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The bioactive lysophospholipid mediator sphingosine-1-phosphate (S1P) promotes the egress of newly formed T cells from the thymus and the release of immature B cells from the bone marrow. It has remained unclear, however, where and how S1P is released. Here, we show that in mice, the S1P transporter spinster homolog 2 (Spns2) is responsible for the egress of mature T cells and immature B cells from the thymus and bone marrow, respectively. Global Spns2-KO mice exhibited marked accumulation of mature T cells in thymi and decreased numbers of peripheral T cells in blood and secondary lymphoid organs. Mature recirculating B cells were reduced in frequency in the bone marrow as well as in blood and secondary lymphoid organs. Bone marrow reconstitution studies revealed that Spns2 was not involved in S1P release from blood cells and suggested a role for Spns2 in other cells. Consistent with these data, endothelia-specific deletion of Spns2 resulted in defects of lymphocyte egress similar to those observed in the global Spns2-KO mice. These data suggest that Spns2 functions in ECs to establish the S1P gradient required for T and B cells to egress from their respective primary lymphoid organs. Furthermore, Spns2 could be a therapeutic target for a broad array of inflammatory and autoimmune diseases.

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The sphingosine-1-phosphate transporter Spns2 expressed on endothelial cells regulates lymphocyte trafficking in mice

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The bioactive lysophospholipid mediator sphingosine-1-phosphate (S1P) promotes the egress of newly formed T cells from the thymus and the release of immature B cells from the bone marrow. It has remained unclear, however, where and how S1P is released. Here, we show that in mice, the S1P transporter spinster homolog 2 (Spns2) is responsible for the egress of mature T cells and immature B cells from the thymus and bone marrow, respectively. Global Spns2-KO mice exhibited marked accumulation of mature T cells in thymi and decreased numbers of peripheral T cells in blood and secondary lymphoid organs. Mature recirculating B cells were reduced in frequency in the bone marrow as well as in blood and secondary lymphoid organs. Bone marrow reconstitution studies revealed that Spns2 was not involved in S1P release from blood cells and suggested a role for Spns2 in other cells. Consistent with these data, endothelia-specific deletion of Spns2 resulted in defects of lymphocyte egress similar to those observed in the global Spns2-KO mice. These data suggest that Spns2 functions in ECs to establish the S1P gradient required for T and B cells to egress from their respective primary lymphoid organs. Furthermore, Spns2 could be a therapeutic target for a broad array of inflammatory and autoimmune diseases.

Introduction

Sphingosine-1-phosphate (S1P) is a bioactive lysophospholipid mediator that plays a crucial role in diverse physiological functions, such as lymphocyte trafficking, vascular development, and inflammation (1–5). S1P exerts biological functions mostly through activating cell-surface G protein–coupled receptors S1P1–S1P5, while intracellular S1P is also known to act as a second messenger to regulate inflammation (6). It remains unclear how intracellular S1P is transported to the outside of the cells to activate S1P receptors expressed on the cells.

The activation of S1P1 signaling in lymphocytes by S1P has been shown to promote the egress of newly formed T cells from the thymus and that of mature T and B cells from secondary lymphoid organs such as spleen and lymph nodes (7–10). An immunosuppressive molecule, FTY720, produces peripheral lymphopenia by blocking the lymphocyte egress from the thymus and lymph nodes. Interestingly, FTY720 was found to elicit the immunosuppressive effect by functionally antagonizing the S1P/S1P1 signaling pathway (8, 11–13). In fact, this compound has been recently approved by the United States Food and Drug Administration for treatment of autoimmune diseases (14). In addition, S1P/S1P1 receptor signals direct the release of immature B cells from the bone marrow to the peripheral blood (15, 16).

The concentration of S1P is abundant in circulatory fluids, such as blood and lymph (~μM), whereas it is normally kept low in the lymphoid tissues (~nM) by S1P-degrading enzymes that include lipid phosphate phosphatase 3 (17, 18). However, it has been suggested that this concentration difference of S1P is required but not sufficient for lymphocyte egress from lymphoid tissues into the circulation (9), implying the significance of S1P gradient made in lymphoid tissues. Consistently, S1P produced by neural crest–derived perivascular cells is required for efficient T cell egress (19). Moreover, lymphatic ECs release S1P, which is necessary for lymphocyte egress from lymph nodes into lymph (20). However, it is still unclear how B and T cell egress from the primary lymphoid organs and which cells release S1P that promotes the egress of these cells.

S1P is generated inside of the cell by phosphorylation of sphingosine in a reaction catalyzed by sphingosine kinase 1 and 2, 2 closely related isozymes, and is exported toward the outside of the cell to stimulate its cell-surface receptors (21, 22). Release of S1P is observed in a variety of cells, such as platelets, erythrocytes, mononuclear cells, neutrophils, mast cells, and ECs (3, 21–28). In vitro analyses have revealed that ABC transporters mediate S1P release in several types of cells, such as mast cells, erythrocytes, platelets, breast cancer cells, and

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Results

Spns2 is essential for trafficking of both T and B cells. To address the physiological functions of Spns2 in mammals, we generated global Spns2-KO (Spns2<sup>−/−</sup>) mice by crossing Spns2<sup>fl/fl</sup> mice, in which exon 2 of the Spns2 gene is flanked with loxP sites, with mice expressing Cre recombinase under the control of cytomegalovirus promoter (Supplemental Figure 1; supplemental material available online with this article; doi:10.1172/JCI60746DS1). RT-PCR analyses of the RNA extracted from the lungs of WT and Spns2<sup>−/−</sup> mice revealed that Spns2<sup>−/−</sup> mice express a mutant mRNA transcript lacking exon 2-derived sequence encoding aa 124–145 of WT Spns2 (Supplemental Figure 2, A and B). This Spns2 mutant protein failed to localize at the plasma membrane and lost the ability to export S1P (Supplemental Figure 2, A and B). Thus, we conclude that Spns2<sup>−/−</sup> mice are indeed functionally disrupted for Spns2.

