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Generation of a Mouse Model Lacking the Non-Homologous End-Joining Factor Mri/Cyren

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Abstract: Classical non-homologous end joining (NHEJ) is a molecular pathway that detects,
processes, and ligates DNA double-strand breaks (DSBs) throughout the cell cycle. Mutations in
several NHEJ genes result in neurological abnormalities and immunodeficiency both in humans and
mice. The NHEJ pathway is required for V(D)J recombination in developing B and T lymphocytes,
and for class switch recombination in mature B cells. The Ku heterodimer formed by Ku70 and Ku80
recognizes DSBs and facilitates the recruitment of accessory factors (e.g., DNA-PKcs, Artemis, Paxx
and Mri/Cyren) and downstream core factor subunits X-ray repair cross-complementing group 4
(XRCC4), XRCC4-like factor (XLF), and DNA ligase 4 (Lig4). Accessory factors might be dispensable
for the process, depending on the genetic background and DNA lesion type. To determine the
physiological role of Mri in DNA repair and development, we introduced a frame-shift mutation in
the Mri gene in mice. We then analyzed the development of Mri−/− mice as well as wild type
and immunodeficient controls. Mice lacking Mri possessed reduced levels of class switch
recombination in B lymphocytes and slow proliferation of neuronal progenitors when compared to
wild type littermates. Human cell lines lacking Mri were as sensitive to DSBs as the wild type
controls. Overall, we concluded that Mri/Cyren is largely dispensable for DNA repair and mouse
development.

Keywords: NHEJ; double-strand breaks; mouse model; lymphocyte; neurodevelopment

1. Introduction

Non-homologous end-joining (NHEJ) is a molecular pathway that recognizes, processes, and
repairs DNA double-strand breaks (DSBs) throughout the cell cycle [1]. Core NHEJ factors Ku70 and
Ku80 form heterodimer (Ku) that is rapidly associated with the DSB sites facilitating recruitment of
downstream factors including core x-ray cross-complementing 4 (XRCC4) and DNA ligase 4 (Lig4).
XRCC4-like factor (XLF) is also a core factor that binds XRCC4 and stimulates Lig4-dependent DNA ligation. A number of accessory NHEJ factors are required for specific DNA end processing and DNA complex stabilization, in other words, DNA-dependent protein kinase, catalytic subunit (DNA-PKcs), nuclease Artemis and structural components, a parologue of XRCC4 and XLF (PAXX), and modulator of retroviral infection (Mri) [2,3]. Mice lacking Ku70, Ku80, DNA-PKcs, or Artemis possess severe combined immunodeficient phenotype (SCID), while inactivation of both alleles of the Xlf gene results in 2–3-fold reduced B and T cell counts [1,4–7]. Mice lacking PAXX or Mri possess no or very modest phenotype due to functional redundancy with XLF [8–12]. In contrast, mice lacking either XRCC4 or Lig4 demonstrate p53- and Ku-dependent embryonic lethality, which correlates with massive neuronal apoptosis in the central nervous system [1,13–17]. Combined inactivation of Xlf and Dna-pkcs results in p53- and Ku70-dependent perinatal lethality in mice [10,18,19]. Moreover, deficiency or haploinsufficiency for Trp53 rescues synthetic lethality between Xlf and Paxx [10]. XLF is also functionally redundant in mouse development with Mri [20], recombination activating gene 2, RAG2 [21], and a number of DNA damage response (DDR) factors including Ataxia telangiectasia mutated (ATM) [6], histone H2AX [6,22], mediator of DNA damage checkpoint protein 1 (MDC1) [10], and p53-binding factor (53BP1) [7,23].

Development of B and T lymphocytes depends on programmed DSBs induced by RAG during the V(D)J recombination and NHEJ pathway, which is used for error-prone DNA repair [1]. Moreover, mature B cells replace constant regions of immunoglobulins during the somatic recombination process known as class switch recombination (CSR), when DSBs are initiated by activation-induced cytidine deaminase (AID) and Uridine-N-glycosylase (UNG), and NHEJ is used for DNA repair [1,24,25]. Furthermore, the NHEJ process is required for neurodevelopment by preventing neuronal apoptosis [1,26].

Mri was initially described as an open reading frame at human chromosome 7 (C7orf49), a factor reversing the resistance to retroviral infection in cell lines [27]. Mri was found to enhance NHEJ [28] and possess an N-terminal Ku-binding motif (KBM) [29]. Later, Mri/Cyren was suggested to inhibit NHEJ at telomeres during the S and G2 phases of the cell cycle [30], and finally confirmed to be a bona fide NHEJ factor, which is functionally redundant with XLF in mouse development including the V(D)J recombination and development of the central nervous system [20]. However, it was not clear whether XLF and Mri functionally overlap during the early stages of neurodevelopment (e.g., supporting proliferation and self-renewal of neuronal stem cells). Moreover, due to the lack of a viable mouse model deficient for both XLF and Mri, the impact of Mri on B and T lymphocyte development in vivo is not fully understood.

Here, we introduced a frame-shift mutation to exon 2 of the murine Mri gene. By interbreeding heterozygous parents, we obtained Mri+/−, Mri+−, and Mri+/+ mice at nearly expected ratios. Mri−/− deficient mice possessed normal body size and number of B and T lymphocytes; however, we detected that stimulated primary mature Mri−/− B cells had reduced levels of IgG1, and Mri+− neurospheres showed a reduced proliferation rate when compared to the Mri+/+ controls.

2. Materials and Methods

2.1. Mouse Models

All experiments involving mice were performed according to the protocols approved by the Animal Resources Care Facility of Norwegian University of Science and Technology (NTNU, Trondheim, Norway). Ung−/− mice were described previously [31]. Mri+− mice were generated on request and described here for the first time.

