Synthetic lethality between murine DNA repair factors XLF and DNA-PKcs is rescued by inactivation of Ku70

Mengtan Xing¹, Magnar Bjørås¹, Jeremy A. Daniel², Frederick W. Alt³,* and Valentyn Oksenych¹,²,³,⁴,*

¹Institute for Cancer Research and Molecular Medicine, Norwegian University of Science and Technology, Laboratory Center, Erling Skjalgssons gate 1, 7491 Trondheim, Norway
²The NNF Center for Protein Research, Faculty of Health and Medical Sciences, University of Copenhagen, Blegdamsvej 3B, 2200 Copenhagen, Denmark
³Howard Hughes Medical Institute; Program in Cellular and Molecular Medicine, Boston Children’s Hospital; Department of Genetics, Harvard Medical School, Boston, MA, 02115
⁴St. Olavs Hospital, Trondheim University Hospital, Clinic of Medicine, Postboks 3250 Sluppen, 7006 Trondheim

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*Corresponding authors:
alt@enders.tch.harvard.edu (Frederick W. Alt)
valentyn.oksenych@ntnu.no (Valentyn Oksenych)
Abstract

DNA double-strand breaks (DSBs) are recognized and repaired by the Classical Non-Homologous End-Joining (C-NHEJ) and Homologous Recombination pathways. C-NHEJ includes the core Ku70 and Ku80 (or Ku86) heterodimer that binds DSBs and thus promotes recruitment of accessory downstream NHEJ factors XLF, PAXX, DNA-PKcs, Artemis and other core subunits, XRCC4 and DNA Ligase 4 (Lig4). In the absence of core C-NHEJ factors, DNA repair can be performed by Alternative End-Joining, which likely depends on DNA Ligase 1 and DNA Ligase 3. Genetic inactivation of C-NHEJ factors, such as Ku70, Ku80, XLF, PAXX and DNA-PKcs results in viable mice showing increased levels of genomic instability and sensitivity to DSBs. Knockouts of XRCC4 or Lig4, on the other hand, as well as combined inactivation of XLF and DNA-PKcs, or XLF and PAXX, result in late embryonic lethality in mice, which in most cases correlate with severe apoptosis in the central nervous system. Here, we demonstrate that inactivation of the Ku70 gene rescues the synthetic lethality between XLF and DNA-PKcs, resulting in triple knockout mice that are indistinguishable from Ku70-deficient littermates by size or levels of genomic instability. Moreover, we find that combined inactivation of Ku70 and XLF results in viable mice. Together, these findings suggest that Ku70 is epistatic with XLF and DNA-PKcs and support a model in which inactivation of Ku70 allows DNA lesions to become accessible to alternative DNA repair pathways.
1. Introduction

Non-Homologous DNA End Joining (NHEJ) is a cellular pathway that recognizes and fixes DNA double-strand breaks (DSBs) throughout the cell cycle. Core NHEJ factors include Ku70 and Ku80 (Ku86), a heterodimer that recognizes DSBs and has binding sites for multiple other NHEJ proteins. Ligation of DNA breaks is mediated by another heterodimer that includes core NHEJ factors, the DNA ligase 4 (Lig4) enzyme and its binding partner, XRCC4. DNA-dependent protein kinase, catalytic subunit (DNA-PKcs) and nuclease Artemis are accessory NHEJ factors, yet all of these proteins are individually required for lymphocyte development [1]. XRCC4-like factor (XLF) [2, 3] and paralog of XRCC4 and XLF (PAXX) [4-6] are NHEJ factors whose function remains to be determined.

Activation of the DNA damage response (DDR) signaling pathway occurs upon induction of DSBs. It consists of multiple enzymes responsible for posttranslational modifications as well as scaffold factors that are recruited to the chromatin areas adjacent to the DSBs. Related protein kinases ataxia telangiectasia mutated (ATM) and DNA-PKcs phosphorylate multiple substrates in response to DSBs, including histone H2AX and most of the NHEJ factors. Scaffold proteins MDC1 and 53BP1, ubiquitin ligases RNF8 and RNF168, and their downstream effectors RIF1 and PTIP then access modified chromatin [1].

