The Mre11 complex (Mre11, Rad50, and Nbs1) is a central component of the DNA damage response (DDR), governing both double-strand break repair and DDR signaling. Rad50 contains a highly conserved Zn$^{2+}$-dependent homodimerization interface, the Rad50 hook domain. Mutations that inactivate the hook domain produce a null phenotype. In this study, we analyzed mutants with reduced hook domain function in an effort to stratify hook-dependent Mre11 complex functions. One of these alleles, Rad50$^{46}$, conferred reduced Zn$^{2+}$ affinity and dimerization efficiency. Homozygous Rad50$^{46}$/46 mutations were lethal in mice. However, in the presence of wild-type Rad50, Rad50$^{46}$ exerted a dominant gain-of-function phenotype associated with chronic DDR signaling. At the organismal level, Rad50$^{+/46}$ exhibited hydrocephalus, liver tumorigenesis, and defects in primitive hematopoietic and gametogenic cells. These outcomes were dependent on ATM, as all phenotypes were mitigated in Rad50$^{+/46}$ Atm$^{+/−}$ mice. These data reveal that the murine Rad50 hook domain strongly influences Mre11 complex-dependent DDR signaling, tissue homeostasis, and tumorigenesis.

**Keywords:** Mre11 complex; double-strand breaks; Rad50; ATM
Structural analyses of the complex suggest a model in which the Mre11 complex’s effect on DSB repair is attributable to its ability to bridge DNA molecules in trans and in cis (de Jager et al. 2001, 2004; van Noort et al. 2003; Moreno-Herrero et al. 2005). In trans, two complexes bind DNA independently by their globular domains and, via association through the Rad50 hook domain, would bridge sister chromatids and promote HDR between them. Indeed, studies in yeast have shown that the dimerization function of the hook domain underlies the influence of this domain on DSB repair, telomere maintenance, and meiotic DSB formation (Wiltzius et al. 2005). In cis, the globular domain could bridge two DNA ends in a manner that would promote NHEJ (Williams et al. 2008). In this mode, end-to-end DNA bridging is effected by adjacent monomeric components of the dimeric globular domain (Lammens et al. 2011; Lim et al. 2011).

Mutations that target the invariant cysteines of the Rad50 hook domain (Hopfner et al. 2002; Wiltzius et al. 2005; Hohl et al. 2011; He et al. 2012) globally disrupt Mre11 complex functions and thus phenocopy the null Rad50 mutation. Hence, the specific contribution of the Rad50 hook domain to the repair and signaling functions of the Mre11 complex has not been examined in isolation from the other domains of the complex.

To gain insight into Rad50 hook domain’s role, our laboratory targeted the residues adjacent to the invariant cysteine residues in S. cerevisiae in an effort to obtain hypomorphic alleles. The alleles obtained were examined and found to encode hook domains with reduced Zn$^{2+}$-dependent dimerization and increased DNA damage sensitivity [M Hohl, C Tous, T Kocharczyk, A Krężel, A Aguilera, and JHJ Petrini, in prep.]. In this study, two of the yeast alleles [rad50-46 and rad50-47] were modeled in mice.

We found that in the absence of the wild-type Rad50 protein, both Rad50$^{46}$ and Rad50$^{47}$ alleles failed to support viability at either the cellular or organismal level, consistent with the view that the Rad50 hook domain is integral to the functions of the Mre11 complex. The Rad50$^{46}$ and Rad50$^{47}$ gene products exerted dominant gains of function in the presence of the wild-type protein.

Heterozygous Rad50$^{46+}$ and Rad50$^{47+}$ cells were relatively proficient in DNA repair but exhibited chronic DDR signaling at the cellular level and severe ATM-dependent phenotypes at the organismal level. The data suggest that the Rad50$^{46}$ and Rad50$^{47}$ proteins promote aberrant heterotypic dimerization at the hook domain interface. As the functional defect imparted by Rad50$^{46}$ and Rad50$^{47}$ predominantly affected Mre11 complex-dependent DDR signaling, these data reveal a previously unrecognized influence of the Rad50 hook domain on Mre11 complex-dependent regulation of the apical DDR kinases.

**Results**

**Rad50 hook domain mutants**

Having previously shown that alteration of the cysteine residues of the Rad50 hook domain was tantamount to complete elimination of the protein (Hopfner et al. 2002), we reasoned that alteration of adjacent residues might result in a partial loss of function that would provide...
insight regarding the functional importance of the Rad50 hook. Using S. cerevisiae, we identified two mutants, designated rad50-46 and rad50-47, that behaved as hook domain hypomorphs and have in common the presence of charged residues in the vicinity of the cysteines that coordinate zinc [Fig. 1A]. As a consequence, both mutants showed reduced hook-mediated dimerization and reduced affinity for Zn$^{2+}$ [M Hohl, C Tous, T Kótačzyk, A Krčzal, A Aguilera, and JHJ Petrini, in prep.].

A 22.7-kDa fragment of bacterially produced mouse Rad50 [Rad50-HK] that contained the hook domain flanked by small segments of the coiled-coil region was purified and examined by size exclusion chromatography and multiangle light scattering (SEC-MALS). Rad50-HK$^\text{WT}$ eluted as a single peak [Fig. 1B,C] with a molecular weight of 44.6 kDa, indicating a dimeric species. Inductively coupled plasma mass spectrometry (ICP-MS) revealed that the protein to Zn$^{2+}$ molar ratio of this species was 2.4, approximating the expected value of 2 for a Zn$^{2+}$-dependent dimer (Fig. 1D). In contrast, Rad50-HK$^{46}$ eluted in three peaks [Fig. 1B]. Peak 2 [P2] was similar to Rad50-HK$^\text{WT}$ in molecular weight and Zn$^{2+}$ content. Peak 1 [P1] and peak 3 [P3] contained very little Zn$^{2+}$ and represent monomers and higher-molecular-weight aggregates, respectively (Fig. 1D). These data suggest that the Rad50$^{46}$ hook domain is structurally compromised but retains some ability to dimerize and bind Zn$^{2+}$.

