The Serine/Threonine Phosphatase PP4 Is Required for Pro-B Cell Development through Its Promotion of Immunoglobulin VDJ Recombination

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Abstract

PP4 phosphatase regulates a number of crucial processes but the role of PP4 in B cells has never been reported. We generated B cell-specific pp4 knockout mice and have identified an essential role for PP4 in B cell development. Deficiency of PP4 in B lineage cells leads to a strong reduction in pre-B cell numbers, an absence in immature B cells, and a complete loss of mature B cells. In PP4-deficient pro-B cells, immunoglobulin (Ig) DJ joining is impaired and Ig µ heavy chain expression is greatly decreased. In addition, PP4-deficient pro-B cells show an increase of DNA double-strand breaks at Ig loci. Consistent with their reduced numbers, residual PP4-deficient pre-B cells accumulate in the G1 phase, exhibit excessive DNA damage, and undergo increased apoptosis. Overexpression of transgenic Ig in PP4-deficient mice rescues the defect in B cell development such that the animals have normal numbers of IgM+ B cells. Our study therefore reveals a novel function for PP4 in pro-B cell development through its promotion of VµDJµ recombination.

Introduction

B cell development initiates in the bone marrow (BM) of adult mice and is a tightly controlled process. Developing B cells can be divided chronologically into six Hardy fractions (Frs.) A to F according to the recombination status of the immunoglobulin (Ig) heavy chain (HC) locus, the Ig light chain (LC) locus, and the expression pattern of particular cell surface markers [1], [2]. The process starts with D-JH recombination in Fr. A cells, followed by Vµ-DJµ and DJµ-Vµ recombination in Fr. B and Fr. C cells [2], [3]. When a differentiating B cell reaches the Fr. D stage, Vµ-Jµ recombination commences [2]. Successful Ig Vµ-DJµ/Vµ-Jµ recombination leads to the expression of a surface IgM-containing BCR complex that enables a B cell to continue to the Fr. E and Fr. F stages [4]. At the molecular level, DJµ/Vµ-DJµ recombination is initiated when two Ig gene segments flanked by recombination signal sequences (RSSs) are paired and cleaved by RAG [5], [6]. The two gene segments are brought together by the cell’s non-homologous end joining (NHEJ) machinery via the sequential recruitment of NHEJ factors. A deficiency of any of these factors results in a failure in DJµ/Vµ-DJµ recombination, an early block in B cell development, and ultimately a shortage of mature B lymphocytes [7].

Protein phosphatase 4 (PP4) belongs to the type 2A protein serine/threonine phosphatase (PP2A) family. In mammals, the catalytic subunit of PP4 (PP4c) selectively binds to one or two of several different regulatory subunits, including R1 [8], [9], R2 [10], R3 [11], R4 [12], Smck [17], to form a PP4 holoenzyme. The composition of the PP4 holoenzyme presumably determines its catalytic activity and also confers its substrate and tissue specificity [18]. At the cellular level, PP4 activity is required for microtubule organization and centrosome maturation via mechanisms that are highly conserved among mammalian species [19], [20], [21]. PP4 is also necessary for DNA repair via the homologous recombination pathway through dephosphorylation of the RPA2 subunit of replication protein A [22], and through dephosphorylation of γH2AX during cell division [18], [23], [24]. Lastly, PP4 has been implicated in multiple signal transduction pathways, including pre-TCR/TGR signaling [25], TNF-α signaling [26], [27], Toll-like receptor 4 signaling [28], and NF-kB signaling [29], [30].

Cell-specific deletion of PP4 in mice leads to a partial block in thymocyte development at the double negative (DN) stage. The Ca2+ mobilization and PLC-γ1 phosphorylation normally induced by anti-CD3 stimulation are impaired in these PP4-deficient cells [25]. Whether PP4 plays an analogous role in B cell development is unknown. In this study, we utilized mb-1/cre mice to delete the pp4c gene specifically in B cells and identified a pivotal role for PP4 in pro-B cell development. Deletion of PP4 severely disrupted pro-B cell differentiation and consequently led to a complete absence of mature B cells. In PP4-deficient pro-B cells, DJµ recombination was greatly reduced and Ig µ HC expression was decreased. We also found that PP4-deficient pre-B cells in vivo accumulated in the G1 phase, showed an elevated level of DNA damage, and
underwent increased apoptosis. Significantly, PP4-deficient pro-B cells transgenically expressing IgM successfully differentiated into normal numbers of IgM+ B cells. Our results therefore reveal the indispensable role of PP4 in promoting the VH-DJH recombination required for continued pro-B cell differentiation and the production of mature B cells.

**Materials and Methods**

**Mice**

PP4Cf/f mice [25], mb-1/cre mice [31], and IgHΔE transgenic mice [32] generated as previously described were maintained in strict accordance with the recommendations of the Guide for the Care and Use of Laboratory Animals of the National Health Research Institutes (NHRI). The IgHΔE transgenic and mb-1/cre mice used in all experiments were heterozygous. All protocols were approved by NHRI’s Institutional Animal Care and Use Committee (Permit Number: 099111-A), and all efforts were made to minimize suffering.

**BrdU Incorporation in vivo and in vitro**

For in vivo BrdU incorporation, mice were injected i.p. with 2 mg BrdU (Sigma) in 200 μl sterile PBS at 16 h and 24 h before sacrifice and cell sorting. For in vitro BrdU incorporation, isolated cells were cultured for 16 h in RPMI 1640 medium containing 10% FCS (Hyclone) and 10 μM BrdU prior to analysis using a BrdU-Flow kit (BD PharmingenTM).

