Acetyltransferases GCN5 and PCAF are required for B cell maturation in mice

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Abstract: B lymphocyte development includes two DNA recombination processes, the V(D)J recombination of immunoglobulin (Igh) gene variable region and class switching when the Igh constant regions are changed from IgM to IgG, IgA, or IgE. The V(D)J recombination is required for successful maturation of B cells from pro-B to pre-B to immature-B and then to mature B cells in the bone marrow. The CSR occurs outside the bone marrow when mature B cells migrate to peripheral lymphoid organs, such as spleen and lymph nodes. Both V(D)J recombination and CSR depend on an “open chromatin” state that makes DNA accessible to specific enzymes, recombination activating gene (RAG), and activation-induced cytidine deaminase (AID). Acetyltransferases GCN5 and PCAF possess redundant functions acetylating histone H3 lysine 9 (H3K9). Here, we generated by complex breeding a mouse model with B cells lacking both GCN5 and PCAF. We found that double-deficient mice possess low levels of mature B cells in the bone marrow and peripheral organs, accumulation of pro-B cells in bone marrow, and reduced CSR levels. We concluded that both GCN5 and PCAF are required for B cell development in vivo.

Keywords: KAT2A; KAT2B; mice; acetyltransferase; B cell; lymphocyte; class switching

1. Introduction

Development of B lymphocytes starts in the bone marrow where progenitor (pro)-B cells using recombination-activating genes (RAG) generate DNA double-strand breaks (DSBs) and initiate the V(D)J recombination. In maturing B cells, the V(D)J recombination process is genetic recombination of variable (V), diversity (D), and joining (J) gene segments arranging into a newly formed VDJ part of immunoglobulin gene (Ig) (1-4). Following the V(D)J recombination, B cells develop from pro-B cells expressing specific markers cluster of differentiation 19 (CD19), B220/CD45, and CD43 (CD19+B220+CD43+) to pre-B cells (CD19+B220+CD43+), immature B (CD19+B220+IgM+, low immunoglobulin M, IgM) and mature B (CD19+B220+IgM+, high IgM) cells in bone marrow (1). Mature B lymphocytes leave the bone marrow and through the blood migrate to periphery populating spleen and lymph nodes.

Then, mature B cells initiate another DNA recombination process to change the constant regions of immunoglobulin genes referred to as class switch recombination (CSR). During the CSR in mice, IgM is replaced by IgG3, IgG1, IgG2a, IgG2b, IgE, and IgA (1). The CSR is initiated by non-productive transcription known as a germ-line transcription (GLT) which is needed to separate two DNA strands. Single-stranded DNA is then targeted by activation-induced cytidine deaminase (AID), a B lymphocyte-specific enzyme...
deaminating cytosine to uracil (C to U). Then uracil DNA N-glycosylase (UNG) removes the uracil from DNA leading to the single-strand break formation (SSB)(5). Two SSBs facing each other then form a DSB and allow recombination (1-4). Both V(D)J and CSR can be regarded as processes following fundamentally similar strategies of genomic recombination (6, 7).

The DSBs formed during the V(D)J recombination and CSR are recognized, processed, and repaired by the non-homologous end-joining pathway (NHEJ) and initiate more complex signaling and chromatin modification known as DNA damage response (DDR) (1-4). The NHEJ is initiated when Ku70 and Ku80 heterodimer (Ku) recognizes and binds the DSBs. Ku serves as a platform for downstream factors including DNA-dependent protein kinase catalytic subunit (DNA-PKcs), X-ray repair complementing protein 4 (XRCC4), XRCC4-like factor (XLF), Paralogue of XRCC4 and XLF (PAXX), a modulator of retrovirus infection (MRJ), XRCC4 and DNA ligase 4 (LIG4) (2-4). There are additional factors that are sometimes optional for NHEJ, including Artemis with nuclease activity required for processing of hairpin-sealed DNA ends and overhangs.

One type of DDR pathway acts downstream of ataxia telangiectasia mutated (ATM) protein kinase, which is activated by DSBs and then phosphorylates multiple substrates, including NHEJ and DDR factors. ATM phosphorylates histone H2AX, which in turn recruits mediator of DNA damage checkpoint 1 (MDC1) and facilitates the accumulation of really interesting new gene (RING) finger motif (RNF) 8 and RNF168 ubiquitin-ligases, and then the p53-binding protein (53BP1). Phosphorylation of H2AX is related to the acetylation of histones, including histone H3K9 (8). Acetylation of histone H3K9 is mediated by GCN5 and PCAF (8, 9).