**Derivation of Rad50$^{46}$ and Rad50$^{47}$ mice**

To analyze the function of the Rad50 hook in vivo, mice expressing analogs of the yeast rad50-46 and rad50-47 alleles were derived. The mice initially retained a LoxP-flanked neo cassette within intron 13 of the mutant Rad50 locus [Supplemental Fig. S1A,B]. These alleles were designated Rad50$^{\text{WT}}$/46 and Rad50$^{\text{WT}/C0}$. Rad50$^{\text{WT}/C0}$ chimeric mice were unable to produce viable offspring. No live Rad50$^{\text{WT}/C0}$ births were noted among 327 pups produced from 27 different chimeras. PCR genotyping of tissues from Rad50$^{\text{WT}/C0}$ chimeras [including isolated sperm] indicated that the Rad50$^{\text{WT}/C0}$ allele was present (Supplemental Fig. S1C). On this basis, we concluded that the Rad50$^{\text{WT}/C0}$ allele exerted a dominant lethal effect. In contrast, Rad50$^{\text{WT}/\text{Neo}46}$ chimeric mice transmitted the Rad50$\times$Neo$^{46}$ allele to viable offspring, and 50% of the embryonic stem cell-derived pups were Rad50$^{\text{WT}/\text{Neo}46}$.

Rad50$^{\text{WT}/\text{Neo}46}$ also exerted a dominant phenotype. The effects observed were more pronounced following deletion of the LoxP-flanked neo cassette [henceforth Rad50$^{\text{WT}/46}$ mice] by crossing to CAG-Cre mice. Rad50$^{\text{WT}/46}$ mice were born at the expected Mendelian ratios but displayed intense skin hyperpigmentation [Fig. 2A], and 27% of them developed hydrocephalus compared with 4% of Rad50$^{\text{WT}/\text{Neo}46}$ [Fig. 2B]. RT–PCR analysis indicated that the expression of Rad50$^{\text{WT}/\text{Neo}46}$ was lower than Rad50$^{\text{WT}/46}$ [Supplemental Fig. S1D]. Hence, the extent of dominance was correlated with the dosage of the Rad50$^{\text{WT}/46}$ gene product.

Rad50$^{\text{WT}/46}$ males were infertile (discussed further below). This precluded interbreeding of heterozygotes to obtain Rad50$^{\text{46}/\text{46}}$ mice. Instead, Rad50$^{\text{46}/\text{46}}$ females were crossed with Rad50$^{\text{46}/\text{46}}$ males [Luo et al. 1999]. No Rad50$^{\text{46}/\text{46}}$ mice were obtained from 90 offspring ($\chi^2$, $P < 10^{-4}$), indicating that the Rad50$^{\text{46}}$ allele does not support embryonic viability.

Rad50$^{\text{46}/\text{46}}$ and Rad50$^{\text{47}/\text{47}}$ mice were crossed with Rad50$^{\text{46}/\text{46}}$ [in which exons 1 and 2 are flanked by loxP sites] [Adelman et al. 2009] and a tamoxifen-inducible Cre [Ventura et al. 2007]. In this setting, Cre-mediated excision of the floxed exons of the Rad50$^{\text{46}}$ allele to generate Rad50$^{\text{46}/\text{46}}$- and Rad50$^{\text{47}/\text{47}}$-cells in vivo was carried out. The induction of Cre is variable in vivo, and so affected tissues were likely to exhibit mosaicism for the deletion genotype. Tamoxifen administered in the food produced excision in the Rad50$^{\text{46}}$ allele in various tissues but was predominant in the intestine [Supplemental Fig. S2B]. Rad50$^{\text{46}/\text{46}}$ and Rad50$^{\text{47}/\text{47}}$-mosaic mice did not live beyond 2 wk due to intestinal failure. Histological analysis of the gut 9 d after beginning the treatment evidenced a profound alteration of Rad50$^{\text{46}/\text{46}}$-intestinal morphology [Supplemental Fig. S2C]. These data indicate that the Rad50$^{\text{46}}$ allele does not support the viability of proliferating cells in vivo.

**DNA repair and checkpoint signaling in Rad50$^{\text{46}/\text{46}}$ cells**

We sought to determine the molecular basis of the Rad50$^{\text{46}/\text{46}}$ phenotype. SV40-immortalized Rad50$^{\text{46}/\text{46}}$ and Rad50$^{\text{47}/\text{47}}$ mouse embryonic fibroblasts (MEFs) showed similar levels of Mre11 complex proteins, and immunoprecipitations indicated that complex integrity was not overtly compromised [Fig. 2C].