Both NHEJ and the DDR are required for DNA repair events that are associated with the assembly of gene segments of immunoglobulins and T lymphocyte receptors, known as V(D)J recombination. During this process, Recombination Activating Gene (RAG) nuclease generates DSBs next to V, D and J gene segments, and NHEJ factors recognize and ligate DNA ends. Mature B lymphocytes change immunoglobulin isotypes in a process called Class Switch Recombination (CSR), when Activation induced Cytosine deaminase (AID) initiates DNA modifications that results in DSBs in the immunoglobulin constant region gene [1, 7]. CSR is also dependent on NHEJ and the DDR, although it can be robust, reaching about a half of wild type level, even in the absence of core NHEJ factors, likely due to the activity of DNA Ligase 1 and DNA ligase 3 [8, 9].
Mutations in several NHEJ factors in human are associated with developmental disorders that include immunodeficiency, neurological abnormalities and cancer [1]. A number of mouse knockouts were generated to model and investigate effects of NHEJ deficiency in vivo. Ku70- [10] and Ku80- [11] deficient mice are live-born, possess blocks in B and T lymphocyte development, and show moderate levels of p53-mediated apoptosis in central nervous system (CNS) [12]. Differently, deficiency for Lig4 or XRCC4 leads to p53-mediated late embryonic lethality in mice associated with high levels of apoptosis in the central nervous system [13-17]. Knockout of DNA-PKcs or Artemis results in live mice with immunodeficiency due to defects in B and T lymphocyte development [18, 19], although a kinase dead (KD) mutation of DNA-PKcs results in embryonic lethality associated with neuronal apoptosis [20]. The XLF knockout results in nearly wild type mice with a mild reduction of lymphocyte count [21, 22], and knockout of PAXX does not lead to any detectable phenotype in mice [23, 24].

Inactivation of several NHEJ genes simultaneously allows studying genetic interactions within this pathway in vivo. For example, inactivation of Ku80 rescued embryonic lethality of Lig4-deficient mice [25] and mice with catalytically dead DNA-PKcs [20]. Inactivation of Ku70 also rescued lethality of Lig4-deficient mice [26, 27]; moreover, combined inactivation of XLF and DNA-PKcs [28] or XLF and PAXX resulted in synthetic lethality [23, 24].

XLF overlaps functionally with multiple factors in mouse development, lymphocyte development and maintenance of genomic stability. We and others have found that XLF-deficient lymphocytes are proficient in V(D)J recombination due to compensatory functions of other NHEJ factors, such as DNA-PKcs [28] and PAXX [23, 24, 29, 30]; the DDR factors, such as ATM and H2AX [31], 53BP1 [32, 33]; and RAG [34]. Combined inactivation of XLF and H2AX leads to early embryonic lethality [31], while mice with double deficiency of DNA-PKcs and XLF [28] or PAXX and XLF [23, 24] possess late embryonic lethality, resembling deficiency for XRCC4 or Lig4 [15, 16]. In the latter cases, the embryonic lethality is associated with massive cell death, including in the CNS. Besides functional
overlap with XLF, DNA-PKcs is also functionally redundant with the related protein kinase ATM in aspects of mouse development, CSR and V(D)J recombination [35-37].

Here, we demonstrated that genetic inactivation of Ku70 rescues embryonic lethality of XLF/DNA-PKcs double-deficient mice; moreover, combined inactivation of Ku70 and XLF results in live mice phenotypically indistinguishable from Ku70-deficient littermates.


2.1. Mouse models

All experiments involving mice were performed according to the protocols approved by the Animal Resources Care Facility of Boston Children’s Hospital (ARCH), University of Copenhagen and Norwegian University of Science and Technology (NTNU). Ku70+/− [10], XLF+/Δ [21], DNA-PKcs+/− [16] and p53+/− [38] mice were previously described.

