Sensing of intermediates in V(D)J recombination by ATM

Eric J. Perkins, Ayyappan Nair, Dale O. Cowley, Terry Van Dyke, Yung Chang, and Dale A. Ramsden

1Curriculum in Genetics and Molecular Biology, 2Lineberger Comprehensive Cancer Center, and 3Department of Biochemistry and Biophysics, University of North Carolina at Chapel Hill, North Carolina 27599, USA; 4Department of Microbiology, Arizona State University, Tempe, Arizona 85287, USA

Ataxia-telangiectasia mutated (ATM) is required for resistance to radiation-induced DNA breaks. Here we use chromatin immunoprecipitation to show that ATM also localizes to breaks associated with V(D)J recombination. ATM recruitment to the recombining locus correlates approximately with recruitment of the break-initiating factor RAG1 and precedes efficient break repair, consistent with localization of ATM to normal recombination intermediates. A product of ATM kinase activity, Ser 18-phosphorylated p53, was detected similarly at these breaks, arguing that ATM phosphorylates target proteins in situ. We suggest routine surveillance of intermediates in V(D)J recombination by ATM helps suppress potentially oncogenic translocations when repair fails.

Received October 25, 2001; revised version accepted November 28, 2001.

Ataxia-telangiectasia is characterized by cerebellar degeneration, immunodeficiency, and a high frequency of malignancy, usually lymphoid in origin [Morell et al. 1986]. The gene mutated in this disease [ataxia-telangiectasia mutated, or ATM] is required for resistance to DNA double-strand break (DSB) inducing agents such as ionizing radiation, and is an important trigger for the cellular response to DSBs (for review, see Rotman and Shiloh 1999). ATM may act as a primary sensor of DSBs, first binding directly to DSBs and consequently activating through phosphorylation several downstream effectors of the DNA damage response, including Nbs1, BRCA1, Chk2, and p53 [for review, see Durocher and Jackson 2001]. For example, after treatment with ionizing radiation, phosphorylation of p53 at Ser 15 [Ser 18 in mice] in cells is largely ATM dependent [for review, see Giaccia and Kastan 1998] and is required for effective p53-dependent responses to this stimulus [Chao et al. 2000].

In contrast to its important role in mediating resistance to exogenous DSB-inducing agents, genetic evidence for a role for ATM in repair of DSB intermediates in V(D)J recombination is less clear. V(D)J recombination is an essential step in the generation of a diverse repertoire of immunoglobulins and T-cell receptors (for review, see Gellert 1997). RAG1 and RAG2 proteins initiate V(D)J recombination by introducing DSBs precisely adjacent to recombination-targeting signals that flank segments of immunoglobulin and T-cell receptor coding sequence. Efficient resolution of broken species requires factors implicated in end-joining DSB repair, including Ku, XRCC4, ligase IV, and the ATM-related kinase DNA-PKcs, but not ATM. Cells from patients with AT support normal levels of V(D)J recombination using an extra-chromosomal substrate assay (Hiiesh et al. 1993), and mature antigen receptor-bearing lymphocytes are readily observed in ATM-deficient mice [Barlow et al. 1996, Elson et al. 1996, Xu et al. 1996].

Are V(D)J recombination intermediates recognized by ATM as DNA damage? The lack of requirement for ATM in V(D)J recombination might suggest that ATM is excluded from breaks associated with this pathway, possibly through masking of ends by RAG1 and RAG2, to avoid counter-productive apoptotic responses during this normal cellular process. However, both AT patients (for review, see Taylor et al. 1996) and ATM-deficient mice (Barlow et al. 1996, Liyanage et al. 2000, Xu et al. 1996) are prone to lymphoid malignancies that harbor translocations involving antigen receptor genes. Moreover, ATM-deficient mice no longer develop tumors with such translocations when V(D)J recombination is blocked, due to deficiency in RAG1 or RAG2 (Liao and Van Dyke 1999, Petiniot et al. 2000). Genetic evidence thus indicates that although ATM is normally not required for V(D)J recombination, this factor does play an important role in protection against tumors caused by aberrant V(D)J recombination.

However, the molecular basis for these latter observations is unknown. We therefore used chromatin immunoprecipitation (for review, see Orlando 2000) to show that both ATM and a product of ATM kinase activity, Ser 18-phosphorylated p53, are recruited to DSBs associated with V(D)J recombination. Our data further provide a biochemical basis for a model in which ATM monitors the repair of intermediates in V(D)J recombination, and subsequently helps protect against aberrant recombination when repair fails.

