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BRIEF DEFINITIVE REPORT

Sarco/endoplasmic reticulum Ca²⁺-ATPase (SERCA) activity is required for V(D)J recombination

Chun-Chin Chen^{1*}, Bo-Ruei Chen^{2,3*}, Yinan Wang¹, Philip Curman^{4,5}, Helen A. Beilinson⁶, Ryan M. Brecht⁶, Catherine C. Liu⁶, Ryan J. Farrell^{7,10}, Jaime de Juan-Sanz⁷, Louis-Marie Charbonnier⁸, Shingo Kajimura⁹, Timothy A. Ryan⁷, David G. Schatz⁶, Talal A. Chatila⁸, Jakob D. Wikstrom^{4,5}, Jessica K. Tyler¹, and Barry P. Sleckman^{2,3}

A whole-genome CRISPR/Cas9 screen identified *ATP2A2*, the gene encoding the Sarco/endoplasmic reticulum Ca²⁺-ATPase (SERCA) 2 protein, as being important for V(D)J recombination. SERCAs are ER transmembrane proteins that pump Ca²⁺ from the cytosol into the ER lumen to maintain the ER Ca²⁺ reservoir and regulate cytosolic Ca²⁺-dependent processes. In preB cells, loss of SERCA2 leads to reduced V(D)J recombination kinetics due to diminished RAG-mediated DNA cleavage. SERCA2 deficiency in B cells leads to increased expression of SERCA3, and combined loss of SERCA2 and SERCA3 results in decreased ER Ca²⁺ levels, increased cytosolic Ca²⁺ levels, reduction in *RAG1* and *RAG2* gene expression, and a profound block in V(D)J recombination. Mice with B cells deficient in SERCA2 and humans with Darier disease, caused by heterozygous *ATP2A2* mutations, have reduced numbers of mature B cells. We conclude that SERCA proteins modulate intracellular Ca²⁺ levels to regulate *RAG1* and *RAG2* gene expression and V(D)J recombination and that defects in SERCA functions cause lymphopenia.

Introduction

All developing lymphocytes must assemble the second exon of genes encoding antigen receptor chains (Bassing et al., 2002). This occurs through V(D)J recombination, a reaction where the RAG endonuclease, composed of the RAG1 and RAG2 proteins, introduces DNA double-strand breaks (DSBs) at the border of RAG recognition sequences (RSs) and variable (V), diversity (D), or joining (J) gene segments undergoing recombination (Fugmann et al., 2000). RAG-mediated DNA cleavage results in a signal end (SE) and coding end (CE) at each DSB, and the nonhomologous end-joining pathway of DNA DSB repair joins the two SEs to form a signal join (SJ) and the two CEs to form a coding join (CJ; Helmink and Sleckman, 2012).

The V(D)J recombination reaction can be studied in mouse preB cells transformed by the viral abl kinase, hereafter referred to as abl preB cells (Bredemeyer et al., 2006; Helmink and Sleckman, 2012). Treating these cells with the abl kinase inhibitor imatinib leads to cell cycle arrest in G0/G1, induction of *RAG1* and *RAG2* gene expression, and V(D)J recombination at the

endogenous immunoglobulin light chain κ (*Igk*) locus and chromosomally integrated retroviral V(D)J recombination substrates (Bredemeyer et al., 2006; Helmink and Sleckman, 2012; Hung et al., 2018; Muljo and Schlissel, 2003). Here we used abl preB cells for a CRISPR/Cas9 whole-genome guide RNA (gRNA) screening approach to identify novel genes encoding proteins required for V(D)J recombination. This approach revealed the Sarco/ER Ca²⁺-ATPase 2 (SERCA2) protein, encoded by the *ATP2A2* gene, as a potential regulator of V(D)J recombination.

SERCA proteins are ATP-dependent Ca²⁺ transporters that move Ca²⁺ from the cytosol into the ER in order to maintain low cytosolic, and high ER, Ca²⁺ concentrations (Stammers et al., 2015; Vandecaetsbeek et al., 2011). There are two additional SERCA family members, SERCA1 and SERCA3, encoded by distinct genes, *ATP2A1* and *ATP2A3*, respectively (Stammers et al., 2015; Vandecaetsbeek et al., 2011). SERCA1 is expressed primarily in skeletal muscle, and SERCA3 is expressed in non-muscle cells, including lymphocytes (Vandecaetsbeek et al., 2011;

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Wu et al., 1995; Wuytack et al., 1994). Mutations in *ATP2A2* result in a rare autosomal dominant skin disorder, Darier disease (Ahn et al., 2003; Burge and Wilkinson, 1992; Leong et al., 2017; Tavadia et al., 2002).

Here we show that in developing B cells, SERCA2 and SERCA3 have redundant functions required to promote *RAG1* and *RAG2* expression and V(D)J recombination. Moreover, SERCA2 deficiency leads to B lymphopenia in mice and humans. We propose that SERCA proteins regulate intracellular Ca^{2+} homeostasis in a way that is required for normal *RAG1* and *RAG2* expression and efficient V(D)J recombination.

Results and discussion

SERCA2 is required for efficient V(D)J recombination

We generated WT abl preB cells containing a doxycycline-inducible *Cas9* cDNA and pMGINV retroviral V(D)J recombination substrate (Fig. 1 A; Hung et al., 2018). pMGINV has an anti-sense GFP cDNA flanked by two RSs that mediate V(D)J recombination by inversion placing the GFP cDNA in the sense orientation leading to GFP expression (Hung et al., 2018). A lentiviral murine whole-genome gRNA library was introduced into these cells followed by doxycycline treatment for 7 d to promote *Cas9* expression and gene inactivation (Tzelepis et al., 2016). These cells were treated with imatinib for 4 d to induce RAG expression and V(D)J recombination of pMGINV (Fig. 1 A). Abl preB cells that had not (GFP⁻) or had (GFP⁺) completed V(D)J recombination were isolated by flow-cytometric cell sorting, and gRNAs were sequenced from both populations.

GFP⁻ cells exhibited significant enrichment in gRNAs to *Rag1* and *Rag2*, and *Lig4* and *Xrcc6*, which encode the DNA ligase 4 and Ku70 proteins, respectively, that are required for the repair of RAG-mediated DSBs, demonstrating the veracity of our approach (Table S1; Helmink and Sleckman, 2012). Several distinct gRNAs to *Atp2a2* were significantly enriched (approximately sixfold) in the GFP⁻ population, suggesting that the SERCA2 may regulate V(D)J recombination (Table S1).

SERCA2 promotes RAG cleavage

To validate our finding, we generated *Atp2a2* “bulk inactivated” abl preB cells by introducing a gRNA to the 5' region of *Atp2a2* into WT abl preB cells containing a doxycycline-inducible *Cas9* cDNA and pMGINV. These cells were treated with doxycycline for 6 d, leading to inactivation of most *Atp2a2* alleles in the bulk population, as evidenced by the loss of SERCA2 protein (Fig. 1 B). *Atp2a2* mRNA splice variants leading to different SERCA2 isoforms are generated by alternative splicing in the 3' region of *Atp2a2* and can be detected by the antibody we use for Western blotting (Lipskaia et al., 2014; Vandecaetsbeek et al., 2011). Thus, we conclude that SERCA2 isoforms are not expressed in the bulk *Atp2a2* inactivated abl preB cells. Bulk *Atp2a2* inactivated abl preB cells exhibited diminished pMGINV V(D)J recombination kinetics, as evidenced by the reduced fraction of GFP⁺ cells at 2 and 4 d after imatinib treatment (23% and 77%, respectively) compared with WT cells (56% and 88%, respectively; Fig. 1, C and D). Several *Atp2a2*^{-/-} abl preB cell clones (c16, c21, c25, c54, c67, c72, c224, and c225) were generated from two distinct lines of

WT abl preB cells, and SERCA2 loss was confirmed by Western blotting (Fig. 1 E and Fig. S1 A). As assessed by flow cytometry, *Atp2a2*^{-/-} abl preB cell clones also exhibited impaired V(D)J recombination kinetics compared with WT abl preB cells (Fig. 1 F and Fig. S1 B).