**Cell Sorting and Cell Culture**

Single-cell suspensions were prepared from the BM of mice by gentle pipetting and sieving through 70-μm nylon mesh filters (Falcon; BD). BM cells were cultured in RPMI 1640 medium containing 10% FCS and 2 mM L-glutamine (Invitrogen), 100 U penicillin-streptomycin (Invitrogen), 100 mM Hepes (Sigma-Aldrich), and 0.055 mM 2-mercaptoethanol (Gibco). For flow cytometric analyses, cells were suspended in FACS buffer (2% BSA in PBS containing 0.1% NaN₃).

**Comet Assays**

To perform comet assays to detect DNA damage, Fr. B and Fr. C cells were sorted by FACSAria (BD) and processed using the OxiSelect™ Comet Assay Kit following the manufacturer’s instructions (Cell Biolabs). Data were analyzed using Comet Assay IV™ v4.3 software.

**Confocal Imaging**

Fr. B and Fr. C cells were sorted, fixed in 4% paraformaldehyde, and permeabilized in 0.2% saponin. After blocking with Image-iT™ FX signal enhancer (Invitrogen), cells were stained with anti-mouse IgM-FTTC (HC-specific antibody (Ab) (BD), seceded on Superfrost® Plus microscope slides (Thermo Scientific), and mounted using Prolong® Gold anti-fade reagent plus DAPI (Invitrogen).

**Ig DJH Recombination Assay and Coding End Assay**

For DJH recombination analysis, genomic DNA from 5 × 10⁵ cells was extracted and subjected to nested PCR using a primer 5’ of the DFL16.1 segment and a primer 3’ of the JH4 segment in the Ig HC locus (Fig. S1). 1 ng of genomic DNA was utilized for first-round PCR using the following primers: dil6i16.15'-CCAGGGCTTTCATGGAAGGATCTAG-TACTGT-3' and jh4.5'-TTTCTCAATTGAGGCCTC-CAAAAGCTC-3'. PCR conditions were as follows: 96°C for 3 min; 2 cycles of 94°C for 40 sec, 65°C for 45 sec, and 72°C for 1 min 45 sec; 3 cycles of 94°C, 63°C and 72°C for these same time intervals; and 10 cycles of 94°C, 60°C and 72°C for these same time intervals. For second-round PCR, primers in-dll16.15'-AAGGGATCTACTAGTGTTTATTAGACGTAAG- TAGCTAC-3' and in-jh4.5'-AGGCTCTGAGATCCCTGAGCACG'-3' were used with PCR conditions as follows: 94°C for 5 min; 3 cycles of 94°C, 67°C and 72°C for the same time intervals as used for first-round PCR, and 24 cycles of 94°C, 64°C and 72°C for these same time intervals. PCR products were separated by 1% agarose gel electrophoresis and capillary-transferred to nylon membranes. Blots were hybridized with 5 ng/ml DIG-labeled PCR probe synthesized with the following primers: jhp: 5’-ATTACATATGCTATGACTACTGGOGTG-CAAAGG-3' and in-jh4.5'-CATATAACTACGGCCTGTCTCATACC-3', and detected using the DIG system following the manufacturer’s instructions (Roche). To detect DNA double-strand breaks at Ig loci, oligonucleotides BW-1.3'-GGCGTGACCAGGAGATCTGAAATTC-3' and BW-2.5'-GAATTCAAGTAC-3' were annealed for the preparation of BW linker as described previously [33]. 1 μg of genomic DNA from sorted Fr. B cells was treated with 0.1 U/50 μl of Klenow at 25°C for 30 min. After Klenow-inactivation, genomic DNA was washed and ligated with 1000 pM/50 μl of BW linker overnight. 500 ng of ligated DNA was utilized for first-round PCR using the primer 16.1b: 5’-GCCCTTCACAAAGAAGAAGAAG-3' at the upstream of DFL16.1 segment and the primer BW-1 (Fig. S2). PCR conditions were as follows: 96°C for 3 min; 3 cycles of 94°C for 1 min, 58°C for 2 min, and 72°C for 1 min 45 sec; 12 cycles of 94°C for 1 min, 62°C for 2 min, and 72°C for 1 min 45 sec. For second-round PCR, the primer 16.1a: 5’-GAAGTCCGCCCAAGAACACGAC-3' at the downstream of primer 16.1b and BM-1 were utilized. PCR conditions were as follows: 94°C for 3 min; 27 cycles of 94°C for 1 min, 62°C for 2 min and 72°C for 1 min. PCR products were separated by 1% agarose gel electrophoresis and capillary-transferred to nylon membranes. Blots were hybridized with 700 ng/ml DIG-labeled PCR probe of 220 bp synthesized with primers 16.1a and COD-R: 5’-CCAGGCCAGCAGGTTAGTTT-3’, and detected as described above.

**Flow Cytometry**

Single-cell suspensions of 1 × 10⁶ cells were washed twice with FACS buffer and maintained in the dark at 4°C throughout the experiment. A CantoII flow cytometer (BD) and FACSDivia software (BD) were used for the acquisition of flow cytometric data, and FlowJo software (Tree Star, Inc.) was used for analysis. For apoptosis assays, cells were stained with B220-APC and Annexin V-FITC using the Apoptosis Detection Kit I (BD). To stain and sort B cells, BM cells were incubated on ice for 15 min with anti-mouse Abs as follows: PE-1FITC/PE, IgG-Pacific Blue™ (both from Biolegend); IgM-FITC/eFluor 450 (eBioscience), B220-PE/PerCP/APC-Cy7, CD21-PE, CD5-PE, CD43-APC, CD25-PE-Cy7, CD24-PE-Cy7, and IgM-APC/PE-Cy7 (all from BD). Immunostained cells were washed twice in ice-cold FACS buffer prior to sorting by FACSAria cell sorter (BD).

