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The Role of Sphingosine-1-Phosphate Transporter Spns2 in Immune System Function

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Sphingosine-1-phosphate (SIP) is lipid messenger involved in the regulation of embryonic development, immune system functions, and many other physiological processes. However, the mechanisms of SIP transport across cellular membranes remain poorly understood, with several ATP-binding cassette family members and the spinster 2 (Spns2) member of the major facilitator superfamily known to mediate SIP transport in cell culture. Spns2 was also shown to control SIP activities in zebrafish in vivo and to play a critical role in zebrafish cardiovascular development. However, the in vivo roles of Spns2 in mammals and its involvement in the different SIP-dependent physiological processes have not been investigated. In this study, we characterized Spns2-null mouse carrying the Spns2<sup>tm1a(KOMP)Wtsi</sup> allele. The Spns2<sup>tm1a/tm1a</sup> animals were viable, indicating a divergence in Spns2 function from its zebrafish ortholog. However, the immunological phenotype of the Spns2<sup>tm1a/tm1a</sup> mice closely mimicked the phenotypes of partial SIP deficiency and impaired SIP-dependent lymphocyte trafficking, with a depletion of lymphocytes in circulation, an increase in mature single-positive T cells in the thymus, and a selective reduction in mature B cells in the spleen and bone marrow. Spns2 activity in the nonhematopoietic cells was critical for normal lymphocyte development and localization. Overall, Spns2<sup>tm1a/tm1a</sup> resulted in impaired humoral immune responses to immunization. This study thus demonstrated a physiological role for Spns2 in mammalian immune system functions but not in cardiovascular development. Other components of the SIP signaling network are investigated as drug targets for immunosuppressive therapy, but the selective action of Spns2 may present an advantage in this regard. The Journal of Immunology, 2012, 189: 000–000.

Lipid messenger sphingosine-1-phosphate (SIP) is essential for normal embryonic development and the functions of the cardiovascular and immune systems (1), and components of the SIP signaling network are widely investigated as drug targets for suppression of transplant rejection, autoimmunity, and sepsis (2). In the extracellular environment SIP acts through five G protein coupled receptors (SIP<sub>1–5</sub>) expressed on a variety of cell types (reviewed in Refs. 3, 4). In particular, the loss of SIP receptor 1 (SIP<sub>R</sub>) results in embryonic lethality with abnormal development of the cardiovascular system, whereas a lymphocyte-specific loss of SIP<sub>R</sub> causes impaired exit of mature T cells out of the thymus (5, 6), B cells out of the bone marrow (7, 8), as well as a severe defect in lymphocyte egress from secondary lymphoid organs during their physiological recirculation (9, 10). SIP<sub>R</sub> is also required for normal localization of marginal zone B cells (11), B1 cells (12), plasma cells (13), and gut intraepithelial T lymphocytes (14), whereas SIP receptor 5 controls NK cell migration (15). Intracellular activities of SIP as a second-messenger molecule are broadly linked to cell survival and growth (16), and in the immune system intracellular SIP promotes inflammatory and antimicrobial activities of mast cells, macrophages, and neutrophils (17–22). In particular, SIP is produced downstream of FcεR crosslinking and stimulates mast cell degranulation (23, 24). It is also produced downstream of TNFR signaling and is required for TNFR-associated factor 2-dependent RIP1 activation and NF-κB signaling (25). These activities of SIP have been implicated in the pathologies of allergic and inflammatory disorders (26, 27).

The activities of SIP as a chemokine and a second messenger are critically dependent on the SIP concentrations in the different cellular compartments, the tissue environment, and the circulation, and these are established through the controlled rates of SIP production, degradation, and transport. SIP is synthesized by the sphingosine kinases, sphingosine kinase (SphK1) and SphK2 (28–30), and it is degraded by the SIP lyase and SIP phosphatases (1, 31). Combined loss of SphK1 and SphK2 in mice results in embryonic lethality with abnormal cardiovascular development, whereas the loss of SIP lyase causes a milder phenotype with impaired lymphocyte egress from lymphoid organs. In contrast, the mechanisms of SIP transport across cellular membranes remain poorly understood. Several proteins of the ATP-binding cassette (ABC) transporter superfamily can mediate SIP secretion in cell culture, for example ABCG1 in mast cells (32), ABCA1 in RBCs (33), and another family member in platelets
Expression of the cystic fibrosis transmembrane conductance regulator on epithelial cell lines was also shown to contribute to S1P transport (35). However, knockout mice for these proteins do not phenocopy the knockouts of known S1P signaling network components, and plasma S1P levels in the mice lacking ABCA1 and ABCC1 are unaltered (36). Some S1P is also produced extracellularly by the extracellular SpnK1 enzyme released from the vascular endothelium (37).

SpnK homolog 2 (Spns2) is a member of the major facilitator superfamily of transmembrane proteins (38). In recent studies, zebrafish Spns2 was shown to mediate S1P secretion, and the loss of Spns2 activity resulted in lethal defects in cardiovascular development (39, 40), similar to the phenotypes of the SpnK1/−/−SpnK2/−/− and S1P1/−/− mouse lines (30, 41). The human Spns2 protein was also shown to mediate the secretion of S1P as well as of S1P receptor agonist and immunosuppressive drug phospho-FTY720 (42). Furthermore, human Spns2 expression could rescue the developmental defects in zebrafish embryos (39). However, the in vivo functions of Spns2 in the mammalian system have not been investigated.