To determine whether the requirement for SERCA2 is at the RAG DNA cleavage or DNA DSB repair step of V(D)J recombination, we assayed pMGINV by Southern blotting, which permits the detection of unrepaired SEs and normally repaired SJs (Fig. 1 A). In *Atp2a2*^{-/-} abl preB cells treated with imatinib, the hybridizing band reflecting pMGINV SJs in *NheI*-digested DNA formed with slower kinetics than that in WT cells (Fig. 1 G and Fig. S1 C). Moreover, while hybridizing bands reflecting unrepaired pMGINV SEs were readily detected in *XbaI*-digested DNA from imatinib-treated *Lig4*^{-/-} abl preB cells, they were not detected in *Atp2a2*^{-/-} abl preB cells (Fig. 1 G and Fig. S1 C). We conclude that loss of SERCA2 leads to diminished V(D)J recombination due to impaired RAG cleavage.

SERCA2 and SERCA3 function redundantly during V(D)J recombination

SERCA3, a less potent ER Ca^{2+} transporter, is often coexpressed with SERCA2, including in lymphoid cells (Chandrasekera et al., 2009; Vandecaetsbeek et al., 2011; Wu et al., 1995; Wuytack et al., 1994). Abl preB cells could compensate for loss of SERCA2 by up-regulating SERCA3 (Lipskaia et al., 2014). Indeed, bulk *Atp2a2* inactivated abl preB cells had increased levels of *Atp2a3* transcripts (Fig. 2 A) and SERCA3 protein (Fig. 2 B). While bulk *Atp2a3* inactivation led to a near complete loss of SERCA3 protein in WT abl preB cells, it caused only partial loss of SERCA3 in *Atp2a2*^{-/-} abl preB cells, suggesting that these cells cannot tolerate the loss of SERCA2 and SERCA3 (Fig. 2 C). Bulk *Atp2a3* inactivation in WT abl preB cells did not lead to significant defects in V(D)J recombination of pMGINV (Fig. 2 D). However, bulk *Atp2a3* inactivation in *Atp2a2*^{-/-} abl preB cells, which led to a reduction in SERCA3 protein, resulted in a near complete block in pMGINV V(D)J recombination (Fig. 2 D).

Southern blot analysis of V(D)J recombination at the *Igk* locus yielded similar results (Fig. 2, E and F). Treatment of WT abl preB cells with imatinib led to a progressive reduction in the intensity of the hybridizing band corresponding to the unrearranged *Igk* locus and the appearance of heterogeneously sized bands reflecting diverse *Vκ* to *Jκ* rearrangements (Fig. 2, E and F). Southern blot analyses of cells with bulk *Atp2a2* inactivation or clonal *Atp2a2*^{-/-} abl preB cells revealed a mild reduction in the kinetics of *Igk* locus rearrangement (Fig. 2, F and G). Strikingly, bulk *Atp2a3* inactivation in *Atp2a2*^{-/-} abl preB cells led to a near complete block of *Igk* rearrangements with persistence of the band reflecting the unrearranged *Igk* locus and undetectable *Vκ* to *Jκ* rearrangements after imatinib treatment for up to 5 d (Figs. 2, F and G). We conclude that SERCA2 and SERCA3 have redundant functions that are critical for the RAG cleavage step of the V(D)J recombination reaction.

Loss of SERCA2 and SERCA3 disrupts Ca^{2+} homeostasis

We assessed ER Ca^{2+} levels in abl preB cells using ER-GCaMP6-150, a low-affinity calcium indicator designed to detect μ M Ca^{2+}

