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Accessibility
Ly9 (CD229) cell-surface receptor is crucial for the development of spontaneous autoantibody production to nuclear antigens

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The Signaling Lymphocyte Activation Molecule Family (SLAMF) genes, which encode cell-surface receptors that modulate innate and adaptive immune responses, lay within a genomic region of human and mouse chromosome 1 that confers a predisposition for the development of systemic lupus erythematosus (SLE). Herein, we demonstrate that the SLAMF member Ly9 arises as a novel receptor contributing to the reinforcement of tolerance. Specifically, Ly9-deficient mice spontaneously developed features of systemic autoimmunity such as the production of anti-nuclear antibodies (ANA), -dsDNA, and -nucleosome autoantibodies, independently of genetic background (B6.129) or (BALB/c.129). In aged (10- to 12-month-old) Ly9−/− mice key cell subsets implicated in autoimmunity were expanded, e.g., T follicular helper (Tfh) as well as germinal center (GC) B cells. More importantly, in vitro functional experiments showed that Ly9 acts as an inhibitory receptor of IFN-γ producing CD4+ T cells. Taken together, our findings reveal that the Ly9 receptor triggers cell intrinsic safeguarding mechanisms to prevent a breach of tolerance, emerging as a new non-redundant inhibitory cell-surface receptor capable of disabling autoantibody responses.

Keywords: SLAMF, Ly9 (CD229, SLAMF3), anti-DNA autoantibodies, disease susceptibility, systemic lupus erythematosus, murine Lupus
detected in lupus patients and lupus-prone mice had suffered isotype switching and somatic hypermutation, which facilitated the binding to self-antigens with high affinity, all indicating the involvement of GC pathways in this disease (21, 22). This, therefore, provides a rationale for determining not only the exact role played by SLAMF receptors as key drivers of Th and GC formation, but also their potential as appealing therapeutic targets for autoimmune-mediated diseases.

While recent reports have identified Slamf6 (Ly108) receptor and its isoforms and Slamf2 (28a) as contributing to the role played by Sle1b in tolerance (23–25), the involvement of other SLAMF members cannot be excluded. Of particular interest is the Ly9 (CD229, Slamf3) molecule, since a comparative analysis between B6 and the autoimmune congenic strain B6.Sle1b revealed significant differences in its isoforms usage, as well as in the extent of polymorphisms and expression levels (9), with the evidence indicating the possible participation of Ly9 in B6.Sle1b autoimmunity. Briefly, Ly9 expression is restricted to hematopoietic cells, including B and T lymphocytes (26). As has been shown in other SLAMF members, Ly9 functions as a homophilic adhesion receptor and its cytoplasmic tail contains two copies of the conserved tyrosine-based switch motif (ITSM), which is a docking site for the adapter molecules SAP and EAT-2 (27, 28). Ly9-deficient mice with a mixed 129 × B6 background exhibited no major T-cell developmental abnormalities and only very mild defects in T-cell responses (29). Recent findings demonstrate the role of Ly9 as a unique inhibitory cell-surface receptor regulating the size of the thymic innate CD8+ T-cell pool and the development of invariant Natural Killer T (iNKT) cells (30). Nonetheless, the functional role of Ly9 in lupus pathogenesis remains unknown. Here, we use Ly9-deficient mice, which were generated with 129-derived ES cells and then backcrossed onto B6 or BALB/c backgrounds, in order to determine the role of the Ly9 receptor in autoantibody development.

**MATERIALS AND METHODS**

**MICE**

Ly9−/− mice (129 × B6), generously provided by Dr. McKeon (29), were backcrossed onto BALB/c background for 12 generations to generate the Ly9−/− (BALB/c.129) strain and onto C57BL/6 (B6) background for 12 generations to generate the Ly9−/− (B6.129) strain. Eight-week-old BALB/c and B6 wild-type mice were purchased from Charles River Laboratories (Saint-Aubin-lès-Elbeuf, France). All mice strains were maintained under specific pathogen-free conditions. All samples were handled simultaneously under the same experimental conditions and results are expressed as OD values.

**IgG ISOTYPE DETECTION**

Basal serum IgG isotypes were determined by ELISA using purified goat anti-mouse IgG (Sigma-Aldrich) coated 96-well plates. 1:100 diluted mouse sera were incubated for 1 h at room temperature. After extensive washing, IgG isotypes were detected using a HRP-conjugated anti-mouse IgG and developed with substrate. All samples were handled simultaneously under the same experimental conditions and results are expressed as OD values.