Results

ATM and Ser 18-phosphorylated p53 localize to V(D)J recombination-associated breaks

We initially used a temperature-sensitive Abelson Murine Leukemia virus transformed (ts-Ab-MuLV) cell line as a source of cells undergoing high levels of V(D)J recombination [Chen et al. 1994; Chang and Brown 1999]. Culture of a ts-Ab-MuLV at the nonpermissive temperature induces high levels of RAG protein expression, which in turn mediates breakage of recombination signals at endogenous immunoglobulin k loci [Igk] [Chen et al. 1994; see also Fig. 3A,B, below]. We analyzed breaks that retain the recombination signal [signal ends], as these breaks are longer lived than the ends of coding segments, but are nevertheless efficiently resolved into
junctions and are thus normal recombination intermediates (Ramsden and Gellert 1995). Moreover, because the majority of functional V gene segments in the κ locus recombine by inversion (Thiebe et al. 1999) resolution of signal ends at this locus is typically required for maintenance of chromosomal integrity.

Proteins that bind to these breaks were recovered by immunoprecipitation with appropriate antibodies, and the associated DNA was detected by PCR analysis. Antibodies to RAG1 and DNA-PKCs efficiently recovered a 300-bp DNA fragment adjacent to Jk1, a site of active V(D)J recombination in this cell line (Fig. 1A, B). A similar localization to Jk1 was observed by use of antibodies specific to other factors implicated in the end-joining pathway [Ku, XRCC4, E.J. Perkins and D.A. Ramsden, unpubl.], but not control antibody preparations [IgG1, normal rabbit serum; all ChIP figures] or antibodies to other ubiquitous DNA-binding proteins (TFIID, Sp1; E.J. Perkins and D.A. Ramsden, unpubl.).

Importantly, antibodies to ATM also recovered the Jk1 species (Fig. 1B). Similar results were obtained with three different ATM monoclonal antibodies when used separately (Fig. 1C; more quantitative recovery of ATM-associated DNA was observed using a pool of two of these monoclonals). All antibody preparations precipitated negligible amounts of DNA species from several different loci not undergoing V(D)J recombination. This included a gene active in transcription but not V(D)J recombination [the RAG1 gene, Fig. 1B], an antigen receptor gene not active for recombination in this cell type (T-cell receptor Jα50, Fig. 1C), as well as a region near Ck1 but 2.5 kb distal to a site of recombination; E.J. Perkins and D.A. Ramsden, unpubl.). We conclude RAG1, ATM, and DNA-PKCs specifically localize near actively recombining DNA.

We further characterized immunoprecipitated DNA by ligation-mediated PCR (LMPCR), which specifically detects blunt DSBs by ligation of a small double-stranded DNA linker to DNA ends (Fig. 1A). Results using LMPCR paralleled the previous analysis using the 5’ of Jk1 product (Fig. 1B, cf. Jk1 DSBs with 5’ of Jk1). Localization of RAG1, DNA-PKcs, and ATM to this region thus occurs specifically at DSBs. The greater specificity of LMPCR for actively recombining DNA also allowed us to determine that Ser 18-phosphorylated p53 [S18P-p53] also localizes to DSBs at Jk1 [Fig. 1B]. Similar results were obtained by use of an antibody that recognizes p53 independent of phosphorylation status (E.J. Perkins and D.A. Ramsden, unpubl.). No p53 consensus binding sites can be found near these breaks, arguing localization is not sequence dependent. Instead, our data is consistent with prior observations indicating p53 associates with ATM (Khanna et al. 1998), and that phosphorylation of p53 on Ser 18 is largely ATM dependent in response to ionizing radiation (for review, see Giaccia and Kastan 1998).

Because localization of ATM to breaks associated with V(D)J recombination could be unique to our transformed cell line model, we assessed the ability of ATM to localize to DSBs at a T-cell receptor α locus recombination signal [Jα50] in thymocytes from 1-week-old mice. Antibodies to ATM recovered levels of Jα50 DSBs comparable with that recovered by the RAG1 antiserum, whereas an isotype-matched control monoclonal antibody did not recover this species [Jα50 DSBs, Fig. 2A]. Again, neither RAG nor ATM antibodies recovered significant amounts of DNA from loci not active in V(D)J recombination, including the RAG1 gene as well as the 5’ of Jk1 region [active only in B cells]. In a parallel experiment performed using thymocytes from ATM-deficient littermates, Jα50 DSBs were still recovered by antibodies to RAG1 but not with antibodies to ATM, further confirming the specificity of the ATM antibodies in this assay [Fig. 2B]. ATM therefore also localizes to V(D)J recombination-associated DSBs at the T-cell receptor α locus in primary thymocytes.