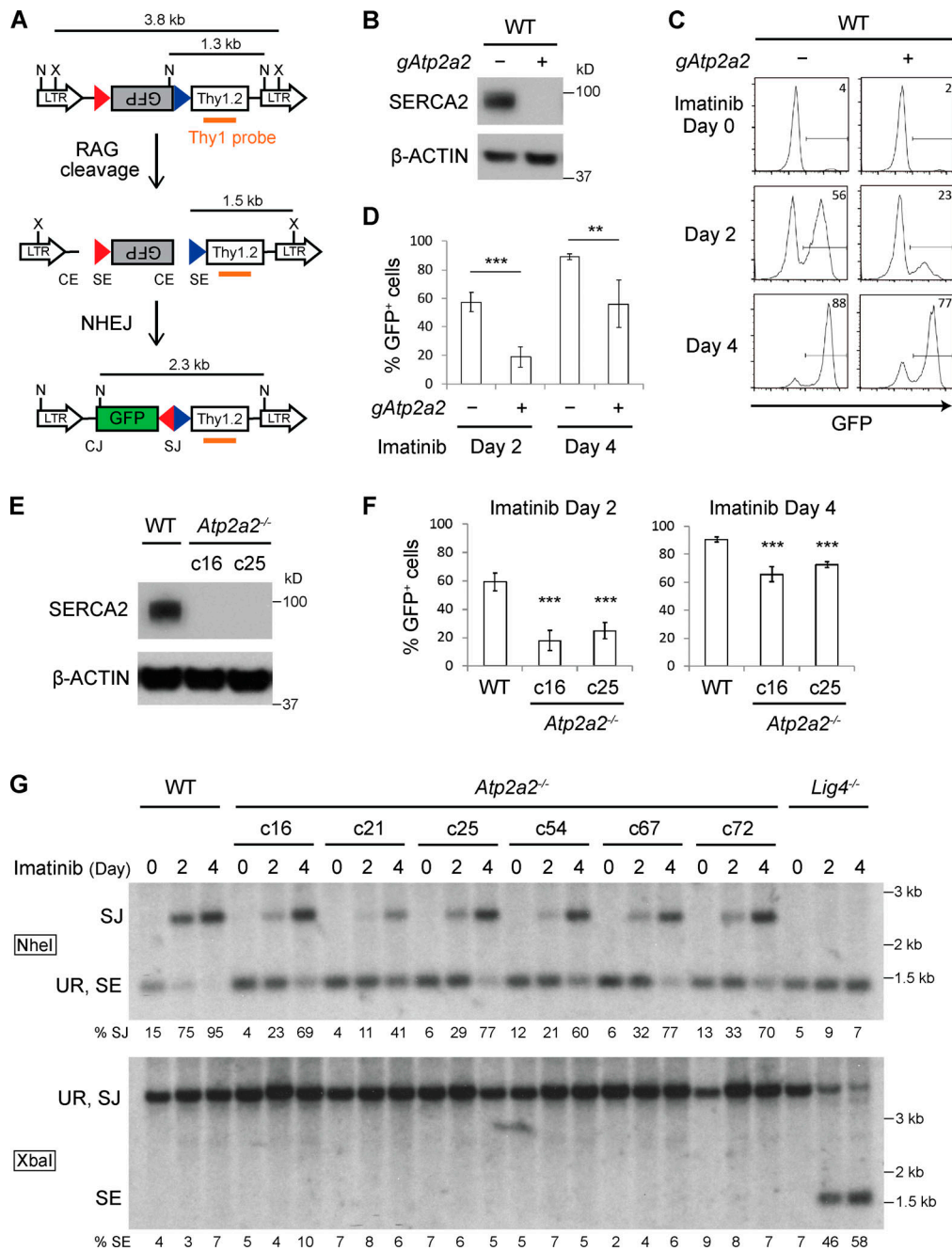


Figure 1. Loss of SERCA2 impairs RAG cleavage. (A) Schematic of pMGINV showing the antisense (gray rectangle) and sense (green rectangle) GFP cDNA, RSs (red and blue triangles), CE, SE, CJ, and SJ. The Thy1.2 cDNA (open rectangle), Thy1 probe (orange line), and *NheI* (N) and *XbaI* (X) sites are indicated. (B) Western blot for SERCA2 and β -actin in WT *abl* preB cells (Tet-On Cas9 clone 302) without (-) an *Atp2a2* gRNA (*gAtp2a2*) and bulk *Atp2a2* inactivated with (+) *gAtp2a2* ($n = 5$). (C) Flow-cytometric analysis for GFP expression in WT (-) or bulk *Atp2a2* inactivated (+) *abl* preB cells with pMGINV that were treated with imatinib for 0, 2, or 4 d. Percentage of GFP-expressing cells is indicated. (D) Mean \pm SD of five experiments performed as in C. P values were calculated using Student's *t* test throughout the figures. **, $P < 0.01$; ***, $P < 0.001$. (E) Western blot analysis of SERCA2 in two *Atp2a2*^{-/-} *abl* preB clones (c16 and c25) and the parent WT *abl* preB cell line (c302) as described in B ($n = 3$). (F) Analysis of pMGINV rearrangement in *Atp2a2*^{-/-} *abl* preB clones and WT *abl* preB cells as described in D (mean \pm SD; $n = 4$). ***, $P < 0.001$ compared with WT. (G) Southern blot analyses of *NheI* (top) or *XbaI* (bottom) digested genomic DNA probed with the Thy1 probe as shown in A. Shown are *Lig4*^{-/-}, WT, and *Atp2a2*^{-/-} *abl* preB clones treated with imatinib for 0, 2, or 4 d. Bands reflecting unrearranged (UR) pMGINV and pMGINV SEs and SJs are indicated. The percentages of pMGINV that have SJs (%SJ) or SEs (%SE) are indicated and quantified as described in Materials and methods ($n = 4$ for SJ analysis in *Atp2a2*^{-/-} c25; $n = 1$ for other *Atp2a2*^{-/-} clones).

concentrations present in the ER lumen (de Juan-Sanz et al., 2017). This indicator contains ER-targeting and retention sequences to ensure high-fidelity measurement of ER Ca²⁺. Upon Ca²⁺ binding, ER-GCaMP6-150 fluorescence changes ~45-fold,

similar to the original GCaMP6 reporter (de Juan-Sanz et al., 2017). ER-GCaMP6-150 was expressed in WT *abl* preB cells, and GFP fluorescence indicative of ER Ca²⁺ could be readily detected by flow cytometry (Fig. 3 A). Treating these cells with the SERCA

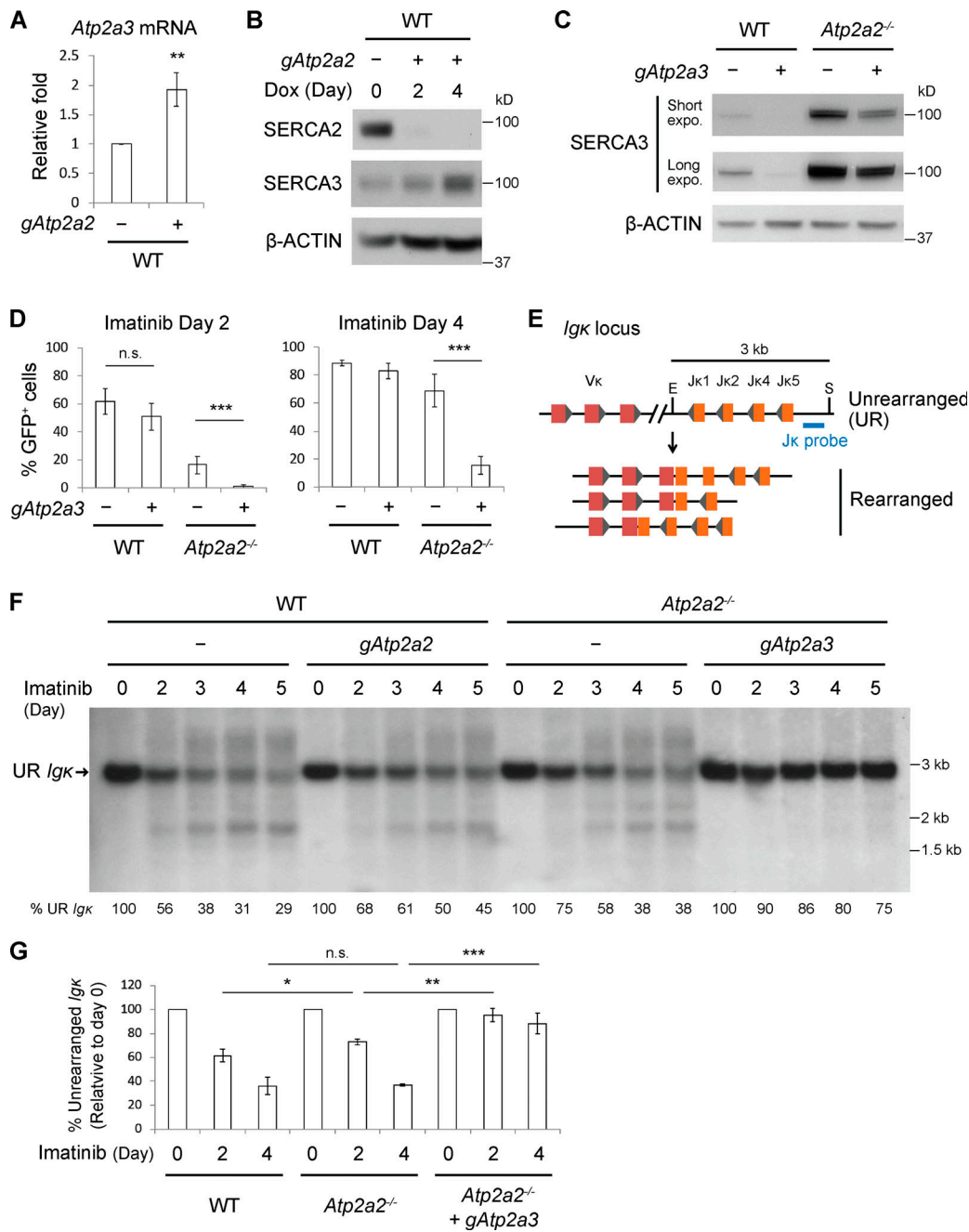


Figure 2. SERCA3 supports V(D)J recombination in SERCA2-deficient cells. (A) Quantitative RT-PCR analysis of *Atp2a3* mRNA in WT *abl* preB cells without (-) *gAtp2a2* and bulk *Atp2a2* inactivated with (+) *gAtp2a2* normalized to β -actin mRNA (mean \pm SD; $n = 3$). **, $P < 0.01$. **(B)** Western blot analysis of SERCA2, SERCA3, and β -actin in WT and bulk *Atp2a2* inactivated *abl* preB cells performed as in Fig. 1B ($n = 1$ for Dox 2 d; $n = 4$ for Dox 4 d or more). **(C)** Western blot analysis of SERCA3 and β -actin in WT and *Atp2a2*^{-/-} *abl* preB cells without (-) *gAtp2a3* and bulk *Atp2a3* inactivated with (+) *gAtp2a3* ($n = 3$). **(D)** Flow-cytometric analysis of pMGINV rearrangement in WT and *Atp2a2*^{-/-} *abl* preB cells (-) and these cells after bulk *Atp2a3* inactivation with (+) *gAtp2a3* (mean \pm SD; $n = 3$). ***, $P < 0.001$. **(E)** Mouse *Igk* locus schematic showing multiple *V κ* and *J κ* gene segments, relative positions of *SacI* (S) and *EcoRI* (E) restriction sites, and *J κ* probe. The 3-kb *J κ* probe-hybridizing *SacI*-*EcoRI* fragment is shown, as are several different *V κ* to *J κ* rearrangements that will give rise to *J κ* probe-hybridizing *SacI*-*EcoRI* fragments of different sizes. **(F)** Southern blot of *SacI* and *EcoRI* digested genomic DNA probed with the *J κ* probe from WT and *Atp2a2*^{-/-} *abl* preB cells (-) and the same cells bulk *Atp2a2* (*gAtp2a2*) or *Atp2a3* (*gAtp2a3*) inactivated and treated with imatinib for the indicated number of days. The fragment generated by the unrearranged (UR) *Igk* locus is indicated. The percentage of lane hybridization that corresponds to the UR *Igk* locus is shown with 0-d imatinib set at 100%. Quantification described in Materials and methods. **(G)** Mean \pm SD of the percentage of UR *Igk* locus quantified as described in F in imatinib-treated WT, *Atp2a2*^{-/-}, and bulk *Atp2a3* inactivated *Atp2a2*^{-/-} *abl* preB cells ($n = 3$). *, $P < 0.05$; **, $P < 0.01$; ***, $P < 0.001$. Dox, doxycycline; expo., exposure; n.s., not significant.