**FLOW CYTOMETRY**

Single-cell suspensions were incubated with 20% heat-inactivated rabbit serum before being stained on ice with fluorophore-labeled antibodies against surface molecules using standard methods. Data was acquired using a FACSCanto II (BD Pharmingen, San Jose, CA, USA) flow cytometer and analyzed with either FACSDiva™ (BD Pharmingen) or FlowJo software (Tree Star, San Carlos, CA, USA). The following anti-mouse mAbs were obtained from BD Pharmingen: CD4-FITC, CD11b-PE, CD21-FITC, CD23-FITC, CD24-FITC, CD43-FITC, CD44-FITC, CD62L-FITC, CD69-FITC, CD154-PE, c-Kit-PE, Ter-119-PE, IgM-biotinylated, and CXCR5-biotinylated. The mAbs CD8-FITC, CD11b-FITC, CD25-PE, CD25-FITC, IgM-FITC, B220-FITC, as well as the isotype-matched control Abs, were acquired from ImmunoTools (Friesoythe, Germany). The following mAbs were obtained from BioLegend (San Diego, CA, USA): CD3-FITC, CD4-Pacific Blue, CD8-PE-Cy5, PD1-PE, PD1-PE-Cy7, B220-Pacific Blue, CD41-FITC, and IgD-APC-Cy7. The mAbs CD3-APC, CD5 PE-Cy7, CD229-APC, Sca-1-APC, and GL-7-FITC were purchased from eBioscience (San Diego, CA, USA). Anti-mouse CD138-APC was obtained from R&D Biosystems.
(R&D System, Wiesbaden, Germany). R-PE labeled murine CD1d tetramer pre-loaded with PBS57 (NIH Tetramer Core Facility, Atlanta, GA, USA) was used to detect iNKT cells according to the manufacturer’s instructions. Streptavidin PERCP-Cy5.5 was obtained from BD Pharmingen and streptavidin PE-Cy5 from eBioscience. For intracellular staining with IFN-γ and IL-17, cells were made permeable with an intracellular staining buffer (eBioscience). Anti-IFN-γ-PE (Clone XMG1.2, BD Pharmingen) or anti-IFN-γ-FTTC (XMG1.2; eBioscience), and anti-IL-17-APC (Clone TC11-18H10.1, BioLegend) were used for intracytoplasmic staining.

**KIDNEY HISTOLOGY AND URINE ASSAYS**

Kidneys were fixed with 4% formalin (PBS), dehydrated, and embedded in paraffin. All sections were counterstained with Gill’s hematoxylin (Panreac, Spain), dehydrated with graded alcohol and xylene, and mounted with DPX (VWR International, Radnor, PA, USA). To evaluate IgG-immunocomplex deposits on kidneys, snap-frozen spleens in OCT media (Sakura Finetek Europe B.V., The Netherlands) were cryosectioned (5 µm), blocked with 6% FCS in PBS, and stained with alexa 488-conjugated anti-mouse IgG (Lifet Technologies Corporation, Invitrogen, Paisley, UK). Images were analyzed using a Nikon Optiphot-2 microscope (Nikon) and acquired with a high-definition color camera (Nikon).

Freshly voided urine samples were tested for proteinuria using Albustix (Siemens Healthcare Diagnostics Inc. Tarrytown, NY, USA).

**IN VITRO CELL ACTIVATION**

Splenic lymphocytes were activated with plate-bound anti-CD3 (2 µg/ml) (145-2C11; BD Pharmingen) combined with purified soluble anti-CD28 (1 µg/ml) (37.51; BD Pharmingen). Splenocytes (100,000 cells/well) were cultured in RPMI 1640 medium supplemented with 10% FBS, 100 IU/ml of penicillin, 100 µg/ml of streptomycin, and 2.5 µM of β-mercaptoethanol in a 96-well plate and activated. Supernatants were collected after 72 h of incubation and IFN-γ levels were measured by ELISA. Additionally, after 24 h of activation, cells were collected and the percentages of activation markers (CD25, CD40L) were analyzed by flow cytometry. Th17 cell polarization was carried out according to BioLegend’s protocol. Briefly, single splenocyte suspensions were activated with 2 µg/ml of plate-bound anti-CD3 (clone 145-11; BioLegend), anti-CD28 (5 µg/ml clone 37.51, BD Pharmingen), TGF-β1 (1 ng/ml, BioLegend), anti-IFN-γ (10 µg/ml clone XMG1.2, BD Pharmingen), anti-IL-4 (10 ng/ml; clone 11B11; BioLegend), IL-6 (50 ng/ml, Immunotools), and IL-23 (5 ng/ml, BioLegend) over 3 days. Cells were then stimulated for 4–5 h with phorbol myristate acetate (PMA; 50 ng/ml; Sigma-Aldrich) and ionomycin (750 ng/ml; Sigma-Aldrich) in the presence of GolgiStop at 1:1500 (Pharmingen).