2.2. Generation of Mri+− Mice

MRI-deficient (M−/−) mice were generated through a CRISPR/Cas9 gene-editing approach in 2017 by Horizon Discovery (Saint Louis, MO, USA) upon request from the Oksenych group (IKOM, Faculty of Medicine and Health Science, NTNU, Trondheim, Norway). Single-guide RNA (sgRNA) GGG CTG TCA TCC AAG AGG GGA GG was designed to target exon 2 of the Mri gene in C57BL/6...
mice. The 14 bp deletion resulted in a premature stop codon (Figure 1A). Cas9 and sgRNAs were injected into single-cell fertilized embryos, which were then transferred back into pseudopregnant females for gestation. Live-born pups were screened for indel mutation by DNA sequencing. Homozygous pups were used for back-crossing with wild type C57BL/6 mice. Heterozygous $Mri^{+/-}$ mice were obtained from Horizon Discovery.

2.3. Mouse Genotyping

Two polymerase chain reactions (PCRs) were designed to determine the mouse genotypes. The first PCR was performed using TCAGGTCTGCCCTACACTGA and GTGGTGGTGCTTCTCTTGTGA primers, detecting both wild type (428 bp) and null (414 bp) alleles (Figure 1B). The second PCR performed with TCAGGTCTGCCCTACACTGA and AGAGGGGAGGACC primers was used to validate the presence of the WT allele (234 bp, Figure 1B). The PCRs were performed using 50 ng of genomic DNA extracted from murine tissues (e.g., ears, tails), in a final reaction volume of 25 μL, using the Taq 2x Master Mix Kit (New England Biolabs® Inc., Ipswich, MA, USA; #M0270L). A 2.5% agarose gel was used to separate 428 bp and 414 bp PCR products during 18 h at 4 °C, 90 V; and 0.7% agarose gel was used to detect the 234 bp PCR product (75 min, room temperature, 124 V). Genomic DNA isolated from the $Mri^{+/-}$ and $Mri^{-/-}$ cells as well as samples with no genomic DNA were used as the PCR controls (Figure 1B).

2.4. Fluorescence-Activated Cell Sorting, Splenocyte, and Thymocyte Count

Fluorescence-activated cell sorting (FACS) analysis was performed as previously described [11,32]. Briefly, spleens and thymi were isolated from 2-month-old mice, and splenocytes and thymocytes were counted using Countess™ Automated Cell Counter (Invitrogen, Carlsbad, CA, United States); the cell suspension was spun down and diluted with PBS to obtain a final cell concentration of 2.5 × 10⁶/mL. The samples of 2.5 × 10⁶ splenocytes or thymocytes were blocked for 15 min at room temperature with Mouse BD fragment crystallizable (Fc) Block™ (1:50 dilution) (BD Biosciences, Franklin Lakes, NJ, USA; #553142). The cells were then incubated with fluorochrome-conjugated antibodies (see below) and sorted.

2.5. Class Switch Recombination

Class switch recombination (CSR) from IgM to IgG1 was performed as previously described [11]. Naïve B lymphocytes were purified from spleens of 2-month-old mice using EasySep™ mouse B cell enrichment kit (STEMCELL Technology, Vancouver, Canada; #19854), according to the manufacturers’ instructions. For each CSR assay, 2 × 10⁴ cells/200 μL were used in duplicates. The cells were stimulated with LPS (lipopolysaccharides, 40 μg/mL; Sigma Aldrich, St. Louis, MO, USA; #437627-5MG) and IL-4 (Interleukin 4, 20 ng/mL; PeproTech, Stockholm, Sweden; #214-14) for 96 h. Then, the cells were blocked with Fc receptor antibody (2.4G2) and normal mouse serum (Invitrogen, Carlsbad, CA, USA; #10410). The cells were washed in PermWash™ (BD Biosciences, NJ, USA; #554723). Intracellular staining was done using fluorescently tagged anti-mouse antibodies (IgG1-APC) (BioLegend, San Diego, CA, USA; #406610) and the succeeding wash was performed in PermWash. The cells were resuspended in 300 μL of CellFix (BD Biosciences, NJ, USA; #340181). Viable CD19+ B lymphocytes were analyzed for IgG1 expression using FlowJo® (Ashland, Oregon, USA) version 7.6 for Windows.

2.6. Double Strand Break Sensitivity Assay

The DSBs sensitivity assay was performed as previously described [10,32,33]. Human nearly-haploid HAP1 cells were generated by the Horizon Discovery Group (Waterbeach, Cambridge, UK, #HZGHC005061c001 and #HZGHC005061c004) and are commercially available. HAP1 cells were cultured according to the manufacturer’s instructions. Doxorubicin (Selleckchem, Houston, TX, USA; #S1208), bleomycin (Selleckchem; #S1214), and etoposide (Sigma-Aldrich, St. Louis, MS, USA; #E1383) were used to induce DSBs, and PrestoBlue™ Cell Viability Reagent (Thermo Fisher,
Waltham, MA, USA; #A13262) was used to estimate cellular metabolism levels. Briefly, 2000 cells per well were seeded into 96-well plates in 100 μL of Iscove Modified Dulbecco Media (IMDM) medium (day 0). On day 1, 50 μL of the medium was replaced with 50 μL of fresh medium containing doxorubicin, bleomycin, or etoposide, when indicated. Each experimental condition was performed in triplicates. On day 4, 11 μL of 10× PrestoBlue reagent was added to the wells and incubated for 30 min at 37 °C. The cellular viability was estimated according to manufacturer’s instructions, using the excitation/emission wavelengths set at 544/590 nm.

2.7. Brain Isolation and Neural Stem Progenitor Cell Culture

The brain was isolated from postnatal day 1 mouse after the cerebellum was removed. The isolated brain was mechanically disrupted in the proliferation medium consisting of Dulbecco Modified Eagle Medium, Nutrient Mixture F12 (DMEM/F12; Thermo Fisher, Waltham, MA, USA; #11330-057), supplemented with penicillin/streptomycin (Thermo Fisher, Waltham, MA, USA; #15140122), B27 without vitamin A (Thermo Fischer Scientific, Waltham, MA, USA; #12587001), EGF (10 ng/mL; PeproTech, Stockholm, Sweden; #AF-100-15), and bFGF (20 ng/mL; PeproTech; #100-18B). Neural stem progenitor cells (NSPC) form free-floating globular structures referred to as neurospheres. The neurospheres were formed during incubation at 37 °C, 5% CO₂ and 95% humidity in order to perform the proliferation and self-renewal assay [34].