In the present work we characterized an Spns2-targeted mouse line Spns2tm1a(Wtsi) generated by the Wellcome Trust Sanger Institute as part of the International Knockout Mouse Consortium, and we demonstrate the requirement for Spns2 for the normal lymphocyte localization and mammalian immune system function but not the other S1P-mediated functions such as embryonic viability and cardiovascular development.

Materials and Methods

Gene targeting and mouse production

The mouse strain carrying the Spns2tm1a(Wtsi) allele was generated by blastocyst injection of embryonic stem cell clone EPD0090_5_B04 obtained from the Knockout Mouse Project resource (43). Prior to microinjection, the identity of the targeted embryonic stem cells was verified by 5′-long-range PCR using a primer external to the targeting vector. Chimeric mice were bred to C57BL/6-Tfg−/− and germline transmission was verified by quantitative PCR (qPCR) to detect the neo transgene included in the mutant allele (single insertion event), as well as by loss of wild-type allele qPCR (correct targeted locus) in the F1 heterozygous mice. The presence of the downstream LoxP site was verified by PCR. The C57BL/6N-Hprt−/−(C57BL/6N-Hprt−/−(C57BL/6N-Hprt−/−(C57BL/6N-Hprt−/−tm1a(Wtsi) and C57BL/6N-Gt(Rosa)26Sor1um1(FLPeR)Dym/Wtsi transgenic lines with systemic expression of Cre and Fip recombinases were previously described (44, 45). The Spns2tm1a(Wtsi) allele was generated by crossing the tm1a allele to the C57BL/6N-Hprt−/−(C57BL/6N-Hprt−/−(C57BL/6N-Hprt−/−tm1a(Wtsi) allele to delete exon 3 and the neo cassette between the LoxP sites, and the Cre allele was bred out of the colony before study. The Spns2tm1a(Wtsi) allele was generated by breeding to C57BL/6N-Gt(Rosa)26Sor1um1(FLPeR)Dym/Wtsi transgenic mice expressing GFP-tagged S1P1. Internalization of S1P1-GFP was quantified using FACS/Kit and the Spot detection algorithm and the “spot total area per object” function on Cellomics ArrayScan VTI system. S1P (Cayman Chemicals) was used to prepare the positive controls.

Recipient mice were immunized by intranasal inhalation of 30 μl PBS containing 10 mg TeCt (gift from Omar Qazi, Imperial College) combined with 1 mg heat-labile toxin of Escherichia coli (gift of Rino Rappuoli, Chiron) adjuvant. Mice were boosted on days 7, 21, and 37. Serum samples were collected on days 36 and 40. Detection of TeCt-specific Abs from sera was performed by ELISA as described above.

Mouse bone marrow transfer experiments

Recipient mice were irradiated with two doses of 4.5 Gy, 3 h apart, and injected i.v. with 3 × 109 donor bone marrow cells. The mice were maintained on clindamycin (250 mg/l) in drinking water for 2 wk and analyzed 8 wk after reconstitution. In one experiment Spns2tm1a(Wtsi) and Spns2tm1a(Wtsi) recipients were reconstituted either with wild-type CD45.1+ marked or with Spns2tm1a(Wtsi) donor bone marrow. In a separate study

ROLES OF Spns2 IN IMMUNE SYSTEM FUNCTION

RNA isolation and qPCR

For the comparisons of Spns2 transcript levels in wild-type and Spns2tm1a(Wtsi) tissues (Fig. 1B) RT-qPCR was performed using an RNA-to-Ct 1-Step kit (Applied Biosystems) in a 10 μl reaction with 1 μl total RNA (20–50 ng depending on tissue type). A TaqMan assay (Mn01249325) spanning the exon flanking the targeted receptor site of the construct was used in a multiplex reaction with a GAPDH endogenous control to normalize for variations between the amounts of input RNA (Applied Biosystems), and amplified in triplicate using a Viia 7 qPCR machine (Applied Biosystems). Analysis was performed using the Viia 7.1 analysis software and the ΔΔCt relative quantification module. For the comparisons of Spns2 transcript levels across different tissues of wild-type mice (Fig. 5A), RNA was isolated using the RNeasy Plus Mini kit (Qiagen) and reverse-transcribed with the Quan- tiTect reverse transcription kit (Qiagen). qPCRs were performed using the Quantitect SYBR Green PCR Kit (Qiagen), using Spns2 QuantiTect primer assay (Qiagen) and Actb primers 5′-CTAAGGCGCAACCGTGAAAAG-3′ (forward) and 5′-ACAGAGGCGATACGGGACA-3′ (reverse) (Sigma-Aldrich). The Spns2 primers spanned the boundaries of exons 5–7 of the Spns2 coding transcript ENSMUST00000045530. The data were acquired on the StepOnePlus real-time PCR system (Applied Biosystems) and analyzed using the ΔΔCt method.