For intracellular staining, cells were fixed and permeabilized using the BrdU-Flow kit (BD Pharmingen™) prior to incubation with μ- vs. HC-FITC or BrdU-Alexa Fluor 647 (all from BD).

**Statistical Analysis**

Data were analyzed by a one-tailed distribution, type 3 Student’s t-test. Differences between treatment groups with p-values ≤0.05 were considered statistically significant.
Figure 1. PP4 deficiency induces a severe block in B cell development. (A) Flow cytometric profiles of: total BM lymphocytes from WT and cKO mice analyzed for B220 vs CD43 expression (left panel); B220+CD43+ BM lymphocytes analyzed for CD24 vs BP-1 (middle panel); and B220+CD43+ BM lymphocytes analyzed for IgM vs IgD (right panel). (B) Quantitative cell count analysis of Frs. A to F cells from WT and cKO mice. (C) Quantitative cell count analysis of splenic B220+ B cells from WT and cKO mice. (D) Flow cytometric profiles of BM, spleen and MLN lymphocytes from WT and cKO mice (n = 4/group) analyzed for B220 vs IgM; B220+ splenocytes analyzed for CD21 vs CD23; and peritoneal lymphocytes analyzed for IgM vs CD5, as indicated. For A-D, results are representative of three independent experiments. For B and C, data are the mean ± SD (n = 4) (*p<0.05; **p<0.005). doi:10.1371/journal.pone.0068804.g001
Results

PP4 Deficiency Leads to a Severe Block in B Cell Development

To study the role of PP4 in B cell development, mb-1/cre mice [31] were bred with pppc-loxp-flanked (floxed) mice [25] to generate mb-1/cre;PP4FP/FP (designated here as cKO) mice. As a control, mb-1/cre mice were bred with pppc WT mice to obtain mb-1/cre;PP4FP/+ (designated here as WT) mice. B cell subsets, including Fr. A (B220CD43+CD24−BP−1); pre-pro-B cells, Fr. B (B220CD43+CD24−BP−1); pre-B cells), Fr. C (B220CD43+CD24−BP−1); pro-B plus large pre-B cells), Fr. D (B220CD43+IgM−IgD−BP−1; small pre-B cells), Fr. E (B220CD43+IgM−IgD−; immature B cells), and Fr. F (B220+CD43+IgM+IgD+/−; mature B cells), from the BM of WT and cKO mice were analyzed by FACS. In cKO mice, combined staining with anti-B220 and anti-CD43 Abs revealed that the B220+CD43+ BM lymphocyte population containing Frs. D to F was greatly reduced (Fig. 1A, left panel). Gating of B220+CD43+ BM lymphocytes on CD24 plus BP-1 expression showed that the percentage of CD24+BP-1+ cells (Fr. C) was strongly reduced in the mutant compared to the control (WT, 31.1% vs cKO, 4.7%) (Fig. 1A, middle panel). Among the residual B220+CD43+ BM lymphocytes in cKO mice, only IgM+IgD− cells (Fr. D) were present (Fig. 1A, right panel). Cell count analyses revealed severe reductions in the Fr. C and Fr. D populations in cKO mice, and the complete absence of the Fr. E and Fr. F subsets (Fig. 1B, Table 1). In the spleen and mesenteric lymph nodes (MLN) of cKO mice, mature B cells were barely detectable (Fig. 1C, 1D, Table 2). In addition, CD5+IgM+ B1a B cells were missing from the peritoneum (Fig. 1D). Thus, B cell development in cKO mice is severely blocked before the Fr. C stage, leading to the complete loss of immature and mature B cells.

To determine the deletion efficiency mediated by mb-1/cre, genomic DNA samples from Frs. A-C cells sorted from WT and cKO mice were subjected to PCR analysis. A scheme illustrating the primers used to detect ppcp deletion efficiency is shown in Fig. 2A. In Frs. A–C from cKO mice, ppcp deletion efficiency was 24.4%, 71.4%, and 75.6%, respectively (Fig. 2B, lanes 4–6). In parallel to our deletion efficiency determinations, we extracted total RNA from WT and cKO Frs. A–C cells to analyze PP4 mRNA by RT-PCR (Fig. 2C). After normalization to HPRT expression, PP4 mRNA from cKO mice was reduced by 50% in Fr. B cells and by 77% in Fr. C cells compared to WT controls (Fig. 2D). Thus, consistent with a previous report [34], our results show that mb-1/cre-mediated ppcp gene deletion initiates in Fr. A cells, with higher deletion efficiencies occurring in Fr. B and Fr. C cells. The finding suggests that the developmental block at Fr. C stage in cKO mice likely correlates with the higher ppcp deletion efficiency in this population.