The DNA repair functions of the Mre11 complex were modestly affected in Rad50$^{\text{46}/\text{46}}$ cells. Rad50$^{\text{46}/\text{46}}$ cells showed increased sensitivity to camptothecin at high doses [Fig. 2D] but were not sensitive to other DNA-damaging agents [Supplemental Fig. S3A]. Accordingly, treatment with clastogens failed to increase the yield of chromosomal aberrations relative to wild-type cells [Supplemental Fig. S3B]. The DR-GFP assay, in which an I-SceI/GFP reporter is integrated at the chromosome (Adelman et al. 2009) and a tamoxifen-inducible Cre [Ventura et al. 2007]. In this setting, Cre-mediated excision of the floxed exons of the Rad50$^{\text{46}}$ allele to generate Rad50$^{\text{46}/\text{46}}$- cells in vivo was carried out. The induction of Cre is variable in vivo, and so affected tissues were likely to exhibit mosaicism for the deletion genotype. Tamoxifen administered in the food produced excision in the Rad50$^{\text{46}}$ allele in various tissues but was predominant in the intestine [Supplemental Fig. S2B]. Rad50$^{\text{46}/\text{46}}$ and Rad50$^{\text{47}/\text{47}}$-mosaic mice did not live beyond 2 wk due to intestinal failure. Histological analysis of the gut 9 d after beginning the treatment evidenced a profound alteration of Rad50$^{\text{46}/\text{46}}$- intestinal morphology [Supplemental Fig. S2C]. These data indicate that the Rad50$^{\text{46}}$ allele does not support the viability of proliferating cells in vivo.

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The Mre11 complex is required for ATM activation [Stracker and Petrini 2011]. Rad50$^{\text{46}/\text{46}}$ cells exhibited indices of spontaneous ATM activation, including pS1987-ATM and γH2AX formation [Fig. 2F]. IR-induced ATM activation in Rad50$^{\text{46}/\text{46}}$ cells was otherwise normal and occurred with wild-type kinetics [Fig. 2F]. Rad50$^{\text{47}/\text{47}}$ skin fibroblasts generated from chimeric mice phenocopied Rad50$^{\text{46}/\text{46}}$ cells in all respects [Supplemental Fig. S4]. These results suggested that Rad50$^{\text{46}/\text{46}}$ [and Rad50$^{\text{47}/\text{47}}$] cells exhibited spontaneous DNA damage. An alternative nonexclusive possibility is that the Rad50$^{\text{46}}$ and Rad50$^{\text{47}}$ alleles are hypermorphic with respect to DNA damage.
signaling, as observed previously in Rad50<sup>S/S</sup> mice (and yeast) (Usui et al. 2006), and thereby exhibited indices of ATM activation at levels disproportional to the extent of DNA damage. This latter interpretation predicts that Rad50<sup>46</sup> phenotypes would be ATM-dependent. The Rad50<sup>+</sup>/46 phenotype is ATM-dependent

Precedent for the hypothesis that Rad50<sup>+</sup>/46 phenotypes could be ATM-dependent comes from Rad50<sup>S/S</sup> mice, which at both the cellular and organismal levels exhibited phenotypes reminiscent of Rad50<sup>+/46</sup> [Bender et al. 2002; Morales et al. 2005]. We found that most aspects of the Rad50<sup>+/46</sup> pathology, including hydrocephalus [Fig. 2B], were rescued in the context of Atm heterozygosity, demonstrating the acute ATM dependence of the Rad50<sup>+/46</sup> phenotype.

Mre11 complex-dependent ATM signaling affects the viability and proliferative behavior of Lin<sup>−</sup> Sca1<sup>+</sup> c-Kit<sup>+</sup> (LSK) cells, which contain primitive hematopoietic components, including hematopoietic stem cells (HSCs) and lineage progenitors [Morales et al. 2008]. Rad50<sup>+/46</sup> mice showed an altered LSK population characterized by an expansion (average increase of 4.7-fold) of the multipotent precursors (MPPs) [Fig. 3A]. This phenotype was also dependent on ATM because it was suppressed in Rad50<sup>+/46</sup> Atm<sup>+/−</sup> mice [Fig. 3A]. The increased MPP population was not correlated with elevated levels of its proximal descendants, the common lymphoid progenitor (CLP) and common myeloid progenitors (CMPs) [Fig. 3B]. CFU-S assays, which measure the ability of MPPs to differentiate and form colonies in the spleen of lethally irradiated mice, were carried out [Spangrude et al. 1988]. We noted a 50% reduction in the colony-forming ability of Rad50<sup>+/46</sup> MPP cells [Fig. 3C]. These data indicate that the expanded Rad50<sup>+/46</sup> MPP population is not fully functional.

A remarkable outcome in Rad50<sup>S/S</sup> Atm<sup>+/−</sup> mice was the suppression of lymphomagenesis associated to ATM deficiency [Morales et al. 2005; Usui et al. 2006]. Similarly, the latency of thymic lymphomas typical of the