Flow cytometry

Cell suspensions of mouse tissues were prepared in RPMI 1640 with 2% (v/v) FCS (Sigma-Aldrich), 100 μg/ml streptomycin, and 100 μg/ml penicillin (all from Invitrogen). Blood was collected into heparin-coated tubes (Kabe Labotecnich) by cardiac puncture, and erythrocytes were lysed using Pharmalyse (BD Biosciences). The cells were stained in PBS with 2% FCS (Sigma-Aldrich) and 0.2% (w/v) sodium azide (Sigma-Aldrich) for 20 min on ice, with the following Abs. Fluorescein-conjugated Abs were against CD4 (clone L3T4), CD8 (53-6.7), CD11b (M1/70), CD21 (7G6), CD86 (GL1), and B220 (RA3-6B2; all from BD Pharmingen). PE-conjugated Abs were against CD8 (clone 53-6.7), CD19 (1D3), CD69 (H1.2F3), CD80 (16-10A1), and IgM (R6-60.2; all from BD Pharmingen). Allophycocyanin-conjugated Abs were against CD4 (RM4-5), CD8 (53-6.7), and CD44 (IM7; all from BD Pharmingen). Allophycocyanin-Cy7 anti-CD45.1 (A20; BioLegend), Alexa Fluor 647-conjugated anti-IgD (clone 11-26-2), and PE-Cy7 anti-CD23 (BioLegend) were also used. Flow cytometric measurements of β-galactosidase activity were performed using FluorReporter lacZ flow cytometry kits (Invitrogen/Molecular Probes). The cells were stained for appropriate combinations of cell surface lineage markers, before loading with fluorescein di-β-D-galactopyranoside and analysis by flow cytometry. The data were acquired on BD FACSaria or LSRII flow cytometers and analyzed with FACSDiva software.

ELISA

For the measurements of Ab levels, mouse blood was collected by tail bleed or cardiac puncture, and serum was prepared and stored at −20°C. For Ag-specific Ab measurements in mouse serum, Nunc MaxiSorp plates were coated overnight at 4°C with 2 mg/ml toxogen fragment C recombinant protein (TestCt) in 0.1 M Na2HPO4 (pH 9.0), blocked with 3% (w/v) BSA in PBS for 1 h, and incubated with 5-fold serial dilutions of mouse serum in PBS with 1% BSA for 1 h. The plates were developed with anti-mouse IgG, IgG1, or IgG2a, HRP-conjugated Abs (BD Pharmingen), followed by o-phenylenediamine substrate tablets (Sigma-Aldrich) dissolved in water. Cytokine ELISA on cell culture supernatants was performed using anti-mouse TNF-α- coating Ab clone IF3F34 and biotin-conjugated detection Ab clone XT3/XT22, followed by avidin HRP (all from eBioscience), and the 3′,3′,5,5′-tetrathymethylbenzilic liquid substrate system (Sigma-Aldrich). Absorbances were measured using the Bio-Rad 680 microplate reader.

Measurements of S1P levels and activity

For the measurements of S1P levels, mouse blood was collected from the retro-orbital sinus. S1P levels in the plasma were measured using the ELISA-based S1P assay kit (Echelon Biosciences) according to the manufacturer’s protocol. Assays of S1P activity in mouse plasma used an S1P, redistribution assay (Thermo Scientific) according to the manufacturer’s instructions. Briefly, the assay measured S1P-induced internalization of S1P, when mouse plasma is added at different dilutions to the U2OS cells stably expressing GFP-tagged S1P1. Internalization of S1P1-GFP was quantified using the spot detection algorithm and the “spot total area per object” function on Cellomics ArrayScan VTI system. S1P (Camyan Chemicals) was used to prepare the positive controls.

Mouse immunization

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**Tissue culture**

Bone marrow-derived macrophages (BMDMs) were generated by culturing bone marrow cells of all hematopoietic lineages of the Spns2

**Tissue staining for β-galactosidase activity**

Mice were fixed by cardiac perfusion with 4% (w/v) paraformaldehyde. Following dissection, the tissues were fixed in 4% paraformaldehyde for a further 30 min, rinsed in PBS, and stained in 0.1% (w/v) 5-bromo-4-chloro-3-indolyl-β-D-galactoside (X-Gal; Invitrogen) for up to 48 h. After a further overnight fixation in 4% paraformaldehyde, the tissues were cleared with 50% (v/v) glycerol and transferred to 70% glycerol for long-term storage. Images were taken using a Leica MZ16A microscope and iMagic software.

**Statistical analyses**

Statistical comparisons were performed with Prism 4.0 software (GraphPad Software) using a two-tailed Student t-test or nonparametric Mann–Whitney U-test for comparisons of two datasets, and ANOVA was used for multiple comparisons.

**Results**

**Spns2 gene targeting and mouse production**

Spns2 gene targeting was carried out as part of the International Knockout Mouse Consortium, in the JM8 embryonic stem cell line on a C57BL/6N genetic background (43). The targeted Spns2

**Abnormal T and B lymphocyte development and localization in Spns2tm1a/ma mice**

To establish the impact of Spns2

**FIGURE 1.**

A. Structure of the Spns2tm1aKOMP/WTSt allele. (B) Average fold reduction in Spns2 transcript levels in Spns2tm1a/ma relative to wild-type tissues (liver, spleen, lymph nodes, and lung), analyzed by qRT-PCR using primers spanning the junction of exons 2–3 of the Spns2 coding transcript ENSMUST00000045303.

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circulation. There were no abnormalities in the numbers of the earlier thymocyte subsets, including CD4+CD8– double-negative thymocytes (subsets 1–4, differentiated by CD44 and CD25 expression) and CD4+CD8+ double-positive thymocytes (data not shown).