2.2. Tail fibroblasts

Primary murine tail fibroblasts were generated and cultured as described [28, 33]. Briefly, mouse tail skin was separated from the rest of the tail, cut into about 4 mm² pieces, and treated with 2 mg/mL collagenase II (Gibco) in DMEM supplemented with 15% (vol/vol) heat-inactivated FCS and antibiotics for 24 h at 37°C, at 5% (vol/vol) CO2 incubator. After being filtered through a 70-μm nylon cell strainer (BD Falcon), cells were washed in DMEM supplemented with 15% (vol/vol) FCS and plated. Fibroblasts from the second or third passages were used for telomere FISH assay.

2.3. Telomere FISH

Metaphases were prepared and chromosomal aberrations were counted as described [28, 33, 39]. The murine tail fibroblasts were synchronized with 100 ng/mL colcemid (KaryoMax, Gibco) for 6h. Then, the cells were treated with trypsin in PBS solution, washed with PBS, and lysed in 75 mM KCl (37°C, 20min). Fibroblasts were fixed in methanol:acetic acid (3:1) as described [39]. Telomeres were visualized with a Cy3-labeled fluorescent CCCTAACCTAACCCTAA probe (Applied
Biosystems). To visualize DNA, the samples were treated with DAPI (Vector Laboratories). Images were obtained with Eclipse microscope (Nikon). Chromosomal breaks were defined as a clear loss of telomere signal from both sister chromatids, and loss of telomere signal from one of the chromatids or clear absence of DAPI in the middle of one chromatid were defined as chromatid breaks.

3. Results

3.1. Generation of Ku70\(^{-}/\)XLF\(^{-}/\) double deficient mice

Ku70 and Ku80 form a heterodimer (Ku) that recognizes DSBs and has affinity to other DNA repair factors, including the catalytic subunit DNA-PKcs forming the DNA-PK holoenzyme. In addition, Ku facilitates recruitment of XLF and PAXX to DSB sites [40, 41]. Mice with individual Ku70, DNA-PKcs or XLF genetic deficiencies are alive and possess different levels of genomic instability. Recently, we found that combined inactivation of XLF and DNA-PKcs leads to synthetic lethality, as XLF and DNA-PKcs have functional redundancy [1, 28] and (Fig. 1). To determine whether XLF functionally overlaps with either the entire DNA-PK complex or functions downstream and is dependent of Ku, we intercrossed mice homozygous for XLF null allele and heterozygous for Ku70 (Ku70\(^{+/}\)XLF\(^{-}/\)). We obtained double knockout Ku70\(^{-}/\)XLF\(^{-}/\) mice at the expected ratio, 25 out of 104 pups analyzed, or 24% (Fig. 1). In control cages, breeding of DNA-PKcs\(^{+/}\)XLF\(^{-}/\) mice resulted in low number of live-born DNA-PKcs\(^{+/}\)XLF\(^{-}/\) double knockout mice (5 out of 104 pups analyzed, or about 5%), and none of them lived longer than 5 days, P5 (Fig. 1). In addition, intercrossing of Ku70\(^{+/}\)DNA-PKcs\(^{+/}\) mice resulted in alive Ku70\(^{+/}\)DNA-PKcs\(^{+/}\) pups that were also born at a proportion close to expected (Fig. 1). We conclude that both XLF and DNA-PKcs function downstream of Ku70. Moreover, we did not observe any evidence of Ku70-independent function of XLF and DNA-PKcs in vivo.