There is a complex genetic interaction associated with the functional redundancy between the NHEJ factors (2, 3), including the following pairs: DNA-PKcs/XLF (4, 10), PAXX/XLF (11-17), and MRI/XLF (18, 19). Moreover, there is a genetic interaction between NHEJ and DDR pathway factors, for example, ATM/XLF and H2AX/XLF (20), MDC1/XLF (21), RNF8/XLF and RNF168/XLF (22), 53BP1/XLF (23, 24), and more (2, 3). Moreover, acetyltransferases GCN5 and PCAF are redundant in promoting histone H3 lysine K9 acetylation (25).

General control non-depressible 5 (GCN5) acetyltransferase is also known as lysine acetyltransferase (KAT) 2A. Germline inactivation of GCN5 in mice resulted in early embryonic lethality due to the role of the protein in neurogenesis (25). GCN5 is functionally redundant with another acetyltransferase, KAT2B, also referred to as p300/CBP-associated factor (PCAF). While inactivation of Pcaf gene in mice has no detectable phenotype, double knockout of Gen5/Pcaf genes resulted in even earlier embryonic lethality (25). Because histone H3K9 acetylation is related to DDR (8), one could propose that GCN5, PCAF, or both enzymes are required for lymphocyte development in vivo. However, embryonic lethality of Gen5- and Gen5- Pcaf- mice (25) challenged the studies. To overcome the obstacle, we developed a complex genetic model, where Pcaf gene was germline-inactivated (25), while floxed Gen5 gene was conditionally-inactivated in B cell lineages by CRE recombinase expressed under Cd19 promoter (26). To sort out CRE-positive and CRE-negative cells, we used Rosa26-stop-YFP knockin which only expressed YFP following the CRE activation (27).

Here, we found that GCN5 and PCAF acetyltransferases are functionally redundant during early B cell maturation, while GCN5 is required for robust CSR.

2. Materials and Methods

2.1. Mice

Gen5f/f (25), Pcaf- (25), Cd19cre+ (26) (# 006785; The Jackson Laboratory, Bar Harbor, ME, USA), Rosa26-stop-YFP+ (27)(# 006148, The Jackson Laboratory, Bar Harbor, ME, USA), and Aid- (28) mice were previously described. Mice used for experiments were between 8-12 weeks of age. All experiments were performed in compliance with the Danish Working Environment Authority, the Danish Animal Experiment Inspectorate, and the Department of Experimental Medicine (University of Copenhagen).
2.2. Flow cytometry

Flow cytometry experiments were performed as we described earlier (18, 21, 29-31). In particular, we used the fluorescent antibodies recognizing the proteins described below: B220 (PE-CF594, FITC, Alexa Fluor 700; all clone RA3-6B2, BD Bioscience, Franklin Lakes, NJ, USA), IgM (PerCP-eFluor 710, APC-eFluor 780, APC, FITC; all clone II/41, eBioscience, Santa Clara, CA, USA), IgG1 (PE, MOPC-21, Biolegend, San Diego, CA, USA), IgG3 (PE, MG3-35, Biolegend, San Diego, CA, USA), CD3-APC (Biolegends, USA, #100312), CD19 (Alexa Fluor 700, APC eFluor 780, both clone 1D3, eBioscience, Santa Clara, CA, USA), CD43 (APC and PE-Cy7, both clones S7, BD Bioscience, Franklin Lakes, NJ, USA).

2.3. Class switch recombination

The CSR was performed as we described earlier (10, 20, 30-34).

2.4. Western blot

The western blot procedures were performed as we described earlier (21, 29, 30, 35). Briefly, the cells were lysed for 30 minutes on ice in Radioimmunoprecipitation assay (RIPA) buffer (Sigma Aldrich, St. Louis, MO, USA, #R0278) supplemented with complete™ EDTA-free Protease Inhibitor Cocktail (Sigma Aldrich, #11873580001). Proteins were analyzed by 4-12% Bis-Tris NuPAGE gels (Invitrogen, Carlsbad, CA, USA, #NP0322), transferred to PDVF membranes (GE Healthcare, Boston, MA, USA, #GE10600023), and probed with indicated antibodies. Rat anti-AID (1:500, Active Motif, Carlsbad, CA, USA, #39886); mouse anti-GCN5, clone A-11 (1:500, Santa Cruz Biotechnology, Dallas, TX, USA, #sc-365321); rabbit anti-PCAF, clone C14G9 (1:1000, Cell Signaling Technology, Leiden, The Netherlands, #3378); rabbit anti-histone H3 (1:1000, Abcam, Cambridge, UK, #ab1791); rabbit anti-histone H3 acetyl K9 (1:500, Abcam, Cambridge, UK, #ab32129).

