The Atr and Atm protein kinases associate with different sites along meiotically pairing chromosomes

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A number of cell-cycle checkpoint genes have been shown to play important roles in meiosis. We have characterized the human and mouse counterpart of the *Schizosaccharomyces pombe* Rad3 protein, named Atr (for ataxia-telangiectasia- and rad3-related), and the protein that is mutated in ataxia-telangiectasia, Atm. We demonstrate that *ATR* mRNA and protein are expressed in human and mouse testis. More detailed analysis of specific cells in seminiferous tubules shows localization of Atr to the nuclei of cells in the process of meiosis I. Using immunoprecipitation and immunoblot analysis, we show that Atr and Atm proteins are ~300 and 350 kD relative molecular mass, respectively, and further demonstrate that both proteins have associated protein kinase activity. Further, we demonstrate that Atr and Atm interact directly with meiotic chromosomes and show complementary localization patterns on synapsing chromosomes. Atr is found at sites along unpaired or asynapsed chromosomal axes, whereas Atm is found along synapsed chromosomal axes. This is the first demonstration of a nuclear association of Atr and Atm proteins with meiotic chromosomes and suggests a direct role for these proteins in recognizing and responding to DNA strand interruptions that occur during meiotic recombination.

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Mitotic cells guard against genetic instability and increase their survival following DNA damage both by direct DNA repair mechanisms and by delaying progression through the cell cycle. Depending on the position of the cell within the cell cycle at the time of irradiation, DNA damage in mammalian cells can effect specific checkpoints that prevent passage from G₁ to S phase, progression through S phase, or passage from G₂ into mitosis. These checkpoints are thought to prevent deleterious events such as replication of damaged DNA or the segregation of fragmented chromosomes during mitosis (Hartwell and Kastan 1994) and therefore assure the accurate transmission of the genetic material. Recently it has been shown that mitotic checkpoint proteins can have a role in meiosis (Kato and Ogawa 1994; Hari et al. 1995; Xu et al., this issue).

In mitotic cells, G₁ arrest following DNA damage is

dependent on the product of the *p53* gene, which is stabilized following DNA damage (Nelson and Kastan 1994). *p53* is a transcription factor that induces production of p21^{CIP1/WAF-1}, a direct inhibitor of cyclin-dependent kinase activity (El-Deiry et al. 1993; Harper et al. 1993; Deng et al. 1995). Loss of G₁ arrest is not associated directly with loss of viability following irradiation in mammalian cells, but appears to be linked to genomic instability, an early event in tumorigenesis (Hartwell and Kastan 1994). Treatment of replicating cells with ionizing radiation transiently prevents progression through S phase by reducing the number of replication initiation events (Lamb et al. 1989). Cells from patients with ataxia-telangiectasia (A-T), an autosomal recessive disease that is characterized by progressive cerebellar degeneration, immune deficiencies, increased incidences of cancer, gonadal abnormalities, sterility, and sensitivity to ionizing radiation (Sedgwick and Boder 1991), have lost the delay in S phase following ionizing radiation and display radiation-resistant DNA synthesis (Painter and Young 1980). The pleiotropic A-T phenotype is thought to be due to an inability of multiple cellular functions to

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be activated in response to certain types of spontaneous and induced DNA damage (for review, see Meyn 1995; Shiloh 1995). For example, A-T cells irradiated in G₁ or S phase accumulate irreversibly in G₂ where they subsequently die from apoptosis, presumably as a consequence of attempting to replicate damaged DNA or entering mitosis with unrepaired chromosomal damage (Meyn 1995; Shiloh 1995). A-T cells irradiated during G₂ fail to pause transiently prior to entering mitosis. Consequently they progress through mitosis with damaged DNA (Beamish and Lavin 1994). In addition, the induction of p53 in irradiated A-T cells is blunted and delayed (Kastan et al. 1992; Canman et al. 1994). Thus, mutations at the A-T locus result in the disruption of at least three checkpoints required for appropriate responses to ionizing radiation and radiomimetic mutagens.