**ATM localization is consistent with recruitment to normal intermediates**

Unresolved signal ends accumulate in both ts-Ab-MuLV cells (Ramsden and Gellert 1995) and primary lymphocytes (Schlissel et al. 1993; Zhu and Roth 1995) upon induction of V(D)J recombination. However, analysis of signal ends in the ts-Ab-MuLV cell model has shown

**Figure 1.** ChIP analysis of Jk1 breaks from ts-Ab-MuLV cells. (A) Diagram of κ locus, with recombination signals [triangles] and coding segments [rectangles] marked. The location of PCR primers [arrows] and PCR products [bars] at bottom. (B) ChIP analysis of extracts from ts-Ab-MuLV cells induced for 2 d. Comparison of PCR amplified, serial dilutions of input DNA template to DNA template immunoprecipitated [IP’d DNA] by antibodies to different proteins. [NRS] Normal rabbit serum control; [S18P-p53] Ser 18-phosphorylated p53; [ATM] a pool of 9E3C8 and 2C1 antibodies. The same DNA samples were used as template to assess levels of three different DNA species, nonspecific locus [RAG1 gene], 5’ of Jk1, and Jk1 DSBs, as noted. (C) ChIP analysis with different ATM antibodies. The ability of different ATM monoclonal antibodies, used separately, to recover Jk1 DSBs or a nonspecific locus (5’ of Jα50) was assessed.
that these ends nevertheless resolve efficiently to signal junctions if cells are returned to pre-induction conditions, and are thus intermediates in the recombination pathway (Ramsden and Gellert 1995). We made use of the ability to control accumulation and resolution of signal ends to distinguish among the following possibilities: (1) ATM localization actually precedes chromosomal breakage, (2) ATM localizes only to breaks long after induction, suggestive of recruitment to a subpopulation of breaks whose resolution has been inhibited, or (3) ATM localization is approximately coincident with break induction and localization of the break-generating factor RAG1, consistent with recruitment to normal DSB intermediates in V(D)J recombination.

Western analysis showed that changes in levels of Ser 18-phosphorylated p53 paralleled the accumulation and resolution of signal ends over the entire time course (Fig. 3, cf. S18Pp53 in A with Jk1 DSBs in B). Localization of ATM in V(D)J recombination

Western analysis shows that total levels of ATM remain constant throughout the time course [Fig. 3A]. Although levels of ATM do not change, this factor nevertheless clearly localizes to Jk1 after 12 h of induction [Fig. 3C]. Thus, there is no pronounced delay in localization of ATM to breaks, relative either to the appearance of breaks or the localization of the break-initiating factor RAG1. We observed a reduction in localization of ATM after 24 h relative to earlier time points, even though localization of RAG1 continues to rise in parallel with the accumulation of breaks. Localization of ATM to DSBs at Jk1 is also no longer observed after return of these cells to pre-induction conditions for 24 h [Fig. 3C]. ATM recruitment does not precede breakage, nor is ATM recruited only long after V(D)J recombination-induced breaks have been generated. Our ChIP analysis does not quantitatively recover ATM-associated DNA, thus we cannot exclude the possibility that ATM is localizing to a rare subpopulation of breaks that cannot be repaired. However, our results are most consistent with recruitment of ATM to normal intermediates in the pathway.

Western analysis showed changes in levels of Ser 18-phosphorylated p53 paralleled the accumulation and resolution of signal ends over the entire time course (Fig. 3, cf. S18Pp53 in A with Jk1 DSBs in B). Localization of ATM in V(D)J recombination.

Figure 2.  ChIP analysis of Jk50 breaks from mouse thymocytes. (A) Comparison of input template to template recovered after immunoprecipitation [IP'd DNA] of thymocyte extract with antibodies to RAG1, ATM (pool IP used both 2C1 and ATXO8 IgG1 antibodies), and an isotype matched control (IgG1). The same DNA samples were used as template to assess levels of nonspecific loci [RAG1 gene, 5′ of Jk1] and Jk50 DSBs. (B) ChIP analysis of ATM-deficient thymocyte extract with antibodies to RAG1 and ATM [using a pool of 2C1 and ATXO8].