Spns2<sup>−/−</sup> mice develop normally, survive to adulthood, and are fertile, although they exhibited symblepharon to a greater or lesser extent (Supplemental Figure 3). In addition, blood biochemical examination revealed no significant differences between WT and Spns2<sup>−/−</sup> mice (Supplemental Figure 4). Notably, hematological analysis showed a significant decrease in white blood cell count in Spns2<sup>−/−</sup> mice compared with control mice, although there were no differences in the other hematological parameters, such as red blood cells, platelets, hemoglobin, hematocrit, mean corpuscular volume, mean corpuscular hemoglobin, and mean corpuscular hemoglobin concentration (Supplemental Figure 5), implying the role of Spns2 in lymphocyte trafficking.

It should be noted that the number and proportion of mature CD4 and CD8 single-positive (SP) T cells was dramatically reduced in the blood of Spns2<sup>−/−</sup> mice (Figure 1A and Supplemental Figure 6A). In addition, immature B cells (CD19<sup>+</sup>CD23<sup>+</sup>IgD<sup>−</sup>IgM<sup>+</sup>), mature recirculating B cells (CD19<sup>+</sup>CD23<sup>−</sup>IgD<sup>+</sup>IgM<sup>+</sup>), and mature recirculating T cells (CD4<sup>+</sup>CD8<sup>+</sup>) in peripheral blood are shown (n = 11). In A and B, bars and circles indicate averages and values for individual mice, respectively.

Figure 1
Mature T and recirculating mature B lymphocytes are remarkably reduced in the peripheral blood of Spns2<sup>−/−</sup> mice. (A and B) Flow cytometric analyses of control (Spns2<sup>+/+</sup>) and global Spns2<sup>−/−</sup> mice. (A) Frequencies (left) and total numbers (right) of CD4 SP (CD4<sup>+</sup>) and CD8 SP (CD8<sup>+</sup>) T cells and mature recirculating B cells (CD19<sup>+</sup>CD23<sup>−</sup>IgD<sup>+</sup>IgM<sup>+</sup>), immature B, and mature recirculating B cells (CD19<sup>+</sup>CD23<sup>+</sup>IgD<sup>−</sup>IgM<sup>+</sup>), Mature rec. B) in peripheral blood are shown (n = 11). In A and B, bars and circles indicate averages and values for individual mice, respectively.
To assess the possibility that lack of mature T cells in the blood of Spns2−/− mice is related to their accumulation in other secondary lymphoid tissues, we further examined mature SP T lymphocytes in peripheral lymph nodes and in the spleen. In contrast with the accumulation of mature T cells in the thymus, the numbers and proportions of mature CD4 and CD8 SP T cells were dramatically reduced in peripheral lymph nodes and in the spleen of Spns2−/− mice, although their structures were normal (Figure 2, C–E, and Supplemental Figure 7). These results show that a decrease in the number of mature T cells in the peripheral blood of Spns2−/− mice is a consequence of impaired T cell egress from the thymus, but not due to the accumulation in the secondary lymphoid organs.

Spns2 regulates egress of immature B cells from the bone marrow into the blood. In the late stage of B cell development in the bone marrow, newly generated immature B cells are exported into the peripheral blood in an S1P/S1P1 signal–dependent manner (15, 16). The immature B cells subsequently undergo maturation in the secondary lymphoid tissues and migrate back to the bone marrow through the blood (recirculating mature B cells). To explore the cause of remarkable reduction of recirculating mature B lymphocytes in the peripheral blood of Spns2−/− mice, we examined the number and proportion of the lymphocytes at different developmental stages. The numbers and frequencies of mature recirculating B cells (B220+IgM+ or CD19+IgM+IgD+) were significantly reduced in the bone marrow of Spns2−/− mice compared with that of control mice (Figure 3, A and B, and Supplemental Figure 9). However, the number of pro–/pre–B cells (B220−IgM− or CD19−IgM−IgD−) was normal in the bone marrow of Spns2−/− mice, although their frequencies were slightly