**MEASUREMENT OF CYTOKINES**

The amounts of IFN-γ in sera or in cell-culture supernatants were evaluated with commercially available ELISA kits (R&D Systems). Intracellular analysis of cytokines produced by CD4+ and CD8+ T cells, and iNKT cells was carried out by FACS analysis. Spleen CD4+ T cells were isolated by using MACS CD4+ T-cell Isolation Kit II (Miltenyi Biotec), following manufacturer’s protocol. Cells were stimulated for 4–5 h with PMA (50 ng/ml), ionomycin (750 ng/ml), and GolgiStop at 1:1500. APC-conjugated anti-CD3 (eBioscience), pacific blue-conjugated anti-CD4 (BioLegend), PE-Cy7-conjugated anti-CD8 (BioLegend), pacific blue-conjugated anti-B220 (BioLegend), and PE-conjugated tetramer-PBS57 (NIH) were used for flow cytometry staining. Cells were then fixed and permeabilized with Foxp3 Staining Buffer (eBioscience) before intracellular staining with alexa fluor 647-conjugated anti-IL-17 (clone TC11-18H10.1; BioLegend), phycoerythrin-conjugated anti-IFN-γ (XMG1.2; BD Pharmingen) or FITC-conjugated anti-IFN-γ (XMG1.2; eBioscience), IL-4 PE-Cy7 (11B11; BD Pharmingen).

**IN VITRO PROLIFERATION AND VIABILITY ASSAYS**

Splenocytes were resuspended in PBS with 5% FCS (106 cells/mL), and stained with CFSE (Invitrogen) (1 µL CFSE for each 106 cells), during 5 min at room temperature. Then cells were washed with cold PBS with 5% FCS and with cold PBS. After the second wash, lymphocytes were resuspended in RPMI media (10% FCS, and 2.5 µM β-mercaptoethanol). Cells were stimulated with 10 µg/mL F(ab')2 anti-mouse IgM (Jackson ImmunoResearch), or complete media as a negative control.

In order to measure viability, lymphocytes were stained with fluorochrome-labeled antibodies against surface antigens using standard methods. Cells were washed twice in azide-free and serum/protein-free PBS, and stained with Live/Dead Fixable far Red (Invitrogen) following manufacturer’s protocol. Then lymphocytes were stained with eFluor 450-conjugated Annexin V (eBioscience) during 15 min at room temperature, and washed with RPMI media. Proliferation and viability were assessed 72 h later.

**STATISTICS**

Mann–Whitney tests were used to calculate p values for all numeric data.

**RESULTS**

**Ly9 DEFICIENT MICE DEVELOP SPONTANEOUS AUTOIMMUNITY**

To assess the role of Ly9 in humoral autoimmune responses, Ly9-deficient mice were generated by mutated 129-derived ES cells (29) and backcrossed with B6 (C57BL/6) or BALB/c mice for 12 generations. First, we analyzed the presence of ANA in sera of 3, 6, 9, and 12 months old Ly9−/− (B6.129) mice by immunofluorescence staining of Hep-2 cells. As shown in Figures 1A,B, an age-associated increase in ANA titers was detected in Ly9−/− (B6.129) mice that by age 12 months was significantly higher than those of their age-matched B6.129 counterparts measured ≤3 months of age. At 12 months of age, 80% of the serum samples from Ly9−/− (B6.129) mice that was significantly higher than those of their age-matched Ly9−/− (wild-type; wt) counterparts. The differences in ANA serum levels between wt and Ly9−/− mice were statistically significant as early as 3 months of age. At 12 months of age, 80% of the serum samples from Ly9−/− (B6.129) showed ANA titers 1:4096. In contrast, 100% of the serum titers from their wt counterparts measured ≤1:512. Further analysis demonstrated that by age 12 months Ly9−/− (B6.129) mice displayed significant increases in their circulating levels of anti-double-stranded (ds) DNA and anti-nucleosome IgG antibodies in comparison to their wt counterparts (Figures 1C,D).

In order to further dissect the role of Ly9 in the humoral autoimmune response, excluding any effects stemming from
ANA titers in Ly9

ANA titers ≥ Ly9 increase in ANA titers was also detected in Frontiers in Immunology | Inflammation, although Figures 1C,D mice from both genetic backgrounds (B6 and Ly9 ground mice, the differences in ANA serum levels between matched wt counterparts. As was observed in the B6 back-

higher ANA titers. This likely reflects the presence of preferential nuclear antigen recognition based on genetic background, which has been also observed in other lupus models (31).

Ly9-deficient mice exhibited a significant increase in IgG2b (Figure 3A) as well as higher ratios of IgG2a, 2b, and IgG3 isotypes vs. IgG1 than wt mice. Due to the high titers of autoantibodies observed on aged Ly9-deficient mice sera, we further investigated the presence of alterations in these animals’ renal physiology and functionality. Twelve-month-old Ly9−/− (BALB/c.129) mice did not exhibit proteinuria (Figure 3B) or differences in their glomerular morphology compared to their wt counterparts. Although we observed a mild increase in IgG-immunocomplex deposits in Ly9-deficient mice (Figure 3C), this initial trait prior to any sign of glomerulonephritis did not trigger a renal pathology.