2.8. Neural Stem Progenitor Cell Proliferation and Self-Renewal Assays

Early passage NSPCs (P3–P10) were used throughout all of the NSPC experiments. A PrestoBlue™ Cell Viability Assay was used to investigate the neurosphere proliferation rates, following the manufacturer’s instructions during each incubation on days 1 to 7. The capacity of neural stem cells to maintain their multipotency ex vivo was assessed by determining the number and two-dimensional size of neurospheres [34]. Single NSPCs were plated onto 6-well suspension plates in the proliferation medium on day 0. During days 8 and 10 in culture, images of the entire wells were captured using an EVOS microscope. Only areas between 50 and 1500 pixels were included in the analyses.

2.9. Antibodies

The following antibodies were used for FACS. Rat anti-mouse anti-CD16/CD32 (Fc Block, BD Biosciences, San Jose, CA, USA; #553141, 1:50); anti-CD4-PE-Cy7 (Thermo Scientific, Waltham, MA, USA, #25-0042-81, 1:100); anti-CD8-PE-Cy5 (BD Biosciences, San Jose, CA, USA, #553034, 1:100); anti-CD19-PE-Cy7 (Biolegends, San Diego, CA, USA, #115520, 1:100); and hamster anti-mouse anti-CD3-APC (Biolegends, USA, #100312, 1:100).

3. Results

3.1. Generation of Mri−/− Mice

To investigate the impact of Mrı on mouse development, we generated a mouse model with 14 bp frame-shift deletion in Mrı exon 2 on a C57BL/6 background (Figure 1A). Purified sgRNA and Cas9 RNA were introduced to fertilized oocytes, resulting in complete inactivation of the Mrı gene. Mrı status (WT, wild type, +/+; heterozygous, +/-; and null, −/−) was confirmed for every experiment by PCR screening (Figure 1B). Mrı−/+, Mrı+/-, and Mrı−/− mice were born from Mrı−/+ parents at ratios close to 1:2:1 (Figure 1C). Thirty-day old Mrı−/+ mice possessed an average body weight of 15.0 g, which was slightly lower, but not significantly different from the Mrı−/+ controls, with a bodyweight of 17.5 g, on average (Figure 1D). The lifespan of Mrı−/+ and Mrı−/+ mice was monitored for up to 12 months, according to the local regulations. During this time frame, both Mrı−/+ and Mrı−/+ mice were fertile and had no cancer incidence, similar to the Mrı−/+ controls.
Figure 1. Generation of Modulator of retrovirus infection\textsuperscript{+} (Mri\textsuperscript{+}) mice. (A) Top. Schematic diagram of murine Mri locus indicating the frame-shift mutation in the exon 2, induced by the single guide RNA (sgRNA) and resulting in a 14 bp deletion. (Bottom) Resulting Mri\textsuperscript{-} locus lacking part of the exon 2. (B) Top. Polymerase chain reaction (PCR)-based genotyping strategy reveals the Mri WT allele (428 bp) and Mri null allele (414 bp). (Bottom) WT gene validation PCR revealed the Mri wild type allele (234 bp). (C) Analyses of 140 pups born from Mri\textsuperscript{+/-} parents revealed expected genetic distribution of Mri\textsuperscript{+/-} (29), Mri\textsuperscript{+/-} (75), and Mri\textsuperscript{-/-} (36) mice, which is close to the Mendelian distribution 1:2:1. (D) Body weight of six to eight week old Mri\textsuperscript{+/-} mice (n = 6) was not distinguishable from Mri\textsuperscript{-/-} mice (n = 7), \( p = 0.4242 \). (E) The weight of spleens isolated from Mri\textsuperscript{+/-} (n = 8) and Mri\textsuperscript{-/-} mice (n = 11) was not significantly different, \( p = 0.8551 \). Spleen size in immunodeficient Dna-pkcs\textsuperscript{-/-} mice (n = 10) was reduced...
when compared to the Mri\(^{+}\) and Mri\(^{-}\) mice, \(p < 0.0001\). (F) Splenocyte count was not affected in the Mri\(^{-}\) mice (\(n = 11\)) when compared to the Mri\(^{+}\) (\(n = 10\), \(p = 0.7713\). A number of splenocytes in immunodeficient Dna-pkcs\(^{-}\) mice (\(n = 6\)) was significantly reduced when compared to Mri\(^{+}\) and Mri\(^{-}\) mice, \(p < 0.0001\). (G) The weight of thymus from Mri\(^{+}\) (\(n = 11\)) and Mri\(^{-}\) (\(n = 11\)) mice was similar, \(p = 0.6796\). Thymus size in immunodeficient Dna-pkcs\(^{-}\) mice (\(n = 7\)) was significantly reduced when compared to Mri\(^{+}\) and Mri\(^{-}\) mice, \(p < 0.0001\). (H) The thymocyte count was nearly identical in Mri\(^{+}\) (\(n = 8\)) and Mri\(^{-}\) (\(n = 6\)) mice, \(p = 0.5285\). A number of thymocytes in immunodeficient Dna-pkcs\(^{-}\) mice (\(n = 6\)) was significantly reduced when compared to Mri\(^{+}\) and Mri\(^{-}\) mice, \(p < 0.0001\).