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**Figure 2**

Egress of mature T cells from the thymus is impaired in Spns2−/− mice. (A–D) Flow cytometric analyses of control (Spns2+/+) and global Spns2−/− mice. (A) A representative flow cytometric analysis of T cells in the thymus. The numbers represent the percentages of CD4 SP, CD8 SP, CD4/CD8 DP T cells, and CD4/CD8 DN thymocytes. (B) Frequencies (left) and total numbers (right) of CD4/CD8 DN (DN), CD4/CD8 DP (DP), CD4 SP (CD4) and CD8 SP (CD8) thymocytes and T cells are shown (n = 11). (C) Frequencies and numbers of CD4 SP (CD4) and CD8 SP (CD8) T cells in peripheral lymph nodes are shown (n = 11). (D) Frequencies and numbers of CD4 SP (CD4) and CD8 SP (CD8) T cells in spleens are shown (n = 11). In B–D, bars and circles indicate averages and values for individual mice, respectively. (E) Spleen sections from control (Spns2+/+) or Spns2−/− mice, stained to detect CD3+ T cells (blue) and B220+ B cells (red). The boxed areas of upper panels are enlarged in lower panels. Scale bars: 200 μm (top row); 50 μm (bottom row).
Immature B cell egress from bone marrow is impaired in Spns2−/− mice. (A–D) Flow cytometric analysis of B cells in control (Spns2+/+) and Spns2−/− mice. (A) A representative flow cytometric analysis of progenitor B (B220+IgM+), immature B (B220+IgM+) and mature recirculating B cells (B220+IgM−) in the bone marrow cavity. Numbers indicate the percentages of IgM− and B220-expressing cells of total lymphocytes. (B) Frequencies (left) and total numbers of pro–/pre– B cells (B220+IgM−), immature B cells (B220+IgM−), and mature recirculating B cells (B220+IgM−) defined as in A are shown (n = 11). (C) Frequencies (left) and numbers (right) of mature recirculating B (mature rec. B) cells (CD19+CD23−IgD+) in peripheral lymph nodes are shown (n = 11). (D) Frequencies (left) and numbers (right) of T1 B cells (CD19−CD21−CD23−), MZ (CD19−CD21−CD23−), and follicular B cells (FO) CD19+CD21+CD23+ in spleens are shown (n = 11). In B–D, bars and circles indicate averages and values for individual mice, respectively.

To further confirm this conclusion, we examined the number and proportion of B lymphocytes in the secondary lymphoid organs. In the peripheral lymph nodes, the number of mature B cells was reduced in Spns2−/− mice, although the frequency of mature B cells was not different from that in control mice (Figure 3C). In the spleen, the numbers and proportions of follicular B cells were significantly decreased in Spns2−/− mice compared with those of control mice, although there was no difference in the number of marginal zone (MZ) and transitional type 1 (T1) and B1 B cells between control and Spns2−/− mice (Figure 3D). These results reveal that the decrease in mature recirculating B cells in the peripheral blood of Spns2−/− mice is not due to their accumulation in the secondary lymphoid organs, although it remains unclear whether Spns2 is involved in the egress of B lymphocytes from the secondary lymphoid organs. Therefore, we conclude that Spns2 is required for the egress of immature B cells from the bone marrow into the blood.

Spns2 is not involved in S1P release from blood cells. Which cells expressing Spns2 are responsible for releasing S1P necessary for lymphocyte trafficking? Blood cells, especially erythrocytes, are known to produce S1P, thereby contributing to high plasma S1P concentration (9, 23–26). Thus, we first investigated whether Spns2 is involved in S1P release from blood cells. In Spns2−/− mice, plasma S1P levels were reduced to 54% of those in control mice (0.39 ± 0.03 μM in control mice; 0.21 ± 0.01 μM in Spns2−/− mice), although plasma sphingosine and glycerolysophospholipids of Spns2−/− mice were comparable to those in control mice (Figure 4, A–H). Most of the plasma S1P is known to be associated with HDL and albumin (40–42). Consistent with the reduced concentration of plasma S1P in Spns2−/− mice, the amount of S1P associated with HDL and albumin was lower in Spns2−/− mice than in WT mice (Supplemental Figure 10).

To further clarify whether the reduction of plasma S1P concentration in Spns2−/− mice is attributed to the decreased S1P release from blood cells, we examined the secretion of S1P from blood cells isolated from either control or Spns2−/− mice. S1P release from blood cells occurred in Spns2−/− mice to an extent similar to that in control mice (Figure 5A). This S1P release was not caused by membrane damage, since no release was observed when the cells were incubated at 4°C (Figure 5A), as previously reported (27). These results suggest that Spns2 is not involved in the release of S1P from blood cells. To further confirm this conclusion, we performed bone marrow reconstitution studies (Supplemental Figure 11). Reconstitution of irradiated Spns2−/− mice with WT bone marrow did not restore the reduced concentration of plasma S1P (Figure 5B). Furthermore, Spns2−/− mice reconstituted with WT bone marrow still exhibited accumulation of mature SP T lymphocytes in the thymus and reduction of mature recirculating B cells in the bone marrow in comparison with WT mice (Figure 5, Figure 5C). Together with the evidence for the decreased number of immature B cells in the peripheral blood of Spns2−/− mice (Figure 1B), these results suggest that the egress of immature B cells from the bone marrow is impaired in Spns2−/− mice. However, the amount of S1P associated with HDL and albumin was lower in Spns2−/− mice than in WT mice (Supplemental Figure 10).
decreased to the level observed in Figure 1). In Spns2-ECKO mice, plasma S1P concentration was ing Cre recombinase under the Tie2 promoter (Supplemental Spns2, we tried to generate mice lacking Spns2 in ECs (Spns2-ECKO: whether Spns2 functions in ECs to regulate lymphocyte traffick

These results suggest that ECs secrete S1P through Spns2. Hence, not in those of kidney and olfactory bulb (Supplemental Figure 12).