Impaired DJH3 Recombination and Increased DNA Double-strand Breaks at Ig loci in PP4-deficient Pro-B Cells

To further characterize the effects of PP4 deficiency during early B cell development, Frs. A–C cells from WT and cKO mice were subjected to intracellular staining to detect Ig μ HC. In WT mice, Ig μ+ cells constituted 0.3%, 21.8%, and 24.6% of Frs. A, B and C cells, respectively, whereas the corresponding percentages in cKO mice were 0.2%, 6.7%, and 7.6% (Fig. 3A, B). Thus, Ig μ HC expression was strongly reduced in the absence of PP4 beginning at the Fr. B stage, an observation confirmed using confocal microscopy to examine Fr. B cells from WT and cKO mice (Fig. 3C). To assess whether this reduction in Ig μ HC was due to impaired Ig DJH recombination, we subjected the genomic DNA of WT and cKO Fr. B cells to an in vitro DJH recombination assay. In both WT and cKO Fr. B cells, PCR products representing rearranged DJH1, DJH2, DJH3 and DJH4, gene segments appeared when the template was diluted 1:1 (Fig. 3D). In WT mice, the rearranged DJH3 gene segment remained detectable until the template dilution reached 1:16. In contrast, the rearranged DJH3 gene segment was detectable up to a dilution of only 1:4 in cKO Fr. B cells, suggesting that PP4 deficiency inhibits DJH3 recombination. These data imply that PP4 is involved in the regulation of Ig DJH/VHDJH recombination such that loss of ppcp impairs Ig μ HC expression.

During VDJ recombination, the hairpin coding ends are generated after the cleavage of recombination signal sequences (RSSs). Before the rejoining process is initiated, hairpin coding ends are opened to form double-strand ends [35]. To address whether the impaired Ig DJH recombination induced by PP4 deficiency is associated with double-strand breaks (DSBs) at RSSs, the genomic DNA of WT and cKO Fr. B cells was extracted and then subjected to coding end assay as described previously [33], [36]. The PCR product of approximate 350 bp, indicating the DSB at the 3′ RSS of DFL16.1 segment (DFL16.1 [3′]), appeared in both WT and cKO Fr. B cells (Fig. 3E and Fig. S2). However, the formation of DFL16.1 [3′]-coding end in cKO mice was

Table 1. Summary of cell counts (×10⁶) of B cell fractions in WT and cKO mice.

| Mice | Total BM | Fr. A | Fr. B | Fr. C | Fr. D | Fr. E | Fr. F |
|------|----------|-------|-------|-------|-------|-------|-------|
| WT   | 57.9 ± 10.63 | 2.89 ± 0.39 | 9.75 ± 3.68 | 6.98 ± 1.79 | 2.96 ± 1.75 | 1.44 ± 1.17 | 2.43 ± 1.62 |
| cKO  | 55.2 ± 1.65 | 4.76 ± 1.11 | 7.76 ± 0.71 | 0.87 ± 0.28 | 0.71 ± 0.05 | 0 ± 0 | 0 ± 0 |
| p    | 0.2992 | 0.0190 | 0.1484 | 0.0006 | 0.0014 | 0.0171 | 0.0049 |

Cell counts of total BM and Frs. A to F cells from WT and cKO mice (n = 4/group) were calculated following immunostaining to detect the appropriate markers and analysis by flow cytometry. Results are representative of three independent experiments.

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Table 2. Cell counts (×10⁶) of total splenocytes and splenic B cells in WT and cKO mice.

| Mice | Splenocytes | Splenic B cells |
|------|-------------|-----------------|
| WT   | 4.54 ± 0.56 | 1.11 ± 0.26 |
| cKO  | 0.50 ± 0.13 | 0.03 ± 0.05 |
| p    | 0.0024      | 0.0085 |

Total cell counts of splenocytes and splenic B cells (B220+ gated lymphocytes) from WT and cKO mice (n = 4/group) were calculated following immunostaining to detect the appropriate markers and analysis by flow cytometry. Results are representative of three independent experiments.

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nearly threefold more than that in WT mice. In addition, the PCR product of approximate 2 kb, indicating the DSB at the 3’ RSS of DSP segment (DSP [3’]), appeared in cKO mice. By contrast, DSP [3’]-coding end in WT mice was barely detectable. These data suggest that PP4 is involved in the regulation of coding end joining during Ig DJH/VHDJH recombination such that loss of pp4c causes DNA DSBs at Ig loci.

PP4 Deficiency is Associated with DNA Breaks and Impaired B Cell Homeostasis

During Ig gene rearrangement, DNA breaks are repaired mainly by factors in the NHEJ pathway [37]. The impaired Ig DJH/VHDJH recombination by PP4 deficiency raised the possibility that this phenomenon was associated with excessive DNA damage. To address this question, WT and cKO Fr. B and Fr. C cells were isolated and subjected to cell-cycle analysis. Cells of the BrdU+7AAD−, BrdU+7AAD+, BrdU+7AAD−, and BrdU+7AAD+ populations were defined as G1, G1/S, S/G2, and G2/M phase cells, respectively. We found that the percentages of Fr. B cells in each phase were comparable between WT and cKO mice (Fig. 4C). However, in cKO Fr. C cells, a 33% increase in G1 phase cells and a corresponding reduction in cells in the G1/S plus S/G2 phases were observed (Fig. 4C, 4D). To confirm these results in vitro, we cultured WT and cKO Fr. B and Fr. C cells for 16 h in the presence of BrdU and analyzed its incorporation. Similar to our in vivo results, no significant difference in the cell-cycle was observed between WT and cKO Fr. B cells (Fig. 4E). However, compared to WT Fr. C cells, cKO Fr. C cells showed a 2-fold increase in the percentage of G1 phase cells, while the percentages of G1/S and S/G2 phase cells were correspondingly diminished (Fig. 4E). Thus, PP4 deficiency affects the cell-cycle progression of Fr. C cells, leading to an accumulation of G1 phase cells and a reduction in S phase cells.