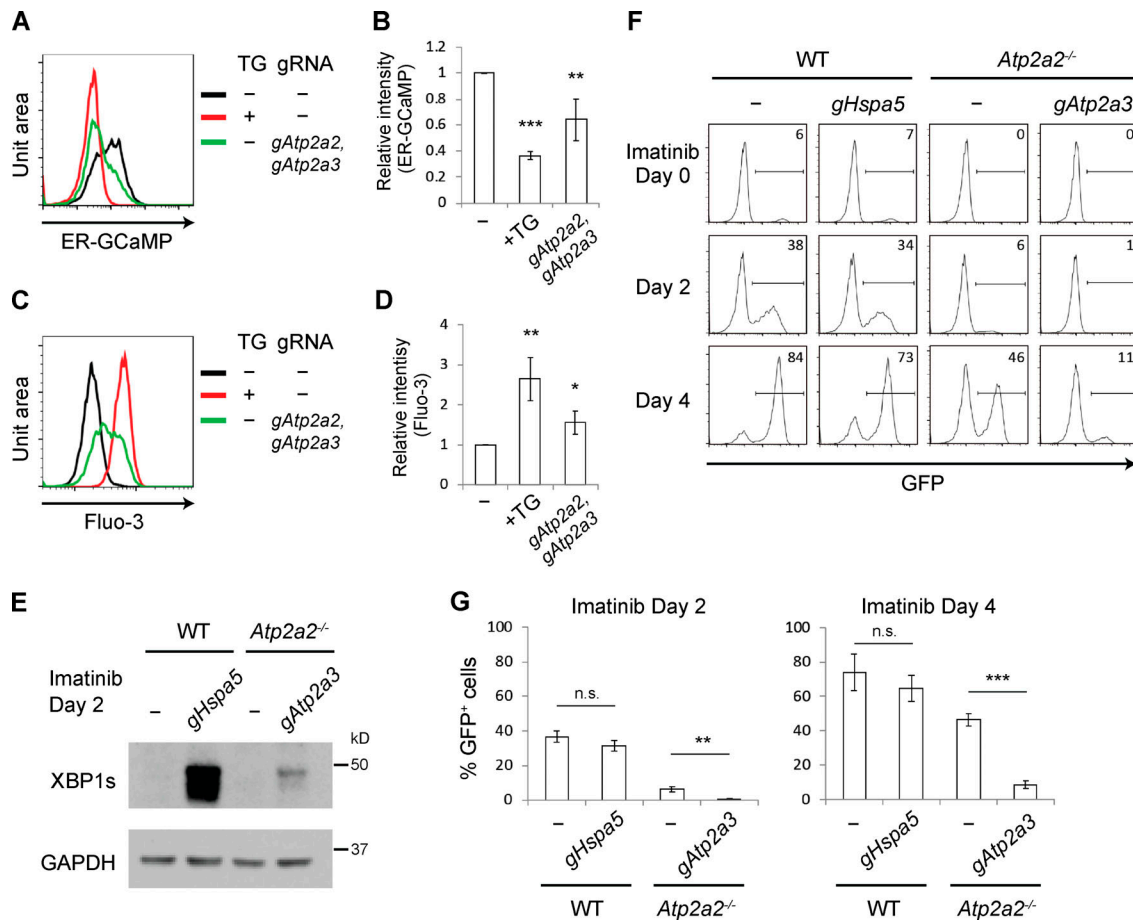


Figure 3. Loss of SERCA2 and SERCA3 disrupts Ca²⁺ homeostasis. (A–D) Flow-cytometric analysis of WT *abl* preB cells expressing ER-GCaMP6-150 (A) or incubated with Fluo-3 AM Ca²⁺ indicator (C) without further treatment (black histogram), treated with thapsigargin (TG; red histogram), or after bulk inactivation of *Atp2a2* and *Atp2a3* genes (green histogram). Median fluorescence intensity was calculated and normalized to that of the black histogram in each experiment to derive the relative intensity of ER-GCaMP6-150 (B; mean ± SD; *n* = 4) and Fluo-3 dye (D; mean ± SD; *n* = 3). *, *P* < 0.05; **, *P* < 0.01; ***, *P* < 0.001 compared with the untreated sample. **(E)** Western blot analysis of XBP1s in imatinib-treated WT *abl* preB cells bulk *Hspa5* (*gHspa5*) inactivated or *Atp2a2*^{-/-} cells bulk *Atp2a3* (*gAtp2a3*) inactivated (*n* = 3). **(F)** Flow-cytometric analysis of GFP expression from pMGINV rearrangement in cells from E treated with imatinib for 0, 2, or 4 d. **(G)** Mean ± SD of three pMGINV experiments performed as in F. **, *P* < 0.01; ***, *P* < 0.001. n.s., not significant.

inhibitor thapsigargin led to a significant reduction in ER-GCaMP6-150 fluorescence, indicative of loss of ER Ca²⁺ (Fig. 3, A and B) and concomitant increase in cytosolic Ca²⁺, as evidenced by Fluo-3 fluorescence (Fig. 3, C and D). Bulk *Atp2a2* and *Atp2a3* inactivation also led to reduced ER-GCaMP6-150 fluorescence (Fig. 3, A and B) and increased Fluo-3 fluorescence (Fig. 3, C and D). We conclude that in *abl* preB cells, loss of SERCA2 and SERCA3 leads to decreased ER Ca²⁺ and increased cytosolic Ca²⁺.

Reduced ER Ca²⁺ promotes IRE1α activation, which leads to expression of the spliced form of XBP1, XBP1s, and activation of the unfolded protein response (UPR; Hetz and Glimcher, 2009). As expected, loss of SERCA2 and SERCA3 led to activation of the UPR, as evidenced by XBP1s expression in *Atp2a2*^{-/-} *abl* preB cells after bulk *Atp2a3* inactivation (Fig. 3 E). BiP, encoded by *Hspa5*, normally binds IRE1α, preventing its activation until the accumulation of unfolded proteins prompts its dissociation from IRE1α and activation of the UPR (Hetz and Glimcher, 2009). Indeed, bulk *Hspa5* inactivation in *abl* preB cells led to high levels of XBP1s expression (Fig. 3 E). However, unlike loss of SERCA2 and SERCA3, loss of BiP had minimal effects on V(D)J

recombination (Fig. 3, F and G). We conclude that pathways other than the UPR are regulated by SERCA2 and SERCA3 to promote efficient V(D)J recombination.

SERCAs are required for RAG gene expression

V(D)J recombination relies on pathways that induce RAG expression and promote accessibility of antigen receptor genes to the RAG proteins (Sleckman et al., 1998). We assayed V(D)J recombination on episomal pMGINV plasmid substrates transiently introduced into *abl* preB cells, but not integrated into the chromosome (Fig. 4 A). pMGINV plasmids were transfected into WT and *Atp2a2*^{-/-} *abl* preB cells treated with imatinib and then recovered 2 d later, and CJ formation was assayed by PCR. These analyses revealed a decrease in pMGINV CJ formation in *Atp2a2*^{-/-} compared with WT *abl* preB cells, suggesting that SERCA proteins have functions regulating the expression or basic biochemical activities of the RAG proteins (Fig. 4 B). Indeed, *Atp2a2*^{-/-} *abl* preB clones had lower levels of *Rag1* and *Rag2* transcripts at 1 and 2 d after imatinib treatment compared with WT cells (Fig. 4, C and D). Strikingly, bulk *Atp2a3* inactivation in