There was also a significant reduction in the number of recirculating B cells in Spns2tm1amtm1a mice (CD45RCD19+; Fig. 3A). This was accompanied by a reduction in the follicular B cell population in the spleen (gated as B220+IgM+IgD– or B220+CD23–CD21low; Fig. 3B–D) and the mature B cell population in the bone marrow (B220+IgM+IgD+; Fig. 3E). In contrast, no abnormalities in earlier B cell developmental subsets were observed, with normal numbers of pro/pre-B cells (B220+IgM– IgD+ ), and immature B cells (B220–IgM– IgD+ ) in the bone marrow and transitional B cells (B220+CD23highCD21low) in the spleen of the Spns2tm1amtm1a mice (Fig. 3D, 3E). Overall, this phenotype resembles the loss of recirculating and mature follicular B cell population previously reported in other mouse models with impaired S1P production or sensing (5, 31, 46). We also observed a reduction in the numbers of B1 cells in the peritoneal cavity of Spns2tm1amtm1a mice (B220+IgM+IgD–; data not shown), consistent with the role of S1P in their control of trafficking (12).

Reduced Ab responses to immunization in Spns2tm1amtm1a mice

To establish whether the defects in lymphocyte development and localization seen in the Spns2tm1amtm1a mice led to impaired Ab responses to immune challenge, the mice were immunized intraperitoneally with tetanus toxoid, boosted at days 7 and 21, and analyzed for Ag-specific Ab titers in the serum at day 36. Significant reductions in Ag-specific total IgG and IgG1 were seen (Fig. 4), indicating impaired humoral immunity in the Spns2tm1amtm1a mouse line. The mice were subsequently given a tertiary boost, and serum Ab levels were measured again 3 d later (day 40), confirming the reduction in Ag-specific total IgG and IgG1 in Spns2tm1amtm1a mice (data not shown). However, the levels of IgG2a were not significantly reduced (Fig. 4), and this might have been due to the low levels and high variation in IgG2a production in both the wild-type and Spns2tm1amtm1a groups.

Normal macrophage responses to inflammatory stimuli in Spns2tm1amtm1a mice

S1P functions not only as a chemoattractant in the extracellular environment, but also as an intracellular messenger required for the normal signaling downstream of TNFR and TLRs (25); furthermore, inhibition of S1P production is protective in animal models of endotoxemia and sepsis (26). To establish whether Spns2tm1amtm1a disrupted normal cellular responses to TNF and NF-κB stimulation, macrophages, generated from Spns2tm1amtm1a and wild-type control mice, were stimulated with LPS (100 ng/ml), IFN-γ (25 ng/ml), and/or TNF-α (25 ng/ml) and analyzed for induction of CD80 and CD86 activation markers and secretion of inflammatory cytokines. No differences in the responses of Spns2tm1amtm1a and wild-type cells were observed (Supplemental Fig. 1B). This suggests that Spns2 does not affect intracellular S1P functions in macrophages downstream of TNF or NF-κB stimulation.

Cell-intrinsic activity of Spns2 is dispensable for lymphocyte development and localization

Many cell types have been shown to produce and release S1P, including erythrocytes, lymphatic and vascular endothelial cells, platelets, and mast cells (32, 34, 46–50). To assess Spns2 expression in a broad range of mammalian cell types, a range of tissues and organs from Spns2tm1amtm1a mice were profiled for the endogenous Spns2 promoter in mice carrying the Spns2tm1amtm1a allele (51). The studies were done as part of the Sanger Mouse Genetics Project high-throughput phenotyping, and the data on the β-galactosidase reporter activity in 39 organs and tissues of Spns2tm1amtm1a mice is summarized in Supplemental Table I, with images available at http://www.sanger.ac.uk/mouseportal/phenotyping/MBNZ/adult-lac-z-expression. The expression of the Spns2 gene was further confirmed in a selection of tissues using wild-type tissues.

**FIGURE 2.** T cell abnormalities in the Spns2tm1amtm1a mice. (A) Percentage of CD4 and CD8 T cells in the blood and numbers of CD4 and CD8 T cells in the spleen of wild-type, Spns2tm1amtm1a, and Spns2tm1amtm1a mice (+, +tm1a, and tm1a, respectively). (B) Percentage of CD4+ and CD8+ single-positive cells in the thymus of wild-type, Spns2tm1amtm1a, and Spns2tm1amtm1a mice (+, +tm1a, and tm1a, respectively). (C) Representative flow cytometry plots of wild-type and Spns2tm1amtm1a thymocytes stained for CD4 and CD8; average percentage of cells within the CD4+ and CD8+ single-positive thymocyte gates; the expression of CD62L and CD69 on CD4+ and CD8+ single-positive thymocytes in wild type (+), Spns2tm1amtm1a (+/tm1a), and Spns2tm1amtm1a (tm1a) mice is indicated. (D) Representative flow cytometry histograms indicating CD24 expression on CD4+ single-positive thymocytes in wild-type and Spns2tm1amtm1a mice. (E) Percentage of CD24low cells within the CD4+ and CD8+ single-positive thymocyte gates; the expression of CD62L and CD69 on CD4+ and CD8+ single-positive thymocytes in wild type (+), Spns2tm1amtm1a (+/tm1a), and Spns2tm1amtm1a (tm1a) mice is indicated. Bars represent means ± SEM. MFI, mean fluorescence intensity. Data are from three to four mice per group and were reproducible in two independent experiments. *p < 0.05, **p < 0.01, ***p < 0.001 using ANOVA with a Bonferroni post hoc test or t test.
and qRT-PCR. Overall, it was demonstrated that there were high Spns2 transcript levels in the liver and lung, lower levels in the lymph nodes, spleen, and bone marrow, and low but detectable levels in the thymus (Fig. 5A).