### 3.2. Mri\(^{-}\) Mice Develop Normal Spleens and Thymi

The NHEJ is required for V(D)J recombination in developing B and T lymphocytes, and for CSR in mature B cells [1]. To determine specific functions of Mri in B and T cell development, we first analyzed spleens and thymi isolated from Mri-deficient and WT mice. The average weights of spleens (91 mg) and thymi (69 mg) as well as the average count of splenocytes (121 million) and thymocytes (173 million) was not affected in Mri\(^{-}\) mice when compared to Mri\(^{+}\) controls (90 mg; 71 mg; 118 million; 186 million, respectively). These numbers were significantly different from the immunodeficient controls, Dna-pkcs\(^{-}\) mice (23 mg; 10 mg; six million; five million, respectively) (Figure 1E–H). Moreover, the proportions of CD19\(^{+}\) B cells in spleens of six-to eight-week old Mri\(^{-}\) mice were on average 60%, which was similar to the proportion of CD19\(^{+}\) in Mri\(^{+}\) mice (55%, \(p = 0.0668\), and significantly higher than the background levels detected in immunodeficient Dna-pkcs\(^{-}\) controls (\(p < 0.0001\); Figure 2A). The average proportion of CD3\(^{+}\) T splenocytes in Mri\(^{-}\) (21%) was also similar to the one observed in the Mri\(^{+}\) controls (22%, \(p = 0.8228\), and higher than in the Dna-pkcs\(^{-}\) controls (1%, \(p < 0.0001\); Figure 2A). Mri\(^{+}\) and Mri\(^{-}\) mice had similar proportions of CD4\(^{+}\) T cells (\(p = 0.8876\)) and CD8\(^{+}\) T cells (\(p = 0.7026\)) in the spleens, while proportions of CD4\(^{+}\) and CD8\(^{+}\) T splenocytes in the Dna-pkcs\(^{-}\) controls were 4–5-fold reduced (\(p < 0.0001\), Figure 2B). In the thymi of six- to eight-week old Mri\(^{+}\) and Mri\(^{-}\) mice, the proportions of CD4\(^{+}\), CD8\(^{+}\), and CD4\(^{+}\)CD8\(^{+}\) T cells were similar (\(p > 0.5589\), while only background levels were detected in the Dna-pkcs\(^{-}\) controls (\(p < 0.0001\), Figure 2C).

### 3.3. Class Switch Recombination to IgG1 Is Reduced in Mri\(^{-}\) Mice

To determine whether Mri deficiency affects CSR, we isolated B cells from the spleens of Mri\(^{+}\) and Mri\(^{-}\) mice and stimulated the cells with LPS and IL-4. After 96 h, we detected that average IgG1 levels were 33% in Mri\(^{-}\) mice, which was significantly lower (\(p = 0.0031\)) than in the Mri\(^{+}\) controls (average 39%; Figure 2D). B lymphocytes isolated from Ung\(^{-}\) mice were used as the negative control and possessed on average only 2% of IgG1 at the end of the experiment (96 h), which was lower than in Mri\(^{+}\) or Mri\(^{-}\) mice (\(p < 0.0001\)).
Figure 2. Lymphocyte development in Mri\(^{+}\) mice. (A) Proportions of T (CD3\(^{+}\)) and B (CD19\(^{+}\)) cells in the spleens of Mri\(^{+}\) (n = 6), Mri\(^{−}\) (n = 3), and Dna-pkcs\(^{−}\) (n = 4) mice. Proportions of T and B cells were similar in Mri\(^{+}\) and Mri\(^{−}\) mice, \(p > 0.0667\), and were only background levels in immunodeficient Dna-pkcs\(^{−}\) mice, \(p < 0.0001\). (B) Proportions of CD4\(^{+}\) and CD8\(^{+}\) T splenocytes in Mri\(^{+}\) (n = 6), Mri\(^{−}\) (n = 3), and Dna-pkcs\(^{−}\) (n = 4) mice. Mri\(^{−}\) mice possessed similar proportions of CD4\(^{+}\) T helper and CD8\(^{+}\) T cytotoxic cells when compared to Mri\(^{+}\) mice, \(p = 0.8876\) and \(p = 0.7026\), respectively. Only background levels of CD4\(^{+}\) and CD8\(^{+}\) T cells are present in immunodeficient Dna-pkcs\(^{−}\) spleens, \(p < 0.0001\). (C) Proportions of CD4\(^{+}\), CD8\(^{+}\), and CD4\(^{+}\)CD8\(^{+}\) thymocytes in Mri\(^{+}\) (n = 6), Mri\(^{−}\) (n = 3), and Dna-pkcs\(^{−}\) (n = 4) mice. Proportions of T cell types in Mri\(^{−}\) mice were similar to the ones detected in Mri\(^{+}\) mice, \(p > 0.5589\), and higher than in Dna-pkcs\(^{−}\) mice, \(p < 0.0001\). (D) CSR to IgG1 in primary B splenocytes isolated from the Mri\(^{+}\) mice (n = 4) was reduced when compared to the cells from the Mri\(^{+}\) mice (n = 3), \(p = 0.0032\). CSR to IgG1 was significantly reduced in the Ung\(^{−}\) B cells (n = 3) when compared to the Mri\(^{+}\) and Mri\(^{−}\), \(p < 0.0001\).
3.4. Lack of Mri Results in the Reduced Proliferation Rate of Neuronal Stem Progenitor Cells

Previous studies have shown that single knockout of NHEJ DNA repair genes (e.g., Xrcc4, Lig4, Ku70) results in impaired nervous system development in mice [1,13,14]. To determine the impact of Mri on the developing nervous system, we used NSPC isolated from Mri+/+ and Mri−/− mice at postnatal day 1. We performed four independent experiments using two cell lines from two mice of each genotype. The average proliferation rate of Mri−/− neurospheres was approximately 35% lower than that in the WT controls, p = 0.0043 (Figure 3B).