Besides the thymus, expression of Spns2 mRNA on the pericytes in the thymus (Figure 6D). Although it has been reported that pericytes covering thymic ECs release S1P to vivo by performing in situ hybridization analyses. Spns2 mRNA was clearly expressed on the ECs in the thymus (Figure 6D). Although it has been reported that pericytes covering thymic ECs release S1P to promote T cell egress from thymus (19), we could not detect expression of Spns2 mRNA on the pericytes in the thymus (Figure 6D). Besides the thymus, expression of Spns2 mRNA was also observed in the ECs of other tissues, such as heart, lung, and hypothalamus, but not in those of kidney and olfactory bulb (Supplemental Figure 12).

These results suggest that ECs secrete S1P through Spns2. Hence, we hypothesized that S1P released from ECs via Spns2 is required for lymphocyte egress from primary lymphoid organs.

ECs regulate thymic egress by releasing S1P through Spns2. To address whether Spns2 functions in ECs to regulate lymphocyte trafficking, we tried to generate mice lacking Spns2 in ECs (Spns2-ECKO: Spns2–/–;Tie2Cre) by crossing the Spns2–/– mice with the mice expressing Cre recombinase under the Tie2 promoter (Supplemental Figure 1). In Spns2-ECKO mice, plasma S1P concentration was decreased to the level observed in Spns2–/– mice (Supplemental Fig-
S1P required for vascular development, since transporters other than Spns2 might also be involved in the release mediated by S1P transporters other than Spns2. In addition, S1P blood cells from thought to be the major cellular source of S1P in plasma (9, 23, 27), with control mice. Although blood cells, especially erythrocytes, are partially but not completely decreased in only transporter of S1P in mammals because plasma S1P levels were Spns2-ECKO mice. On the other hand, Spns2 appears not to be the cells, since symblepharon was observed in Spns2–/– mice, but not in Spns2-ECKO mice. On the other hand, Spns2 appears not to be the only transporter of S1P in mammals because plasma S1P levels were partially but not completely decreased in Spns2–/– mice compared with control mice. Although blood cells, especially erythrocytes, are thought to be the major cellular source of S1P in plasma (9, 23, 27), blood cells from Spns2–/– mice still retained the ability to release S1P (Figure 5A). Thus, S1P release from blood cells appears to be mediated by S1P transporters other than Spns2. In addition, S1P transporters other than Spns2 might also be involved in the release of S1P required for vascular development, since Spns2–/– mice did not show the defects in vascular development that can be observed in S1P-deficient mice (48).

This study clearly reveals that Spns2-dependent S1P release from ECs is important for egress of mature T cells from the thymus into the peripheral blood. Thymic egress is strictly controlled by the S1P/S1P1 signaling pathway (1, 2, 4, 5). In the thymus, thymocytes differentiate into mature T cells, and subsequently express S1P1 through upregulation of Krüppel-like factor 2 (5, 49). S1P1-expressing mature T cells acquire responsiveness to S1P, thereby exiting from the thymus into the peripheral blood. Until recently, it has been assumed that the S1P gradient between the thymus and the blood is required for egress of mature T cells. However, recent evidence has suggested that plasma S1P is insufficient to promote thymic egress (9, 19). Importantly, egress of mature T cells from the thymus was impaired in Spns2–/– mice even though the plasma contains enough concentration of S1P to stimulate lymphocyte S1P1 in vitro (10). Similarly, Zachariah and Cyster have recently reported that neural crest–derived pericytes covering the ECs release S1P responsible for thymic egress without influencing plasma S1P concentration (19). Therefore, mature T cells might be recruited to the abluminal present study on Spns2 largely contributes to the understanding of S1P signaling, which is used in the egress of lymphocytes from primary lymphoid organs (1, 2, 4, 5).

Spns2 is the first S1P transporter functioning in mammals. Intracellularly generated S1P has to be transported out of the cell to stimulate its cell-surface receptors. Several lines of evidence obtained from in vitro studies have suggested the involvement of the ABC family of transporters in S1P release from several types of cells (22, 24, 29–32). However, their biological significance in vivo has remained unclear. In this study, we demonstrate Spns2 as a key S1P transporter that regulates lymphocyte trafficking in mammals. Although Spns2 regulates lymphocyte trafficking by inducing the release of S1P from ECs, this transporter may also function in other cells, since symblepharon was observed in Spns2–/– mice, but not in Spns2-ECKO mice. On the other hand, Spns2 appears not to be the only transporter of S1P in mammals because plasma S1P levels were partially but not completely decreased in Spns2–/– mice compared with control mice. Although blood cells, especially erythrocytes, are thought to be the major cellular source of S1P in plasma (9, 23, 27), blood cells from Spns2–/– mice still retained the ability to release S1P (Figure 5A). Thus, S1P release from blood cells appears to be mediated by S1P transporters other than Spns2. In addition, S1P transporters other than Spns2 might also be involved in the release of S1P required for vascular development, since Spns2–/– mice did not show the defects in vascular development that can be observed in S1P-deficient mice (48).
exported into the peripheral blood (50–52). Sinusoidal entry of immature B cells is thought to be a key step in bone marrow egress. Recently, it has been reported that S1P/S1P1 signaling promotes the movement of immature B cells from parenchyma to sinusoid, thereby facilitating egress of immature B cells from bone marrow (15, 16). Thus, bone marrow sinusoidal ECs may attract immature B cells from the parenchyma by producing S1P through Spns2 and thereby promoting the immature B cell egress into the peripheral blood.