3.2. Inactivation of Ku70 gene rescues perinatal lethality of XLF\(^{-}/\)DNA-PKcs\(^{-}/\) mice
Upon interbreeding mice homozygous for XLF null allele and heterozygous for both DNA-PKcs and Ku70 (Ku70<sup>+/−</sup>DNA-PKcs<sup>−/−</sup>XLF<sup>−/−</sup>), triple null Ku70<sup>+/−</sup>DNA-PKcs<sup>−/−</sup>XLF<sup>−/−</sup> pups were surprisingly born at expected ratios, 17 out of 308, or about 5.5% (Fig. 2). These triple knockout mice were indistinguishable by size and lifespan from Ku70<sup>−/−</sup> or Ku70<sup>−/−</sup>XLF<sup>−/−</sup> littermates (Fig. 3) and were alive up to 12 months of age. Meanwhile, there is a reduced number of live-born XLF<sup>−/−</sup>DNA-PKcs<sup>−/−</sup> mice that rapidly died postnatally (by P5) when carrying one or two wild type alleles of Ku70 (Fig. 2). In particular, there were about 2.6% of Ku70<sup>+/−</sup>DNA-PKcs<sup>−/−</sup>XLF<sup>−/−</sup> (8 out of 308) and 4.2% of Ku70<sup>−/−</sup>DNA-PKcs<sup>−/−</sup>XLF<sup>−/−</sup> pups born (Fig. 2). The average size of Ku70<sup>−/−</sup> mice at day P30 (8.2g) was similar to the ones of Ku70<sup>+/−</sup>XLF<sup>−/−</sup> (8.5g), p=0.9987 and the size of triple knockout Ku70<sup>−/−</sup>DNA-PKcs<sup>−/−</sup>XLF<sup>−/−</sup> mice (9.4g) was similar to double knockout Ku70<sup>−/−</sup>XLF<sup>−/−</sup> (p=0.8328) and single knockout Ku70<sup>−/−</sup> (p=0.6506). See Figure legend for the full statistical comparisons between groups. We conclude that perinatal lethality of XLF<sup>−/−</sup>DNA-PKcs<sup>−/−</sup> mice is Ku-dependent.

3.3. Ku70 is epistatic with XLF and DNA-PKcs in maintaining genomic stability

Combined deficiency for XLF and DNA-PKcs leads to high levels of genomic instability in murine fibroblasts measured as proportion of metaphases with chromosomal and chromatid breaks [28]. Should XLF have Ku-independent functions in maintaining genomic stability, we would expect an increased number of aberrant metaphases in Ku70<sup>−/−</sup>XLF<sup>−/−</sup> cells compared to single Ku70<sup>−/−</sup> and XLF<sup>−/−</sup> cells. Here, we found that levels of genomic instability in XLF<sup>−/−</sup> and DNA-PKcs<sup>−/−</sup> primary murine tail fibroblasts are higher than in WT controls (Fig. 4). Moreover, the levels of genomic instability in Ku70<sup>−/−</sup> (25%), Ku70<sup>−/−</sup>XLF<sup>−/−</sup> (28%) and Ku70<sup>−/−</sup>DNA-PKcs<sup>−/−</sup>XLF<sup>−/−</sup> (32%) fibroblasts are indistinguishable (p>0.2746), which suggests that Ku70 is epistatic with both XLF and DNA-PKcs in maintaining genomic stability, and likely as a result of their functions in the NHEJ pathway. The levels of genomic instability in DNA-PKcs<sup>−/−</sup>XLF<sup>−/−</sup> fibroblasts (26%) were similar to Ku70<sup>−/−</sup> cells, indicating that NHEJ is similarly abrogated in these two distinct genetic deficiencies (Fig. 4).
3.4. Haploinsufficiency for p53 rescues embryonic lethality of XLF−/−DNA-PKcs−/− mice

By intercrossing mice homozygous for XLF null allele and heterozygous for both DNA-PKcs and p53 (DNA-PKcs−/−XLF−/−p53−/+), we found that inactivation of only one allele of p53 results in live DNA-PKcs−/−XLF−/−p53−/+ mice. Eight mice of this genotype were detected at 30 days postnatally in our studies. DNA-PKcs−/−XLF−/−p53−/+ mice possessed reduced life span, which varied from 97 to 202 days, as well as reduced body weight being in the range of Ku70-deficient mice (data not shown). Further analysis is required to determine how haploinsufficiency for p53 rescues perinatal lethality of DNA-PKcs−/−XLF−/− mice.