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**Figure 2.** Rad50<sup>46</sup> allele is dominant. (A) Representative image showing hyperpigmentation in tail and paws in a Rad50<sup>+/46</sup> mouse compared with a Rad50<sup>+/+</sup> littermate. (B) Incidence of hydrocephalus in Rad50<sup>+/46</sup>, Rad50<sup>+/Neo46</sup>, and Rad50<sup>+/46</sup> Atm<sup>+/−</sup> mice and representative X-ray images of an 8-wk-old Rad50<sup>+/46</sup> mouse with hydrocephalus and a Rad50<sup>+/+</sup> littermate. (C) Rad50, Mre11, and Nbs1 protein levels in Rad50<sup>+/+</sup> and Rad50<sup>+/46</sup> SV40-immortalized MEFs and Nbs1 immunoprecipitation to assess Mre11 complex integrity in mock-treated cells (−) or 1 h after 5 Gy IR. Mre11<sup>ATLD1/ATLD1</sup> cells were used as control for cells with reduced Mre11 complex levels [Theunissen et al. 2003]. (D) SV40-immortalized MEFs were treated with camptothecin for 24 h, and colonies were assessed 10 d later. Experiments were done three times in triplicate using two independent cell lines per genotype. Error bars depict standard deviation (SD). (E) Primary mouse fibroblasts Rad50<sup>+/−</sup> Pim1<sup>+/DR-GFP</sup> and Rad50<sup>+/46</sup> Pim1<sup>+/DR-GFP</sup> were infected with SceI-IRES-hCD4 retrovirus, and GFP<sup>+</sup> cells were assessed 72 h after infection. The efficiency of GFP<sup>+</sup> is measured relative to the infected-only (CD4<sup>+</sup>) cells. Experiments were done in triplicate with two independent cell lines per genotype. P-value was determined by Wilcoxon rank-sum test. (F) SV40-immortalized MEFs of the indicated genotypes were left untreated (t = 0) or irradiated with 2 Gy IR, and the levels of pS1987-ATM and γH2AX were assessed in the indicated time points.
**Figure 3.** ATM-dependent hematopoietic alterations in Rad50<sup>+/46</sup> mice. (A) Quantification of long-term hematopoietic stem cells [LT-HSC], short-term hematopoietic stem cells [ST-HSC], and MPPs in bone marrow from mice of the indicated genotypes. Bars denote the average ± standard error of mean (SEM). Rad50<sup>+/46</sup>, n = 8; Rad50<sup>−/46</sup>, n = 8; Rad50<sup>+/46 Atm<sup>−/-</sup></sup>, n = 2. (B) Quantification of CMP, granulocyte macrophage progenitor (GMP), megakaryocyte erythroid progenitor (MEP), and CLP. Bars denote the average ± SEM. Rad50<sup>+/46</sup>, n = 5; Rad50<sup>−/46</sup>, n = 5; Rad50<sup>+/46 Atm<sup>−/-</sup></sup>, n = 2. P-values were determined by unpaired t-test. (C) Representative images and quantification of CFU-S assay at day 13 using bone marrow cells of the indicated genotype. Bars denote the average ± SEM of all the spleens obtained from three independent experiments. Rad50<sup>+/46</sup>, n = 21; Rad50<sup>−/46</sup>, n = 20; Rad50<sup>+/46 Atm<sup>−/-</sup></sup>, n = 5. P-values were determined by unpaired t-test. (D) Kaplan-Meier survival curves of Atm<sup>−/-</sup>, Rad50<sup>+/46</sup>, Rad50<sup>−/46 Atm<sup>−/-</sup></sup> [previously published, Morales et al. 2005], Rad50<sup>+/46</sup>, and Rad50<sup>−/46 Atm<sup>−/-</sup></sup> mice. Rad50<sup>+/46</sup> mouse survival was not assessed beyond 18 mo, and thus the events were censored at that age.

The mammalian Rad50 hook domain

*Atm<sup>−/-</sup>* genotype was markedly increased in Rad50<sup>+/46</sup> *Atm<sup>−/-</sup>* mice (Fig. 3D). In light of recent data suggesting a role for the Mre11 complex in ATR activation [Duursma et al. 2013; Shiotani et al. 2013], the suppression of lymphomagenesis in Rad50<sup>+/46 Atm<sup>−/-</sup></sup> mice may reflect compensatory Mre11 complex-dependent ATR activation.

**Germ cell dysfunction in Rad50<sup>+/46</sup> mice**

Rad50<sup>+/46</sup> mice exhibited severe fertility defects. All males and 27% of the females were infertile, and the remaining Rad50<sup>+/46</sup> females produced fewer and smaller litters than Rad50<sup>+/4</sup> controls during the breeding period [Supplemental Fig. SSA,B]. Rad50<sup>+/46</sup> exhibited ovarian and testicular atrophy [Fig. 4A,B], and the seminiferous tubules displayed severely reduced cellularity [Fig. 4C]. As with the hematopoietic phenotype, testicular defects were ATM-dependent. Testes from Rad50<sup>+/46 Atm<sup>−/-</sup></sup> mice were twofold larger than those from Rad50<sup>+/4</sup> [Fig. 4A], and the cellularity of the seminiferous tubules was increased [Fig. 4C].

Immunohistochemical staining for the synaptonemal complex protein SYCP3 revealed that Rad50<sup>+/46</sup> mice showed an eightfold reduction in meiotic prophase cells per tubule [Fig. 4C,D]. Similar analyses revealed a 23-fold reduction in SYCP3-positive cells from 14-dpp Rad50<sup>+/46</sup> mice [Supplemental Fig. S5C,D]. TUNEL staining of Rad50<sup>+/46</sup> tubules from adult and juvenile mice was unremarkable [Fig. 4C,D, Supplemental Fig. S5C,D], suggesting that the reduction in Rad50<sup>+/4</sup> meiotic cells was not due to apoptotic attrition. The reduction in cellularity did not appear to reflect a stage-specific block in meiotic progression, as chromosome spreads from the rare meiotic cells present revealed spermatocytes in all stages of meiosis [Supplemental Fig. S5E].

Spo11 protein initiates meiotic recombination by forming DSBs through a covalent protein–DNA intermediate. Spo11 is cleaved from DSB ends to liberate Spo11–oligonucleotide complexes [Neale et al. 2005] by a process that in yeast requires activity of the Mre11 complex [Neale et al. 2005; Milman et al. 2009]. In S. cerevisiae rad50-46 mutants, DSBs are formed, but Spo11 remains covalently bound to genomic DNA [M Hohl, C Tous, T Kochan´ czyk, A Krz´ ezl, A Aguilera, and JHJ Petrini, in prep.). We tested whether a defect in removal of Spo11 from DSB ends accounted for the infertility of Rad50<sup>+/46</sup>
mice. Spo11–oligonucleotide complexes were nearly absent in adult and juvenile Rad50<sup>+/46</sup> mice (Supplemental Fig. S5F). The reduction in Spo11–oligonucleotide complexes in Rad50<sup>+/46</sup> testes was not simply a consequence of testicular atrophy because they were evident in Dmc1<sup>−/−</sup> testes, which, like Rad50<sup>+/46</sup>, are severely atrophic (Supplemental Fig. S5F). These data suggested that in Rad50<sup>+/46</sup> mice, meiotic DSBs are not created, or Spo11 is not released following DSB formation. We could not exclude the possibility that this effect is due to the paucity of meiotic cells in Rad50<sup>+/46</sup> testes. To directly address whether defects in the formation or repair of meiotic DSBs were the cause of infertility of Rad50<sup>+/46</sup> mice, we generated Rad50<sup>+/46</sup> Spo11<sup>−/−</sup> mice. Seminiferous tu-