The lymphocyte egress from secondary lymphoid organs such as lymph nodes and spleen also depends on S1P/S1P1 signaling. Pham et al. have recently reported that lymphatic ECs are an in vivo source of S1P required for lymphocyte egress from lymph nodes and Peyer patches (20). Since Spns2 is expressed not only in vascular ECs but also in lymphatic ECs, Spns2 may also regulate lymphocyte egress from lymph nodes by inducing the release of S1P from lymphatic ECs. However, to address this hypothesis, we need to analyze mice lacking Spns2 specifically in the lymphatic ECs because lymphocyte egress from primary lymphoid organs is severely impaired in global Spns2−/− mice. It also remains elusive whether Spns2 is involved in lymphocyte egress from spleen. Thus, this will be a subject for future studies.

In conclusion, we demonstrate that Spns2 is a key S1P transporter involved in lymphocyte trafficking and further indicate that vascular ECs are the major source of S1P in vivo responsible for lymphocyte egress from the thymus and the bone marrow. Thus, this study not only reveals the crucial role of Spns2 as an S1P transporter in mammals, but also contributes to our understanding of molecular mechanisms of S1P-mediated lymphocyte trafficking. Since S1P signaling is profoundly involved in the inflammatory and autoimmune diseases, such as multiple sclerosis, psoriasis, asthma, and rheumatoid arthritis, as well as in transplantation, Spns2 can be a potential therapeutic target for these diseases.

**Methods**

*Generation of Spns2fl/fl mice. Spns2flfl mice (acc. no. CDB0705K; http://www.cdb. riken.jp/arg/mutant%20mice%20list.html), in which exon 2 is flanked by 2 loxP sites, were generated (Supplemental Figure 1A). TT2 ES cells derived from an F1 hybrid of C57BL/6 and CBA mice (53) were transfected with the targeting vector, selected in the presence of G418, and screened for homologous recombination by PCR and Southern blotting. Two ES clones were introduced into host embryos to generate chimeric mice. Chimeric mice with a high ES cell contribution were bred with the CMV-Cre mice (C57BL/6 strain background) expressing Cre recombinase under the control of cytomegalovirus promoter to generate heterozygous Spns2+/− mice (Supplemental Figure 1). Spns2+/− mice were intercrossed to obtain Spns2−/− mice (75% C57BL/6 and 25% CBA genetic background). The chimeric mice were also crossed with the CMV-Flp mice (C57BL/6 strain background) expressing Flp recombinase under the control of cytomegalovirus promoter to generate heterozygous Spns2+/− mice (Supplemental Figure 1). Spns2−/− mice were bred with Tie2-Cre mice (C57BL/6 strain background), which carry the Cre recombinase driven by the Tie2 promoter (45, 46) and were provided by T.N. Sato (Nara Institute of Science and Technology, Nara, Japan) and M. Yanagisawa (University of Texas Southwestern Medical Center, Dallas, Texas, USA) (Supplemental Figure 1). For confirmation of cor-

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**Figure 6**

Spns2 is involved in S1P release from ECs. (A) Expression of Spns2 in ECs. RT-PCR analysis was performed to examine the expression of Spns2 in HUVECs, HMVECs, HAECs, HDLECs, HeLa, and HEK293 cells as indicated at the top. PCR was performed using specific primers for either Spns2 (upper panel) or GAPDH (lower panel). To verify the absence of contaminating genomic DNA, RT-PCR was also performed in the absence of reverse transcriptase (−). (B and C) S1P release from Spns2-depleted ECs. (B) Release of S1P by ECs transfected without (−) or with either control siRNA (control) or 2 independent siRNAs targeting Spns2 (Spns2#1 and Spns2#2). (C) Real-time RT-PCR analysis to assess the efficiency of siRNA-mediated Spns2 knockdown. In B and C, data are expressed relative to those observed in the untransfected cells and shown as mean ± SD of 3 independent experiments. (D) In situ hybridization for Spns2 mRNA in thymus. Antisense probe was hybridized to thymus section (Spns2: purple). Serial sections were also stained with anti-CD31 (CD31: brown) and anti-α-SMA (brown) antibodies to identify ECs and pericytes, respectively. The boxed areas of upper panels are enlarged in lower panels. Arrows and white arrowheads indicate ECs and pericytes, respectively. Scale bars: 50 μm (upper panels); 10 μm (lower panels).
rect targeting, Southern blot analysis was performed with the probe located outside of the regions used in the targeting vector (Supplemental Figure 1). For the genotyping of mice, PCR was performed using a forward primer, 5'-AGGCTCATTTCATGGCTGAT-3', and a reverse primer, 5'-AGCCCT-TGTGCTCTCTGTGTG-3', producing products of 552-bp fragment for WT allele, 842-bp fragment for floxed allele, and 316-bp fragment for deleted allele. All mice were housed under specific pathogen–free conditions.