To establish whether Spns2 was expressed in the lymphocyte populations affected by the Spns2<sup>tm1a/tm1a</sup> phenotype, Spns2 promoter-driven β-galactosidase reporter activity was analyzed in Spns2<sup>+/tm1a</sup> lymphocytes using flow cytometry with a fluorescent β-galactosidase substrate. The data showed no significant β-galactosidase activity over the background level in the different subpopulations of thymocytes or in splenic T cells (Fig. 5B), indicating that the Spns2 gene is not expressed in these cell types. The Spns2 gene was also not expressed in splenic B cells (Fig. 5B). As a positive control, high levels of β-galactosidase reporter activity were detected in hematopoietic cells of another mouse line Mysm1<sup>+/tm1a</sup> (52).

The requirement for Spns2 expression and activity in lymphocytes for their normal development and localization was further tested using bone marrow chimeras. Lethally irradiated Spns2<sup>+/+<sup> Rag1</sup></sup> recipients were reconstituted with a 50:50 mix of wild-type CD45.1<sup>+</sup>-marked and Spns2<sup>tm1a/tm1a</sup> bone marrows and analyzed at 8 wk after reconstitution. No significant differences in the development and localization of wild-type and Spns2<sup>tm1a/tm1a</sup> lymphocytes were observed in this study. For example, ~50% of CD4 and CD8 T cells in the blood and spleen of the chimeric mice were CD45.1<sup>2</sup> Spns2<sup>tm1a/tm1a</sup> (Fig. 5C). This demonstrated that Spns2<sup>tm1a/tm1a</sup> lymphocytes could develop and migrate normally when placed in a wild-type environment, and therefore that Spns2 expression in lymphocytes was dispensable for their normal development and localization. Additionally, an adoptive i.v. transfer of 10<sup>7</sup> Spns2 wild-type GFP-expressing splenocytes into either wild-type or Spns2<sup>tm1a/tm1a</sup> recipients demonstrated a significant reduction in the transferred CD4 T cells in the blood of Spns2<sup>tm1a/tm1a</sup> as compared with wild-type mice at 48 h, suggesting the role of the Spns2<sup>tm1a/tm1a</sup> environment in affecting CD4 T cell localization; however, no reduction in CD8 T cells and B cells was observed (data not shown).

Spns2 activity in the nonhematopoietic stromal cells is essential for normal immune function

The requirement for Spns2 expression and activity on different cell types for normal lymphocyte development and trafficking was

![FIGURE 3. Reduction in the numbers of recirculating B cells in the blood and mature B cells in the spleen and bone marrow of Spns2<sup>tm1a/tm1a</sup> mice. (A) Reduction in the percentage of B cells (gated as CD45<sup>+</sup>CD19<sup>+</sup>) in the blood of Spns2<sup>tm1a/tm1a</sup> mice. (B) Reduction in the absolute number of mature B cells (gated as B220<sup>+</sup>IgM<sup>+</sup>IgD<sup>+</sup>) in the spleen of Spns2<sup>tm1a/tm1a</sup> mice. (C) Representative flow cytometry plots of splenocytes, stained for B220, IgM, and IgD and gated on the B220<sup>+</sup> B cell population. Gates indicate IgM<sup>+</sup>IgD<sup>+</sup> transitional and IgM<sup>+</sup>IgD<sup>+</sup> mature B cells; the average percentage of cells within each gate for all mice in the group is indicated. (D) Reduction in the numbers of mature follicular B cells in the spleen of Spns2<sup>tm1a/tm1a</sup> mice confirmed by B220<sup>+</sup> CD21<sup>+</sup>CD23<sup>+</sup> staining. (E) Reduction in mature B cells (B220<sup>+</sup>IgM<sup>+</sup>IgD<sup>+</sup>) in the bone marrow of Spns2<sup>tm1a/tm1a</sup> mice. All bars represent means ± SEM. Data are from three mice per group and were reproducible in two independent experiments; bone marrow cell counts are per one tibia and femur. **p < 0.01, ***p < 0.001 using ANOVA with a Bonferroni post hoc test.