Identification of *ATM*, the gene mutated in A-T, showed that the Atm protein is similar to a family of phosphatidylinositol 3-kinase (PIK)-like kinases (Savitsky et al. 1995). This family, which includes the products of the *TOR1*, *TOR2*, *FRAP*, *TEL1*, *MEC1*, *rad3*, *mei-41*, *ATR* genes, and the DNA-dependent protein kinase catalytic subunit (DNA-PK_{CS}), shows greatest similarity throughout the protein kinase domain (Heitman et al. 1991; Seaton et al. 1992; Cafferkey et al. 1993; Kunz et al. 1993; Helliwell et al. 1994; Kato and Ogawa 1994; Weinert et al. 1994; Greenwell et al. 1995; Hari et al. 1995; Hartley et al. 1995; Bentley et al. 1996). All of these proteins appear to act as early sensors for defects in various cell-cycle transitions or aberrations in chromosomal mechanisms (Carr 1996). The *Saccharomyces cerevisiae* rapamycin-binding proteins, Tor1p and Tor2p, and their mammalian counterparts, Frap and Raft1, play an important role in mediating the G₁ to S transition (Heitman et al. 1991; Cafferkey et al. 1993; Kunz et al. 1993; Brown et al. 1994; Sabatini et al. 1994). Frap, which autophosphorylates on serine, plays a role in downstream signaling to p70^{S6} kinase and cyclin-dependent kinases (Brown et al. 1994).

The product of the *TEL1* gene, which is most highly related to Atm in primary structure (56% identity through the kinase and carboxy-terminal domains), is required for proper telomere maintenance in *S. cerevisiae* (Greenwell et al. 1995) and plays a partially redundant role with the *MEC1* gene product in conferring resistance to DNA damage (Morrow et al. 1995). DNA-PK is required for V(D)J recombination and for double-stranded break repair after ionizing radiation (Jackson and Jeggo 1995). Mice lacking DNA-PK have severe combined immunodeficiency (SCID; Blunt et al. 1995), and a similar equine syndrome is also the result of a DNA-PK deficiency (Wiler et al. 1995). Mec1p from *S. cerevisiae*, Rad3 from *S. pombe*, and mei-41 from *Drosophila melanogaster* appear to be structural and functional homologs that play important roles in mitotic and meiotic cell-cycle checkpoints and DNA repair. The most closely related mammalian protein to these model system checkpoint proteins is Atr. Unlike Frap and DNA-PK, biochemical activity for Atr and Atm previously has been unidentified.

The Rad3, Mec1p, and Mei-41 group mediate several checkpoint responses in mitotic cells and are important for meiosis. Rad3 is involved in S phase and G₂-M cell-cycle arrest following DNA damage. *rad3* mutants fail to arrest in G₂ after UV or ionizing irradiation and continue into mitosis even if DNA synthesis is incomplete (Al-Khodairy and Carr 1992; Jimenez et al. 1992; Rowley et al. 1992). Rad3 possesses an associated protein kinase activity, and site-directed mutations in the protein kinase domain that render the kinase inactive do not complement *rad3* mutant defects, indicating that Rad3 protein kinase activity is required for checkpoint function (Bentley et al. 1996). Similarly, *MEC1* is required for checkpoints at G₂-M (Weinert et al. 1994) and S-phase transitions (Paulowich and Hartwell 1995) and is essential for meiosis (Kato and Ogawa 1994). Finally, *mei-41* mutant flies are highly sensitive to DNA damaging agents such as ionizing radiation, UV radiation, chemical mutagens, and hydroxyurea (Baker et al. 1976; Boyd et al. 1976; Banga et al. 1986), and cells from these mutants have defects in mitotic checkpoints (Hari et al. 1995) and in meiotic recombination (Baker and Carpenter 1972).