Figure 3. Characterization of neurogenesis in Mri−/− mice. For each experiment, four independent cell lines isolated from two mice were used (n = 8). (A) Neurosphere isolation diagram from Mri+/+ and Mri−/− mice at postnatal day 1. (B) Neurosphere proliferation isolated from the Mri−/− mice was reduced when compared to the Mri+/+ mice, p = 0.0043. (C) Number of neurospheres formed in cell culture for eight days. Mri+/+ and Mri−/− neurospheres possessed similar self-renewal capacity, p = 0.7254. (D) Neurosphere size isolated from Mri+/+ and Mri−/− mice were similar, p = 0.4915. The surface of neurospheres, pxl. Areas between 50 and 1500 pixels were included in the analyses. Four independent experiments using two cell lines of each genotype were performed in all experiments (A–C). p values were calculated using the unpaired t-test. The horizontal bars represent the average.

3.5. Normal Self-Renewal Capacity of Mri-Deficient Neuronal Stem Progenitor Cells

To analyze the capacity of NSPCs to maintain the features of stem cells throughout cell divisions and numerous propagations (self-renewal capacity), we counted the number of neurospheres formed in cell culture. In four independent experiments, we plated 10,000 single NSPCs and cultured for eight days. In total, we counted 5123 neurospheres that originated from Mri+/+, and 4608 from Mri−/− mice. On average, there were 256 neurospheres in each of the 20 Mri++ samples analyzed, and 230
neurospheres in each of the 20 $Mri^+$ samples ($p = 0.7254$, n.s., Figure 3C). In addition, images of the neurospheres were collected and the surface was calculated using ImageJ software. Inactivation of $Mri$ did not affect the average diameter of neurospheres, which was 461 px$^2$ on average in $Mri^{+/+}$ and 427 px$^2$ in $Mri^{-/-}$ neurospheres, $p = 0.4915$ (Figure 3D). We concluded that $Mri$ is dispensable for the self-renewal capacity of NSPC.

3.6. Human HAP1 Cells Lacking $Mri$ Possess Normal Levels of Sensitivity to DNA Double-Strand Breaks

Deficiency for XRCC4, LIG4, XLF, or DNA-PKcs results in hypersensitivity to DSBs in human HAP1 cells [10,32,33]. To determine the effect of $Mri$ on DSB sensitivity, we obtained two independent clones of $MRI^{-/-}$ HAP1 cells as well as WT and $XRCC4^{-/-}$ controls. We exposed the HAP1 cells to DSB-inducing agents bleomycin (0 to 0.4 μM), doxorubicin (0 to 4 nM), and etoposide (0 to 160 nM), and evaluated the cell viability four days later (Figure 4). We observed no hypersensitivity of HAP1 cells lacking $Mri$ when compared to WT controls. However, cells lacking $XRCC4$ were hypersensitive to all indicated compounds, bleomycin, doxorubicin, and etoposide ($p < 0.0001$, Figure 4).

![Figure 4](image-url)

**Figure 4.** Sensitivity to DSBs in $Mri$-deficient HAP1 cells. Sensitization of WT, two independent $Mri$-deficient clones, $MRI^{+/-}$ and $MRI^{-/-}$, and $XRCC4^{+/-}$ HAP1 cells to bleomycin (A), doxorubicin (B), and etoposide (C) at indicated concentrations. Results are from the mean (SD) of three repeats. Cell viability (%) represents the relative proportion of the fluorescence normalized to untreated cells. Comparisons between every two groups were made using one-way ANOVA, GraphPad Prism 8. WT, $MRI^{+/-}$, and $MRI^{-/-}$ vs. $XRCC4^{+/-}$: $p < 0.0001$ (****); WT vs. $MRI^{+/-}$: $p = 0.9983$ (n.s); WT vs. $MRI^{-/-}$: $p = 0.1295$ (n.s); $MRI^{+/-}$ vs. $MRI^{-/-}$, $p = 0.1791$ (n.s).

4. Discussion

We have generated a new knockout mouse model with 14 bp deletion in exon 2 of the $Mri$ gene, $Mri^+$. While we were characterizing our mouse model, another group reported an independently-generated $Mri$-deficient mouse [20], which possessed a similar phenotype. Thus, our observations are confirmatory to the findings observed by the Sleckman group [20].

The mice lacking $Mri$ were live-born at expected ratios and demonstrated normal growth and development of lymphoid organs. $Mri^+/-$, $Mri^-/-$, and $Mri^+/+$ mice possessed similar sizes of spleens and thymi, a similar number of splenocytes and thymocytes, and proportions of B and T cells (Figure 1). Similar to $Mri$-deficient mice, $Paxx^+/-$ mice did not have a detectable phenotype [8–12]. However, inactivation of other NHEJ factors resulted in a reduced number of $B$ and $T$ cells (Figures 1, 2), and block in $B$ and $T$ cell development (e.g., $Artemis^+/-$ [35], $Dna-pkcs^+/-$ [36], $Ku70^-/-$ [37], $Ku80^-/-$ [38]; or even embryonic lethality in $Xrcc4^-/-$ [39] and $Lig4^-/-$ [40]).

The CSR to IgG1 was reduced in primary $B$ cells isolated from $Mri^-/-$ mice when compared to WT controls (Figure 2), which suggests that $Mri$ is involved in specific DNA repair-mediated event. Furthermore, we isolated neuronal stem progenitor cells from $Mri^-/-$ brains and found that these cells proliferate slower when compared to $Mri^+/-$ controls. Reduced proliferation rates of $Mri$-deficient neuronal stem progenitor cells could be explained, as one option, by lower expression or lack of factors functionally redundant with $Mri$ in these cell types. Future studies would be required to address this option. Moreover, future studies may address questions such as neurological defects and...
cognitive functions in mice lacking Mri as well as whether the Mri-deficient mice are prone to infections. In addition, we found that human nearly haploid HAPI cell lines lacking Mri possessed no proliferation defect or hypersensitivity to DNA damaging agents such as etoposide, doxorubicin, and bleomycin (Figure 4). Previously, it was reported that murine embryonic fibroblasts (MEF) lacking Mri were hypersensitive to ionizing radiation when compared to WT controls, although the sensitivity was less obvious than in XLF-deficient MEFs [20]. The discrepancy between our and previously published data could be due to the usage of different cell types, the difference between species as well as distinct ways to induce DNA damages (e.g., chemicals vs. irradiation). Further studies involving various cell type models originated from different species, and using various DNA damaging strategies would deepen our understanding of the specific functions of Mri in DNA repair in mammalian cells. Overall, we concluded that the lack of Mri has a rather minor effect on murine and human cells.