**RT-PCR and real-time RT-PCR.** To check the expression of Spns2 mRNA in Spns2−/− mice, total RNA was extracted from the lungs using TRizol reagent (Invitrogen) and reverse transcribed by random hexamer primers using Superscript II (Invitrogen) according to the manufacturer’s instructions. PCR amplification was carried out with the following primer sets: PCR1, 5′-AAAGGTTGACAGCTGTTCC-3′ and 5′-CCACAGCTGAGGATCATT-3′, for exons 1–3 of the mouse Spns2; and PCR2, 5′-ATGGATGTGCTTGAAATGC-3′ and 5′-TCAGACTTTACGGGATGCA-3′, for complete coding sequence of mouse Spns2. To determine the expression of Spns2 in ECs, RT-PCR was performed using the gene-specific primers for human SPNS2 (5′-ACTTTGGGGTCAAGGACC-3′ and 5′-ATATCCCTTCTGTGAAGGG-3′). Amplification of GAPDH was also performed using the gene-specific primers for human GAPDH (5′-ATGGGGAAGGTGAAGGTCG-3′ and 5′-GGGAGTACCTTCAAC-3′) as a control.

To assess the efficiency of siRNA-mediated knockdown of Spns2, total RNA was extracted from HUVECs transfected without or with either control siRNA or 2 independent siRNAs targeting Spns2 and subjected to quantitative real-time RT-PCR analysis using the Quantifast SYBR Green RT-PCR Kit (Qiagen) as described (54). For each reaction, 100 ng of total RNA was transcribed for 10 minutes at 50°C, followed by a denaturation step at 95°C for 5 minutes, 40 cycles of 10 seconds at 95°C, and 30 seconds at 60°C. Fluorescence data were collected and analyzed using MASTERCycler ep realplex (Eppendorf). For normalization, expression of human GAPDH was determined in parallel as an endogenous control. The gene-specific primers used to amplify human SPNS2 and GAPDH were the same as described above.

**Cell culture, transfection, and siRNA-mediated gene silencing.** HUVECs, human microvascular ECs (HMVECs), and human aortic ECs (HAECs) were purchased from Kurabo and maintained as described previously (55). Human dermal lymphatic ECs (HDLECs) were obtained from Lonza and maintained in EC growth medium EGM-2 (Lonza). Hela and HEK293 cells were cultured in DMEM (Nissui) supplemented with 10% fetal bovine serum and antibiotics (100 μg of streptomycin/ml and 100 U of penicillin/ml).

Stealth siRNAs targeted to human Spns2 (HS515133S and HS515133E) were purchased from Invitrogen. As a control, siRNA duplexes with irrelevant sequences were used. HUVECs were transfected with 20 nM siRNA duplexes using Lipofectamine RNaI MAX reagent (Invitrogen). After incubation for 48 hours, the cells were used for the experiments.

**Detection of subcellular localization of GFP-tagged Spns2.** cDNAs encoding WT and mutant Spns2 were amplified using cDNAs derived from the lungs of WT and Spns2−/− mice by RT-PCR, and cloned into pEGFP-N1 vector to construct the expression plasmids encoding WT and mutant Spns2 with a C-terminal GFP tag, respectively. HUVECs were transfected with the plasmid encoding either WT or mutant Spns2-GFP or with myristoylated GFP–encoding plasmid. GFP and phase contrast images were obtained using an IX81 inverted microscope (Olympus) equipped with a pE-1 LED excitation system (CoolLED).

**SIP release from cultured cells.** HEK293 cells were plated in 24-well plates (5 × 104 cells/well), cultured for 24 hours, and transfected with the expression plasmids indicated in the legend of Supplemental Figure 2D using Lipofectamine 2000 reagent (Invitrogen). After incubation for 24 hours, cells were incubated in 250 μl of serum-free DMEM containing 0.5 % fatty acid–free bovine serum albumin, 10 mM sodium glycerophosphate, 5 mM sodium fluoride, and 1 mM semicarbazide for 24 hours. To determine the role of Spns2 in SIP release from ECs, HUVECs transfected without or with either control siRNA or 2 independent Spns2 siRNAs were detached, replated in collagen-coated 24-well plates (2.5 × 105 cells/well), and cultured for 12 hours. The cells were then washed twice with Medium 199 (Invitrogen) and incubated in 200 μl of Medium 199 containing 20 mM Heps, pH 7.4, 10 mM sodium glycerophosphate, 5 mM sodium fluoride, 1 mM semicarbazide, 0.5% fatty acid free bovine serum albumin, 40 ng/ml vascular endothelial growth factor, 40 ng/ml fibroblast growth factor–2, and 400 ng/ml angioptoin-1 for 12 hours. After incubation, conditioned medium was collected and centrifuged at 15,000 × g for 5 minutes at 4°C to remove cell debris. SIP levels in the conditioned medium were determined as described below.
Spns2 expressed in ECs is required for immature B cell egress from bone marrow. (A–C) Flow cytometric analyses of lymphocytes of control (Spns2+/+) or Spns2-ECKO (Spns2−/-;Tie2Cre) mice. (A) Frequencies (left) and numbers (right) of pro–pre-B cells (B220−IgM−), immature B cells (B220hiIgM−), and mature recirculating B cells (B220hiIgM+) in bone marrow are shown (n = 11). (B) Frequencies (left) and numbers (right) of mature recirculating B (Mature rec. B) cells (CD19hiCD23−IgD−) in peripheral blood are shown (n = 11). (C) Frequencies (left) and numbers (right) of T1 and B1 B cells (T1+B1), MZ, and follicular B cells in spleens are shown (n = 11). T1 and B1 B cells, MZ B cells, and follicular B cells were phenotypically defined as described in the legend of Figure 3D. In A–C, bars and circles indicate averages and values for individual mice, respectively.