The human counterpart of Rad3, Mec1p, and Mei-41, named Atr (for ataxia telangiectasia and *rad3*-related), has been identified (Bentley et al. 1996; Cimprich et al. 1996). Through the conserved kinase and carboxyl terminus, Atr is 53% identical to *Schizosaccharomyces pombe* Rad3, 48% identical to Mei-41, 41% identical to Mec1p, 31% identical to Atm, and 32% identical to Tel1p (Bentley et al. 1996). The sequence of *ATR* is identical to *FRP1*, a clone isolated previously from a Jurkat cDNA library (Cimprich et al. 1996). In *S. pombe*, Rad3 is in a homomultimeric complex (Bentley et al. 1996). Evidence that supports the notion that Atr is a functional Rad3 homolog has been suggested from the observation that Atr can heteromultimerize with Rad3 in vivo and that Atr can complement for *mec1/esr1* UV sensitivity (Bentley et al. 1996).

Signal transduction cascades, initiated by DNA damage or aberrant DNA replication, are thought to mediate the checkpoint responses (for review, see Carr 1996). Checkpoint pathways require three distinct functions: a detector molecule that recognizes abnormal DNA structures, a signaling cascade that amplifies and transmits the information, and effectors that interact with the cell-cycle machinery (Carr 1996). Several candidate proteins for stages of this cascade have been identified, and most of these have mutant sensitivities to DNA-damaging agents. The Atm protein and the Rad3/Mec1p/Mei-41/Atr proteins are likely candidates as detectors of DNA damage or abnormal DNA replication in both mitotic and meiotic cells. Signal amplification could occur through the Rad53p/Cds1 protein kinase (Allen et al. 1994; Sanchez et al. 1996) and effectors might include proteins like p53 for G₁-S transition; replication protein A (RPA) for S phase (Liu and Weaver 1993); and Chk1/Rad27 for G₂-M (Walworth et al. 1993; Al-Khodairy et al. 1994; Walworth and Bernards 1996).

In addition to playing a role in sensing DNA damage or

completion of DNA synthesis in mitotic cells, *MEC1* and *mei-41* play important roles in meiosis (Baker and Carpenter 1972; Carpenter 1979; Kato and Ogawa 1994). Meiosis segregates a complete haploid set of chromosomes to each gamete, and this process requires the negotiation of a complex series of dependent events. Homologous chromosomes must associate, recombine, synapse, and segregate to opposite poles at meiosis I. DNA double-stranded breaks (DSBs) have been linked to the initiation of meiotic chromosomal synapsis in yeast (Petes et al. 1991). However, the events are much less clear in multicellular organisms. Both *mec1* and *mei-41* mutants are defective in meiotic processes (Baker and Carpenter 1972; Kato and Ogawa 1994). Likewise, the gonads of A-T patients and *ATM*-deficient mice have a paucity of germ cells (Sedgwick and Boder 1991; Barlow et al. 1996; Xu et al., this issue), and spermatocytes from *ATM*^{-/-} mice show meiotic arrest in early prophase with severe chromosome fragmentation in aberrant pachytene nuclei (Xu et al., this issue). These observations suggest a link between functions carried out by mitotic checkpoint proteins and functions required for meiosis.

Here we report studies of *Atr* and *Atm*. We demonstrate that *ATR* mRNA and protein are expressed at high levels in mouse testes. Our further studies show that *Atr* is localized to the nuclei of primary spermatocytes, cells that are undergoing meiosis I. We demonstrate that both *Atr* and *Atm* proteins have associated protein kinase activity, consistent with their primary structures. In addition, *Atr* and *Atm* show specific association with chromosomes in cells that are in early meiosis I as demonstrated by antibody localization on surface-spread spermatocytes. Both the *Atr* and *Atm* proteins are present at pairing forks in meiotic prophase as chromosomes synapse; however, they do not colocalize. Instead, they occupy complementary positions: *Atr* localizes along unsynapsed chromosome axes and *Atm* interacts with synapsed axes. This is the first demonstration that these cell-cycle checkpoint protein homologs specifically associate with chromosomal DNA and suggests a role for these *Atm*-like proteins in recognition of meiotic DNA configurations associated with meiotic synapsis.