Table 3, suggesting that DNA damage is more severe in Fr. C cells lacking PP4.

To understand the effect of PP4 deficiency on cell-cycle progression, Fr. B and Fr. C cells were isolated and subjected to cell-cycle analysis. Cells of the BrdU+7AAD−, BrdU+7AAD+, BrdU+7AAD−, and BrdU+7AAD+ populations were defined as G1, G1/S, S/G2, and G2/M phase cells, respectively. We found that the percentages of Fr. B cells in each phase were comparable between WT and cKO mice (Fig. 4C). However, in cKO Fr. C cells, a 33% increase in G1 phase cells and a corresponding reduction in cells in the G1/S plus S/G2 phases were observed (Fig. 4C, 4D). To confirm these results in vitro, we cultured WT and cKO Fr. B and Fr. C cells for 16 h in the presence of BrdU and analyzed its incorporation. Similar to our in vivo results, no significant difference in the cell-cycle was observed between WT and cKO Fr. B and Fr. C cells (Fig. 4E). However, compared to WT Fr. C cells, cKO Fr. C cells showed a 2-fold increase in the percentage of G1 phase cells, while the percentages of G1/S and S/G2 phase cells were correspondingly diminished (Fig. 4E). Thus, PP4 deficiency affects the cell-cycle progression of Fr. C cells, leading to an accumulation of G1 phase cells and a reduction in S phase cells.
To investigate if PP4 deficiency was associated with increased apoptosis, freshly sorted WT and cKO Fr. B and Fr. C cells were subjected to AnnexinV plus 7AAD staining. While percentages of early apoptotic (AnnexinV$^+$7AAD$^+$) and late apoptotic (AnnexinV$^+$7AAD$^+$) cells were comparable between WT and cKO Fr. B cells, significant increases in early and late apoptotic cells were observed in cKO Fr. C cells compared to WT controls (Fig. 4F, 4G). To confirm this effect on cell viability \textit{in vitro}, WT and cKO

**Figure 3.** Reduced Ig$\mu$ HC expression and impaired DJH recombination in PP4-deficient pro-B cells. (A) Flow cytometric profiles of Frs. A, B and C cells from WT and cKO mice (n = 3/group) analyzed for B220 vs intracellular Ig$\mu$ HC (upper panel), or B220 vs isotype control (lower panel). Results are representative of two independent experiments. (B) Quantitation of percentages of the Fr. A, B and C cells in (A) expressing intracellular Ig$\mu$ HC. Data are the mean ± SD (n = 2). (C) Confocal images of intracellular Ig$\mu$ HC staining in Fr. B cells from WT and cKO mice (n = 2/group). DAPI, nuclei. Results are representative of three independent experiments. (D) Southern blot of genomic DNA from Fr. B cells of WT and cKO mice (n = 2/group) subjected to a DJH$_{\gamma}$ recombination assay. Genomic DNA was serially diluted from 1:1 to 1:64 prior to nested PCR. Bands were visualized using DIG-labeled southern blotting. CD14, loading control. Results are representative of two independent experiments. (E) Southern blot of genomic DNA from Fr. B cells of WT and cKO mice (n = 2/group) subjected to the coding end assay. Genomic DNA was filled by Klenow, ligated with double-strand oligo linker of 5$'$-overhang, and subjected to nested PCR. Bands were visualized using DIG-labeled southern blotting. CD14, loading control. Ten mice per group were used in this experiment, and two independent experiments were performed.

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Fr. B and Fr. C cells were cultured in RPMI 1640 medium containing 10% FCS for 16 h followed by 7AAD staining. The percentage of viable cells in the cKO Fr. B culture was drastically reduced compared to the control (WT; 49.4% vs. cKO, 21.7%), as was true for the cKO Fr. C culture (WT, 53.7% vs. cKO, 10.6%) (Fig. 4H). Taken together, these analyses indicate that PP4 deficiency severely interferes with the cell-cycle of Fr. C cells and induces their apoptosis, thereby disrupting B cell homeostasis.

### Discussion

In this study, we describe the essential role of PP4 in early B cell development through its regulation of Ig DJH/VµDHJH recombination. Utilizing the mb-1/cre-flox system to generate cKO mice, we have shown that cre activity under the control of the mb-1 promoter initiates at the pre-pro-B cell stage (Fr. A). However, the deletion efficiency of the pp-ff allele in Fr. A cells was poor, precluding us from investigating the function of PP4 in this population. Fortunately, acceptable deletion efficiencies were achieved in cKO Fr. B and Fr. C cells, permitting us to determine the effects of PP4 deficiency on these subsets. With respect to Fr. B cells, loss of PP4 leads to impaired Ig DJH recombination accompanied by reduced µ HC expression. These defects occur in the absence of abnormal cell-cycle progression or excessive apoptosis. During WT B cell development, secondary DJH rearrangements, which employ a more 5’ D segment to replace the existing DJH rearrangement, occur frequently in pro-B and pre-B cells [40]. Hence, it is likely that PP4-deficient Fr. B cells, which fail to complete DJH/VµDHJH recombination, undergo a secondary DJH/VµDHJH rearrangement rather than apoptosis. Consistent with that, we found that PP4-deficient Fr. B cells generate more coding ends at 3’ RSS sites of DFL16,1 and DSP. It suggests that the impaired DJH recombination by PP4 deficiency increases the usage of more 5’ D segments for a secondary rearrangement. Very importantly, the analyses of comet scores and apoptosis in Fr. B cells (shown in Fig. 4A and 4F) did not show any significant difference between WT and cKO mice. The data strongly suggest that PP4 regulates Ig DJH/VµDHJH recombination at the joining phase such that loss of pp-ff causes DSBs at Ig loci.