![FIGURE 4. Impaired humoral immune response in the Spns2<sup>tm1a/tm1a</sup> mouse line. Wild-type (+/+) and Spns2<sup>tm1a/tm1a</sup> (tm1a/tm1a) mice were immunized intranasally with TetC, boosted at days 7 and 21, and analyzed for Ag-specific Ab titers in the serum at day 36. Bars represent means ± SEM. Data are from six mice per group. *p < 0.05, **p < 0.01 using a Student t test.](http://www.jimmunol.org/)
S1P levels in the plasma of CD45.1+-marked and reconstituted with wild-type (CD45.1+) or with Spns2 tm1b/tm1b (CD45.1−) bone marrow and analyzed by flow cytometry at 8 wk after the reconstitution. The results indicated that the Spns2 genotype of the nonhematopoietic cells was of primary importance for normal lymphocyte development in the chimeras. Thus, when the wild-type hematopoietic system was reconstituted into Spns2 tm1a/tm1a hosts, the numbers of T cells were depleted in the blood and spleen to the same extent as in Spns2 tm1a/tm1a mice reconstituted with Spns2 tm1a/tm1a bone marrow (Fig. 6). Similarly, mature B cells were depleted to an equal extent in both groups of mice in the blood, spleen, bone marrow (Supplemental Fig. 2). In contrast, in the chimeric mice with selective loss of Spns2 function in the hematopoietic compartment, there was a trend toward decreased lymphocyte numbers but this did not reach statistical significance (Fig. 6, Supplemental Fig. 2), further indicating that Spns2 is primarily functional in the nonhematopoietic cells. Overall, these data suggested that Spns2 expression and function on the cells of the nonhematopoietic stroma had a primary role in the maintenance of normal lymphocyte development and immune system function.

No significant reduction in the plasma S1P levels in Spns2 tm1b/tm1b mice

S1P levels in the plasma of Spns2 tm1a/tm1a and wild-type mice were measured using two methods, an ELISA-based S1P assay (Echelon Biosciences; Fig. 7A) and the S1P redistribution assay that measures S1P-induced internalization of S1P1 (Thermo Scientific; Fig. 7B, 7C). In the latter assay, the plasma of wild-type and Spns2 tm1a/tm1a mice was added at different dilutions to U2OS cells expressing GFP-tagged S1P1, and the internalization of S1P1 GFP was quantified using a Cellomics Array-Scan VTI high-throughput cell imaging system. No significant differences in S1P concentrations or activity were observed between Spns2 tm1a/tm1a and wild-type mice in either assay (Fig. 7A, 7C). This indicated that Spns2-independent mechanisms exist for maintaining overall S1P levels in the blood of Spns2 tm1a/tm1a mice, but did not rule out the possibility that S1P levels are reduced in certain localized environments and that this was responsible for the altered lymphocyte distribution and immune function in Spns2 tm1a/tm1a mice. Normal viability and lack of gross developmental defects in the Spns2 tm1a/tm1a mice are also consistent with localized rather than systemic defects in S1P export. S1P concentrations in the lysates of spleen and thymus tissues were below the limit of detection of the Echelon Biosciences ELISA assay for both wild-type and Spns2 tm1a/tm1a mice (<0.06 μM; data not shown).

Characterization of Spns2 knockout Spns2 tm1b/tm1b mice

To further confirm that the Spns2 tm1a/tm1a mice were phenotypically equivalent to Spns2 knockout animals, the Spns2 tm1a/tm1a line was crossed to the C57BL/6N-Hprt tm1b/tm1b mice. The resulting allele structure was designated Spns2 tm1a/tm1a, and the homozygous Spns2 tm1b/tm1b mice were viable with no signifi-cant increase in embryonic mortality. Flow cytometry analysis of lymphoid organs demonstrated a reduction in CD4 and CD8 T cells and mature B cells in the spleen of Spns2 tm1b/tm1b mice (Supplemental Fig. 3B), comparable to Spns2 tm1a/tm1a mice (Figs. 1A, 2A). There was also an increase in the proportion of CD4+CD8− and CD4−CD8+ T cells in the thymus of Spns2 tm1b/tm1b mice (Figs. 1A, 2A).
mice, and these cells expressed higher levels of CD62L and lower levels of CD24, indicating their more mature status (Supplemental Fig. 3B), as seen previously in \( \text{Spns2}^{tm1a/tm1a} \) mice (Fig. 1B–E).

Overall, this confirmed that the \( \text{Spns2}^{tm1a/tm1a} \) phenotype is equivalent to the \( \text{Spns2}^{ knockout} \) phenotype. Additionally, to further confirm that the \( \text{Spns2}^{tm1a/tm1a} \) phenotype resulted from the gene-trap cassette in the \( \text{Spns2} \) locus, the \( \text{Spns2}^{tm1a/tm1a} \) animals were crossed to a transgenic line with systemic expression of Flp recombinase \( \text{C57BL/6N-Gt(Rosa)}^{26Sortm1(FLP1)Dym/Wtsi} \) (45), causing germline excision of the gene-trap cassette (Supplemental Fig. 3A). The resulting allele was designated \( \text{Spns2}^{tm1c(KOMP)Wtsi} \), and the \( \text{Spns2}^{tm1c/tm1c} \) mice showed a rescue of lymphocyte numbers in the spleen, bone marrow, and thymus (Supplemental Fig. 3C and data not shown), confirming that the immune phenotype of the \( \text{Spns2}^{tm1a/tm1a} \) line was caused by the \( \text{Spns2}^{tm1a} \) gene-trap cassette.