Quantification of S1P using LC-MS/MS. Quantification of S1P was performed according to previously described methods (56) with minor modifications. Briefly, plasma and conditioned medium were mixed and sonicated with 10-fold volume of methanol and an internal standard (C17-S1P). Similarly, plasma and conditioned medium were mixed and sonicated with methanol and analyzed by the LC-MS/MS analysis. Then 20 μl of methanol extract was injected and separated by Nanospace LC (Shiseido), using a C18 CAPCELL PAK ACR column (1.5 × 250 mm; Shiseido), using a gradient of solvent A (5 mM ammonium formate in water) and solvent B (5 mM ammonium formate in 95% [v/v] acetonitrile). Elution was sequentially ionized with an ESI probe, and the parent ion (m/z 380.2) and the fragment ion (m/z 264.2) were monitored in the positive mode by a Quantum Ultra Triple Quadrupole Mass Spectrometer (Thermo Fisher Scientific). Similarly, other lysophospholipids, including lysophosphatidic acid (LPA), lysophosphatidylcholine (LPC), lysophosphatidylethanolamine (LPE), lysophosphatidylglycerol (LPG), lysophosphatidylinositol (LPI), and lysophosphatidylserine (LPS), were extracted with methanol and analyzed by the LC-MS/MS system. For each lysophospholipid class, 12 acyl chains (14:0, 16:0, 16:1, 18:0, 18:1, 18:2, 18:3, 20:3, 20:4, 20:5, 22:5, and 22:6) were monitored.

For quantification of HDL- and albumin-bound S1P, mouse plasma was subjected to size-exclusion chromatography according to previously described methods (57) with some modifications. Briefly, 100 μl of plasma was loaded onto a Superose 12 column (GE Healthcare) using an AKTA Explorer System (GE Healthcare) and eluted with PBS at 0.25 ml/min at 4°C. Fractions were collected every 2 minutes (0.5 ml). S1P concentration in each fraction was determined by LC-MS/MS as described above.

Biochemical and hematological test of blood. Blood was collected from WT (Spns2+/+, n = 4) and Spns2−/- (Spns2−/-;Tie2Cre, n = 4) mice via the abdominal aorta under inhalation anesthesia (isoflurane) using EDTA as an anticoagulant. Blood biochemistry parameters (total protein, total bilirubin, aspartate aminotransferase, alanine aminotransferase, triglycerol, glucose, blood urea nitrogen, and albumin) were determined by using a blood biochemical analyzer, Fuji DRI-CHEM 5500V (Fuji Film). Hematology and blood clotting parameters (white blood cells, red blood cells, platelets, hemoglobin, hematocrit, mean corpuscular volume, mean corpuscular hemoglobin, and mean corpuscular hemoglobin concentration) were determined by using an XT-1800iv hematology analyzer (Sysmex).

Antibodies and flow cytometric analysis. Unless otherwise stated, all anti-mouse monoclonal antibodies were obtained from ebioscience Inc. Antibodies used for cell-surface staining were PE-conjugated anti-CD19 (eBio1D3), anti-CD8 (53-6.7), and anti-CD23 (B3B4); FITC-conjugated anti-B220 (RA3-6B2), anti-CD19 (ebio1D3), anti-CD23 (B3B4); PE-Cy7-conjugated anti-IgM (II/41); and PerCP-Cy5.5 conjugated anti-CD62L (MEL-14) (BioLegend); Pacific Blue–conjugated anti-IgD (11-26) and anti-CD4 (RM4-5) (BD Biosciences); APC-conjugated anti-CD62L (MEL-14) (BioLegend); Pacific Blue–conjugated anti-IgD (11-26) and anti-CD4 (RM4-5) (BD Biosciences); PeCy7-conjugated anti-IgM (II/41); and PerCP-Cy5.5 conjugated anti-CD21/CD35 (7E9) (BioLegend). Single-cell suspensions of freshly isolated thymus, spleen, peripheral lymph nodes (inguinal, axillary, and brachial), and total bone marrow cells of femur and tibia were subsequently incubated with anti-CD16/CD32 for 10 minutes, followed by staining with a combination of conjugated antibodies in FACs buffer (PBS + 4% heat-inactivated FCS + 2 mM EDTA) for 10 minutes. Before antibody staining, 250 μl of freshly isolated blood was treated with heparin solution, and red blood cells were lysed with BD Pharm Lyse solution (BD Biosciences). For immunohistochemistry, paraffin was then removed from the sections by placing the sections in a pressure cooker for 3 minutes. Endogenous
Blood was collected from anesthetized WT mice via the inferior vena cava using heparinized syringes and transferred to tubes containing EDTA as an anticoagulant. Blood cells were separated from plasma by centrifugation at 1,200 × g for 5 minutes at 4°C and washed twice with ice-cold PBS to remove plasma residues. The cells were resuspended in the ice-cold incubation buffer containing 20 mM Hepes, pH 7.4, 138 mM NaCl, 3.3 mM NaH2PO4, 2.9 mM KCl, 1.0 mM MgCl2, 1 mg/ml glucose, and 1% fatty acid-free bovine serum albumin at a cell density of 5 × 10^6 cells/ml. Then 500 μl of blood cell suspensions (2.5 × 10^5 cells) was incubated at 4°C or at 37°C for 90 minutes. After incubation, the cells were pelleted by centrifugation at 1,200 × g for 5 minutes at 4°C. The SIP levels in the supernatants were determined as described above. To quantify the total amount of SIP in the blood cells, cells were collected from 500 μl of cell suspensions by centrifugation at 1,200 × g for 5 minutes at 4°C and homogenized in 100 μl of methanol.