In summary, our findings reveal that the absence of the Ly9 receptor per se initiates the progressive development of autoantibodies, independently of any epistatic interactions.

episodic interactions, we determined autoantibodies in the serum of Ly9−/− (BALB/c.129) mice. Once again, an age-associated increase in ANA titers was also detected in Ly9−/− (BALB/c.129) mice that was significantly higher than those of their age-matched wt counterparts. As was observed in the B6 background mice, the differences in ANA serum levels between wt and Ly9−/− (BALB/c.129) mice were statistically significant as early as 3 months of age, reaching its highest level at age 12 months when 91.6% of Ly9−/− (BALB/c.129) mice presented ANA titers ≥ 1:256 compared with 72.7% of their wt counterparts whose titers measured ≤ 1:64 (Figures 2A,B). Notably, ANA titers in Ly9−/− (B6.129) mice were always higher than those of Ly9−/− (BALB/c.129) at the evaluated time points, most likely due to the additional effect of the epistatic interactions, which induced spontaneous loss of immune tolerance to nuclear antigens (25). Comparable levels of anti-dsDNA and anti-nucleosome were detected on Ly9-deficient mice from both genetic backgrounds (Figures 2C,D and Figures 1C,D), although Ly9−/− (B6.129) mice displayed much
**Ly9−/− AGED MICE EXHIBITED SPLENOMEGALY AND ALTERED KEY CELL SUBSETS RELATED TO SELF-TOLERANCE**

The role of Ly9 as an inhibitor molecule in the development of spontaneous autoimmunity, excluding any effects dictated by epistatic interactions, is further supported by the observation that 12-month-old Ly9−/− (BALB/c.129) mice exhibited splenomegaly, which is a feature often present in SLE-prone mice (32, 33). The mean and SEM of spleen weight in wt (n = 7) and Ly9−/− (BALB/c.129) mice (n = 11) were 95.53 ± 3.40 and 147.7 ± 10.36 mg, respectively (p = 0.0021). Consistently, an increase of 75.8% in Ly9−/− (BALB/c.129) spleen cell numbers, compared with wt spleens, was observed (Figures 4A,B).

A deeper analysis of cell subsets by flow cytometry demonstrated altered B- and T-cell peripheral homeostasis in Ly9-deficient mice. We first inspected the B-cell signature of the disease in 12-month-old Ly9−/− (BALB/c.129) mice (Table 1). Notably, the most significant difference was found in GC B cells, with a percentage of 0.64 ± 0.19 in wt and 1.28 ± 0.30 in Ly9-deficient mice (p = 0.0004), which was also evident due to the striking expansion of GC cell numbers (Figure 4C and Table 1). Although a major percentage of GC was observed in Ly9-deficient mice, only a slight decrease in the percentage of Follicular B cells was detected: 88.39 ± 3.15 (Ly9+/+) and 78.01 ± 7.25 (Ly9−−), which is indicative of follicular B-cell areas with a major proportion of GC B cells. Transitional 1 (T1) B cells in wt and Ly9−/− (BALB/c.129) mice showed percentages of 4.48 ± 1.08 and 9.47 ± 3.12, respectively. Marginal zone (MZ) B cells from Ly9-deficient mice also showed increased percentages: 4.54 ± 1.78
(Ly9<sup>+/+</sup>) and 8.70 ± 2.97 (Ly9<sup>−/−</sup>) (Table 1). Thus, the previously mentioned B-cell subsets, most of them involved in autoimmune diseases (34, 35), showed a near twofold percentage increase in Ly9<sup>−/−</sup>/mice compared to their control counterparts.

No altered proportion was found in the studied peritoneal B-cell subsets from the 12-month-old mice (data not shown). Our data also revealed a slightly increased frequency in the occurrence of the most immature B-cell lineages, multipotent progenitor (MMP) and Pro-B cells in the bone marrow of Ly9-deficient mice (data not shown), which is a variation that has been observed in other lupus-prone mice (36).

In examining the T-cell signature of Ly9 deficiency-mediated autoimmunity, the most remarkable difference we observed was in Tfh cells, which showed a threefold percentage increase and cell number expansion (Figure 4D). Importantly, excessive Tfh-cell numbers have been linked to a positive-selection defect in GC, which would account for such differences in the autoantibody generation (21). As previously mentioned, Ly9-deficient mice also exhibited higher ratios of IgG2a,b and IgG3 isotypes vs. IgG1, suggesting that the peripheral tolerance checkpoint that controls GC and Tfh cells becomes altered by the absence of Ly9 molecule (Figure 3A). In the Ly9-deficient spleen, we found a slight increase in the ratios occurring between CD4<sup>+</sup> and CD8<sup>+</sup> T cells. We also observed an increase in effector CD4<sup>+</sup> and CD8<sup>+</sup> T-cell subsets, although it was not significant. Interestingly, the nKRT cell pools in these animals were also enlarged: 3.53 ± 1.23% wt and 7.67 ± 2.80% Ly9<sup>−/−</sup> mice (Table 1).