**Discussion**

In this study, we have characterized an \( \text{Spns2} \)-targeted mouse line and demonstrated that \( \text{Spns2} \) is required for normal lymphocyte development and localization, as well as for normal humoral immune response to immunization. Overall the changes in lymphocyte subpopulations in \( \text{Spns2}^{tm1a/tm1a} \) and \( \text{Spns2}^{tm1b/tm1b} \) mice closely mimicked the phenotypes of partial S1P deficiency and impaired S1P-dependent lymphocyte trafficking, including the depletion of lymphocytes in circulation, increase in the mature single-positive T cells in the thymus, and a selective reduction in the mature B cell population in the spleen and bone marrow (5, 6, 10, 31, 46). Although we did not detect a reduction in S1P levels in \( \text{Spns2}^{tm1a/tm1a} \) mouse plasma, the phenotypic data presented in this study together
with the previous in vitro demonstrations that human Spns2 can transport S1P and S1P mimic FTY720 (39, 42), suggest that the Spns2\textsuperscript{tm1a/	extltm1a} phenotype may arise from localized disruptions in S1P concentrations at certain restricted physiological locations.

Critically, this work indicates that Spns2 functions are limiting for lymphocyte trafficking with some degree of specificity, as the viability and lack of developmental defects in Spns2\textsuperscript{tm1a/	extltm1a} mice contrast with lethality and defects in cardiovascular and neural development in Spns2-mutant zebrafish (39, 40). Importantly, this is not due to different requirements for S1P production between the two species, as knockout mice lacking sphingosine kinases SphK1\textsuperscript{−/−} SphK2\textsuperscript{−/−} or S1P receptor S1P\textsubscript{1}\textsuperscript{−/−} are also embryonic lethal with abnormal cardiovascular development (30, 41). In comparison, Spns2\textsuperscript{tm1a/	extltm1a} mice more closely mimic the phenotypes of partially reduced S1P production, such as the single-null knockouts for SphK1 or SphK2 (53–55), or the knockout for S1P lyase with normal S1P production but disrupted S1P concentration gradients (56). Overall, this indicates that sufficient S1P levels are maintained in correct anatomical and cellular locations in the Spns2\textsuperscript{tm1a/	extltm1a} mice to allow normal embryonic development, and it suggests significant divergence in the expression and functions of Spns2 between mice and zebrafish. We can speculate that alternative mechanisms of S1P release may operate during embryonic development in mouse but not in zebrafish species. These may include other proteins that have been shown to transport S1P in vitro in mammalian cells, such as ABCG1 and ABCA1 (32, 33), or extracellular S1P production by secreted SphK1 enzyme (37), or even the two Spns2 paralogs Spns1 and Spns3.

This work further demonstrated that Spns2 activity in the nonhematopoietic cells of the stroma is of key importance for normal immune system function. In contrast, previous studies showed that plasma S1P levels in the mouse are maintained by hematopoietic cells such as erythrocytes (46), and ABC family transporters have been implicated in mediating S1P export from erythrocytes (32) and platelets (34). This suggests that the primary role of Spns2 is to maintain appropriate S1P concentrations at other in vivo locations, consistent with the unaltered S1P levels in the plasma of Spns2\textsuperscript{tm1a/	extltm1a} mice. Lymph is one of the sites where analysis of S1P concentrations would be particularly interesting (46), as S1P in the lymph was shown to be derived from nonhematopoietic cells, in particular the lymphatic endothelium (47). Overall, this warrants further investigation of Spns2 expression in different nonhematopoietic cell types and tissues, including lymphatic endothelium (46, 47).

S1P also acts as an intracellular messenger, and in mast cells and APCs increased S1P production is associated with cell activation, degranulation, and inflammatory cytokine production (18, 20, 23, 24). However, the present work indicates that in contrast to the SphK1 knockdown cells, Spns2\textsuperscript{tm1a/	extltm1a} macrophages respond normally to TNFR and TLR stimulation. This indicates that Spns2 likely does not affect intracellular S1P levels, at least in this cell type. Whether the activity of Spns2 and other S1P transporters can affect intracellular S1P levels in other cell types, by altering S1P secretion or uptake, remains to be addressed in future research.

Components of the S1P signaling pathways are targets for therapies aimed at treating autoimmune, transplant rejection, inflammatory diseases, and cancer (2). The S1P receptor agonist FTY720 is the most advanced of such therapies and was recently approved for the treatment of multiple sclerosis (57). Other pharmaceutical agents are under development and have shown efficacy in animal models of inflammatory diseases, sepsis, and cancer (27, 58, 59). Knowledge of the mechanisms regulating S1P concentrations in vivo in a mammal is essential for the future development of such pharmaceutical agents and may lead to better targeted therapies. For example, S1P-targeting therapies for the treatment of autoimmunity and transplant rejection aim to achieve immunosuppression without inhibiting other S1P activities. S1P-targeting cancer therapies suppress vascularization, cell growth, and migration, but ideally they aim to maintain full immune system function. In contrast, therapies for systemic inflammatory disorders primarily aim to suppress inflammation and intravascular coagulation while retaining the protective activities of S1P on endothelial barrier integrity (60, 61). Given the diverse roles of S1P in many physiological processes, understanding the mechanisms regulating its bioavailability in different tissues and under different conditions is essential for the development of such therapies. Transporter proteins have proven highly effective drug targets in other areas, particularly neuropharmacology (62). The demonstration that Spns2 deficiency selectively impaired lymphocyte functions and Ag-specific immune responses, without affecting vascular and neural development, highlights Spns2 as a possible drug target with potential for the treatment of autoimmunity and transplant rejection.

While this manuscript was under review, another study was published that substantially agrees with many of the findings presented in our study (63).