Figure 3. Correlation of V(D)J recombination activity with factor localization. Cells were harvested before induction (L), after 12 h (H12) and 24 h (H24) of induction, and after 24 h of induction followed by 24 h without induction (HL). [H20] A PCR control with water substituted for template. [A] Western analysis of various proteins, as noted at left. [B] PCR analysis of different input DNA species, as noted at right. [S18Pp53] Vj1 signal junctions. Note that only a small subpopulation of total signal junctions are detected by this PCR as sequences 3′ of the ~100 different Vβ segments are not conserved. [C] ChIP analysis of the localization of different proteins or the nonspecific NRS control, as noted at left. PCR reactions in B and C were shown to be in the linear range by serial dilution of the H24 sample (E.J. Perkins and D.A. Ramsden, unpubl.).
Ser 18 p53 to Jκ1 DSBs, as determined by ChIP analysis, roughly paralleled that of ATM, in that it was observed only during induction (Fig. 3C).

Phosphorylation of p53 at Ser 18 is enhanced in a scid cell line

ATM and DNA-PKcs both localize to signal ends [Fig. 1B] and both phosphorylate p53 at Ser 18 in vitro [for review, see Giaccia and Kastan 1998], so either kinase could potentially mediate this phosphorylation event at V(D)J recombination intermediates. We therefore examined levels of Ser 18-phosphorylated p53 in ts-Ab-MuLV cells derived from a scid mouse, deficient in DNA-PKcs [Chang and Brown 1999], and compared these findings with our previous results from wild-type ts-Ab-MuLV cells. Upon induction of V(D)J recombination, levels of phosphorylation at Ser 18 of p53 were >10-fold higher in the scid cell line [Fig. 4A]. Recruitment of Ser 18-phosphorylated p53 to breaks in scid cells was also much more readily detectable relative to previous experiments with wild-type cells [cf. levels of S18P p53 relative to ATM or RAG1 in Fig. 1B with Fig. 4B]. DNA-PKcs is therefore not required for this event, as phosphorylation of p53 at Ser 18 instead appears to be much more efficient in the DNA-PKcs-deficient cell line. Similar results were obtained previously by comparing responses of wild-type and DNA-PKcs-deficient cells in response to ionizing radiation [Araki et al. 1999], arguing that the enhanced phosphorylation of p53 at Ser 18 in scid cells observed here is not unique to this cell line model.

Discussion

We observe localization of RAG1 to signal ends generated at an endogenous locus active in V(D)J recombination. This supports previous work, using artificial recombination substrates, that shows RAG1/RAG2 complexes remain bound to signal ends after they complete signal-targeted cleavage [Agrawal and Schatz 1997; Hiom and Gellert 1998]. DNA-PKcs, an end-joining factor primarily linked to resolution of coding end intermediates in V(D)J recombination, was also shown to localize to signal ends. However, localization of DNA-PKcs to signal ends is not completely unexpected, as deficiency in DNA-PKcs also results in a mild, but significant, impairment of signal end resolution [Lieber et al. 1988; Bogue et al. 1998; Fukumura et al. 2000].

Most importantly, we observed ATM at V(D)J recombination-associated breaks, both in the Igκ locus in a pre-B cell line, as well in the T-cell receptor α locus in primary mouse thymocytes. Recruitment of ATM to the recombining locus in the pre-B cell line was roughly coincident with both break induction and recruitment of RAG1, and preceded efficient resolution of breaks to repaired junctions. These results are consistent with recruitment of ATM to normal recombination intermediates.

We also observed Ser 18-phosphorylated p53 at breaks associated with V(D)J recombination. p53 physically associates with ATM [Khanna et al. 1998] and phosphorylation of p53 at Ser 18 is ATM dependent in response to other sources of double-strand breaks [for review, see Giaccia and Kastan 1998]. Our data, taken together with these previous observations, argues that ATM is not only recruited to V(D)J recombination intermediates, but also responds to these breaks by phosphorylating associated molecules of p53. This is consistent with models in which ATM is a direct sensor of DNA damage [for review, see Durocher and Jackson 2001], and suggests that transduction of ATM-dependent signals may typically be initiated at the site of DNA damage.