2.5. Statistics

We performed statistical analyses with one-way ANOVA using GraphPad Prism 8.0.1.244 (San Diego, CA, USA). In the tests, p values less than 0.05 were defined as significant, i.e. *p<0.05; **p<0.01; ***p<0.001; and ****p<0.0001.

3. Results

3.1. Generation of mice lacking GCN5 and PCAF in B cells

Combined inactivation of Gcn5 and Pcaf genes in mice results in embryonic lethality (25). To overcome this challenge, we designed a complex genetic model when floxed Gcn5 gene is conditionally inactivated in B cell lineages by CRE enzyme under the Cd19 promoter (Cd19Cre+) (26). To sort out the cells with activated CRE, we used a model with knocked-in yellow fluorescent protein gene (YFP) into ROSA-26 locus. The YFP is inactive until CRE removes “STOP” signal (Rosa-26-YFP+) (27). Thus, we obtained Gcn5+/Pcaf+/Cd19Cre+/YFP+ mice and simpler controls. Further in the text, we will skip Cd19Cre+ and YFP+ for simplicity in most of the cases, and will refer to the mice based on the status of Gcn5 and Pcaf genes, i.e. as Gcn5+/Pcaf+, Gcn5+, Pcaf+ and WT. When the CRE is active and describing sorted B cells, we indicate Gcn5−/−, a knockout status of the gene. Lack of GCN5 and PCAF was also validated by western blot (Figure S1) and quantitative PCR.
3.2. Mice lacking GCN5 and PCAF in B cells are alive and possess small spleens

We obtained alive mice with germline inactivation of *Pcaf* gene and conditional inactivation of *Gcn5* in B cells under the *Cd19* promoter (Figure 1). We found that germline inactivation of *Pcaf* gene alone has no detectable effect on mouse development, in line with the previous observation (25). Conditional inactivation of *Gcn5* gene in B cells had no visible effect on size of WT and *Pcaf*−/− mice, which were 15 to 19 g on average (p>0.1433) (Figure 1A). However, inactivation of *Gcn5* resulted in smaller spleens in mice (*Gcn5*−/−, 54 mg), when compared to WT (69 mg) and *Pcaf*−/− (72 mg) mice. Combined inactivation of *Pcaf* and *Gcn5* in B cells resulted in even smaller spleens (*Gcn5*−/−*Pcaf*−/−, 29 mg, p<0.0001). Spleens of mice without CRE activity with *Gcn5* gene being floxed and functional (*Gcn5*f/f *Pcaf*−/−, 67 mg), were comparable in size to the ones of WT (Figure 1B, C).

Figure 1. Generation of mice with germline inactivation of *Pcaf* and conditional inactivation of *Gcn5* gene in B cell lineages. (A) Sizes of 8 weeks-old mice of indicated genotype are similar (p=0.1433). (B) Size of spleens from mice of indicated genotypes. WT vs *Pcaf*−/−, n.s. p=0.9559; WT vs *Gcn5*−/−, *p=0.0192; WT vs *Gcn5*−/−*Pcaf*−/−, ****p<0.0001; *Pcaf*−/− vs *Gcn5*−/−, *p=0.0119; *Pcaf*−/− vs *Gcn5*−/−*Pcaf*−/−, **** p<0.0001; *Gcn5*−/− vs *Gcn5*−/−*Pcaf*−/−, ***p=0.0008. (C) Example of spleens of indicated genotypes. *Gcn5*−/− indicates *Cd19*Cre-dependent inactivation of *Gcn5* in B cell lineages.