We conclude that Ly9 gene ablation in a BALB/c background results in the disturbance of B and T cell subsets involved in autoimmunity, with major differences occurring in both Tfh cells and GC B cells.

**Ly9 RECEPTOR MODULATES IFN-γ SECRETION BY CD4<sup>+</sup> T CELLS**

To determine the lymphocyte subsets that facilitate the ignition of SLE-related pathology in the absence of Ly9, we searched for any altered peripheral cell populations in 8- to 12-week-old Ly9<sup>−/−</sup> (BALB/c.129) mice. Slight differences were observed in the T-splenic compartment with a small increase in percentage of CD4<sup>+</sup> T and iNKT. Notably, most of the alterations were displayed by B-cell subsets, with Transitional T1 B cells presenting the major difference (Table 2). A deeper examination of T1 and MZ B cells including IgM as a cell marker (T1:CD23<sup>−</sup> CD21<sup>−</sup> IgM<sup>+</sup>, MZ: CD23<sup>+</sup> CD21<sup>+</sup> IgM<sup>+</sup>) (37) demonstrates that Ly9-deficient mice possessed approximately a threefold increase of T1 subset and also an expanded MZ B subset (Figure S1 in Supplementary Material). Based on these results, we conclude that Ly9-deficiency alters the development of B-cell subsets which may be involved in the generation of autoantibodies.

Abnormalities in BCR signaling could shape the splenic B-cell populations and predispose to autoimmune disease. Therefore, we assessed the proliferation and survival of splenic B cell from Ly9<sup>−/−</sup> mice after IgM stimulation. The proliferation and apoptotic responses of Ly9-deficient B cells were similar to those observed in the wt B cells, with the exception of a slight decrease in percentage of late apoptotic cells in the Ly9-deficient mice (Figure 5).

Since earlier studies have demonstrate the critical implication of SLAMF receptors in Th1/Th2/Th17 polarization (14, 26, 38), we hypothesized that Ly9-deficient splenocytes in 8- to 12-week-old mice would foster an alteration in T-cell cytokine production prior to disease onset, thereby enabling autoantibody production at older ages. First, we asked whether the ablation of murine Ly9 could modulate IL-17 secretion by CD4<sup>+</sup> T cells under Th17 polarizing conditions, since earlier reports have implicated the human Ly9 receptor in IL-17 T-cell secretion (38, 39). No significant differences were detected in the percentage of IL-17 secreting CD4<sup>+</sup> T cells (Figure 6A). On the other hand, our group has previously shown that monoclonal antibodies against Ly9 negatively regulate TCR signaling, thereby inhibiting ERK phosphorylation and IFN-γ secretion (26, 40). Herein, we investigated the role of Ly9 in IFN-γ modulation by activating splenic T cells with anti-CD3 and anti-CD28. A significant increase in IFN-γ secretion, as well as an increase in the expression of CD40L in CD4<sup>+</sup> T cells was detected in the absence of Ly9 compared with wt cells (Figures 6B, C). A similar result was obtained when evaluating the expression of the CD25 activation marker (data not shown).
addition, PMA/ionomycin activation of splenic cells revealed that Ly9-deficient mice foster CD4+ T, CD8+ T, and iNKT cells capable of secreting major quantities of IFN-γ prior to the development of autoimmunity (Figure 6D). Furthermore, isolated Ly9-deficient CD4+ T cells consistently showed an increased IFN-γ production after PMA/ionomycin activation as compared with wt mice (Figure 6E). In contrast, no significant difference in percentage of IL-4 producing CD4+ T cells was observed (data not shown). In accordance with these results, 12-month-old Ly9−/− (BALB/c.129) mice exhibited an increased percentage of IFN-γ producing CD4+ T cells, which correlated with the high ANA titers detected in serum (Figure 6F). Nevertheless, we could not detect IFN-γ in the serum of these mice (data not shown).

Thus, in the absence of the Ly9 molecule occurs an increase in peripheral T and iNKT IFN-γ secretion, a key cytokine in the pathogenesis of SLE which has been previously shown to trigger the accumulation of pathogenic Tfh and GC B cells (41).

### Discussion

Despite extensive research, the mechanisms by which susceptibility and effector genes initiate and promote autoantibody production and tissue damage remain poorly defined. In the present study, we have shown that Ly9 deficiency results in a spontaneous loss of tolerance, reflected principally in the development of autoantibodies, a process which is thought to underlie the initiation of SLE syndrome.