Combined inactivation of Mri and Xlf [20], however, revealed an important role of Mri in mouse development, which was overlooked due to its functional redundancy with XLF. Synthetic lethality between Mri and Xlf complicated studies of genetic interaction between these factors in vivo. There are several potential ways to overcome this challenge. One option is to use conditional knockouts of Xlf or Mri genes. Moreover, there might be another genetic-based approach. Inactivation of one or two alleles of Trp53 partially rescued synthetic lethality between Xlf and Dna- pkcs [10, 18, 19] and Xlf and Paxx [10]. In addition, deficiency or haploinsufficiency for Trp53 rescued embryonic lethality of Lig4−/− and Xrcc4−/− mice [13, 14]. Inactivation of the Atm gene rescued embryonic lethality of Lig4−/− mice [41]. Finally, inactivation of both alleles of Ku80 rescued embryonic lethality of Lig4−/− mice [17], and inactivation of Ku70 rescued synthetic lethality between Xlf and Dna-pkcs [19]. Based on these data, one can speculate that inactivation of Trp53, Atm, Ku70, or Ku80 will rescue synthetic lethality between Xlf and Mri. Moreover, given the critical roles of Ku70 and Ku80 in the initiation of classical NHEJ, one could propose that mice lacking all known NHEJ factors (e.g., Ku70−/− Ku80−/− Dna-pkcs−/− Artemis−/− Xlf−/− Paxx−/− Mri−/− Xrcc4−/− Lig4−/−) will be viable, indistinguishable from Ku-deficient mice, and serve as a suitable in vivo model to investigate alternative end-joining, A-EJ.

5. Conclusions

A new Mri-deficient mouse model was generated. Mri-deficient mice possessed normal body size and number of B and T lymphocytes; however, Mri is required for an efficient class switch recombination process in mature B cells. Mri−/− neurospheres showed a reduced proliferation rate, but similar self-renewal capacity when compared to the Mri+/+ controls.

Author Contributions: V.O. designed and performed experiments, contributed key reagents, and analyzed the data. A.S. performed the CSR assay using primary B cells. M.X. performed the DSBs sensitivity assay using human HAPI cells. S.C.-Z., C.H., Ø.R., Q.Z., A.A., J.W., and N.B.L. performed the lymphocyte analyses. Ø.R., W.W., and P.J. performed and analyzed the neurosphere-based experiments. R.G.-F. developed the Mri−/− genotyping strategy. M.B. and V.O. interpreted the results. The paper was written by Ø.R., S.C.-Z., and V.O.; all authors read and approved the manuscript.

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Abbreviations:

ATM  Ataxia-telangiectasia mutated  
CSR   Class switch recombination
DDR   DNA damage response
DNA-PKcs DNA-dependent protein kinase
DSBs  DNA double-strand breaks
GFAP  Glial fibrillary acid protein
HAP1  A near-haploid human cell line derived from KBM-7 cell line
IL-4  Interleukin 4
Lig4  DNA ligase IV
LPS   Lipopolysaccharides
Mri   Modulator of retroviral infection
NHEJ  Non-homologous end-joining
NSPC  Neuronal stem progenitor cells
PAXX  Paralogue of XRCC4 and XLF
PCR   Polymerase chain reaction
XLF   XRCC4-like factor
XRCC4  X-ray repair cross-complementing protein 4

References

1. Kumar, V.; Alt, F.W.; Oksenych, V. Functional overlaps between XLF and the ATM-dependent DNA double strand break response. DNA Repair (Amst) 2014, 16, 11–22, doi:10.1016/j.dnarep.2014.01.010.
2. Pannunzio, N.R.; Watanabe, G.; Lieber, M.R. Nonhomologous DNA end-joining for repair of DNA double-strand breaks. J. Biol. Chem. 2018, 293, 10512–10523, doi:10.1074/jbc.TM117.000374.
3. Scully, R.; Panday, A.; Elango, R.; Willis, N.A. DNA double-strand break repair-pathway choice in somatic mammalian cells. Nat. Rev. Mol. Cell Biol. 2019, 10.1038/s41580-019-0152-0, doi:10.1038/s41580-019-0152-0.
4. Li, G.; Alt, F.W.; Cheng, H.L.; Brush, J.W.; Goff, P.H.; Murphy, M.M.; Franco, S.; Zhang, Y.; Zha, S. Lymphocyte-specific compensation for XLF/cernunnos end-joining functions in V(D)J recombination. Mol. Cell 2008, 31, 631–640, doi:10.1016/j.molcel.2008.07.017.
5. Vera, G.; Rivera-Munoz, P.; Abramowski, V.; Malivert, L.; Lim, A.; Bole-Feyssot, C.; Martin, C.; Florkin, B.; Latour, S.; Revy, P.; et al. Cernunnos deficiency reduces thymocyte life span and alters the T cell repertoire in mice and humans. Mol. Cell Biol. 2013, 33, 701–711, doi:10.1128/MCB.01057-12.
6. Zha, S.; Guo, C.; Boboila, C.; Oksenych, V.; Cheng, H.L.; Zhang, Y.; Wesemann, D.R.; Yuen, G.; Patel, H.; Goff, P.H.; et al. ATM damage response and XLF repair factor are functionally redundant in joining DNA breaks. Nature 2011, 469, 250–254, doi:10.1038/nature09604.
7. Oksenych, V.; Alt, F.W.; Kumar, V.; Schwer, B.; Wesemann, D.R.; Hansen, E.; Patel, H.; Su, A.; Guo, C. Functional redundancy between repair factor XLF and damage response mediator 53BP1 in V(D)J recombination and DNA repair. Proc. Natl. Acad. Sci. 2012, 109, 2455–2460, doi:10.1073/pnas.1121458109.
8. Abramowski, V.; Etienne, O.; Elsaid, R.; Yang, J.; Berland, A.; Kermasson, L.; Roch, B.; Musilli, S.; Moussu, J.P.; Lipson-Ruffert, K.; et al. PAXX and Xlf interplay revealed by impaired CNS development and immunodeficiency of double KO mice. Cell Death Differ. 2018, 25, 444–452, doi:10.1038/cdd.2017.184.
9. Balmus, G.; Barros, A.C.; Wijnhoven, P.W.; Lescale, C.; Hasse, H.L.; Boroviak, K.; le Sage, C.; Doe, B.; Speak, A.O.; Galli, A., et al. Synthetic lethality between PAXX and XLF in mammalian development. Genes Dev. 2016, 30, 2152–2157, doi:10.1101/gad.290510.116.
10. Castaneda-Zegarra, S.; Xing, M.; Gago-Fuentes, R.; Saeterstad, S.; Oksenych, V. Synthetic lethality between DNA repair factors Xlf and Paxx is rescued by inactivation of Trp53. *DNA Repair (Amst)* **2019**, *73*, 164–169, doi:10.1016/j.dnarep.2018.12.002.