Generation of bone marrow chimeras. Bone marrow chimeras were generated with 5 × 10^6 freshly isolated total bone marrow cells from femur and tibia of WT and Spns2 floxed mice (donor). Isolated cells were injected i.v. into lethally irradiated (900 cGy) Spns2–/– and Spns2-ECKO mice (host). Hema-topoietic reconstitution of lymphoid organs of hosts by donor-derived cells was controlled 6 weeks after bone marrow transfer by genotyping of total bone marrow cells. Furthermore, lymphoid organs of reconstituted mice were FACS analyzed as described above.

In situ hybridization and immunohistochemistry on serial tissue sections. In situ hybridization and immunohistochemistry on serial tissue sections was performed by Genostaff. Briefly, the thymus, heart, lung, brain, and kidney of 8-week-old mice were dissected after perfusion, fixed with Tissue Fixative (Genostaff), and then embedded in paraffin by proprietary procedures and sectioned at 6 μm.

For in situ hybridization, tissue sections were de-waxed with xylene and rehydrated through an ethanol series and PBS. The sections were fixed with 4% paraformaldehyde in PBS for 15 minutes and then washed with PBS. The sections were treated with 8 μg/ml Proteinase K in PBS for 30 minutes at 37°C, washed with PBS, refixed with 4% paraformaldehyde in PBS, again washed with PBS and placed in 0.2 N HCl for 10 minutes. After washing with PBS, the sections were acetylated by incubation in 0.1 M tri-ethanolamine–HCl, pH 8.0, 0.25% acetic anhydride for 10 minutes. After washing with PBS, the sections were dehydrated by incubation in 1 M ethanolamine–HCl, pH 8.0, 0.25% acetic anhydride for 10 minutes. After washing with PBS, the sections were dehydrated through a series of ethanol. The cDNA templates for Spns2 were 535-bp and 634-bp fragments corresponding to bases 1629–2163 and 2291–2924 of mouse Spns2 cDNA (GenBank NM_153060.2). Sense and antisense riboprobes for Spns2 mRNA were synthesized using a digoxigenin RNA labeling kit (Roche) according to the manufacturer's protocol. Hybridization was performed with probes at concentrations of 300 ng/ml in the Probe Diluent-1 (Genostaff) at 60°C for 16 hours. After hybridization, the sections were washed in 5x HybridWash (Genostaff), equal to 5x SSC, at 50°C for 20 minutes and then in 50% formamide, 2x HybridWash at 50°C for 20 minutes, followed by RNase treatment in 50 μg/ml RNase A in 10 mM Tris-HCl, pH 8.0, 1 M NaCl, and 1 mM EDTA for 30 minutes at 37°C.

Then the sections were washed twice with 2x HybridWash at 50°C for 20 minutes, twice with 2x HybridWash at 50°C for 20 minutes, and once with TBST (0.1% Tween 20 in TBS). After treatment with 0.5% blocking reagent (Roche) in TBST for 30 minutes, the sections were incubated with anti-DIG AP conjugate (Roche) diluted 1:1,000 with TBST for 2 hours at RT. The sections were washed twice with TBST and then incubated in 100 mM NaCl, 50 mM MgCl2, 0.1% Tween 20, and 100 mM Tris-HCl, pH 9.5. Coloring reactions were performed with NBT/BCIP solution (Sigma-Aldrich) overnight and then washed with PBS. The sections were counterstained with Kernechtrot Stain Solution (Mutoh), dehydrated, and mounted with Malinol (DBS).

For immunohistochemistry, serial tissue sections were deparaffinized with xylene and rehydrated through an ethanol series and PBS. Endogenous peroxidase activity was blocked with 0.3% H2O2 in methanol for 30 minutes. For CD31 staining, the sections were treated with Protein Block (Dako) and avidin/biotin blocking kit (Vector), and incubated with 0.1 μg/ml of anti-CD31 rabbit polyclonal antibody (Spring Bioscience) at 4°C overnight. Immunocomplexes were detected with biotin-conjugated goat anti-rabbit IgG (Dako) and peroxidase-conjugated streptavidin (Nichirei). For γ-SMA staining, sections were treated with Blocking Reagent A (Nichirei) and incubated with anti-γ-SMA mouse monoclonal antibody (Dako) at 4°C overnight. The sections were then blocked with Blocking Reagent B (Nichirei) and incubated with Simple Stain Mouse MAX-PO (M) (Nichirei). Peroxidase activity was visualized by diaminobenzidine. The sections were counterstained with Mayer's hematoxylin, dehydrated, and mounted with Malinol (Muto).

Statistics. Data were analyzed using GraphPad Prism software (GraphPad Software Inc.). Statistical significance was determined using a 2-tailed Mann-Whitney U test for paired samples or 1-way ANOVA and nonparametric tests for multiple groups. P < 0.05 was considered statistically significant.

Study approval. All animal experiments were approved by the animal committee of the National Cerebral and Cardiovascular Center and performed according to the regulations of the National Cerebral and Cardiovascular Center.

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