Figure 4. Germ cell dysfunction in Rad50<sup>+/46</sup> mice. (A) Representative images of testis from mice of the indicated genotypes and average testes weight ± SEM (Rad50<sup>+/+</sup>, n = 7; Rad50<sup>+/46</sup>, n = 8; Rad50<sup>−/−</sup> Atm<sup>−/−</sup>, n = 5). Bar, 2.5 mm. (B) Representative images of anti-VASA-stained mid-ovary sections from 3-wk-old mice and quantification of the number follicles. Bar, 0.4 mm. Ovaries analyzed: n = 10. P-value was determined by unpaired t-test. (C) Hematoxylin and eosin (H&E), TUNEL, SYCP3, and PLZF immunohistochemical staining of testis sections from adult mice of the indicated genotypes. Bar, 50 μm. (D) Quantification of immunohistochemical staining from C. The TUNEL-positive area is expressed as a ratio to the total testis area. Testis analyzed: Rad50<sup>+/+</sup>, n = 8; Rad50<sup>−/−</sup>, n = 10; Rad50<sup>−/−</sup> Atm<sup>−/−</sup>, n = 5. P-values were determined by unpaired t-test.
bules from Rad50+/−/ mice showed reduced cellularity compared with Spo11−/− [Supplemental Fig. S5G]. Analysis of sections from Rad50+/−/ Spo11−/−/ mice revealed that Rad50+/−/ was epistatic to Spo11−/−. This result indicated that the Rad50+/−/ phenotype was independent of Spo11 and thus excluded the possibility of impaired Spo11 removal as a cause of Rad50+/−/ infertility.

These data suggest that the Rad50+/−/ defect was manifest prior to initiation of the meiotic program. Supporting that interpretation, Rad50+/−/ tubules displayed 1.6-fold more cells positive for PLZF [a marker of undifferentiated spermatogonia] [Buas et al. 2004; Costoya et al. 2004] compared with Rad50+/+ mice [Fig. 4C,D]. This expansion of primitive precursors together with the reduction of meiotic cells and the lack of increased cell death was consistent with a block in spermatogonial differentiation, analogizing the block observed in primitive hematopoietic precursors.

Mre11 complex and ATM activation in Rad50+/−/ cells

Rad50+/−/ and Rad50−/− MEFs and ear fibroblast (EF) cultures were established. Lentiviral-mediated delivery of Cre resulted in excision within the Rad50+/−/ allele in 3 d [Supplemental Fig. S6A] to produce Rad50+/−/ and Rad50−/− cells that could be maintained in culture for 7–8 d after Cre infection. This provided a window of time to assess the outcome of limiting cells to Rad50−/− as the sole source of Rad50 protein. As shown previously in Rad50+/−/ cell extracts [Adelman et al. 2009], Rad50+/−/ cells displayed decreased levels of Rad50, Mre11, and Nbs1 [Fig. 5A], suggesting that Mre11 complex proteins were less stable in Rad50+/−/ cells.

Acute disruption of the Mre11 complex results in a dramatic increase in chromosomal aberrations [Demuth et al. 2004; Frappart et al. 2005; Reina-San-Martin et al. 2005; Yang et al. 2006; Adelman et al. 2009]. Excision in the Rad50+/−/ allele in Rad50+/−/ and Rad50+/−/ cells produced a similar outcome, with 53% of Rad50+/−/ and 42% of Rad50+/−/ cells showing chromosomal aberrations compared with 12% of Rad50+/−/ [Fig. 5B]. Cells with multiple aberrations (three or more) were not seen in Rad50+/−/ controls but represented up to 30% and 11.8% in Rad50+/−/ and Rad50+/−/, respectively [Supplemental Fig. S6B].

Figure 5. Mre11 complex and ATM activation in Rad50+/−/ cells. (A) Analysis of Mre11 complex levels in Rad50+/−/ and Rad50+/−/ SV40-immortalized MEFS 8 d after the lentiviral delivery of Cre or GFP and Nbs1 immunoprecipitation to assess the Mre11 complex integrity in the same cells. Kap1 was used as loading control. (B) Analysis of chromosomal aberrations in Rad50+/−/ cells showing chromosomal aberrations. (C) Analysis of the G2/M checkpoint in Rad50+/−/ and Rad50+/−/ SV40-immortalized MEFS 8 d after the lentiviral delivery of Cre or GFP. Shown is average ± SD of three independent experiments. Twenty-five or more metaphases were scored per genotype and experiment. (D) Analysis of pS824-Kap1 and Chk2 phosphorylation in Rad50+/−/ and Rad50+/−/ SV40-immortalized MEFS 8 d after the lentiviral delivery of Cre or GFP after 2 Gy of γ radiation. Time 0 corresponds to untreated cells. Note that Chk2 migrates as a higher-molecular-weight species when phosphorylated (pChk2). Ku70 levels were used as loading control.
Rad50Δ−/− MEFs were unable to activate ATM in response to IR, as evidenced by reduced phosphorylation of Kap1 and Chk2 following IR treatment (Fig. 5D). Rad50Δ−/− MEFs were also impaired in ATM activation but to a lesser extent. Whereas phosphorylation of Kap1 and Chk2 was undetectable in Rad50Δ−/− MEFs, we could consistently detect it in Rad50Δ−/− MEFs (Fig. 5D). This suggests that Rad50Δ−/− cells retain a degree of Mre11 complex function; however, this observation may reflect the presence of residual wild-type Rad50 protein.