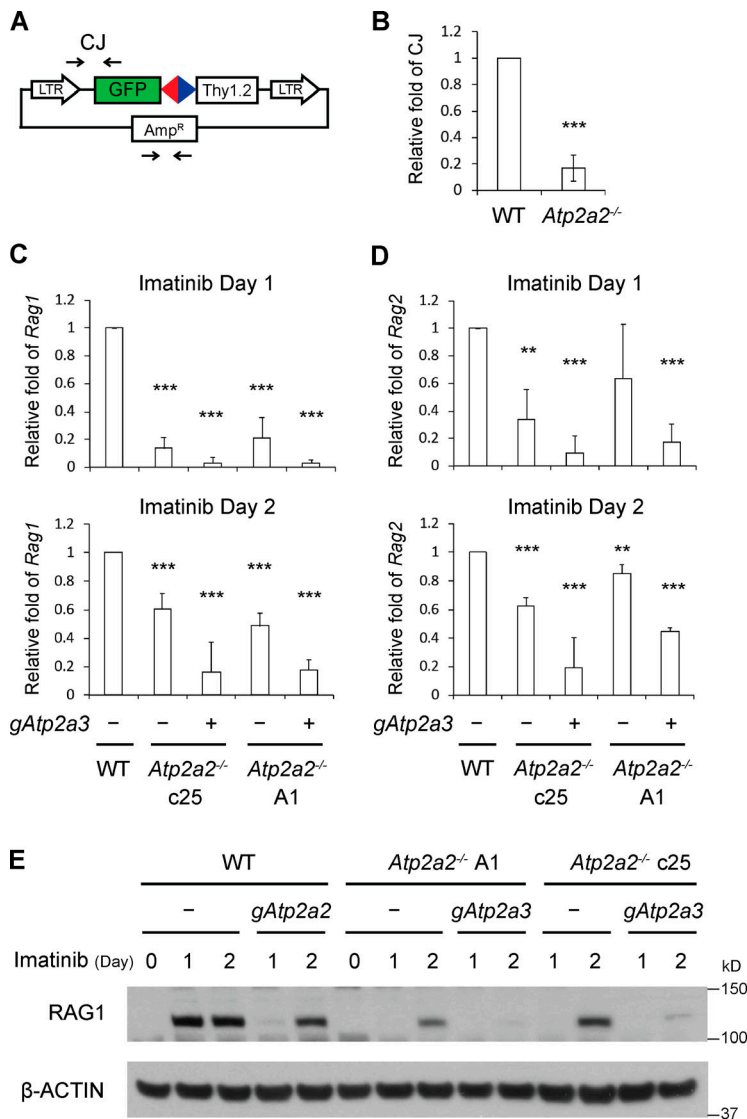


Figure 4. Loss of SERCA2 and SERCA3 leads to diminished Rag expression. (A) Schematic of pMGINV with locations of PCR primers (arrows) used to amplify CJ and the plasmid backbone (Amp^R). (B) Quantitative PCR analysis of pMGINV CJ formation in plasmids recovered from WT and *Atp2a2*^{-/-} abl preB cells treated with imatinib (mean ± SD; n = 3). ***, P < 0.001. (C and D) Levels of *Rag1* (C) and *Rag2* (D) mRNA assayed by quantitative RT-PCR in abl preB cells treated with imatinib for 1 or 2 d. WT and *Atp2a2*^{-/-} abl preB clones (-) were assayed, as were *Atp2a2*^{-/-} abl preB clones with bulk *Atp2a3* inactivation (+). *Rag1* and *Rag2* mRNA levels were normalized to β -actin mRNA and relative to the WT value, which was set at 1. *Rag1* and *Rag2* mRNAs were not detected at significant levels in cells not treated with imatinib (mean ± SD; n = 3). **, P < 0.01; ***, P < 0.001 compared with WT. (E) Western blot analysis for RAG1 and actin in WT and *Atp2a2*^{-/-} abl preB cell clones (-) and cells bulk *Atp2a2* (*gAtp2a2*) or *Atp2a3* (*gAtp2a3*) inactivated and treated with imatinib for 1 or 2 d (n = 1 for 1 d, n = 2 for 2 d).

Atp2a2^{-/-} abl preB cell clones led to a profound reduction in *Rag1* and *Rag2* transcripts after imatinib treatment (Fig. 4, C and D). In agreement, RAG1 protein levels were lower in bulk *Atp2a2* inactivated abl preB cells, *Atp2a2*^{-/-} abl preB cell clones, and after bulk *Atp2a3* inactivation in *Atp2a2*^{-/-} abl preB cell clones (Fig. 4 E). We conclude that SERCA2 and SERCA3 function redundantly to promote *Rag1* and *Rag2* expression.

SERCA2 deficiency leads to B lymphopenia in mice and humans

There are at least four splice variants of SERCA2, with SERCA2a and SERCA2b being the main variants (Fig. S2 A; Gélébart et al., 2003; Stammers et al., 2015; Vandecaetsbeek et al., 2011). Analyses of mRNA from proB, preB, and immature B cells purified from the bone marrow of WT mice revealed SERCA2a, SERCA2b, and SERCA3 mRNA in all of these cells (Fig. S2 B). As *Atp2a2*^{-/-} mice exhibit embryonic lethality, we generated mice carrying a conditional *Atp2a2* floxed allele (*Atp2a2*^{fllox}) and the *Cd79a/mb1-cre* allele (*mb1-cre*), which leads to Cre expression in the early proB cells (Hobeika et al., 2006; Ikeda et al., 2017; Periasamy et al., 1999). SERCA2 protein expression was

effectively abolished in B cells from *Atp2a2*^{fllox/fllox}:*mb1-cre* mice (Fig. 5 A). *Atp2a2*^{fllox/fllox}:*mb1-cre* mice exhibited an increase in the percentage of bone marrow proB cells compared with *Atp2a2*^{fllox/fllox} or *mb1-cre* mice (Fig. 5 B and Fig. S2 C). Moreover, *Atp2a2*^{fllox/fllox}:*mb1-cre* mice exhibited a decreased percentage of bone marrow preB cells (Fig. 5 C and Fig. S2 C) and a decrease in the percentage of mature splenic B cells compared with *Atp2a2*^{fllox/fllox} or *mb1-cre* mice (Fig. 5 D and Fig. S2 C). These findings are consistent with potential defects in antigen receptor gene assembly in developing B cells.

Darier disease is an autosomal-dominant disease caused by heterozygous ATP2A2 mutations that span the entire gene, consistent with the notion that the disease is caused by ATP2A2 haploinsufficiency (Korosec et al., 2006; Leong et al., 2017). We assayed the peripheral blood of 16 Darier disease patients. Remarkably, six patients exhibited a reduction in the percentage of B cells in peripheral blood compared with the normal reference ranges for healthy individuals (Fig. 5 E). The remaining 10 patients could have SERCA2 levels that can cause the disease but are adequate for normal B cell development. Darier disease

