation pattern. In the Djungarian hamsters (Phodopus sungorus), hibernation was induced by shortening the L:D cycle to 8 h:16 h for ∼14 wk at an ambient temperature of 21 ± 1 °C. Daily torpor was determined by observation at the start of the light phase (usual torpor phase) and a single body temperature measurement at the time of decapitation. All experiments were approved by the Institutional Animal Ethical Committees of the University Medical Center Groningen and University of Aberdeen.

**Forced Hypothermia.** Summer-etheric Syrian hamsters and Wistar rats housed at an L:D cycle of 12 h:12 h were anesthetized by injecting 200 mg/kg ketamine and 1.5 mg/kg diazepam i.p. Spontaneously breathing rats were cooled. Spontaneously breathing hamsters and ventilated rats were cooled and rewarmed. Animals were cooled by applying ice-cold water to their fur and were rewarmed using a water-based heating mattress; both processes were performed at a rate of ∼1 °C of body temperature per 3 min. A catheter was inserted into the jugular vein for blood sampling, rectal temperature was measured continuously, and heart rate (ECG) was monitored on the anesthesia monitor Cardiocap 5 (Datex Ohmeda).

**Splenectomies.** Splenectomies were performed on summer-etheric and torpid Syrian hamsters. Immediately after induction of anesthesia (2–2.5% isofluoran/o₃), a blood sample was drawn by cardiac puncture, and 4 mg/kg fluinixin-meglumien (Finadine; Schering-Plough) was given s.c. as analgesia. Summer-etheric animals that underwent splenectomy recovered in a warm room (L:D cycle 8 h:16 h). After induction of hibernation, animals were killed during their third torpor bout, which was 60.3 ± 8.1 d following splenectomy. During surgery in the third torpor bout, torpid animals were kept at <10 °C with ice-packs and were recovered in the climate-controlled room. Animals were killed upon reaching euthermia.

**Statistical Analysis and Data Presentation.** Data are presented as mean ± SEM. Statistical analysis was performed by student’s t-test or one-way ANOVA with post hoc least significant difference (SPSS 16.0 for Windows), with P < 0.05 considered significantly different. Correlations were calculated using Pearson’s correlations. Sigmaplot 11.0 was used to produce the graphs shown in this article.

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Supporting Information

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SI Materials and Methods

Blood Sample Analysis. After animals were killed by i.p. pentobarbital, 250 μL of blood collected via cardiac puncture into EDTA-coated cups (microlon EDTA-K3; Greiner) was analyzed on the Sysmex XE-2100, an automated hematology analyzer (1, 2). Differential leukocyte counts were validated manually using Wright-Giemsa–stained blood smears. The remainder of the blood was collected in a polypropylene tube, mixed 1:10 (vol/vol) with a solution containing prostaglandin E1 (94 nmol/L) (Sigma-Aldrich), Na2CO3 (0.65 mmol/L) (Sigma-Aldrich), EDTA (90 mM) (Titriplex; Sigma-Aldrich), and theophylline (10 mM) (Sigma-Aldrich) to minimize platelet activation. Samples were centrifuged for 30 min at 17,000 × g at 4 °C, snap-frozen in liquid nitrogen, and stored at −80 °C.

Labeling of Lymphocytes. To obtain lymphocytes for autologous transfusion, Syrian hamsters were splenectomized as described in Materials and Methods. A cannula was placed into the jugular vein connected to an s.c. access port (Soluport; Instech Solomon) implanted between the scapulas for infusion of cells during torpor. The spleen was cut into small pieces and washed on a70-μm cell strainer (Greiner) with sterile saline to obtain a suspension of splenocytes. Cells were spun down for 10 min at 800 × g at 4 °C and were resuspended onto a layer of Lympholyte Mammal (Cedarlane Laboratories). Lymphocytes then were separated from other cells by centrifuging for 20 min at 800 × g at 4 °C. The lymphocyte fraction was washed, centrifuged for 10 min at 800 × g at 4 °C, and resuspended in salmine supplemented with 10% DMSO followed by storage in liquid nitrogen. When animals showed torpidity (around 10 wk after splenectomy), lymphocytes were thawed; purity was assessed by analyzing a Wright-Giemsa–stained smear, and viability was measured with a manual cell count after addition of Trypan blue (Sigma-Aldrich). Cells were fluorescently labeled by incubation in 25 μM carboxyfluorescein diacetate succinimidyl ester in saline (Invitrogen) for 15 min at 37 °C. After the labeled lymphocytes were infused during torpor, arousal was induced by gently handling the animals. The animals were killed 2 d after entrance into the subsequent torpor bout.