3.3. Mice lacking GCN5 and PCAF in B cells are alive and possess small spleens

To detect mature *Gcn5*−/−*Pcaf*−/− B cells, we identified B220+IgM+ cells in the spleen using flow cytometry (Figure 2 A, B). Inactivation of *Pcaf* gene alone did not affect B cell proportions in the spleen (58%) when compared to WT mice (52%, p=0.4777) (Figure 2A). Inactivation of *Gcn5* alone resulted in an insignificant reduction of mature splenocytes when compared to WT (*Gcn5*−/−, 46%, p<0.2532), and there were the least of *Gcn5*−/−*Pcaf*−/− B cells in the spleen (21%, p<0.0001). Similarly, the number of *Gcn5*−/−*Pcaf*−/− B splenocytes was the lowest (3.4 million), while the number of *Gcn5*−/− B cells (27 million) was also reduced when compared to *Pcaf*−/− (54 million, *p=0.0065) and WT (44 million, *p=0.0214) controls (Figure 2B).
3.4. Inactivation of Gcn5 and Pcaf results in a reduced proportion of B cells in the blood

To detect mature B cells in the blood, we used B220 markers (Figure 2C, D). Inactivation of Pcaf alone resulted in 62% of B cells after red blood cells were lysed, which was comparable to WT mice with 65% of B cells in the blood (p=0.92). Inactivation of Gcn5 gene alone resulted in a modest reduction of B cell proportion to 56% (p=0.12), while combined inactivation of Gcn5 and Pcaf led to very low B cell levels in blood (22%, p=0.0001). Levels of B cells in blood of control mice without CRE recombinase expression, when Gcn5 gene was functional (Gcn5<sup>+/+</sup>Pcaf<sup>+/+</sup>) were comparable to WT mice (78%). We concluded that GCN5 and PCAF are both required and functionally redundant for B cell development in mice.

One reason of low B cell count in spleen and blood of Gcn5<sup>−/−</sup>Pcaf<sup>−/−</sup> mice could be cell death following normal development of B cells in bone marrow and migration to periphery. Another option could be blocked or delayed maturation of B cells in bone marrow during the earlier stages. To test the latter possibility, we analyzed B cell development in bone marrow of the mice (Figure 3).

3.5. Inactivation of Gcn5 and Pcaf results in accumulation of pro-B cells in bone marrow

To characterize B cell maturation in bone marrow, we followed the expression of B220 (B220+IgM<sup>+</sup>) and IgM (B220+IgM<sup>+</sup>) on the lymphocyte surface. Inactivation of Gcn5 or Pcaf resulted in an insignificant decline of B220+IgM<sup>+</sup> population (26%, 5-6 million) when compared to WT (32%, 8 million, p=0.44)(Figure 3 A, B). Combined inactivation of Gcn5 and Pcaf resulted in an additional reduction of mature B cells in bone marrow (3 million, 16%) (Figure 3 A, B).

We further focused on B220+IgM<sup>+</sup> populations by determining the CD43+ (pro-B cells) and CD43- (pre-B cells). Proportion of early stage pro-B cells increased from WT mice.
(20% on average) and *Pcaf*−/− mice (26%, *p*=0.78) to *Gcn5*−/− (39%, **p=0.0022) to *Gcn5*−/−*Pcaf*−/− (55%, *p<0.0001) (Figure 3 C, D).

We concluded that GCN5 and PCAF are required for the maturation of B cells from the pro-B cell stage to pre-B and later to mature B cells.

![Flow cytometry analyzes of developing B lymphocytes in bone marrow of 8-12-weeks-old mice of indicated genotypes.](image)

Figure 3. Flow cytometry analyzes of developing B lymphocytes in bone marrow of 8-12-weeks-old mice of indicated genotypes. (A) Examples of B220+IgM− (pro-B and pre-B), B220+IgM−low (immature B) and B220+IgM−high (mature B) cell populations. (B) Summary of several experiments from (A) indicating B220+IgM− cells. WT vs *Pcaf*−/−, n.s. *p*=0.6445; WT vs *Gcn5*−/−, n.s., *p*=0.4432; WT vs *Gcn5*−/−*Pcaf*−/−, **p=0.0008; *Pcaf*−/− vs *Gcn5*−/−, n.s., *p=0.9997; *Pcaf*−/− vs *Gcn5*−/−*Pcaf*−/−, *p<0.0138; *Gcn5*−/− vs *Gcn5*−/−*Pcaf*−/−, *p<0.0211 (C) CD43+ (pro-B cells) and CD43− (pre-B cells) gated from B220+IgM− populations in (A). (D) Summary of several experiments detecting B220+IgM−CD43+ (pro-B) cells. WT vs *Pcaf*−/−, n.s. *p=0.7825; WT vs *Gcn5*−/−, **p=0.0022; WT vs *Gcn5*−/−*Pcaf*−/−, **p<0.0001; *Pcaf*−/− vs *Gcn5*−/−, n.s., *p=0.0505; *Pcaf*−/− vs *Gcn5*−/−*Pcaf*−/−, **p<0.0001; *Gcn5*−/− vs *Gcn5*−/−*Pcaf*−/−, *p<0.0322 *Gcn5*−/− and *Gcn5*−/−*Pcaf*−/− indicate *Cd19*−/−-dependent inactivation of *Gcn5* in B cell lineages.