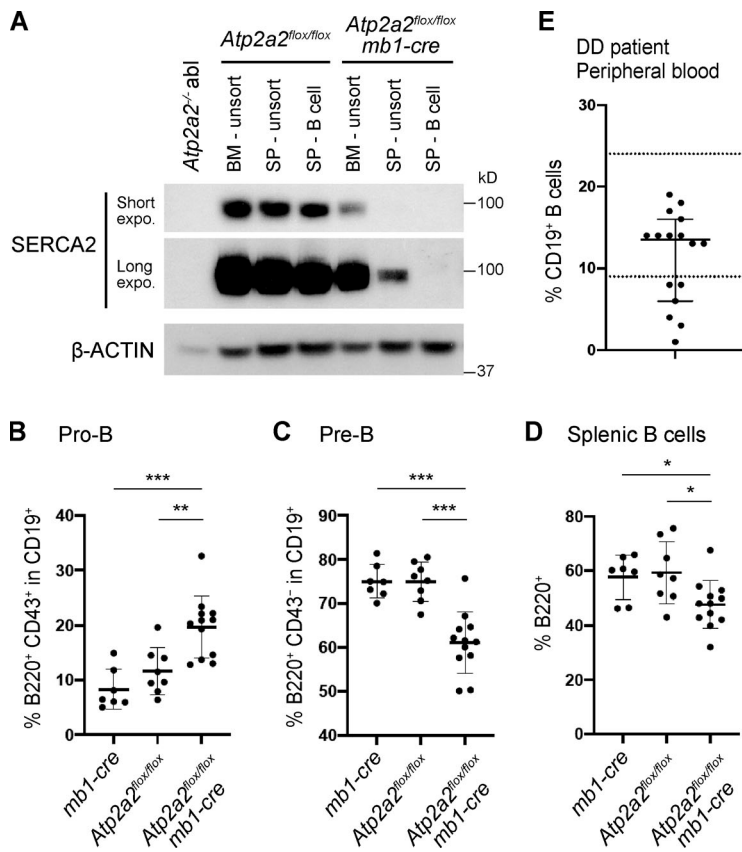


Figure 5. SERCA2 deficiency impairs B cell development in vivo. (A) Western blot analysis of SERCA2 and actin in cells isolated from *Atp2a2^{flox/flox}* and *Atp2a2^{flox/flox};mb1-cre* mice. Bone marrow (BM), whole spleen (SP - unsorted), and purified splenic B cells (SP - B cell) were analyzed (*n* = 1). (B–D) Mean ± SD of the percentages of proB (B), preB (C), and mature splenic B (D) cells assayed by flow-cytometric analysis of BM (B and C) and spleen (D) from *mb1-cre* (*n* = 7), *Atp2a2^{flox/flox}* (*n* = 8), and *Atp2a2^{flox/flox};mb1-cre* (*n* = 12) mice. *, *P* < 0.05; **, *P* < 0.01; ***, *P* < 0.001. (E) Percentage of CD19⁺ B cells in the peripheral blood of Darier disease (DD) individuals (median with 95% CI; mean, 11.4%; *n* = 16). The normal laboratory range (9–24%) for the percentage of B cells in peripheral blood is indicated by the dotted lines. expo., exposure.

patients are prone to skin infections, likely due to disrupted skin integrity, but could also be impacted by concomitant immune dysfunction (Tayabali et al., 2019; von Eiff et al., 2001; Wheeland et al., 1985).

We have shown that loss of SERCA functions in *abl* preB cells leads to decreased ER Ca²⁺ and increased cytosolic Ca²⁺, which are accompanied by diminished RAG expression and a block in V(D)J recombination. This is not due to activation of the UPR. Rather, it likely reflects the perturbation in signaling pathways required for RAG gene expression (Kuo and Schlissel, 2009). In this regard, the binding of Ca²⁺ to calmodulin leads to the activation of calcineurin, a phosphatase that can activate NFAT, which can inhibit RAG expression (Amin and Schlissel, 2008; Clapham, 2007; Patra et al., 2006). Thus, we speculate that the increased cytosolic Ca²⁺ in SERCA-deficient B cells could negatively regulate RAG expression through the activation of NFAT or other transcriptional pathways. Changes in intracellular Ca²⁺ levels could also modulate RAG activity as Ca²⁺ inhibits RAG enzymatic activity *in vitro* (Hiom and Gellert, 1997). Increased Ca²⁺ can also promote chromatin condensation, which would reduce RAG accessibility to antigen receptor genes (Phengchat et al., 2016; Schatz and Ji, 2011).

Materials and methods

Mice

All mice were maintained on B6 background, and all animal protocols were approved by the Weill Cornell Institutional Animal Care and Use Committee or the University of Alabama at

Birmingham Animal Resources Program. *Atp2a2^{flox}* (*tmc* allele) mice were obtained from S. Kajimura (Ikeda et al., 2017). Genotyping was performed by PCR using Platinum Taq Polymerase (Invitrogen; 11304011) with the following primers: *Atp2a2* Fwd: 5'-CACCTTGTTTAGCCTAGCTTTTAC-3'; and *Atp2a2* Rev: 5'-GTTGCACACTCTTTCTGTCTG-3'. The PCR program is as follows: 94°C/2 min → [94°C/15 s → 58°C/30 s → 68°C/40 s] × 34 cycles → 68°C/3 min. The product size is 589 bp for WT and 700 bp for *flox* allele. To detect the knockout allele after crossing with *Cre* mice, a third primer (*Atp2a2* 3'LOXP.R1: 5'-ACTGATGGCGAGCTCAGACC-3') was added together with the Fwd/Rev primer set. The product size for *Cre*-deleted allele is 200 bp.

Mb1-cre allele genotyping was performed using the above PCR conditions with the following primers: *Mb1* Fwd: 5'-CTGCGGTAGAAAGGGGGTC-3'; *Mb1* Rev: 5'-CCTTTCGAGGTCAGGGAGCC-3'; *Mb1-Cre* Fwd: 5'-ACCTCTGATGAAGTCAGGAAGAAC-3'; and *Mb1-Cre* Rev: 5'-GGAGATGTCCTTCACTCTGTTCT-3' (Yen et al., 2019). The expected product size is 400 bp for WT and 500 bp for the *Cre* allele.

Cell culture

abl preB cells were cultured in DMEM-HG (Gibco; 11960044) supplemented with 10% FBS (Gemini; 100106), nonessential amino acids, 2 mM L-glutamine, 1 mM sodium pyruvate, 100 U/ml penicillin-streptomycin, and 60 μM β-mercaptoethanol. For viral infection with gRNA expression virus, 1 × 10⁶ *abl* preB cells were mixed with viral supernatant and Polybrene (Santa Cruz; sc134220; 10 μg/ml final concentration) in 6-well plates and centrifuged at 1,800 rpm for 1.5 h at room temperature.

Cells were purified by flow-cytometric cell sorting on day 3 after infection based on the expression of blue fluorescent protein (BFP) or human CD2 from the pKLV-U6gRNA lentiviral vector (Addgene; 50946) using a BD FACSAria Fusion Sorter. gRNAs used were as follows (protospacer adjacent motif, or PAM, sequence underlined): *gAtp2a2*: ATGGGCAAAGTGTATCGACAGG; *gAtp2a3*: TGATCCGGTCTGACCGCAAGGG; *gHspa5*: CTCGGCGT GAGGTAGAAAAGG; and *gRosa26* control: CTCCAGTCTTTCTAG AAGATGG. Abl preB cell lines containing pMGINV were generated as previously described (Hung et al., 2017). Cells were treated with 3 μ M imatinib (Sigma-Aldrich; CDS022173) at the density of 1×10^6 cells/ml to induce G1 arrest. GFP signals arising from rearranged pMGINV were assayed on a BD LSRII flow cytometer, and the data were analyzed using FlowJo.

CRISPR/Cas9 genetic screening

The WT abl preB cell line carrying a Tet-On Cas9 transgene (clone 302) was infected with a lentiviral gRNA library containing 90,230 gRNAs targeting 18,424 mouse genes (Addgene; 67988; Tzelepis et al., 2016). 140×10^6 cells were infected to achieve >500-fold coverage using the infection described above at low multiplicity of infection. 3 d after infection, cells expressing BFP indicative of positive infection were isolated by flow-cytometric cell sorting and treated with 3 μ g/ml doxycycline for 1 wk to induce Cas9 expression. Cells were then treated with 3 μ M imatinib for 4 d as described above to induce V(D)J recombination of pMGINV. Cells that had (GFP⁺) and had not (GFP⁻) undergone pMGINV recombination were purified by flow-cytometric cell sorting, and genomic DNA was isolated from these cells. The gRNAs from each population were amplified using nested PCR and sequenced on an Illumina HiSeq 2500 platform performed in the Weill Cornell Epigenomics Core. The sequencing primer for the pKLV2 gRNA plasmid used in the library was 5'-GGCTTTATATATCTTGTGGAAAGGACGAAACACCG-3'.

The gRNA sequence regions were retrieved from FASTQ files using Seqkit (Shen et al., 2016). The derived sequences were mapped to the original gRNA sequences in the library (Tzelepis et al., 2016). The number of reads of each gRNA was normalized as follows:

$$\text{Normalized reads per gRNA} = \frac{\text{Reads per gRNA}}{\text{Total reads for all gRNAs in a sample}} \times 10^6 + 1$$

(Shalem et al., 2014). The enrichment score of a gRNA is calculated as a ratio of normalized reads of the gRNA between two samples (GFP⁻ versus GFP⁺).

Southern blotting

Southern blot analyses of pMGINV were performed as previously described (Hung et al., 2018). To quantify the percentage of SJs and SEs in pMGINV, the background of the scanned blot was subtracted in ImageJ, and the measured intensity of SJ and SE bands was divided by the sum of SJ plus unrearranged (UR)/SE bands and SE plus UR/SJ bands, respectively, in the same lane. To quantify *Igk* rearrangement, the intensity of the

germline UR *Igk* band was measured after background subtraction and then divided by the intensity of whole lane (1–5 kb) that represents different *Igk* rearrangement products hybridizing to Jk probe. The derived values were normalized to that of day 0 of imatinib treatment in each genotype to indicate the remaining levels of UR *Igk* locus.