In contrast to Fr. B cells, cells in Fr. C that fail to accomplish a secondary or tertiary DJH/VµDHJH rearrangement do undergo apoptosis. It has been proposed that such premature death is invoked to prevent the survival of a cell with oncogenic potential [41]. This hypothesis might explain why WT Fr. C cells display more apoptosis than WT Fr. B cells. Even so, apoptosis in cKO Fr. C cells is more severe than in WT Fr. C cells, consistent with the drastic reduction in Fr. C cell counts in the mutant mouse. Because the sole defect in DNA repair during V(D)J recombination would not lead to the formation of visible comets, it is likely that the severe DNA fragmentation in PP4-deficient Fr. C cells is due to other mechanisms, such as apoptosis [42]. Nevertheless, cKO Fr. C cells were observed to accumulate in the G1 phase, indicating a disruption of cell-cycle progression. It is noteworthy that VµDHJH recombination is normally restricted at the G1 phase to ensure the repair of recombination-liberated DNA ends prior to re-entry of the cell-cycle [33]. The accumulation of G1 phase Fr. C cells in cKO mice, together with the strong induction of coding ends-associated DSBs in Fr. B cells, likely reflects the fact that Fr. C cells lacking PP4 cannot complete VµDHJH rearrangement and thus undergo apoptosis. Consequently, PP4 deficiency results in a severe reduction of Fr. C cells and developmental arrest. We were able to restore IgM+ BM B cells to WT levels in PP4-deficient mice through transgenic Ig expression. This finding strongly supports our hypothesis that PP4 is indispensable for Ig VH/DH recombination. It also reveals that the transgenic Ig transmitted functional BCR signaling that overcame the PP4 deficiency and supported the differentiation of IgM+ B cells.

### Rescue of the Development of PP4-deficient Pro-B Cells by Ig Transgene Expression

Based on our findings above, we hypothesized that PP4 is required for pro-B cell development because it promotes Ig DJH/VµDHJH recombination. To test this theory, we crossed our cKO mice to IgHEL transgenic mice [32] to generate cKO/IgHEL mice. The IgHEL transgene drives constitutive BCR expression, which promotes the differentiation of IgM+ cells in cKO/IgHEL mice [39]. Thus, the near-absence of Fr. C and Fr. D cells in cKO/IgHEL mice was likely mediated by their expression of the Ig transgene. We can therefore conclude that the Ig transgene was able to drive PP4-deficient B cell development by promoting the differentiation of IgM+ B cells.

### Table 3. Comet scores of Fr. B and Fr. C cells from WT and cKO mice.

| Frs. | Mice | Tail % intensity | Tail length/head length |
|------|------|-----------------|------------------------|
| Fr. B | WT (n = 36) | 13.82 ± 14.62 | 1.00 ± 0.31 |
|       | cKO (n = 57) | 10.33 ± 9.66 | 1.03 ± 0.38 |
| P    | 0.1082 | 0.3495 |
| Fr. C | WT (n = 61) | 10.79 ± 12.97 | 0.97 ± 0.85 |
|       | cKO (n = 41) | 16.50 ± 15.43 | 1.84 ± 1.25 |
| P    | 0.0275 | 0.0001 |

Tail % intensity, tail intensity expressed as a percentage of the comet’s total intensity; tail length, the horizontal distance from the centre of the head to the end of the tail; head length, the horizontal distance from the start of the head to the end of the head. Two mice per group were used in this experiment, and two independent experiments were performed.
Figure 4. PP4 deficiency causes excessive DNA damage and impairs B cell homeostasis. (A) Microscopic images of comet assays of Fr. B and Fr. C cells from WT and cKO mice (n = 2/group). Results are representative of two independent experiments. (B) Enlarged microscopic images of the comet assays of the WT and cKO Fr. C cells in (A). (C) In vivo BrdU incorporation assay. WT and cKO mice (n = 3/group) were i.p. injected with BrdU at 16 h and 24 h before isolation of Fr. B and Fr. C cells by cell sorting. Cells were subjected to intracellular staining with 7AAD plus anti-BrdU and analyzed by flow cytometry. Numbers in quadrants represent the percentage of the indicated population relative to the total. Results are representative of three independent experiments. (D) Quantification of the percentages of G1, G1/S, and S/G2 phase cells among the Fr. B and Fr. C cells in (C). Data are the mean ± SD (n = 3). (E) In vitro BrdU incorporation assay. Fr. B and Fr. C cells were sorted from WT and cKO mice (n = 2/group),
cultured for 16 h in the presence of BrdU, and subjected to intracellular staining and flow cytometric analysis as for (C). Results are representative of two independent experiments. (F) In vivo apoptosis assay. Fr. B and Fr. C cells from WT and cKO mice (n = 4/group) were stained to detect AnnexinV plus 7AAD and analyzed by flow cytometry. Numbers in the lower right quadrant indicate the percentage of cells in early apoptosis, whereas the values in the upper right quadrant indicate the percentage of cells in late apoptosis. Results are representative of two independent experiments. (G) Quantitation of Fr. B and Fr. C cells in early or late apoptosis as calculated from the results in (F). Data are the mean ± SD (n = 4–6/group). (H) In vitro viability assay. Fr. B and Fr. C cells were sorted from WT and cKO mice (n = 2/group) and cultured for 16 h under standard conditions. Cells were surface-stained with 7AAD and the percentage of viable cells was calculated. Data are the mean ± SD (n = 2). Results are representative of two independent experiments (*p<0.05, **p<0.005, ***p<0.0005).