3.6. GCN5 and PCAF are required for robust class switch recombination

The CSR depends on the ATM-dependent DDR (1, 3, 4, 20). Because H3K9 acetylation works downstream of H2AX phosphorylation, and GCN5/PCAF might work downstream of ATM/ATR/DNA-PKcs, we tested if the CSR depends on GCN5 and PCAF (Figure 4). We purified B splenocytes from 8 to 12 weeks old mice and stimulated the CSR from IgM to IgG3 using established protocols (32, 34). We focused on matched pairs of *Gcn5*+/− (the functional equivalent of WT cells) and *Gcn5*−/−, as well as *Gcn5*−/−*Pcaf*−/− (the functional equivalent of *Pcaf*−/−) and *Gcn5*−/−*Pcaf*−/− cells. We used *Aid*−/− cells as a CSR-deficient control to detect an experimental background (Figure 4). Inactivation of *Pcaf* alone had no effect on CSR levels (WT vs *Gcn5*−/−*Pcaf*−/−, *p>0.96). Contrary, the inactivation of *Gcn5* resulted in a reduction of CSR levels from about 14% in WT and *Gcn5*+/− cells to 6% in *Gcn5*−/− cells, *p<0.0008 (Figure 4). Combined deletion of *Pcaf* and *Gcn5* resulted in a similar reduction from 12% in *Gcn5*−/−*Pcaf*−/− cells to 6% in *Gcn5*−/−*Pcaf*−/− cells, **p=0.0080. We concluded that GCN5 is required for robust CSR to IgG3, because additional inactivation of *Pcaf* did not affect CSR levels when compared *Gcn5*−/− and *Gcn5*−/−*Pcaf*−/− B cells, *p=0.9999 (Figure 4).
4. Discussion

Both GCN5 and PCAF are involved in chromatin modification and DDR response, which made them relevant candidates to facilitate lymphocyte development (8, 9). One challenge was the lack of a relevant in vivo model because GCN5 and PCAF have certain redundant functions in acetylating H3K9, and because of the germline inactivation of Gcn5 results in early embryonic lethality in mice (25). Here, we generated and analyzed a complex mouse model which allowed studying of double-deficient Gcn5–PCAF– B cells development in vivo and ex vivo. We used a germline knockout of Pcaf (25), a conditional knockout of Gcn5f/f, a knockin of CRE recombinase expressed under the B cell-specific Cd19 promoter (26), and a knockin of YFP to track the activity of CRE recombinase (27).

For such a complex mouse model (Gcn5f/fPcaf–Cd19cre- Rosa-26-YFP), multiple controls were used. In one line of the controls, the mice lacking PCAF, having floxed Gcn5 gene but expressing no CRE recombinase were used (Gcn5f/fPcaf–Rosa-26-YFP), Figure S2). The Gcn5-deficient and Gcn5/PCAF double-deficient B cells possessed developmental delay with lower levels of mature B cells in spleen and blood, and accumulation of progenitor B cells in bone marrow (Figures 1-4). Contrary, the control mice without CRE expression demonstrated WT levels of B cell development in all the groups, i.e. WT levels of B220+IgM+ mature B cells in the spleen (Figure S2 A and B) and blood (Figure S2 C and D). In addition, these mice possessed high levels of B220+IgM+ cells (Figure S2 E, F), as well as stable and low levels of pro-B cells (B220+IgM-CD43+) in bone marrow (Figure S2, G and H).