4. Discussion

4.1. Models to explain genetic interaction between Ku70, DNA-PKcs and XLF

The loss-of-function principle is widely used in current genetic studies of the NHEJ pathway. Traditionally, only one gene is inactivated; yet recent studies reveal that simultaneous inactivation of several genes is necessary to study functional overlap between proteins and to draw the most precise models of cellular pathways. Mice with inactivated Ku70 and Ku80 are alive but of smaller size than wild type littermates [10, 11]. Deficiency for DNA-PKcs or XLF leads to live-born mice with modest DNA repair defects [18, 21, 22]. DNA-PKcs forms the DNA-PK holoenzyme with the heterodimer Ku70/Ku80, and while the function of the catalytic subunit depends on the presence of Ku, inactivation of either the Ku70 or Ku80 gene will disrupt the whole DNA-PK complex [1, 42, 43]. Combined inactivation of XLF and DNA-PKcs leads to perinatal lethality in mice and abrogates NHEJ [28]. Here, we proposed and experimentally verified two models that would explain the genetic interaction between Ku, DNA-PKcs and XLF (Fig. 5). In one model, XLF has functions in mouse development and DNA repair independent of Ku and potentially outside of NHEJ. In this case, XLF has functional redundancy with the entire DNA-PK holoenzyme, and combined inactivation of Ku and XLF would result in similar perinatal lethality as described for DNA-PKcs−/−XLF−/− mice earlier [28]. In the other model, Ku recruits the downstream factors, including XLF and DNA-PKcs [1, 42, 43] and the function
of XLF in mouse development and DNA repair completely depends on Ku. In this case, combined inactivation of Ku70 and XLF genes would result in mice with a phenotype identical to those with single Ku70 deficiency (Fig. 5, left panel). Indeed, our results strongly support the latter model. We found that the triple knockout Ku70−/−XLF−/−DNA-PKcs−/− mice are alive, and are nearly identical to Ku70−/− littermates by size and levels of genomic instability (Fig. 2-4). While the current manuscript was in the preparation, our colleagues found that combined inactivation of Ku80 and XLF results in alive double knockout Ku80−/−XLF−/− mice that resembles Ku80-knockout littermates [24].

4.2. Why does deficiency for Ku70 or p53 rescues perinatal lethality of DNA-PKcs−/−XLF−/− mice?

Knockout of the upstream NHEJ genes Ku70 or Ku80 rescues embryonic lethality of mice with inactivated Lig4 [25-27]. Here, we demonstrated a similar effect with inactivation of Ku70 rescuing otherwise perinatally lethal DNA-PKcs−/−XLF−/− double knockout mice. It is tempting to speculate that inactivation of Ku70 or Ku80 might also rescue the synthetic lethality between XLF and PAXX [23, 24], although this hypothesis remains to be tested. When one of the Ku genes is inactivated simultaneously with a downstream factor (Lig4, XLF, DNA-PKcs, etc.), the resulting phenotype reassembles Ku70−/− single knockout (Fig. 3, 4 and [25]). This suggests that the functions of downstream NHEJ factors is likely dependent on Ku. Indeed, there is no clear evidence of any Ku-independent function of these factors (XLF, PAXX, DNA-PKcs, Artemis, XRCC4 and Lig4) in vivo. With classical NHEJ is inactive in the absence of Ku70 or Ku80, the DSB sites become available for the alternative end-joining pathway(s). Although no significant difference exists between the genomic instability in Ku70−/− and DNA-PKcs−/−XLF−/− fibroblasts (Fig. 4 and [28]), rescue of the lethal phenotype can be explained by a critical function of alternative end-joining in other organs, such as brain. For example, inactivation of Lig4 or XRCC4 can be more detrimental for the viability of neurons than inactivation of Ku [10, 13-16, 44].

Inactivation of one or two alleles of Trp53 gene (p53) is also known to rescue the embryonic lethality associated with knockouts of downstream NHEJ factors XRCC4 and Lig4 [13, 14]. In this
case, reduction of p53-dependent apoptosis may keep cells with accumulated DNA damages alive and extend the time necessary for alternative end joining to identify and fix a likely critical number of DSBs above a threshold that is compatible with life.