Western blotting

$3\text{--}5 \times 10^6$ abl preB cells were lysed in 50 μ l buffer (10 mM Tris, pH 8, 1 mM EDTA, 10% glycerol, 0.5% NP-40, 400 mM NaCl, 1 mM dithiothreitol, and $1\times$ protease/phosphatase inhibitor cocktails) on ice for 30 min and centrifuged at 14,000 rpm for 15 min to collect supernatants. 20 μ g of the lysates were boiled in $1\times$ lithium dodecyl sulfate sample buffer (Invitrogen) for 5 min and separated on NuPAGE 4–12% Bis-Tris mini gels (Invitrogen). Proteins were transferred to a nitrocellulose membrane in transfer buffer (25 mM Tris, 192 mM glycine, and 20% methanol) at 100 V for 90 min at room temperature using a wet transfer apparatus (Bio-Rad). The membrane was blocked in 5% milk/Tris buffered saline-Tween20 (TBST) at room temperature for 1 h. Primary antibody was diluted in 2.5% milk/TBST and probed overnight at 4°C. The membrane was then washed in TBST and probed with HRP-conjugated secondary antibodies (Promega) at room temperature for 1 h followed by TBST wash and visualization with enhanced chemiluminescence reagents (Thermo Fisher Scientific; 32109). Primary antibodies used were anti-SERCA2, rabbit polyclonal, 1:2,000 (Cell Signaling Technology; 4388); anti-SERCA3, rabbit polyclonal, 1:5,000 (GeneTex; GTX102381); anti-RAG1, rabbit monoclonal, 1:1,000 (Abcam; ab172637); anti-XBPs, rabbit monoclonal, 1:1,000 (Cell Signaling Technology; 12782); anti- β -actin, mouse monoclonal, 1:20,000 (Sigma-Aldrich; A2228); and anti-GAPDH, mouse monoclonal, 1:5,000 (Sigma-Aldrich; G8795).

RNA analysis

RNA was extracted from 5×10^6 abl preB cells using the RNeasy Mini Kit (Qiagen; 74104). 1 μ g RNA was used for cDNA synthesis using SuperScript III Reverse transcription (Invitrogen; 18080044) with oligo-dT primer. The quantitative PCR reaction was set up with 0.5 μ l cDNA, 2 μ l 10 mM forward/reverse primer mix, and 5 μ l LightCycler 480 SYBR Green I Master Mix (Roche; 04707516001) to a total volume of 10 μ l and analyzed on a Roche LightCycler 480. The Ct value of a target transcript was normalized to that of the β -actin transcript of the same sample to calculate Δ Ct, which was then normalized to the Δ Ct of control sample to analyze fold changes ($2^{-\Delta\Delta\text{Ct}}$). The forward/reverse primers were designed to bind different exons to minimize cross-amplification from genomic DNA. Primer sequences for mouse mRNAs were as follows: *Rag1* Fwd: 5'-AGGCTGTGGAGCAAGGTA-3'; *Rag1* Rev: 5'-AGGATCTCACCTAAACAGC-3'; *Rag2* Fwd: 5'-AGGATTCAGAGAGGGATAAGC-3'; *Rag2* Rev: 5'-CCATCTGCAGGGACATTTTTG-3'; β -actin Fwd2: 5'-CATGGCATTGTTACCAACTGG-3'; β -actin Rev2: 5'-ATGGCTACGTACATGCTGG-3'; *Atp2a3* Fwd: 5'-TTGGAGTGTATGTAGGCTG-3'; and *Atp2a3* Rev: 5'-TCAGAGAGCTGTTGAGGG-3'. The following primers were used to amplify *Atp2a2* transcripts: all forms Fwd: 5'-TCGACAGGACAGAAAGAGTGTG-3'; all forms Rev: 5'-GTA

TGCTTGATGACGGAGACAG-3'; isoform-a Fwd: 5'-GCTCATTTC
CCAGATCACACCG-3'; isoform-a Rev: 5'-GTTACTCCAGTATTG
CGGGTTG-3'; isoform-b Fwd: 5'-ACCTTTGCCGCTCATTTTCCA
G-3'; and isoform-b Rev: 5'-AGGCTGCACACTCTTTACC-3'.

Calcium indicator assay

The ER-GCaMP6-150 plasmid, provided by T.A. Ryan (de Juan-Sanz et al., 2017), is a variant of GCaMP6 Ca²⁺ indicator containing the signal peptide of calreticulin and KDEL ER retention motif suitable for detecting Ca²⁺ in the ER. WT abl preB cells carrying Tet-On Cas9 transgene were infected with lentivirus carrying ER-GCaMP6-150. Cells emitting green fluorescence were isolated by flow-cytometric cell sorting and single-cloned. The responsiveness of ER-GCaMP6-150 to Ca²⁺ fluctuation in the ER was confirmed by flow-cytometric analysis following treatments with 50 nM SERCA inhibitor thapsigargin for 30 min. The levels of ER-GCaMP6-150 fluorescence in the cycling abl preB cells expressing gRNA were analyzed on day 3 or 4 of doxycycline treatment. For cytosolic Ca²⁺ analysis, abl preB cells (without ER-GCaMP6-150) were treated with 2 μM Fluo 3-AM (Sigma-Aldrich; 39294) at 37°C for 30 min in growth medium. The stained cells were washed once in prewarmed growth medium and then aliquoted for thapsigargin treatment before flow cytometry or analyzed directly.

Episomal V(D)J recombination assay

Abl preB cells were pretreated with 3 μM imatinib overnight to induce G1 arrest. 15 × 10⁶ cells were transfected with 8 μg pMGINV plasmid using Amaxa Nucleofector (Program X-001; Lonza). The cells were incubated in media containing imatinib for 2 d, and the plasmid was isolated from these cells. pMGINV CJ formation was quantified by quantitative PCR using the primers 23RS-Fwd2: 5'-TCTCCCCTTGAACCTCCTC-3' and CJ-GFP-Strat: 5'-GAACAGCTCCTCGCCCTTG-3'. The CJ products were normalized to the levels of Amp^R gene on the vector backbone to control for transfection efficiency (primers: AmpR Fwd1: 5'-ATAAACCAGCCAGCGGAAG-3'; and AmpR Rev1: 5'-AACCGGAGCTGAATGAAGC-3').

Lymphocyte development

Bone marrow and spleen cells from 4- to 5-wk-old mice were incubated with mouse Fc Block (BD; 553141; 1:100 dilution) on ice for 15 min. Cells were incubated with fluorophore-conjugated primary antibody (1:500 dilution) at 4°C for 30 min, then analyzed on a BD LSRII flow cytometer. For bone marrow, viable CD19⁺ cells were analyzed for B220 and CD43 expression. For spleen samples, viable cells were analyzed for B220 and Thy1.2 expression. Antibodies used were PacificBlue-anti-h/mB220 (BioLegend; 103227), APC-anti-mCD43 (BioLegend; 143208), FITC-anti-mCD19 (BioLegend; 115506), APC-anti-mCD90.2/Thy1.2 (BioLegend; 105312), and PE/Cy7-anti-mIgM (BD; 552867).

Human peripheral blood analysis

Ethical permit for blood sampling of Darier disease patients was granted by the regional ethical review board in Stockholm, Sweden (dnr 2017/1098-32). Prior to sampling, patients gave

informed written consent. Patients were treated according to the principles of the Helsinki declaration. CD19⁺ B cells were analyzed in peripheral blood by flow cytometry (<https://www.karolinska.se/KUL/Alla-anvisningar/Anvisning/10106>). Reference data are based on healthy adult individuals at the Karolinska University Hospital clinical laboratory (Stockholm, Sweden) and comprise values between the 5th and 95th percentiles.

Statistical analysis

Statistical comparisons between two samples were analyzed by unpaired Student's *t* test. P values are indicated in the figure legends.