Reduced ATM activation in Rad50Δ−/− cultures was associated with impairment of the DNA damage-dependent G2/M checkpoint. One hour after 5 Gy of IR, cells with at least one copy of wild-type Rad50 showed an 80%–90% reduction in the mitotic cells, indicative of a proficient G2/M checkpoint (Fig. 5C). Excision in the Rad50Δ allele to produce Rad50Δ−/− or Rad50Δ−/− cells caused defects in the G2/M checkpoint, with cells exhibiting only a 36% and 52% reduction in the proportion of mitotic cells, respectively, after IR (Fig. 5C).

Rad50Δ− promotes tumorigenesis

The Mre11 complex has been implicated in the suppression of malignancy in several contexts [Stracker and Petrini 2011; Foster et al. 2012; Gupta et al. 2013]. Accordingly, cohorts of 30 Rad50Δ−/Δ, 31 Rad50Δ−/Neo46, and 34 Rad50Δ− mice were followed for 18 mo. Although no signs of morbidity were evident, multiple liver neoplasms were noted upon necropsy. The incidence was 26% in Rad50Δ−/Δ mice, 37% in Rad50Δ−/Δ mice, and 3% in Rad50Δ− mice (Fig. 6A). Most of the affected mice presented hyperplastic nodules and/or adenomas, but 10% of Rad50Δ− mice developed malignant hepatoblastoma (Fig. 6A,B). Unlike adenomas, which arise from mature hepatocytes, hepatoblastomas arise from hepatic progenitor cells of the portal areas [Marquardt and Thorgeirsson 2010]. This indicates that both differentiated hepatocytes and immature progenitors are susceptible to Rad50Δ−-induced malignancy.

Discussion

We showed previously that alteration of either invariant cysteine of the S. cerevisiae Rad50 hook domain results in complete inactivation of Rad50, limiting the ability to distinguish hook-specific from general Mre11 complex functions [Hopfner et al. 2002]. The residues between the cysteines are also nearly invariant. The first is either Pro (85%) or Tyr (10%), and the second position contains Leu or Val in 80% of the cases [Stracker and Petrini 2011]. In this study, the functional significance of the Rad50 hook domain in the mammalian Mre11 complex was examined through the generation of mouse mutants affecting those residues. Neither of the hook mutants encoded gene products that support viability at the organismal or cellular level, indicating that, as in S. cerevisiae, the hook domain is required for Mre11 complex function in mammals. In the heterozygous configuration, we found that Rad50Δ− and Rad50Δ− behaved as dominant gain-of-function alleles that promoted chronic DDR signaling. These outcomes at the molecular level were associated with severe phenotypes in vivo, including embryonic lethality, hydrocephalus, defects in the development and maintenance of primitive cells of the bone marrow and germine, and increased incidence of liver neoplasms.

Rad50Δ− is not a loss-of-function allele

Despite the phenotypic severity observed, the data are inconsistent with Rad50Δ− being a null allele. First, the phenotypes in Rad50Δ−/Δ cells are milder than Rad50Δ−/Δ. Although perdurance of wild-type Rad50 protein following Cre-mediated deletion in Rad50Δ−/Δ cultures cannot be excluded, Rad50Δ− may retain limited functionality. Second, Rad50Δ− exerts dose-dependent dominance; the phenotype of Rad50Δ−/Neo46, in which expression of the mutant gene is lower than in Rad50Δ−/Δ, was less severe. In contrast, Rad50Δ−/Δ mice and cells are indistinguishable from wild-type [CF Bender, CA Adelman, and JHJ Petrini, unpubl.]. Hence, the observed phenotypes do not simply reflect reduced dosage of wild-type Rad50. Finally, genetic ablation of Rad50 in the liver is not associated with detectable pathology [Adelman et al. 2009], whereas liver neoplasia and liver tumors were common in Rad50Δ−/Δ mice.

Given that mutations in Rad50Δ− and Rad50Δ− affect one of the Mre11 complex’s dimerization domains, it is likely that the dominant behavior of these alleles results from interaction of the mutant gene product with the wild-type Rad50 protein. Supporting this hypothesis, we found that Rad50WT and Rad50Δ− hook domains interact in vitro [Supplemental Fig. S7]. In principle, Rad50Δ−/Δ...
cells could contain wild-type homodimers, mutant/wild-type heterodimers, and mutant homodimers. The Rad50+/46 phenotypes observed are likely attributable to heterotopic complexes. Several lines of evidence indicate that alterations of the hook domain exert global effects on Mre11 complex function [Hopfner et al. 2002; Wiltzius et al. 2005; Hohl et al. 2011]. The data presented here further suggest that these effects can be exerted in trans within the dimeric assembly of the complex.

Chronic DDR signaling

Rad50+/46 and Rad50+/47 cells exhibit constitutive γH2AX formation. However, Rad50+/46 cells exhibited only a modest reduction in HDR efficiency and a slight sensitivity to camptothecin and did not show increased spontaneous chromosomal aberrations. These facts are inconsistent with spontaneous DNA damage as the cause of γH2AX formation and suggest the possibility that Rad50+/46 behaves as a hypermorph with respect to DNA damage signaling, as previously observed for the Rad50S allele [Usui et al. 2006]. Although the underlying mechanisms of Mre11 complex-dependent DDR regulation remain unclear, the increased latency of lymphomas in Rad50+/46 Atm−/− mice supports this interpretation.