4.3. DNA-PKcs deficiency combined with XLF knockout

There are several mouse models with mutations in DNA-PKcs gene. The mice homozygous for catalytic dead DNA-PKcs (DNA-PKcs<sup>KD/KD</sup>) possess embryonic lethality with high levels of genomic instability in affected embryos. This embryonic lethality is however rescued by combined inactivation of Ku80 or Ku70, leading to alive Ku80<sup>−/−</sup>DNA-PKcs<sup>KD/KD</sup> and Ku70<sup>−/−</sup>DNA-PKcs<sup>KD/KD</sup> mice. Moreover, inactivation of one or two alleles of Trp53 gene also lead to alive p53<sup>−/−</sup>DNA-PKcs<sup>KD/KD</sup> and p53<sup>+/−</sup>DNA-PKcs<sup>KD/KD</sup> mice [20]. Complete knockout of DNA-PKcs [18] leads to live mice with no expressed DNA-PKcs protein; and a spontaneous mutation derived from the in bred classical SCID mouse line also results in alive mice [45]. This latter mouse model carries minor modification at the C-terminus of the large DNA-PKcs subunit, which results in reduced amount of protein in the cells. In addition, this protein lacks its kinase activity. Thus, the SCID mouse model is distinct from the DNA-PKcs-null model, as it has residual amount of protein, albeit enzymatically inactive. Earlier, we have demonstrated that combined inactivation of XLF and DNA-PKcs results in perinatal lethality of DNA-PKcs<sup>−/−</sup>XLF<sup>−/−</sup> mice. Moreover, chemical inhibition of DNA-PKcs kinase in XLF-deficient but not in WT pro-B cells completely blocks the joining of blunt signal ends (DSBs) at the chromosomal level [28].

There remains no mechanistic explanation of such a functional overlap between the DNA-PKcs and XLF. Whether or not inactivating XLF in the SCID mice would result in a similar synthetic lethality is an intriguing question. Such a double deficient model could be used in the future to explain whether functional overlap between XLF and DNA-PKcs is due to the structural or catalytic features of the DNA-PKcs.

Author contributions
MX, MB, JAD, FWA and VO designed and analyzed the experiments, the majority of which were performed by VO. MX and VO analyzed mouse genotypes and performed the T-FISH. VO wrote the manuscript. All the authors read and approved the final version of the manuscript.

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Figure legends

**Fig. 1. Double knockout Ku70<sup>−/−</sup>XLF<sup>−/−</sup> mice are viable.**

The number of thirty-day-old mice (P30) of indicated genotypes. **Top.** Combined inactivation of XLF and DNA-PKcs leads to perinatal lethality. No DNA-PKcs<sup>−/−</sup>XLF<sup>−/−</sup> mice survived longer than day P5 (*). **Middle.** Mice with combined inactivation of Ku70 and XLF are live-born and develop as Ku70-deficient littermates. **Bottom.** Mice with combined inactivation of Ku70 and DNA-PKcs are live-born and develop as Ku70-deficient littermates.

**Fig. 2. Triple knockout Ku70<sup>−/−</sup>DNA-PKcs<sup>−/−</sup>XLF<sup>−/−</sup> mice are viable.**

The number of thirty-day-old mice (P30) of indicated genotypes. Combined inactivation of XLF and DNA-PKcs leads to perinatal lethality of Ku70<sup>+/+</sup> and Ku70<sup>+/−</sup> mice with no mice survived longer
than day P5 (*). Combined inactivation of XLF, DNA-PKcs and Ku70 results in live-born mice that develop as Ku70 single deficient littermates.

Fig. 3. Double knockout Ku70\textsuperscript{−/−}XLF\textsuperscript{−/−} and triple knockout Ku70\textsuperscript{−/−}DNA-PKcs\textsuperscript{−/−}XLF\textsuperscript{−/−} mice have similar size as Ku70\textsuperscript{−/−} mice. Body weights of thirty-day-old mice (P30) of the indicated genotypes in grams.

The average weight of at least nine mice of each genotype is indicated. No XLF\textsuperscript{−/−}DNA-PKcs\textsuperscript{−/−} were alive at day P30 (n=0; n/a=not available). The average size of wild type mice was 20.3g; 18.8g for XLF\textsuperscript{−/−} mice and 15.0g for DNA-PKcs\textsuperscript{−/−} mice. All Ku70-deficient mice were smaller. The average size of Ku70\textsuperscript{−/−} mice (8.2g) was similar to the ones of Ku70\textsuperscript{−/−}XLF\textsuperscript{−/−} (8.5g). The average size of triple knockout Ku70\textsuperscript{−/−}DNA-PKcs\textsuperscript{−/−}XLF\textsuperscript{−/−} mice (9.4g) was similar to Ku70\textsuperscript{−/−}XLF\textsuperscript{−/−} and Ku70\textsuperscript{−/−}.