The mice lacking PCAF and with conditional knockout of Gcn5 in B cells were of normal size and visually not different from the WT and PCAF-deficient littermates (Figure 1A). One clear feature the Gcn5f/fPcaf–Cd19cre- Rosa-26-YFP mice had was a small spleen (Figure 1 B and C) which was also the case in mice lacking Gcn5 in B cells. The small spleen could indicate a defect in B cell development, and we indeed found low numbers of mature B cells in the spleen, blood, and bone marrow. One could propose that mature B cells lacking GCN5 or both GCN5 and PCAF possess low proliferation speed or tend to trigger apoptosis. Alternatively, GCN5 and PCAF might be required for the V(D)J recombination. This option could be tested by, for example, using vAbl pre-B cell lines as we and others did before, e.g. (3, 10, 11, 16, 20-24). Another intriguing question is whether the physical presence or enzymatic activity of GCN5 and PCAF are required for the observed phenotypes, i.e. abrogated B cell maturation and reduced levels of CSR. To investigate this question one could use specific inhibitors of GCN5 and PCAF enzymes, or enzyme-dead mutations introduced to the Gcn5 and Pcaf genes.

The CSR levels were reduced in B cells lacking GCN5 (Figure 4). The challenge in this set of experiments was that the mice of Gcn5f/fPcaf–Cd19cre- Rosa-26-YFP genotype were rather rare and possessed a very low number of suitable B splenocytes (Figures 1 and 2). Although our data on IgG3 is sufficient, one could extend the study in the future.
by generating knockout cell lines lacking GCN5 and PCAF and suitable for CSR. One possible model system is CH12F3 cells capable to support CSR to IgA (36), which were used in the past for this kind of experiment (17, 29, 34). The CSR itself is a complex multistage process. Generating relevant cell lines will also provide tools to determine specific stages of CSR affected in GCN5-deficient mice, i.e. germline transcription, AID recruitment, generation of DSBs, or DNA repair.

5. Conclusions

Acetyltransferases GCN5 and PCAF possess redundant functions in B cell maturation. GCN5 is required for class switch recombination in vivo.