Localization of an activated form of p53 at breaks associated with V(D)J recombination is consistent with genetic data. p53 deficiency increases the frequency of malignancy linked to aberrant V(D)J recombination in mice also deficient in genes required for end-joining [Vanasse et al. 1999; Difilippantonio et al. 2000, Frank et al. 2000]. However, comparison of mice deficient in either p53 or ATM as well as mice deficient in both genes indicates the roles of these genes in suppressing tumorigenesis are overlapping, but not congruent [Westphal et al. 1997; Xu et al. 1998]. Two other putative targets of ATM kinase activity, NBS1 and the histone H2AX, have been associated with chromatin actively undergoing V(D)J recombination [Chen et al. 2000]. A significant role for these factors [or other ATM targets] in addition to p53 may be required for ATM's full protective effect.

How does ATM distinguish between breaks that are normal intermediates from breaks that are precursors to aberrant rearrangements? ATM may be recruited to all V(D)J recombination DSB intermediates, but ATM kinase activity (and consequent activation of its downstream effectors) is normally attenuated when these breaks use the end-joining pathway for repair (Fig. 5). However, the enhanced phosphorylation of p53 in DNA-PKcs-deficient cells [Fig. 4, Araki et al. 1999] suggests that if a complete end-joining complex is not assembled, the ATM kinase remains active, resulting in more effective triggering of downstream effectors [p53, NBS1, H2AX, etc.]. This model can explain genetic data indicating that ATM is dispensable for normal V(D)J recom-

Figure 4.  p53 phosphorylation in scid cells.  (A) Samples from a wild-type/scid heterozygous ts-Ab-MuLV cell line [wt/scid] and a scid homozygous ts-Ab-MuLV cell line [scid/scid] both before (L) and after induction of V(D)J recombination for 2 d [H48] were compared by Western analysis. [S18P-p53] Ser 18-phosphorylated p53.  (B) Extracts from scid ts-Ab-MuLV cells were analyzed by ChIP after 2 d of V(D)J recombination induction, as also performed for wild-type cells in Fig. 1C. The same DNA samples were used as template to assay levels of nonspecific loci [RAG1 gene, 5' of Jκ50, Cκ], and Jκ1 DSBs, as noted.
ATM in V(D)J recombination

Materials and methods

Antibodies
Three mouse IgG1 monoclonals recognizing ATM were used as follows: 9E3C8 (Ab-8, NeoMarker), ATX08 (Ab-8, NeoMarker), and 2C1 (GeneTex). A rabbit polyclonal antiserum was used to detect Ser 18-phosphorylated p53 (Cell Signaling Technology) and the monoclonal Ab421 [Ab-1, Oncogene Research] was used to detect p53δ independent of phosphorylation status. We verified the specificity of these antibodies in parallel immunoprecipitation experiments using extracts from the same cells, the same antibody formulations, and using the same conditions as used for ChIP analysis, except adapted for visualization of protein by Western analysis (E.J. Perkins and D.A. Ramsden, unpubl.). With respect to p53 antibodies, with this antibody, roughly equivalent amounts of p53δ were recovered from an induced scid cell extract by both PAb421 and the antibody specific to Ser 18-phosphorylated p53, whereas only Ab421 recovered p53δ from an extract sample treated with λ serine phosphatase (New England Biolabs; E.J. Perkins and D.A. Ramsden, unpubl.). A pool of monoclonals recognizing DNA-PKcs (Neomarkers), RAG-1 (PharMingen), and Ku (the gift of David Schatz, Yale University, New Haven, CT) were also used. Purified Mouse IgG1 (Pharmingen), normal rabbit serum (Sigma), antisera recognizing Sp1 [PEP2, Santa Cruz], a monoclonal recognizing TFIID (SI-1, Santa Cruz), and a monoclonal recognizing β tubulin [TUB 2.1, Sigma] were used as controls.