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Figure 5. The defect in B cell development caused by PP4 deficiency can be rescued by Ig transgene expression. (A) Flow cytometric profiles of: total BM lymphocytes from WT, cKO and cKO/IgHEL mice (n = 3–5/group) analyzed for B220 vs IgM (far left panel) or B220 vs CD43 (middle left panel); or B220^CD43^ BM lymphocytes analyzed for CD24 vs BP-1 (middle right panel); or B220^CD43^ BM lymphocytes analyzed for IgM vs IgD (far right panel). (B) Quantitative cell count analysis of Frs. A to C cells from the mice in (A). (C) Quantitative cell count analysis of Frs. D to F cells from the mice in (A). For B and C, data are the mean ± SD (n = 3–5/group). Results are representative of two independent experiments (*p<0.05, **p<0.005; ns, not significant).

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than the effect of mb-1/cre mediated PP4 deficiency in B cells. Although the mechanism for VDJ recombination is in general highly conserved between B and T lymphocytes, gene-targeting of NHEJ factors in some cases can lead to unequal severities in B and T cell developments. For example, in Ku70 knockout (KO) mice and DNA polymerase Pol γ (PGK1 KO mice), VDJ recombination at Ig loci is drastically impaired, whereas TCRβ/α gene rearrangement is functional [43] [44] [45]. A comprehensive recombination assay in PP4-deficient thymocytes would be necessary to address whether PP4 is also involved in TCR gene rearrangement.

PP4 has been implicated in two DNA repair pathways: homologous recombination-mediated repair of DNA breaks via dephosphorylation of γH2AX [18], [24], and NHEJ-mediated end-joining by dephosphorylation of KAP1 (KRAB-associated protein 1) [46]. Whether the disruption of γH2AX or KAP1 phosphorylation status can lead to abrupt developmental arrest at the pro-B cell stage requires further investigation. In any case, the results of our study provide the first genetic evidence establishing a critical role for PP4 in pro-B cell development through its promotion of Ig VDJ recombination.

Supporting Information

Figure S1 Schematic diagram of southern blotting strategy used to detect DJH recombination. (TIFF)

Figure S2 Schematic diagram of southern blotting strategy used to detect coding ends at 3′ of D segments. (TIFF)

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Author Contributions

Conceived and designed the experiments: YwS THT. Performed the experiments: YwS VpC MYC. Analyzed the data: YwS. Contributed reagents/materials/analysis tools: YwS MR THT. Wrote the paper: YwS.