Fluorescent Microscopy. Frozen samples were embedded in Tissue Tek (Sakura), sectioned into 200-μm-thick slices, and counter stained using TOTO-3 iodide (Invitrogen). Images were taken at 200× magnification (Leica SP2 AOBS) and processed using Imaris 6.4.

Real-Time PCR. RNA isolation was performed using the RNA Isolation Kit (Bioké) according to the manufacturer’s instructions. RNA concentration was determined spectrophotometrically at 260 nm (NanoDrop ND-1000; NanoDrop Technologies), and purity was checked on 1% agarose gel. One microgram of RNA was mixed with 4 μL RT buffer, 0.2 μL dNTP, 0.5 μL Rnasin, 1 μL reverse transcriptase, 1 μL random hexamers (Promega), and H2O in 20 μL. cDNA was produced on a C1000 Thermal Cycler (Bio-Rad Laboratories). Oligonucleotide primer sequences (Biolegio) are shown in Table S1. Amplified products (CFX 384 Real-Time System; Bio-Rad Laboratories) were checked by obtaining melting curves and verification on 1% agarose gels.

Liquid Chromatography-Electrospray Tandem Mass Spectrometry. Sphingolipids were extracted and analyzed by liquid chromatography-electrospray tandem mass spectrometry on a PE-Scex API 3000 triple-quadrupole mass spectrometer equipped with a turbo ion-spray source as described previously (3, 4). HPLC separation was performed as described previously (5), with the following changes: An Alltima C-18 column (2.1 × 150 mm, 5 μm; Grace Davson Discovery Sciences) was used at a flow rate of 200 μL/min. N2 was used as the nebulizing gas and drying gas for the turbo ion-spray source. The ion-spray needle was held at 5,500 V; the orifice temperature was set to 500 °C. N2 was used to induce dissociations collisionally in Q2. Multiple reaction-monitoring scans were acquired by setting Q1 and Q3 to pass the precursor and product ions of the most abundant sphingolipid molecular species. Multiple reaction-monitoring transitions were optimized for each individual component (C-17SoP: 366.2/250.4; C-17SaP: 368.2/270.4; C-18SoP: 380.2/264.4; C-18SaP: 382.2/284.4; C-17So: 286.2/238.1; C-17Sa: 288.2/240.1; C-18So: 300.2/252.3; C18Sa: 302.2/254.2). Quantitation was achieved by spiking the samples before extraction with sphingosine (d17:1), sphinganine (d17:0), sphingosine-1-phosphate (S1P) (d17:1), and sphinganine-1-phosphate (d17:0) (Avanti Polar Lipids, Inc.).

Blood Rewarming ex Vivo. Washed erythrocytes (25 μL) derived from torpid Syrian hamsters (4 °C) were pipetted into small polypropylene tubes (Eppendorf) containing DMEM/F12 cell culture medium (Invitrogen) supplemented with 40% Probumin (Millipore). An inhibitor for the ATP-binding cassette (ABC)-A1 transporters (glyburide, 1 mM; Sigma-Aldrich), the ABC-C1 transporter (MK571, 50 μM; Cayman Chemicals), or both was added to the samples. Samples then were incubated at 37 °C for 30 min. Another sample was left at 4 °C for 30 min. After incubation, samples were centrifuged for 30 min at 17,000 × g at 4 °C. A negative control sample was centrifuged before incubation at 37 or 4 °C. Supernatant was snap-frozen in liquid nitrogen and stored at −80 °C.