Multiple comparisons between 15 groups with one-way ANOVA, GraphPad Prizm 7.03:

WT vs XLF\textsuperscript{−/−}, p=0.3814 (n.s.); WT vs DNA-PKcs\textsuperscript{−/−}, p<0.0001 (**); WT vs Ku70\textsuperscript{−/−}, p<0.0001 (****); WT vs Ku70\textsuperscript{−/−}XLF\textsuperscript{−/−}, p<0.0001 (****); WT vs Ku70\textsuperscript{−/−}DNA-PKcs\textsuperscript{−/−}XLF\textsuperscript{−/−}, p<0.0001 (****); XLF\textsuperscript{−/−} vs DNA-PKcs\textsuperscript{−/−}, p<0.0001 (****); XLF\textsuperscript{−/−} vs Ku70\textsuperscript{−/−}, p<0.0001 (****); XLF\textsuperscript{−/−} vs Ku70\textsuperscript{−/−}XLF\textsuperscript{−/−}, p<0.0001 (****); XLF\textsuperscript{−/−} vs Ku70\textsuperscript{−/−}DNA-PKcs\textsuperscript{−/−}XLF\textsuperscript{−/−}, p<0.0001 (****); DNA-PKcs\textsuperscript{−/−} vs Ku70\textsuperscript{−/−}, p<0.0001 (****); DNA-PKcs\textsuperscript{−/−} vs Ku70\textsuperscript{−/−}XLF\textsuperscript{−/−}, p<0.0001 (****); DNA-PKcs\textsuperscript{−/−} vs Ku70\textsuperscript{−/−}DNA-PKcs\textsuperscript{−/−}XLF\textsuperscript{−/−}, p<0.0001 (****); Ku70\textsuperscript{−/−} vs Ku70\textsuperscript{−/−}XLF\textsuperscript{−/−}, p=0.9987 (n.s.); Ku70\textsuperscript{−/−} vs Ku70\textsuperscript{−/−}DNA-PKcs\textsuperscript{−/−}XLF\textsuperscript{−/−}, p=0.6506 (n.s.); Ku70\textsuperscript{−/−}XLF\textsuperscript{−/−} vs Ku70\textsuperscript{−/−}DNA-PKcs\textsuperscript{−/−}XLF\textsuperscript{−/−}, p=0.8328 (n.s.)

Fig. 4. Genomic instability in double knockout Ku70\textsuperscript{−/−}XLF\textsuperscript{−/−} and triple knockout Ku70\textsuperscript{−/−}DNA-PKcs\textsuperscript{−/−}XLF\textsuperscript{−/−} mice are similar to Ku70\textsuperscript{−/−} mice. Analyses of genomic instability in primary tail fibroblasts.

Fibroblasts were isolated from live-born mice of indicated genotypes. Top. Number (n) and proportion (%) of metaphases with detected chromosomal or chromatid breaks in fibroblasts of indicated genotypes. Fifty metaphases per mouse and three to five mice per genotype were analyzed. Bottom. Examples of chromatid break, intact chromosomes and chromosomal breaks (left and middle). Schematic view of intact chromosome, chromatid break and chromosomal break (right).
Fig. 5. Two models of functional interaction between the Ku, DNA-PKcs and XLF NHEJ factors.

Left. Ku70/Ku80 heterodimer is recruited to the DNA break sites and facilitates recruitment of both XLF and DNA-PKcs. XLF has functional redundancy with DNA-PKcs, and inactivation of Ku70 gene will rescue the perinatal lethality of XLF−/−DNA-PKcs−/− double knockout mouse (this model is supported by our results). Right. Ku70/Ku80 is recruited to the DSB site and binds DNA-PKcs subunit to form a DNA-PK holoenzyme. Even though XLF recruitment to DSB is Ku-dependent, it could have Ku-independent function in mouse development, and thus functionally redundant with entire DNA-PK complex. Genetic inactivation of Ku70 will not rescue the embryonic lethality of XLF−/−DNA-PKcs−/− double-knockout mice (this model is not supported by our results).