11. Gago-Fuentes, R.; Xing, M.; Saeterstad, S.; Sarno, A.; Dewan, A.; Beck, C.; Bradamante, S.; Bjoras, M.; Oksenych, V. Normal development of mice lacking PAXX, the parologue of XRCC4 and XLF. *FEBS Open Bio* **2018**, *8*, 426–434, doi:10.1002/2211-5463.12381.

12. Liu, X.; Shao, Z.; Jiang, W.; Lee, B.J.; Zha, S. PAXX promotes Ku accumulation at DNA breaks and is essential for end-joining in XLF-deficient mice. *Nat. Commun.* **2017**, *8*, 13816, doi:10.1038/ncomms13816.

13. Boboila, C.; Jankovic, M.; Yan, C.T.; Wang, J.H.; Wesemann, D.R.; Zhang, T.; Fazeli, A.; Feldman, L.; Nussenzweig, A.; Nussenzweig, M., et al. Alternative end-joining catalyzes robust IgH locus deletions and translocations in the combined absence of ligase 4 and Ku70. *Proc. Natl. Acad. Sci. USA* **2010**, *107*, 3034–3039, doi:10.1073/pnas.0915067107.

14. Boboila, C.; Yan, C.; Wesemann, D.R.; Jankovic, M.; Wang, J.H.; Manis, J.; Nussenzweig, A.; Nussenzweig, M.; Alt, F.W. Alternative end-joining catalyzes class switch recombination in the absence of both Ku70 and DNA ligase 4. *J. Exp. Med.* **2010**, *207*, 417–427, doi:10.1084/jem.20092449.

15. Karanjawala, Z.E.; Adachi, N.; Irvine, R.A.; Oh, E.K.; Shibata, D.; Schwarz, K.; Hsieh, C.L.; Lieber, M.R. The embryonic lethality in DNA ligase IV-deficient mice is rescued by deletion of Ku: Implications for unifying the heterogeneous phenotypes of NHEJ mutants. *DNA Repair (Amst)* **2002**, *1*, 1017–1026.

16. Oksenych, V.; Kumar, V.; Liu, X.; Guo, C.; Schwer, B.; Zha, S.; Alt, F.W. 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. USA* **2013**, *110*, 2234–2239, doi:10.1073/pnas.1222573110.

17. Xing, M.; Bjoras, M.; Daniel, J.A.; Alt, F.W.; Oksenych, V. Synthetic lethality between murine DNA repair factors XLF and DNA-PKcs is rescued by inactivation of Ku70. *DNA Repair (Amst)* **2017**, *57*, 133–138, doi:10.1016/j.dnarep.2017.07.008.

18. Hung, P.J.; Johnson, B.; Chen, B.R.; Byrum, A.K.; Bredemeyer, A.L.; Yewdell, W.T.; Johnson, T.E.; Lee, B.J.; Deivasigamani, S.; Hindi, I., et al. MRI Is a DNA Damage Response Adaptor during Classical Non-homologous End Joining. *Mol. Cell* **2018**, *71*, 332–342 e338, doi:10.1016/j.molcel.2018.06.018.

19. Lescate, C.; Abramowski, V.; Bedora-Faure, M.; Murignieux, V.; Vera, G.; Roth, D.B.; Revy, P.; de Villartay, J.P.; Deriano, L. RAG2 and XLF/Cernunnos interplay reveals a novel role for the RAG complex in DNA repair. *Nat. Commun.* **2016**, *7*, 10529, doi:10.1038/ncomms10529.

20. Chen, B.R.; Quinet, A.; Byrum, A.K.; Jackson, J.; Berti, M.; Thangavel, S.; Bredemeyer, A.L.; Hindi, I.; Mosammaparast, N.; Tyler, J.K., et al. MRI and H2AX function in series to promote replication fork stability. *J. Cell Biol.* **2019**, *218*, 2113–2123, doi:10.1083/jcb.201808134.

21. Liu, X.; Jiang, W.; Dubois, R.L.; Yamamoto, K.; Wolner, Z.; Zha, S. Overlapping functions between XLF repair protein and 53BP1 DNA damage response factor in end joining and lymphocyte development. *Proc. Natl. Acad. Sci. USA* **2012**, *109*, 3903–3908, doi:10.1073/pnas.1120160109.

22. Alt, F.W.; Zhang, Y.; Meng, F.L.; Guo, C.; Schwer, B. Mechanisms of programmed DNA lesions and genomic instability in the immune system. *Cell* **2013**, *152*, 417–429, doi:10.1016/j.cell.2013.01.007.