Cells and ChIP analysis
The wild-type/scid heterozygote SP9 and scid/scid homozygote S4ts-Ab-9 mice were further processed for ChIP analysis as described previously (Chang and Brown 1999). Western analysis verified that levels of DNA-PKcs were >50-fold lower in the scid cell line compared with the wild-type cell line, whereas levels of ATM were equivalent (E.J. Perkins and D.A. Ramsden, unpubl.). Levels of DNA-PKcs recruited to J1 breaks in scid cells as determined by ChIP analysis, although detectable, were also reduced >10-fold relative to wild-type cells. Thymocytes were harvested from 7-day-old mice, and single cell suspensions prepared by mincing and stretching through a 100-µm mesh. Mice were either reared from 7-day-old mice, and single cell suspensions prepared by mincing and stretching through a 100-µm mesh. Mice were either

Acknowledgments
We thank M. Gellert, Y. Xiong, B. Weissman, A. Sancar, and M. Backlund for critical reading of this manuscript, as well as D. Schatz (Yale University) for the gift of antibodies that recognize mouse Ku. This work was supported by NIH grant CA 84442-01, ACS grant RPG-99-192-01-GMC, a Leukemia and Lymphoma Society fellowship to D.C. and a Searle Scholar grant to D.A.R.

The publication costs of this article were defrayed in part by payment of page charges. This article must therefore be hereby marked “advertisement” in accordance with 18 USC section 1734 solely to indicate this fact.
both signal and coding joint formation. *Proc. Natl. Acad. Sci.* 95: 15539–15544.
Chang, Y. and Brown, M.L. 1999. Formation of coding joints in V(D)J recombination-inducible severe combined immune deficient pre-B cell lines. *Proc. Natl. Acad. Sci.* 96: 191–196.
Chao, C., Saito, S., Anderson, C.W., Appella, E., and Xu, Y. 2000. Phosphorylation of murine p53 at ser-18 regulates the p53 responses to DNA damage. *Proc. Natl. Acad. Sci.* 97: 11936–11941.
Chen, Y.Y., Wang, L.C., Huang, M.S., and Rosenberg, N. 1994. An active v-abl protein tyrosine kinase blocks immunoglobulin light-chain gene rearrangement. *Genes & Dev.* 8: 688–697.
Difilippantonio, M.J., Zhu, J., Chen, H.T., Meffre, E., Nussenzweig, M.C., Max, E.E., Ried, T., and Nussenzweig, A. 2000. DNA repair protein Ku80 suppresses chromosomal aberrations and malignant transformation. *Nat. Genet.* 25: 510–514.
Durocher, D. and Jackson, S.P. 2001. DNA-PK, ATM and ATR as sensors of DNA damage: Variations on a theme? *Curr. Opin. Cell Biol.* 13: 225–231.
Elson, A., Wang, Y., Daugherty, C.J., Morton, C.C., Zhou, F., Campos-Torres, J., and Leder, P. 1996. Pleiotropic defects in ataxia-telangiectasia protein-deficient mice. *Proc. Natl. Acad. Sci.* 93: 13084–13089.
Frank, K.M., Sharpless, N.E., Gao, Y., Sekiguchi, J.M., Ferguson, D.O., Zhu, C., Manis, J.P., Horner, J., DePinho, R.A., and Alt, F.W. 2000. DNA ligase IV deficiency in mice leads to defective neuorgenesis and embryonic lethality via the p53 pathway. *Mol. Cell.* 5: 993–1002.
Fukumura, R., Araki, R., Fujimori, A., Tsutsui, Y., Kurimasa, A., Li, G.C., Chen, D.J., Tatsumi, K., and Abe, M. 2000. Signal joint formation is also impaired in DNA-dependent protein kinase catalytic subunit knockout cells. *J. Immunol.* 165: 3883–3889.
Gellert, M. 1997. Recent advances in understanding V(D)J recombination. *Adv. Immunol.* 64: 39–64.
Giacca, A.J. and Kastan, M.B. 1998. The complexity of p53 modulation: Emerging patterns from divergent signals. *Genes & Dev.* 12: 2973–2983.
Hiom, K. and Gellert, M. 1998. Assembly of a 12/23 paired signal complex: A critical control point in V(D)J recombination. *Mol. Cell.* 1: 1011–1019.