The Rad50+ and Rad50− mutations affect domains widely separated yet engender similar DDR signaling phenotypes [Rad50S is the K20M substitution]. AFM studies have shown that DNA binding by the globular domain promotes spatial reorganization of the coiled-coil and hook domain [Moreno-Herrero et al. 2005]. Conversely, Rad50+/46 and Rad50− mutations as well as mutations that truncate the Rad50 coiled-coil domain affect functions that are likely specified within the globular domain [Hohl et al. 2011]. These data illustrate the functional interdependence of the hook and globular domains within the Mre11 complex.

The Rad50+/46 phenotype and progenitor cells

Consistent with the interpretation that the Rad5046 allele chronically activates ATM, the phenotypic outcomes in Rad50+/46 mice are dependent on Atm. Again, this situation is comparable with Rad500− mice, and the tissues primarily affected are the same: primitive cells in the hematopoietic and germline lineages. These observations indicate that the Mre11–ATM-dependent signaling pathway is important for the maintenance and function of primitive cells in the bone marrow and germline [Morales et al. 2008; Takubo et al. 2008; Maryanovich et al. 2012]. The Mre11 complex–ATM signaling axis likely affects progenitor cells in other organs systems [Shiloh and Ziv 2013]. For example, hyperpigmentation, common to both Rad50S/S and Rad50+/46, has also been observed in mice with chronic DDR signaling in the skin due to telomere dysfunction [Hockemeyer et al. 2008; He et al. 2009; Martinez et al. 2010] or DNA repair defects [Zhang et al. 2011]. In those instances, melanocyte expansion appears to occur in response to epidermal stem cell depletion [Stout and Blasco 2009].

Tumorigenesis in Rad50+/46 mice

Roughly a quarter (26.7%) of Rad50+/46 mice developed hepatocellular adenomas that were frequently coincident with additional preneoplastic lesions, primarily hyperplastic nodules and areas of disorganized architecture. The Mre11 complex–ATM pathway has been previously implicated in liver cancer, as Atm is required for diethylnitrosamine-induced liver carcinogenesis [Teoh et al. 2010]. Notably, almost one-third of the tumor-bearing mice developed hepatoblastoma, a rare tumor that in humans primarily affects infants and children. The blastic and undifferentiated character of this tumor suggests that it arises from hepatic progenitor cells of the portal areas [Marquardt and Thorgerisson 2010]. This outcome represents an additional instance in which Mre11 complex–ATM signaling affects homeostasis by affecting progenitor cells.

Mre11 complex mutations have been identified in ~4% of human tumors [data from cBioPortal; Cerami et al. 2012]. The observation here of dominant Rad50 mutations promoting tumorigenesis supports the possibility that heterozygous Mre11 complex mutations may underlie tumor development, a possibility that should be considered in the genomic analysis of tumors.

Materials and methods

Rad50-HK expression, purification, and analysis

6xHis-SMT3-tagged Rad50-HK proteins [amino acids 591–773] were expressed in Escherichia coli. Soluble extracts were prepared in HK buffer (200 mM NaCl, 50 mM Tris at pH 7.5, 0.2% NP-40, 10% glycerol, 0.5 mM DTT) plus protease and phosphatase inhibitors and was purified by nickel-chelating affinity chromatography followed by Ulp1 protease treatment and a second nickel-chelating affinity purification step to separate the 6xHis-SMT3 tag. Rad50-HK proteins were analyzed by SEC using a 24-μL Superdex200 column equilibrated with 20 mM Tris [pH 7.6], 500 mM NaCl, 0.1 mM EDTA, and 2 mM DTT.

SEC-MALS was performed as described [Rambo and Tainer 2010] with a size exclusion column equilibrated with 20 mM Tris [pH 7.6], 500 mM NaCl, 0.1 mM EDTA, and 1 mM tris[2-carboxyethyl]phosphine. MALS experiments were performed using an 18-angle DAWN HELEOS light-scattering detector connected in tandem to an Optilab refractive index concentration detector [Wyatt Technology]. Metal concentrations were measured using a quadrupole-based ICP-MS [7500ce, Agilent Technologies] equipped with a MicroMist nebulizer (Agilent Technologies).

Gene targeting and mice

The targeting vectors were constructed using the recombineering method previously described [Liu et al. 2003]. Targeting was carried out via electroporation into the embryonic stem cell AC1 that had a 129/SvEv-background [gift of G. Olz and E. Rhuley]. Selection, clonal expansion, and analysis were performed using previously published procedures [Luo et al. 1999]. Chimeras were generated at the SKI Transgenic Facility using standard methods, and male chimeras were crossed with C57BL6 mice to detect germline transmission. Rad50−Neu46 mice were mated with CAG-Cre transgenic mice [Sakai and Miyazaki 1997] to generate...
the \textit{Rad50}^{+}\textit{Neo}^{+} allele. All mice were maintained on mixed 129/SvEv and C57BL6 background.

Mice were housed in ventilated rack caging in a pathogen-free facility. The Institutional Animal Care and Use Committee of Memorial Sloan-Kettering Cancer Center approved animal use protocols. Atrm^{−/−} and Spo11^{−/−} mice were previously described (Barlow et al. 1996, Baudat et al. 2000).

\textbf{Cellular assays}

MEFs and EFs were generated, cultured, and immortalized as described (Bender et al. 2002). \textit{Rad50}^{−}\textit{Neo}^{−} chimeric mice by growing the cells in medium containing 1 mg/mL G418 followed by lentiviral delivery of Cre and clonal selection.