We first found that the absence of the Ly9 gene in mice with a B6 and BALB/c background resulted in the development of autoantibodies. The loss of tolerance to nuclear antigens detected in Ly9−/− (B6.129) mice was reminiscent of those previously observed in B6.Sle1b and B6.129Chr1b congenic mouse strains (6, 42). We found that Ly9−/− (B6.129) mice developed autoantibodies as early as 12 weeks of age. In fact, by 9 months of age all mice were autoantibody positive, a pattern similar to that found in SLAMF receptor Slamf1-deficient (B6.129) mice (25). Most SLAMF receptor-deficient mice have been generated by altering those SLAMF genes located in chromosome 1 via the homologous recombination of 129-derived embryonic stem cells (25, 29, 43, 44). Once the resultant mice are backcrossed onto B6 mice [e.g., Slamf1−/− (B6.129)], the deficient mice are affected by the epistatic interaction that occurs between the 129 gene segment and the B6 genome. Thus, epistatic interactions between 129-derived and B6 genes in Ly9−/− (B6.129) mice greatly contribute to the autoantibody response. Exceptionally, the disruption of the Ly108 gene in congenic mice [Ly108−/− (B6.129)] mitigates the generation of autoantibodies, indicating that SLAMF members carry out opposing functions (e.g., Ly108 vs. Ly9).

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**Table 1 | Flow cytometry analysis of B- and T-splenic cell subsets from 12-month-old wt and Ly9−/− (BALB/c.129) mice.**

<table>
<thead>
<tr>
<th>Parental</th>
<th>Subset</th>
<th>Marker</th>
<th>Ly9+/+</th>
<th>Ly9−/−</th>
</tr>
</thead>
<tbody>
<tr>
<td>B220+</td>
<td>T1</td>
<td>B220+CD21+CD23−</td>
<td>4.48 ± 1.08</td>
<td>** 9.47 ± 3.12</td>
</tr>
<tr>
<td></td>
<td>Follicular-B</td>
<td>B220+CD21+CD232lymph</td>
<td>88.39 ± 3.15</td>
<td>** 78.01 ± 7.25</td>
</tr>
<tr>
<td></td>
<td>MZ-B</td>
<td>B220+CD21+CD23−/low</td>
<td>4.54 ± 1.78</td>
<td>** 8.70 ± 2.97</td>
</tr>
<tr>
<td>Lymphocytes</td>
<td>GC B cells</td>
<td>B220+GL7+</td>
<td>0.64 ± 0.19</td>
<td>*** 1.28 ± 0.30</td>
</tr>
<tr>
<td></td>
<td>Plasma B cells</td>
<td>B220−low CD138+</td>
<td>0.19 ± 0.03</td>
<td>0.24 ± 0.03</td>
</tr>
<tr>
<td></td>
<td>CD4</td>
<td>CD4−/− CD8+</td>
<td>73.55 ± 4.00</td>
<td>76.65 ± 5.66</td>
</tr>
<tr>
<td>T- AND NK-CELL LINEAGE</td>
<td>CD8 SP CD4+ CD4−/− CD8+</td>
<td>21.03 ± 4.08</td>
<td>16.28 ± 3.2</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Effector CD4</td>
<td>CD3−/− CD4+ CD8−/−</td>
<td>42.87 ± 4.26</td>
<td>50.50 ± 10.01</td>
</tr>
<tr>
<td></td>
<td>Effector CD8</td>
<td>CD3−/− CD8+ CD4−/−</td>
<td>17.53 ± 12.18</td>
<td>35.27 ± 15.02</td>
</tr>
<tr>
<td></td>
<td>Reg T cells</td>
<td>CD3−/− CD4+CD25+</td>
<td>17.16 ± 1.62</td>
<td>20.10 ± 15.02</td>
</tr>
<tr>
<td></td>
<td>Tth</td>
<td>CD3−/− CD4+PD1+</td>
<td>3.27 ± 1.10</td>
<td>9.90 ± 3.09</td>
</tr>
<tr>
<td></td>
<td>Tefh</td>
<td>CD4−/−CD4−/−PD1+CD62L−/low</td>
<td>3.06 ± 1.42</td>
<td>5.29 ± 2.00</td>
</tr>
<tr>
<td></td>
<td>iNKT</td>
<td>CD3−/− CD8+</td>
<td>3.53 ± 1.23</td>
<td>** 7.67 ± 2.8</td>
</tr>
<tr>
<td></td>
<td>CD1d Tetra</td>
<td>CD3−/− CD8+</td>
<td>3.53 ± 1.23</td>
<td>** 7.67 ± 2.8</td>
</tr>
<tr>
<td></td>
<td>NK</td>
<td>CD3−/− DX5+</td>
<td>3.55 ± 1.02</td>
<td>3.60 ± 0.74</td>
</tr>
</tbody>
</table>