23. Yeap, L.S.; Meng, F.L. Cis- and trans-factors affecting AID targeting and mutagenic outcomes in antibody diversification. *Adv. Immunol.* **2019**, *141*, 51–103, doi:10.1016/bs.ai.2019.01.002.

24. Alt, F.W.; Schwer, B. DNA double-strand breaks as drivers of neural genomic change, function, and disease. *DNA Repair (Amst)* **2018**, *71*, 158–163, doi:10.1016/j.dnarep.2018.08.019.

25. Agarwal, S.; Harada, J.; Schreifels, J.; Lech, P.; Nikolai, B.; Yamaguchi, T.; Chanda, S.K.; Soma, N.V. Isolation, characterization, and genetic complementation of a cellular mutant resistant to retroviral infection. *Proc. Natl. Acad. Sci. USA* **2006**, *103*, 15933–15938, doi:10.1073/pnas.0602674103.

26. Slavoff, S.A.; Heo, J.; Budnik, B.A.; Hanakahi, L.A.; Saghatelian, A. A human short open reading frame (sORF)-encoded polypeptide that stimulates DNA end joining. *J. Biol. Chem.* **2014**, *289*, 10950–10957, doi:10.1074/jbc.C113.533968.
29. Grundy, G.J.; Rulten, S.L.; Arribas-Bosacoma, R.; Davidson, K.; Kozik, Z.; Oliver, A.W.; Pearl, L.H.; Caldecott, K.W. The Ku-binding motif is a conserved module for recruitment and stimulation of non-homologous end-joining proteins. Nat. Commun. 2016, 7, 11242, doi:10.1038/ncomms11242.

30. Arnoult, N.; Correia, A.; Ma, J.; Merlo, A.; Garcia-Gomez, S.; Maric, M.; Tognetti, M.; Benner, C.W.; Boulton, S.J.; Saghatelian, A., et al. Regulation of DNA repair pathway choice in S and G2 phases by the NHEJ inhibitor CYREN. Nature 2017, 549, 548–552, doi:10.1038/nature24023.

31. Nilsen, H.; Rosewell, I.; Robins, P.; Skjelbred, C.F.; Andersen, S.; Stupphaug, G.; Daly, G.; Krokan, H.E.; Lindahl, T.; Barnes, D.E. Uracil-DNA glycosylase (UNG)-deficient mice reveal a primary role of the enzyme during DNA replication. Mol. Cell 2000, 5, 1059–1065.

32. Dewan, A.; Xing, M.; Lundbaek, M.B.; Gago-Fuentes, R.; Beck, C.; Aas, P.A.; Liabakk, N.B.; Saeterstad, S.; Chau, K.T.P.; Kavli, B.M., et al. Robust DNA repair in PAXX-deficient mammalian cells. FEBS Open Bio. 2018, 8, 442–448, doi:10.1002/2211-5463.12380.

33. Xing, M.; Oksenyych, V. Genetic interaction between DNA repair factors PAXX, XLF, XRCC4 and DNA-PKcs in human cells. FEBS Open Bio. 2019, 10.1002/2211-5463.12681, doi:10.1002/2211-5463.12681.

34. Wang, W.; Osenbroch, F.; Skinnes, R.; Esbensen, Y.; Bjoras, M.; Eide, L. Mitochondrial DNA integrity is essential for mitochondrial maturation during differentiation of neural stem cells. Stem Cells 2010, 28, 2195–2204, doi:10.1002/stem.542.

35. Rooney, S.; Sekiguchi, J.; Zhu, C.; Cheng, H.L.; Manis, J.; Whitlow, S.; DeVido, J.; Foy, D.; Chaudhuri, J.; Lombard, D., et al. Leaky Scid phenotype associated with defective V(D)J coding end processing in Artemis-deficient mice. Mol. Cell 2002, 10, 1379–1390, doi:10.1016/s1097-2765(02)00755-4.

36. Gao, Y.; Chaudhuri, J.; Zhu, C.; Davidson, L.; Weaver, D.T.; Alt, F.W. A targeted DNA-PKcs-null mutation reveals DNA-PK-dependent functions for Ku in V(D)J recombination. Immunity 1998, 9, 367–376.

37. Gu, Y.; Seidl, K.J.; Rathbun, G.A.; Zhu, C.; Manis, J.P.; van der Stoep, N.; Davidson, L.; Cheng, H.L.; Sekiguchi, J.M.; Frank, K., et al. Growth retardation and leaky SCID phenotype of Ku70-deficient mice. Immunity 1997, 7, 653–665.

38. Nussenzweig, A.; Chen, C.; da Costa Soares, V.; Sanchez, M.; Sokol, K.; Nussenzweig, M.C.; Li, G.C. Requirement for Ku80 in growth and immunoglobulin V(D)J recombination. Nature 1996, 382, 551–555, doi:10.1038/382551a0.

39. Gao, Y.; Sun, Y.; Frank, K.M.; Dikkes, P.; Fujiwara, Y.; Seidl, K.J.; Sekiguchi, J.M.; Rathbun, G.A.; Swat, W.; Wang, J., et al. A critical role for DNA end-joining proteins in both lymphogenesis and neurogenesis. Cell 1998, 95, 891–902.

40. Frank, K.M.; Sekiguchi, J.M.; Seidl, K.J.; Swat, W.; Rathbun, G.A.; Cheng, H.L.; Davidson, L.; Kangaloo, L.; Alt, F.W. Late embryonic lethality and impaired V(D)J recombination in mice lacking DNA ligase IV. Nature 1998, 396, 173–177, doi:10.1038/24172.

41. Sekiguchi, J.; Ferguson, D.O.; Chen, H.T.; Yang, E.M.; Earle, J.; Frank, K.; Whitlow, S.; Gu, Y.; Xu, Y.; Nussenzweig, A., et al. Genetic interactions between ATM and the nonhomologous end-joining factors in genomic stability and development. Proc. Natl. Acad. Sci. 2001, 98, 3243–3248, doi:10.1073/pnas.051632098.