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Fig. S1. Average body temperature during (A) torpor entry and (B) arousal in hibernating Syrian hamsters. Graphs demonstrate the time-dependent changes in mean body temperature from 12 torpor–arousal cycles measured by temperature loggers in the animals’ peritoneum.

Fig. S2. Hibernation phase does not affect the number of circulating erythrocytes. Bars represent mean ± SEM of four to six animals per group. Data were analyzed using one-way ANOVA with post hoc least significant difference.

Fig. S3. The number of circulating lymphocytes is reduced significantly in aroused (winter-euthermic) Syrian hamsters compared with summer-euthermic Syrian hamsters. Bars represent mean ± SEM of 11 animals per group. Groups were compared using two-tailed independent samples student’s t-test. *P < 0.05.

Fig. S4. Forced hypothermia of anesthetized rats induces lymphopenia without affecting the number of circulating erythrocytes. (A) Forced hypothermia of anesthetized rats does not affect the number of circulating erythrocytes. (B) Body temperature-dependent decrease in circulating lymphocytes in forced hypothermia in anesthetized spontaneously breathing rats. (C) Body temperature-dependent decrease in circulating lymphocytes followed by restoration upon rewarming (30 min after reaching 37 °C) in anesthetized intubated rats. Bars represent mean ± SEM of five to seven animals per group. Groups were compared using one-way ANOVA and post hoc least significant difference. **P < 0.01.
Fig. S5. Complementary levels of (A) S1P and (B) sphingosine in erythrocytes during different periods of hibernation. Bars represent mean ± SEM of four to six animals per group. Data were analyzed using one-way ANOVA with post hoc least significant difference. */** indicates significant difference at * \( P < 0.05 \); ** \( P < 0.01 \).

Fig. S6. Proposed model of S1P release from erythrocytes governing lymphocyte dynamics during hibernation or hypothermia. Sphingosine is taken up by erythrocytes and is phosphorylated intracellularly by sphingosine kinase to S1P, which is transported into the plasma by ATP-dependent ABC transporters. During torpor, in response to lower temperature, release of S1P from erythrocytes is inhibited. The inhibition lowers S1P plasma levels, affecting the S1P gradient from blood to lymph node (indicated by solid line). In turn, egress of lymphocytes from lymphoid organs is inhibited, resulting in profound lymphopenia. In addition, the low body temperature during torpor may affect ATP-dependent phosphorylation of sphingosine (indicated by dotted line).

Table S1. Oligonucleotides designed for real time PCR

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<th>Transcript</th>
<th>Orientation</th>
<th>Sequences</th>
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<td>CD3ε</td>
<td>Forward</td>
<td>AAGGCCAAGGCCAGCCTGTGAC</td>
</tr>
<tr>
<td></td>
<td>Reverse</td>
<td>GGCTCATAGTCTGGTTGGGA</td>
</tr>
<tr>
<td>CD20</td>
<td>Forward</td>
<td>GCATTCTGTCGGTGATCTCTC</td>
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<tr>
<td></td>
<td>Reverse</td>
<td>CTCCAGCTGACAGCAGAACATT</td>
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Oligonucleotides were developed using Primer Designer 4.0 for Windows, based on regions of homology in the sequences of rat (Rattus norvegicus) and mouse (Mus musculus) that were determined using Nucleotide search and BLAST (NCBI Entrez).

Table S2. Expression levels of CD3ε and CD20 in the spleen during torpor and arousal

<table>
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<th>Transcript</th>
<th>Torpor</th>
<th>Arousal</th>
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<tbody>
<tr>
<td>CD3ε (marker for T lymphocytes)</td>
<td>1.00 ± 0.16</td>
<td>0.87 ± 0.10</td>
</tr>
<tr>
<td>CD20 (marker for B lymphocytes)</td>
<td>1.00 ± 0.24</td>
<td>0.87 ± 0.24</td>
</tr>
</tbody>
</table>

The table shows fold mRNA expression ± SEM of aroused animals (n = 5) compared with torpid animals (n = 5). Expression of CDs was normalized to GAPDH as housekeeping gene. No significant differences were found.