References

Figure 1

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*Live born pups that lived less than five days

Figure 2

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<td>Ku70(^+/+)/DNA-PKcs(^+/+)/XLF(^/-)</td>
<td>23</td>
<td>19</td>
</tr>
<tr>
<td>Ku70(^+/+)/DNA-PKcs(^+/+)/XLF(^/-)</td>
<td>36</td>
<td>38</td>
</tr>
<tr>
<td>Ku70(^+/+)/DNA-PKcs(^+/+)/XLF(^/-)</td>
<td>17</td>
<td>19</td>
</tr>
</tbody>
</table>

*Live born pups that lived less than five days
Figure 3

Mouse weight at P30, g

<table>
<thead>
<tr>
<th></th>
<th>Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>WT</td>
<td></td>
</tr>
<tr>
<td>XLF⁻⁻⁻</td>
<td></td>
</tr>
<tr>
<td>DNA-PKcs⁻⁻⁻</td>
<td></td>
</tr>
<tr>
<td>XLF⁻⁻⁻ DNA-PKcs⁻⁻⁻</td>
<td></td>
</tr>
<tr>
<td>Ku70⁻⁻⁻ XLF⁻⁻⁻</td>
<td></td>
</tr>
<tr>
<td>Ku70⁻⁻⁻ DNA-PKcs⁻⁻⁻</td>
<td></td>
</tr>
<tr>
<td>n=12</td>
<td>n=12</td>
</tr>
<tr>
<td>n=12</td>
<td>n=0 (n/a)</td>
</tr>
<tr>
<td>n=10</td>
<td>n=12</td>
</tr>
<tr>
<td>n=9</td>
<td></td>
</tr>
</tbody>
</table>

p<0.3814, n.s.
p<0.0001, ****
p<0.0001, ****
p<0.8328, n.s.
p=0.9987, n.s.

Figure 4

Genomic instability in tail fibroblasts

<table>
<thead>
<tr>
<th>Genotype</th>
<th>Number of mice, n</th>
<th>Total metaphases, n</th>
<th>Abnormal metaphases, n</th>
<th>Chromosomal breaks, n</th>
<th>Chromatid breaks, n</th>
</tr>
</thead>
<tbody>
<tr>
<td>Wild type</td>
<td>5</td>
<td>250</td>
<td>14</td>
<td>9</td>
<td>1</td>
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<tr>
<td>XLF⁻⁻⁻</td>
<td>5</td>
<td>250</td>
<td>30</td>
<td>24</td>
<td>8</td>
</tr>
<tr>
<td>DNA-PKcs⁻⁻⁻</td>
<td>5</td>
<td>250</td>
<td>23</td>
<td>36</td>
<td>4</td>
</tr>
<tr>
<td>XLF⁻⁻⁻ DNA-PKcs⁻⁻⁻</td>
<td>5</td>
<td>250</td>
<td>66</td>
<td>104</td>
<td>9</td>
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<tr>
<td>Ku70⁻⁻⁻</td>
<td>3</td>
<td>150</td>
<td>38</td>
<td>49</td>
<td>12</td>
</tr>
<tr>
<td>XLF⁻⁻⁻ Ku70⁻⁻⁻</td>
<td>5</td>
<td>250</td>
<td>69</td>
<td>93</td>
<td>17</td>
</tr>
<tr>
<td>XLF⁻⁻⁻ DNA-PKcs⁻⁻⁻ Ku70⁻⁻⁻</td>
<td>4</td>
<td>200</td>
<td>65</td>
<td>97</td>
<td>15</td>
</tr>
</tbody>
</table>

Intact chromosome
Chromatid break
Chromosomal break
Intact chromosome
Chromatid break
Chromosomal break
Figure 5

Model 1
Inactivation of Ku rescues synthetic lethality of XLF<sup>-/-</sup>DNA-PKcs<sup>-/-</sup> mice

Model 2
Inactivation of Ku does not rescue synthetic lethality of XLF<sup>-/-</sup>DNA-PKcs<sup>-/-</sup> mice