Hsieh, C.L., Arlett, C.F., and Lieber, M.R. 1993. V(D)J recombination in ataxia telangiectasia, Bloom’s syndrome, and a DNA ligase I-associated immunodeficiency disorder. *J. Biol. Chem.* 268: 20105–20109.
Khanna, K.K., Keating, K.E., Kozlov, S., Scott, S., Gatei, M., Hobson, K., Taya, Y., Gabrielli, B., Chan, D., Lees-Miller, S.P. et al. 1998. ATM associates with and phosphorylates p53: Mapping the region of interaction. *Nat. Genet.* 20: 398–400.
Liao, M.J. and Van Dyke, T. 1999. Critical role for Atm in suppressing V(D)J recombination-driven thymic lymphoma. *Genes & Dev.* 13: 1246–1250.
Lieber, M.R., Hesse, J.E., Lewis, S., Bosma, G.C., Rosenberg, N., Mizuuchi, K., Bosma, M.J., and Gellert, M. 1988. The defect in murine severe combined immune deficiency: Joining of signal sequences but not coding segments in V(D)J recombination. *Cell.* 55: 7–16.
Liyawan, M., Weaver, Z., Barlow, C., Coleman, P., Pankratz, D.G., Anderson, S., Wynshaw-Boris, A., and Ried, T. 2000. Abnormal rearrangement within the alpha/delta T-cell receptor locus in lymphomas from Atm-deficient mice. *Blood* 96: 1940–1946.
Morrell, D., Cromartie, E., and Swift, M. 1986. Mortality and cancer incidence in 263 patients with ataxia-telangiectasia. *J. Natl. Cancer Inst.* 77: 89–92.
Orlando, V. 2000. Mapping chromosomal proteins in vivo by formaldehyde-crosslinked-chromatin immunoprecipitation. *Trends Biochem. Sci.* 25: 99–104.
Petiotin, L.K., Weaver, Z., Barlow, C., Shen, R., Eckhaus, M., Steinberg, S.M., Ried, T., Wynshaw-Boris, A., and Hodes, R.J. 2000. Recombinase-activating gene (RAG) 2-mediated V(D)J recombination is not essential for tumorigenesis in Atm-deficient mice. *Proc. Natl. Acad. Sci.* 97: 6664–6669.
Ramsden, D.A. and Gellert, M. 1995. Formation and resolution of double-strand break intermediates in V(D)J recombination. *Genes & Dev.* 9: 2409–2420.
Rotman, G. and Shiloh, Y. 1999. ATM: A mediator of multiple responses to genotoxic stress. *Oncogene* 18: 6135–6144.
Schlissel, M., Constantinescu, A., Morrow, T., Baxter, M., and Peng, A. 1993. Double-strand signal sequence breaks in V(D)J recombination are blunt, 5’-phosphorylated, RAG-dependent, and cell cycle regulated. *Genes & Dev.* 7: 2520–2532.
Taylor, A.M., Metcalf, J.A., Thick, J., and Mak, Y.F. 1996. Leukemia and lymphoma in ataxia telangiectasia. *Blood* 87: 423–438.
Thiebe, R., Schable, K.F., Bensch, A., Brensing-Kuppers, J., Heim, V., Kirschbaum, T., Mitloehner, H., Ohnrich, M., Pourrajabi, S., Roschenthaler, F., et al. 1999. The variable genes and gene families of the mouse immunoglobulin κ locus. *Eur. J. Immunol.* 29: 2072–2081.
Vanasse, C.J., Hallbrook, J., Thomas, S., Burgess, A., Hoekstra, M.F., Distecher, C.M., and Willerford, D.M. 1999. Genetic pathway to recurrent chromosome translocations in murine lymphoma involves V(D)J recombinase. *J. Clin. Invest.* 103: 1669–1675.
Westphal, C.H., Rowan, S., Schmaltz, C., Elson, A., Fisher, D.E., and Leder, P. 1997. atm and p53 cooperate in apoptosis and suppression of tumorigenesis, but not in resistance to acute radiatio toxicity. *Nat. Genet.* 16: 397–401.
Xu, Y., Ashley, T., Brainerd, E.E., Bronson, R.T., Meyn, M.S., and Baltimore, D. 1996. Targeted disruption of ATM leads to growth retardation, chromosomal fragmentation during meiosis, immune defects, and thymic lymphoma. *Genes & Dev.* 10: 2411–2422.
Xu, Y., Yang, E.M., Brugarolas, J., Jacks, T., and Baltimore, D. 1998. Involvement of p53 and p21 in cellular defects and tumorigenesis in Atm−/− mice. *Mol. Cell. Biol.* 18: 4385–4390.
Zhu, C. and Roth, D.B. 1995. Characterization of coding ends in thymocytes of scid mice: Implications for the mechanism of the V(D)J recombination. *Immunity* 2: 101–112.