Western blots were carried out on 40 μg of protein extracted with NTEN (20 mM Tris at pH 8, 150 mM NaCl, 1 mM EDTA, 0.5% NP-40) plus protease and phosphatase inhibitors. For H2AX blots, we used 3 μg of the histone fraction extracted with 0.1 M HCl after a pre-extraction with NTEN. All of the antibodies were incubated overnight at 4°C in 5% milk. The antibodies used were rabbit anti-Rad50 polyclonal (custom Petrini laboratory antibody m84-7), rabbit anti-Nbs1 polyclonal (custom Petrini laboratory antibody 92-5), rabbit anti-Mre11 polyclonal (custom Petrini laboratory antibody 42-5), ATM (Cell Signaling), ATM pS1981 (Cell Signaling), γH2AX (Millipore), H2AX (Bethyl Laboratories), Kap1 pS824 (Abcam), Kap1 (Novus Biologicals), Chk2 (Clone 7, Millipore), and Ku70 (Santa Cruz Biotechnology).

For Nbs1 immunoprecipitation, cell extracts of SV40-immortalized MEFS were prepared with lysis buffer A (300 mM NaCl, 20 mM Tris at pH 8.0, 0.5% sodium deoxicholate, 1% Triton X-100, 5 mM MgCl2 supplemented with protease and phosphatase inhibitors). Total protein (0.5 mg) was incubated with 2 μL of anti-Nbs1 antiserum and protein A-agarose beads (Amersham) for 2 h at 4°C, washed extensively, and eluted with Laemmli’s buffer.

For colony-forming assays, cells were treated with the indicated clastogens and plated in triplicate, and colonies were assessed 10 d later by staining with crystal violet.

Metaphases were prepared from cultures treated with 2 x 10^{-7} M colcemid for 1 h. Cells were harvested, hypotonically swollen with 0.075 M KCl for 15 min at 37°C, fixed, washed in ice-cold 3:1 methanol:acetic acid, and dropped on slides. Slides were stained with 5% Giemsa (Sigma) for 10 min and rinsed with distilled water, and coverslips were mounted with Permount (Fisher). Images were captured using an Olympus IX60 microscope and imaged with a Hamamatsu CCD camera. More than 25 spreads were scored for each sample.

G2/M checkpoint was done as previously described (Theunissen and Petrini 2006).

Lentiviral production, concentration, and titering were carried out using previously described methods (Dull et al. 1998; Lois et al. 2002). For lentivirus-Cre and lentivirus-GFP infections, 5 x 10^{5} cells were resuspended in 3 mL of DMEM supplemented with 10% CCS containing 5 μg/mL polybrene and lentivirus at a multiplicity of infection (MOI) 50. Twenty-four hours after infection, fresh medium was added, and cells were trypsinized and replated every 3 d to avoid confluency.

\textbf{DR-GFP assay}

\textit{Pim}^{1}\textit{DR-GFP}\textit{CRE} mice (Kass et al. 2013) were crossed with \textit{Rad50}^{−}\textit{GFP} and heterozygous \textit{Pim}^{1}\textit{DR-GFP} \textit{Rad50}^{−}\textit{GFP} and \textit{Pim}^{1}\textit{DR-GFP} \textit{Rad50}^{−}\textit{GFP} primary skin fibroblast cultures were established. To deliver I-SceI and monitor only the I-SceI-expressing cells, we inserted and IRES-hCD4 cassette into the retroviral vector pMX-Iscel (gift from Fred Alt). I-SceI-hCD4-expressing retrovirus produced in Phoenix cells were used to infect primary cultures in the presence of 5 μg/mL polybrene for 24 h. Seventy-two hours after the infection, cells were harvested, stained with anti-CD4 PE-Cy5 (BD-Pharmingen), and analyzed by flow cytometry. The number of GFP-positive cells was expressed as a ratio to the CD4-positive [infected] cells.

\textbf{Hematopoietic lineage analysis}

Hematopoietic lineages were determined as previously described (Brenet et al. 2013). Briefly, bone marrow single-cell suspensions prepared from both back legs were depleted of lineage-positive cells using a lineage cell depletion kit (Miltenyi Biotech) and simultaneously stained with different antibodies. Hematopoietic populations were identified as follows: long-term HSCs, LKS CD34+ Flk2+; short-term HSCs, LKS CD34+ Flk2−; MPPs, LKS CD34− Flk2−; CMPs, LKS CD34− Flk2−; granulocyte macrophage progenitors, LKS CD34− FcγR−; and megakaryocyte erythroid progenitors, LKS CD34− FcγR+. The CLP population was defined as LKS Flk2+, IL7Rα+. DAPI was used to exclude dead cells during flow cytometric analysis.

MPPs were enumerated using the day 13 CFU-S assay. In brief, recipient mice were lethally irradiated and injected with bone marrow cells. Spleens were isolated 13 d after transplantation and fixed in Bouin’s solution, and macroscopic colonies were counted.

\textbf{Histological sample preparation, staining, and analysis}

Tissue samples for histological analyses were fixed overnight at 4°C with 4% paraformaldehyde, rinsed, stored at 4°C in 70% ethanol, and then processed for paraffin embedding. Eight-micrometer sections were prepared, and slides were processed and stained at the Memorial Sloan-Kettering molecular cytogenetics core facility for hematoxylin and eosin (H&E), TUNEL, anti-VASA, anti-SYCP3, and anti-PLZF.

H&E-stained and immunohistochemically stained slides were digitally scanned using a Micra scanner. TUNEL was quantified relative to the total area using standardized color filter settings in the Metamorph image analysis software. Anti-PLZF- and anti-SYCP3-positive cells per tubule were manually scored from at least 25 tubules per testis.

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