**T1, transitional T cells; MZ-B, marginal zone B cells; GC B cells, germinal center B cells; SP, single positive; Reg, regulatory; Tfh, follicular helper T cells; iNKT, invariant natural killer T cells; NK, natural killer; p-value significance; **p < 0.01, ***p < 0.001. Results represent the mean ± SEM of 7 wt and 11 Ly9−/− (BALB/c.129) mice for each subset analyzed.
We next assessed the role of Ly9 in autoimmune disease without the confounding influence of mixed haplotypes by utilizing Ly9−/− (BALB/c.129) mice. Although autoantibody titers were lower compared to Ly9+/− (B6.129) mice, this strain clearly developed an autoimmune response based on the significant increases of ANA, anti-dsDNA, and anti-chromatin antibodies compared to their wt counterparts. By contrast, other SLAMF receptor-deficiencies embedded in the BALB/c genome, such as Slamf1−/− (BALB/c.129) and Slamf2−/− (BALB/c.129), do not develop any autoimmune response (25), underscoring the role played by Ly9 as a negative regulator in the pathogenesis of lupus. Consequently, among the various SLAMF receptors, both Ly9 and 2B4 rise as unique factors contributing to the reinforcement of tolerance (24).

As a consequence of breaching tolerance, a plethora of disorders can develop a posteriori; e.g., T and B cells are reportedly involved in the amplification and perpetuation of the autoimmune response, resulting in inflammation and cytokine dysregulation (43). This proved to be the case in our 12-month-old Ly9-deficient mice in which various B and T cells subsets underwent alterations. Notably, the most noteworthy differences in older mice were found in Tfh and GC B cells, two cell types which have been shown to preferentially express the Ly9 receptor (46).

In order to begin to understand why Ly9-deficient mice developed spontaneous autoimmunity, we search for any abnormal peripheral B and T cell development prior to full autoantibody disorder on 8- to 12-week-old Ly9-deficient mice. We observed alterations in the B-cell splenic compartment, with the most prominent expansion displayed by Ly9-deficient transitional T1 B cells. Interestingly, SLE patients present increased numbers of T1 cells, although their role in lupus is still ill defined (35). In addition, 8- to 12-week-old Ly9-deficient mice displayed an increase in MZ B cells. Even though many autoreactive antibodies appear to be the product of GC reactions, major evidence begins to reveal that MZ B cells play a key role in homeostasis and tolerance. Notably, the MZ B-cell expansion has been directly implicated in lupus pathogenesis in some murine models (47–50), but not others (51, 52). Autoimmunity mediated by B cell is usually linked to a B-cell hyperresponsiveness upon BCR stimulation, change B-cell SHP-1, FcRII, CD22, Cbl-b, or overexpressing CD19, leads to a cytokine imbalance prior to disease onset. As a consequence of breaching tolerance, a plethora of disorders can develop a posteriori; e.g., T and B cells are reportedly involved in the amplification and perpetuation of the autoimmune response, resulting in inflammation and cytokine dysregulation (43). This proved to be the case in our 12-month-old Ly9-deficient mice in which various B and T cells subsets underwent alterations. Notably, the most noteworthy differences in older mice were found in Tfh and GC B cells, two cell types which have been shown to preferentially express the Ly9 receptor (46).

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Table 2 | Flow cytometry analysis of B- and T-splenic cell subsets from 8- to 12-week-old wt and Ly9−/− (BALB/c.129) mice.

<table>
<thead>
<tr>
<th>Parental Subset</th>
<th>Marker</th>
<th>Ly9+/−</th>
<th>Ly9−/−</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>B-CELL LINEAGE</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>B220+</td>
<td>T1</td>
<td>B220+CD21−CD23low</td>
<td>10.87 ± 0.71</td>
</tr>
<tr>
<td></td>
<td>Follicular-B</td>
<td>B220+CD21+CD23high</td>
<td>79.96 ± 0.84</td>
</tr>
<tr>
<td></td>
<td>MZ</td>
<td>B220+CD21+CD23low</td>
<td>4.76 ± 0.21</td>
</tr>
<tr>
<td><strong>Lymphocytes</strong></td>
<td>GC B cells</td>
<td>B220+GL7+</td>
<td>4.65 ± 0.65</td>
</tr>
<tr>
<td></td>
<td>Plasma B cells</td>
<td>B220−CD138+</td>
<td>0.27 ± 0.06</td>
</tr>
<tr>
<td><strong>T-AND NK-CELL LINEAGE</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>CD3+</td>
<td>CD4 SP</td>
<td>CD4+CD8−</td>
<td>65.93 ± 6.27</td>
</tr>
<tr>
<td></td>
<td></td>
<td>CD4−CD8−</td>
<td>16.81 ± 4.78</td>
</tr>
<tr>
<td>CD4+</td>
<td>Effector CD4</td>
<td>CD3+CD4+CD8−</td>
<td>28.58 ± 1.66</td>
</tr>
<tr>
<td></td>
<td>Effector CD8</td>
<td>CD3+CD8+CD4−</td>
<td>7.50 ± 2.80</td>
</tr>
<tr>
<td></td>
<td>Reg T cells</td>
<td>CD3+CD4−CD25+</td>
<td>8.77 ± 1.22</td>
</tr>
<tr>
<td></td>
<td>Tfh</td>
<td>CD3+CD4−PD1+</td>
<td>3.40 ± 0.44</td>
</tr>
<tr>
<td></td>
<td>iNKT</td>
<td>B220−CD3+</td>
<td>4.22 ± 0.53</td>
</tr>
<tr>
<td></td>
<td></td>
<td>CD1d Tetr+</td>
<td></td>
</tr>
<tr>
<td>CD3−</td>
<td>NK</td>
<td>CD3−DX5−</td>
<td>0.82 ± 0.32</td>
</tr>
</tbody>
</table>