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References

1. F. W. Alt, Y. Zhang, F. L. Meng, C. Guo, B. Schwer, Mechanisms of programmed DNA lesions and genomic instability in the immune system. Cell 152, 417-429 (2013).
2. X. S. Wang, B. J. Lee, S. Zha, The recent advances in non-homologous end-joining through the lens of lymphocyte development. DNA Repair (Amst) 94, 102874 (2020).
3. S. Castaneda-Zegarra et al., Genetic interaction between the non-homologous end-joining factors during B and T lymphocyte development: In vivo mouse models. Scand J Immunol 92, e12936 (2020).
4. V. Kumar, F. W. Alt, V. Oksenych, Functional overlaps between XLF and the ATM-dependent DNA double strand break response. DNA Repair (Amst) 16, 11-22 (2014).
5. D. L. Bordin, L. Lirussi, H. Nilsen, Cellular response to endogenous DNA damage: DNA base modifications in gene expression regulation. DNA Repair (Amst) 99, 103051 (2021).
6. X. Zhang et al., Fundamental roles of chromatin loop extrusion in antibody class switching. Nature 575, 385-389 (2019).
7. Y. Zhang et al., The fundamental role of chromatin loop extrusion in physiological V(D)J recombination. Nature 573, 600-604 (2019).
8. H. S. Lee, J. H. Park, S. J. Kim, S. J. Kwon, J. Kwon, A cooperative activation loop among SWI/SNF, gamma-H2AX and H3 acetylation for DNA double-strand break repair. EMBO J 29, 1434-1445 (2010).
9. Q. Jin et al., Distinct roles of GCN5/PCAF-mediated H3K9ac and CBP/p300-mediated H3K18/27ac in nuclear receptor transactivation. EMBO J 30, 249-262 (2011).
10. V. Oksenych et al., Functional redundancy between the XLF and DNA-PKcs DNA repair factors in V(D)J recombination and nonhomologous DNA end joining. Proc Natl Acad Sci U S A 110, 2234-2239 (2013).
11. V. Abramowski et al., PAXX and Xlf interplay revealed by impaired CNS development and immunodeficiency of double KO mice. Cell Death Differ 25, 444-452 (2018).
12. G. Balmus et al., Synthetic lethality between PAXX and XLF in mammalian development. Genes Dev 30, 2152-2157 (2016).
13. S. Castaneda-Zegarra, M. Xing, R. Gago-Fuentes, S. Saeterstad, V. Oksenych, Synthetic lethality between DNA repair factors Xlf and Paxx is rescued by inactivation of Trp53. DNA Repair (Amst) 73, 164-169 (2019).
14. C. Lescale et al., Specific Roles of XRCC4 Paralogs PAXX and XLF during V(D)J Recombination. Cell Rep 16, 2967-2979 (2016).
15. X. Liu, Z. Shao, W. Jiang, B. J. Lee, S. Zha, PAXX promotes KU accumulation at DNA breaks and is essential for end-joining in XLF-deficient mice. Nat Commun 8, 13816 (2017).
16. P. J. Hung et al., Deficiency of XLF and PAXX prevents DNA double-strand break repair by non-homologous end joining in lymphocytes. Cell Cycle 16, 286-295 (2017).
17. V. Kumar, F. W. Alt, R. L. Frock, PAXX and XLF DNA repair factors are functionally redundant in joining DNA breaks in a G1-arrested progenitor B-cell line. Proc Natl Acad Sci U S A 113, 10619-10624 (2016).
18. S. Castaneda-Zegarra et al., Leaky severe combined immunodeficiency in mice lacking non-homologous end joining factors XLF and MRI. Aging (Albany NY) 12, 23578-23597 (2020).
19. P. J. Hung et al., MRI Is a DNA Damage Response Adaptor during Classical Non-homologous End Joining. Mol Cell 71, 332-342 e338 (2018).
20. S. Zha et al., ATM damage response and XLF repair factor are functionally redundant in joining DNA breaks. Nature 469, 250-254 (2011).
21. C. Beck, S. Castaneda-Zegarra, C. Huse, M. Xing, V. Oksenyych, Mediator of DNA Damage Checkpoint Protein 1 Facilitates V(D)J Recombination in Cells Lacking DNA Repair Factor XLF. Biomolecules 10 (2019).
22. B. R. Chen et al., The RNF8 and RNF168 Ubiquitin Ligases Regulate Pro- and Anti-Resection Activities at Broken DNA Ends During Non-Homologous End Joining. DNA Repair (Amst) 108, 103217 (2021).
23. X. Liu et al., Overlapping functions between XLF repair protein and 53BP1 DNA damage response factor in end joining and lymphocyte development. Proc Natl Acad Sci U S A 109, 3903-3908 (2012).
24. V. Oksenyych et al., Functional redundancy between repair factor XLF and damage response mediator 53BP1 in V(D)J recombination and DNA repair. Proc Natl Acad Sci U S A 109, 2455-2460 (2012).
25. W. Xu et al., Loss of Gcn5l2 leads to increased apoptosis and mesodermal defects during mouse development. Nat Genet 26, 229-232 (2000).
26. R. C. Rickert, J. Roes, K. Rajewsky, B lymphocyte-specific, Cre-mediated mutagenesis in mice. Nucleic Acids Res 25, 1317-1318 (1997).
27. S. Srinivas et al., Cre reporter strains produced by targeted insertion of EYFP and ECFP into the ROSA26 locus. BMC Dev Biol 1, 4 (2001).
28. M. Muramatsu et al., Class switch recombination and hypermutation require activation-induced cytidine deaminase (AID), a potential RNA editing enzyme. Cell 102, 553-563 (2000).
29. A. Dewan et al., Robust DNA repair in PAXX-deficient mammalian cells. FEBS Open Bio 8, 442-448 (2018).
30. R. Gago-Fuentes et al., Normal development of mice lacking PAXX, the parologue of XRCC4 and XLF. FEBS Open Bio 8, 426-434 (2018).
31. S. Castaneda-Zegarra et al., Generation of a Mouse Model Lacking the Non-Homologous End-Joining Factor Mri/Cyren. Biomolecules 9 (2019).
32. L. M. Starnes et al., A PTIP-PA1 subcomplex promotes transcription for IgH class switching independently from the associated MLL3/MLL4 methyltransferase complex. Genes Dev 30, 149-163 (2016).
33. D. Su et al., PTIP chromatin regulator controls development and activation of B cell subsets to license humoral immunity in mice. Proc Natl Acad Sci U S A 114, E9328-E9337 (2017).
34. C. Boboila et al., Robust chromosomal DNA repair via alternative end-joining in the absence of X-ray repair cross-complementing protein 1 (XRCC1). Proc Natl Acad Sci U S A 109, 2473-2478 (2012).
35. R. Gago-Fuentes, V. Oksenyych, Non-Homologous End Joining Factors XLF, PAXX and DNA-PKcs Maintain the Neural Stem and Progenitor Cell Population. Biomolecules 11 (2020).
36. M. Nakamura et al., High frequency class switching of an IgM+ B lymphoma clone CH12F3 to IgA+ cells. Int Immunol 8, 193-201 (1996).