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Disclosures
The authors have no financial conflicts of interest.

References


Supplementary Figures and Table
The Role of Sphingosine-1-phosphate Transporter Spns2 in Immune System Function
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Figure S1. Characterization of Spns2<sup>tm1a/tm1a</sup> mouse phenotype. (A) Reduced white blood cell counts in Spns2<sup>tm1a/tm1a</sup> mice. Bars represent means ± SD, shaded areas represent 95% reference range for all wild type mice of same strain and sex. Blood was collected from terminally anaesthetized mice at 14 weeks of age into EDTA-coated tubes via the retro-orbital sinus, and analyzed on a ScilVet Animal Blood Counter. This data, together with the measurements of the percentages of the different lymphoid subpopulation in the blood (Figures 2A and 3A), indicated ~9.3 fold reduction in the absolute numbers of CD8 T cells, ~7.8 fold reduction in CD4 T cells, and ~3.9 fold reduction in B cells (CD19<sup>+)</sup> in the blood of Spns2<sup>tm1a/tm1a</sup> versus wild type mice. (B) Normal responses of Spns2<sup>tm1a/tm1a</sup> bone marrow derived macrophages (BMDM) to challenges with inflammatory and microbial stimuli. Spns2<sup>tm1a/tm1a</sup> and wild type cells were stimulated with LPS (100 ng/ml), IFNγ (25 ng/ml), and/or TNFα (25 ng/ml), over a 48 hour time-course. TNFα production was measured by ELISA at 4 hours of stimulation; expression of activation markers CD80 and CD86 was measured by flow cytometry at 4, 24, and 48 hours and no differences were seen at any time-points, data presented is from 48 hours. Bars represent means ± SEM, from 4 mice per group; MFI, mean fluorescence intensity.
Figure S2. Bone marrow chimeras experiments indicate that Spns2 expression in the non-hematopoietic cells of the stoma is required for normal B cell development and localization. Lethally irradiated (2 x 4.5Gy) recipients of Spns2^{tm1a/tm1a} and Spns2^{+/tm1a} genotypes were reconstituted either with wild type (+) or with Spns2^{tm1a/tm1a} (tm1a) donor bone marrow, and the numbers of lymphocyte subsets analyzed by flow cytometry at 8 weeks following reconstitution. (A) Numbers of transitional (B220\(^+\)IgM\(^+\)IgD\(^-\)) and mature (B220\(^+\)IgM\(^+\)IgD\(^+\)) B cells in the spleen of the chimeric mice. (B) Numbers of pre/pro-B cells (B220\(^+\)IgM\(^+\)IgD\(^+\)), immature (B220\(^+\)IgM\(^+\)IgD\(^-\)), and mature B cells (B220\(^+\)IgM\(^+\)IgD\(^+\)) in the bone marrow of the chimeric mice, per one tibia and femur. Means ± SEM from 3 mice per group are shown; statistical comparisons using ANOVA with Bonferroni’s post-hoc test to compare each dataset to the control group; * p<0.05, **p<0.01.
Figure S3. Characterization of Spns2 tm1b/tm1b and Spns2 tm1c/tm1c mouse phenotypes. (A) Spns2 tm1a allele structure showing: (left) the LoxP-flanked region that is deleted by Cre-recombinase to produce the Spns2 tm1b allele, lacking exon 3 and predicted to cause a frame-shift in the transcript and full loss of gene function; (right) the FRT-flanked region that is deleted by Flp-recombinase to produce the Spns2 tm1c allele, lacking the splice-acceptor cassette and predicted to allow normal expression of wild type Spns2-transcript. (B) Numbers of T and B cell populations in the spleen, percentages of single positive CD4 and CD8 thymocytes in the thymus, and the expression of CD62L and CD24 markers on the single-positive thymocytes of wild type (+/+) and Spns2 tm1b/tm1b (b/b) mice. (C, left) Numbers of CD4 and CD8 T cells in the spleen of Spns2 tm1c/tm1c (c/c) mice; numbers in the control groups are also shown, including wild type (+/+), Spns2 tm1a/tm1a (a/a), and heterozygous (+/a and +/c) animals. (C, right) Numbers of pre/pro-B cells (B220'IgM'IgD'), immature B cells (B220'IgM'IgD'), and mature B cells (B220'IgM'IgD') in the bone marrow and spleen of Spns2 tm1c/tm1c (c/c), as compared to wild type (+/+) and heterozygous (+/c) mice. Equivalent data from Spns2 tm1a/tm1a (a/a) animals is presented in Figures 3B,E. Bars represent means ± SEM from ≥1493 mice per group; bone marrow cell counts per one tibia and femur; statistical comparisons using t-test or ANOVA with Bonferroni’s post-hoc test, *p<0.05, **p<0.01, ***p<0.001; MFI – mean fluorescence intensity.
Table S1. Activity of the Spns2-promoter-driven β-galactosidase reporter in the tissues and organs of Spns2\textsuperscript{tm1a/} mice; data from three animals. β-galactosidase staining cannot be quantitatively compared across tissues and animals, and therefore “+” and “-” are used to indicate presence or absence of staining. The term 'no data' was assigned if the organ was not available. The term 'ambiguous' indicates uncertainty as to the validity of the observed pattern (staining may be very faint, background, or an artifact due to trapping).

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