T1, transitional 1 B cells; MZ-B, marginal zone B cells; GC B, germinal center B cells; SP, single positive; Reg, regulatory; Tfh, follicular helper T cells; iNKT, invariant natural killer T cells; NK, natural killer; p-value significance; *p < 0.05, ***p < 0.0001. Results represent the mean±SEM of wt and Ly9−/− (BALB/c.129) mice (n = 5) for each subset analyzed.
under Th17 polarizing conditions results in an increase in IL-17 (38, 39). No alterations in IL-17 producing T lymphocytes were observed once Ly9-deficient mice splenic cells were activated under Th17 polarizing conditions. Previous reports showed a diminishing IL-4 production and no altered IFN-γ secretion by Ly9-deficient CD4 T cells (29). In contrast, in the absence of Ly9, we observed an increase in IFN-γ producing T cells with no significant alteration in IL-4 secretion. Importantly, IFN-γ has long been associated with lupus (54). This apparent contradiction could be explained by the influence of epistatic interactions as well as the mice background, since previous studies were performed in Ly9−/− mice with a mixed background (B6 × 129) mice. Recent reports have demonstrated that the overproduction of IFN-γ induces an aberrant accumulation of Tfh and GC cells (41). We also found that these subsets underwent expansion in Ly9-deficient mice, when the influence of confounding epistatic interactions was absent. These observations suggest that the Ly9 molecule may play an inhibitory role in the expansion of these subsets. In addition, Ly9-deficient mice showed a skewed isotype switching toward IgG2a/b, an isotype induced by a Th1 response that requires T cell-stimulated B lymphocytes. Although IgG2a antibodies are the most pathogenic class of immunoglobulin (55,
Ly9 receptor mitigates autoimmune responses

José de Salort performed most of the experiments and analyzed the data; Marta Cuenca performed the B-cell functional experiments; Pablo Engel and Xavier Romero designed the experiments and supervised the study; José de Salort, Pablo Engel, and Xavier Romero wrote the manuscript; Cox Terhorst provided input into the conceptual development of the experiments and edited the manuscript. We thank Adriana Lázaro for technical assistance. We also thank the NIH Tetramer Core Facility for providing R-Pe labeled murine CD1d tetramer pre-loaded with PBS57. This work was supported by the Ministerio de Educación y Ciencia through Grants SAF 2009-07071/SAF 2012-39536 (to Pablo Engel) and by a grant from the NIH (2P01AI65687-06A1 to Pablo Engel and Cox Terhorst). José de Salort was supported by Ayuda Personal Investigador en Formación (APIF), a grant from the Universitat de Barcelona. Marta Cuenca was supported by Ministerio de Educación, Cultura y Deporte (AP2010-1754). Xavier Romero was supported by Beatriz de Pinós (2010 BP-B).

SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at http://www.frontiersin.org/Inflammation/10.3389/fimmu.2013.00225/abstract

Figure S1 | Splenic T1 and MZ1 cells are expanded in Ly9−/− (BALB/c;129) mice. (A) Splenocyte lymphocytes from 8- to 12-week-old wt (n = 5) and Ly9−/− (BALB/c;129) mice (n = 5) were stained using CD23, CD21, and IgM. (B) Representative dot plots from wt and Ly9−/− splenic cells. The gating strategy to characterize transitional and marginal zone (MZ) B cells is shown. (B) Quantitative analysis of the CD23 percentage of MZ and Transitional 1 (T1) B cells as well as (C) MZ and T1-B cells perispleen are shown. SEM and statistical significances are shown. References

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Ly9 receptor mitigates autoimmune responses

Conflict of Interest Statement: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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