Online supplemental material

Fig. S1 shows supporting information for Fig. 1: loss of SERCA2 impairs RAG cleavage. Fig. S2 shows supporting information for Fig. 5: SERCA2 deficiency impairs B cell development in vivo. Table S1 shows CRISPR/Cas9 screening results of candidate genes regulating V(D)J recombination.

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Supplemental material

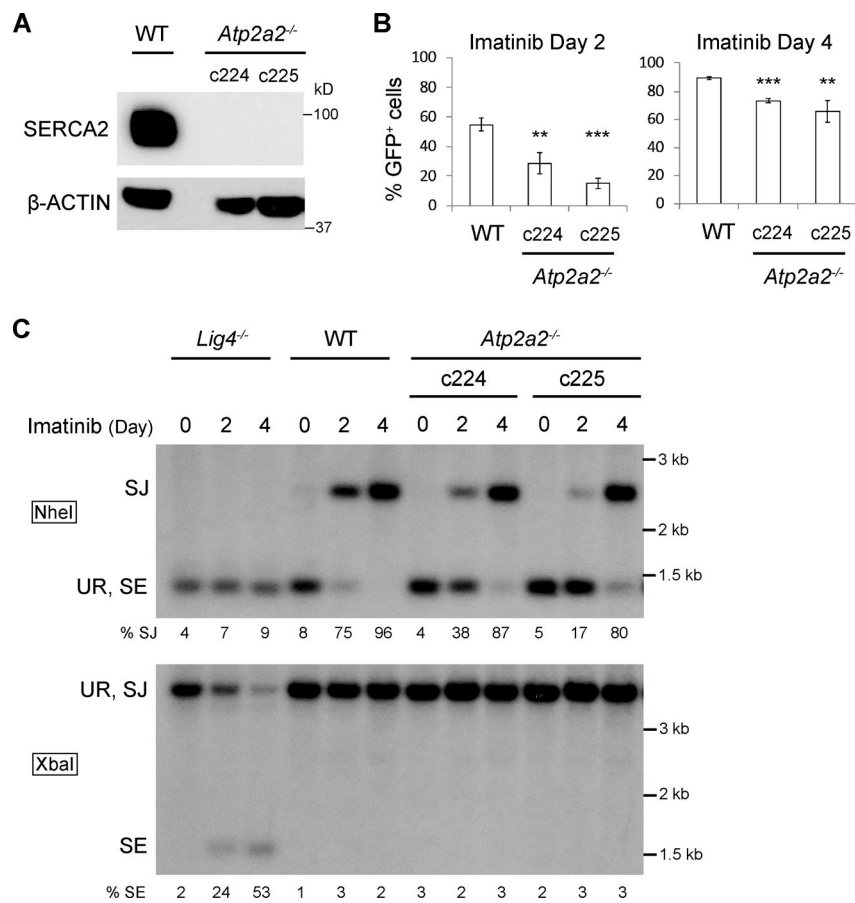


Figure S1. **SERCA2 deficiency impairs V(D)J recombination. (A)** Western blot analysis performed as described in Fig. 1 B in *Atp2a2*^{-/-} abl preB clones (c224 and c225) derived from a distinct WT abl preB cell line (BA600; *n* = 1). **(B)** Analysis of pMGINV rearrangement performed as described in Fig. 1 D (mean ± SD; *n* = 3). **, *P* < 0.01; ***, *P* < 0.001 compared with WT. **(C)** Southern blot analyses of pMGINV in abl preB cells described in A performed as described in Fig. 1 G (*n* = 1).

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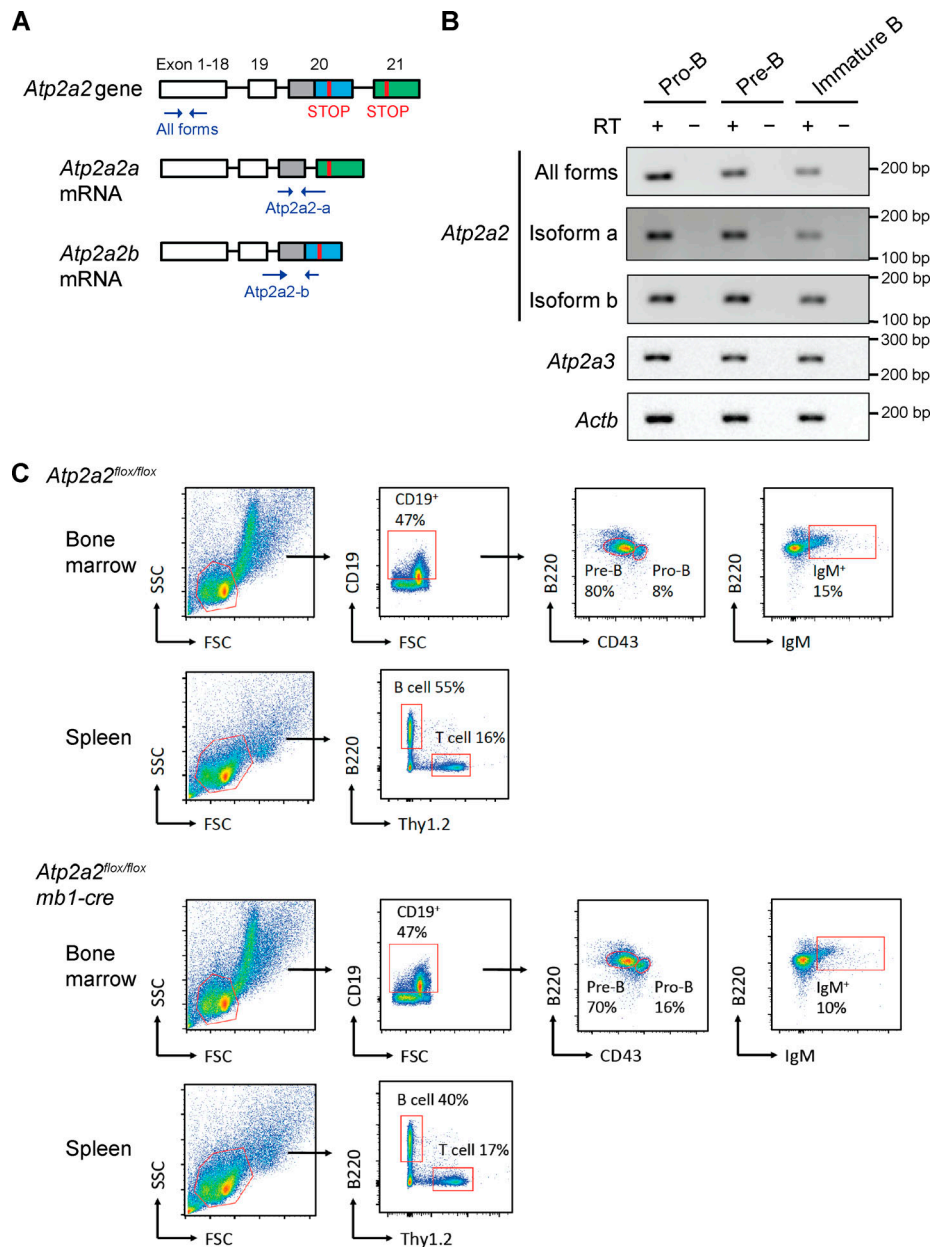


Figure S2. **Analysis of SERCA2-deficient primary B cells.** (A) PCR strategy used to distinguish the *Atp2a2a* and *Atp2a2b* mRNA isoforms. (B) RT-PCR analysis of mRNA representing all *Atp2a2* isoforms and *Atp2a2b* or *Atp2a2a* isoforms in addition to *Atp2a3* transcripts in proB (B220⁺CD43⁺), preB (B220⁺CD43⁻), and immature B (B220⁺IgM⁺) cells purified from bone marrow of WT B6 mice ($n = 2$). (C) Gating strategies for the flow-cytometric analysis of B cell development in mouse bone marrow and spleen shown in Fig. 5, B–D. The analyses shown are of 4-wk-old littermates of *Atp2a2^{flox/flox}* and *Atp2a2^{flox/flox};mb1-cre* mice. SSC, side scatter; FSC, forward scatter.

Table S1 is provided online as a separate Excel file and shows CRISPR/Cas9 screening results of candidate genes regulating V(D)J recombination.