References

1. Hardy RR, Carmack CE, Shinton SA, Kemp JD, Hayakawa K (1991) Resolution and characterization of pro-B and pre-pro-B cell stages in normal mouse bone marrow. J Exp Med 173: 1215–1225.
2. Li YS, Hayakawa K, Hardy RR (1993) The regulated expression of B lineage associated genes during B cell differentiation in bone marrow and fetal liver. J Cell Sci 107: 953–960.
3. Alt FW, Nanculescu GD, Blackwell TK, Wood C, Thomas E, et al. (1984) Ordered rearrangement of immunoglobulin heavy chain variable region segments. EMBO J 3: 1209–1219.
4. Pike KA, Racchielle MJ (2002) Cell surface immunoglobulin receptors in B cell maturation. Semin Immunol 14: 351–358.
5. Bassing CH, Swat W, Alt FW (2002) The mechanism and regulation of chromosome VDJ recombination. Cell 109 Suppl: S45–S55.
6. Jung D, Giaullouasaki G, Mostolsavsky R, Alt FW (2006) Mechanism and control of VDJ recombination at the immunoglobulin heavy chain locus. Annu Rev Immunol 24: 541–570.
7. Puebla-Osorio N, Zhu C (2008) DNA damage and repair during lymphoid development: antigen receptor diversity, genomic integrity and lymphomagenesis. Immuno Res 41: 103–122.
8. Kloeker S, Wadzinski BE (1999) Purification and identification of a novel subunit of protein serine/threonine phosphatase 4. J Biol Chem 274: 5329–5347.
9. Wada T, Miyata T, Iinagi R, Nakanou M, Wagaotuma M, et al. (2001) Cloning and characterization of a novel subunit of protein serine/threonine phosphatase 4 from mesangial cells. J Am Soc Nephrol 12: 2601–2608.
10. Venkataranam DV, Fulton JB, Andreetti AH, Johansen KM, Johansen J (2006) Mapping the Ca2+–dependent binding of an invertebrate homolog of protein phosphatase 4 regulatory subunit 2 to the small EF-hand protein, calbindin. Biochim Biophys Acta 1765: 322–329.
11. Gingras AC, Caballero M, Zarks M, Sanchez A, Hahmun TR, et al. (2005) A novel, evolutionarily conserved protein phosphatase complex involved in clapin sensitivity. Mol Cell Proteomics 4: 1725–1740.
12. Chen GI, Tsehaykon S, Jorgensen C, D’Ambrosio LM, Gourdie RD, et al. (2008) PPM4K1A/A/ A91622 forms a novel stable cytosolic complex with phosphoprotein phosphatase 4. J Biol Chem 283: 29273–29282.
13. Chen J, Peterson KT, Schreiber SL (1998) α4 associates with protein phosphatase X interacts with c-Rel and stimulates c-Rel/nuclear factor-kappa B signaling in T lymphocytes. Immunity 37: 697–708.
14. Hobeika E, Thiemann S, Storch B, Junaza H, Nielsen JE, et al. (2002) Protein phosphatase 4 is involved in tumor necrosis factor-alpha-induced activation of c-Jun N-terminal kinase. J Biol Chem 277: 6301–6308.
15. Mihindukulasariya KA, Zhou G, Qiu J, Tan TH (2004) Protein phosphatase 4 interacts with and down-regulates insulin receptor substrate 4 following tumor necrosis factor-alpha stimulation. J Biol Chem 279: 46588–46594.
16. Chen L, Dong W, Zhou T, Ouyang L, He G, et al. (2008) Protein phosphatase 4 negatively regulates LFS cascade by inhibiting ubiquitination of TRAF6. FEBS Lett 582: 2843–2849.
17. Hu MC, Hung MC, Tan TH (2007) Conditional knockout mice reveal an essential role of protein phosphatase 4 in thymocyte development and pre-T-cell receptor signaling. Mol Cell Biol 27: 79–91.
18. Huang CY, Chou YP, Yang YJ, Huang CH, Yang YJ, et al. (2002) Alpha 4 associates with protein phosphatase 4 from mesangial cells. J Am Soc Nephrol 12: 2601–2608.
19. Sliz P, Hartley SP, Downward J, Wu Z, Zhai S, et al. (2006) PP4 regulates NF-kappa B and BCR-induced proliferation and survival of B lymphocytes. Blood 108: 5717–5726.
20. Scherer CF, Bergeron SG, Lander ES, Gilman S (2004) Protein phosphatase 4 catalytic subunit interacts with and down-regulates insulin receptor substrate 4 following tumor necrosis factor-alpha stimulation. J Biol Chem 279: 46588–46594.
21. Chen L, Dong W, Zhou T, Ouyang L, He G, et al. (2008) Protein phosphatase 4 negatively regulates LFS cascade by inhibiting ubiquitination of TRAF6. FEBS Lett 582: 2843–2849.
22. Hu MC, Tang-Oxley Q, Qiu WR, Wong YP, Mihindukulasariya KA, et al. (1998) Protein phosphatase X interacts with c-Rel and stimulates c-Rel/nuclear factor-kappa B activity. J Biol Chem 273: 33561–33565.
23. Brechmann M, Mock T, Niekles D, Kießling M, Weit N, et al. (2012) A PP4a holoenzyme balances physiological and oncogenic nuclear factor-kappa B signaling in T lymphocytes. Immunity 33: 679–687.
24. Hobeika E, Thiemann S, Storch B, Junaza H, Nielsen JE, et al. (2006) Testing gene function early in the B cell lineage in mb1-cre mice. Proc Natl Acad Sci U S A 103: 13789–13794.
25. Goodnow CC, Crozher J, Adelstein S, Lavoie TB, Smith-Gill SJ, et al. (1988) Altered immunoglobulin expression and functional silencing of self-reactive B lymphocytes in transgenic mice. Nature 334: 676–692.
37. Dudley DD, Chaudhuri J, Bassing CH, Alt FW (2005) Mechanism and control of V(D)J recombination versus class switch recombination: similarities and differences. Adv Immunol 86: 43–112.

38. Chang Y, Bosma GC, Bosma MJ (1995) Development of B cells in scid mice with immunoglobulin transgenes: implications for the control of V(D)J recombination. Immunity 2: 607–616.

39. Spanopoulou E, Roman CA, Corcoran LM, Schlissel MS, Silver DP, et al. (1994) Functional immunoglobulin transgenes guide ordered B-cell differentiation in Rag-1-deficient mice. Genes Dev 8: 1030–1042.

40. Reth MG, Jackson S, Alt FW (1986) VHDJH formation and DJH replacement during pre-B differentiation: non-random usage of gene segments. EMBO J 5: 2131–2138.

41. Guidos CJ, Williams CJ, Grandal I, Knowles G, Huang MT, et al. (1996) V(D)J recombination activates a p53-dependent DNA damage checkpoint in scid lymphocyte precursors. Genes Dev 10: 2038–2054.

42. Enari M, Sakahira H, Yokoyama H, Okawa K, Iwamatsu A, et al. (1998) A caspase-activated DNase that degrades DNA during apoptosis, and its inhibitor ICAD. Nature 391: 43–50.

43. Ouyang H, Nussenzweig A, Kurimasa A, Soares VC, Li X, et al. (1997) Ku70 is required for DNA repair but not for T cell antigen receptor gene recombination In vivo. J Exp Med 186: 921–929.

44. Gu Y, Scidl KJ, Rathbun GA, Zhu C, Manis JP, et al. (1997) Growth retardation and leaky SCID phenotype of Ku70-deficient mice. Immunity 7: 653–665.

45. Bertocci B, De Smet A, Berek C, Weill JC, Reynaud GA (2003) Immunoglobulin kappa light chain gene rearrangement is impaired in mice deficient for DNA polymerase mu. Immunity 19: 203–211.

46. Liu J, Xu L, Zhong J, Liao J, Li J, et al. (2012) Protein phosphatase PP4 is involved in NHEJ-mediated repair of DNA double-strand breaks. Cell Cycle 11: 2643–2649.