Results

Expression of ATR mRNA is elevated in testis

In order to understand the role of *Atr*, the expression pattern was established by Northern blot analysis. All human tissues screened expressed a major transcript of ~8.5 kb. Expression levels were highest in testis with lower, but detectable levels, in all other tissues examined (Fig. 1A). As reported previously (Cimprich et al. 1996), *ATR* also shows low-level expression in a variety of cell lines (data not shown). An identical pattern of expression was seen upon probing of a multiple-tissue Northern blot containing mouse tissues with a mouse cDNA probe that showed 95% homology (Fig. 1B). These results demonstrated that the *ATR* mRNA was elevated in testis, and this elevated expression pattern was conserved in mouse and man.

Cell type-specific expression of Atr in testis

To examine the protein product of *Atr* in greater detail, we generated *Atr*-specific antibodies. Antibodies directed toward the protein kinase domain or the carboxyl terminus recognized a protein in G361 and 293T cells (human melanoma and embryonic kidney cell lines, respectively) that migrated at an approximate molecular weight of 250–300 kD (data not shown), consistent with the predicted molecular mass of *Atr*. Both antibodies recognize the same high molecular mass protein on immunoblots and immunoprecipitations, and reactivity with this protein was blocked by preincubation of the antibodies with the antigen against which it was directed (data not shown).

Since mouse testes were more readily obtained than human testes and the protein kinase domain of human and mouse *Atr* are identical (data not shown), the antisera were tested for cross reactivity on an adult mouse testes cell extract. Both antisera recognized a protein of the expected molecular weight (Fig. 2). These results confirm that the *Atr* protein was expressed in mouse testis.

To determine whether *Atr* expression in mouse testis was the result of nonspecific induction or reflected ex-

Figure 1. *ATR* mRNA is elevated in human and mouse testis. (A) Northern blot containing poly(A)⁺ RNA from various human tissues probed with a human *ATR* probe. (B) Northern blot containing poly(A)⁺ RNA from various mouse tissues and mRNA from total embryo at certain times in development probed with a mouse *ATR* probe. (d7) Day 7; (d11) day 11; (d15) day 15; (d17) Day 17. A 5-kb band is the result of incomplete removal of an unrelated probe from previous use of the Northern blot. The lower bands are likely to represent degradation products of *ATR* mRNA because they vary depending on the lot of Northern blot. Approximate molecular weight standards in kb are shown at the left.

Figure 2. Atr protein is present in mouse testis. Two different rabbit antisera generated against Atr recognize the same protein in mouse testis. Immunoblot analysis of mouse testis cell lysate with rabbit antisera 4747 and 4779. (P) Preimmune. Approximate molecular weight standards in kD are shown at the left.

pression in a specific cell type or stage in testis, we probed an immunoblot containing protein isolated from purified populations of adult mouse testis with the anti-Atr antisera (Fig. 3; Table 1). Atr was expressed most highly in pachytene spermatocytes, cells that are undergoing meiosis I, and its level decreased in purified cells from later meiotic stages. Low levels of Atr were detected in the round spermatid fraction. No expression was detected in the condensing spermatids or mature sperm. We interpret these results to suggest that Atr protein is expressed in the early meiotic steps and is being turned over as spermiogenesis proceeds. The small amount of Atr in the round spermatid fraction could be due to bona fide expression in this cell type, to residual protein from an earlier meiotic stage, or to contamina-

Figure 3. Atr protein level varies with the stage of meiosis. Immunoblot analysis of purified populations of spermatogenic cells. Pachytene spermatocytes and round chromatids were at least 85% pure. The condensing spermatid population was 40–50% pure (contaminating material primarily anucleated residual bodies and some round spermatids). Sperm were obtained from the cauda epididymides. Molecular weight standards are shown at the left (in kD).

Table 1. Meiotic expression pattern of mammalian Atr protein

Cell type	Meiotic stage or function	Atr protein
Leydig cells	interstitial cells	–
Basement membrane	surrounds seminiferous tubule	–
Sertoli cell	compartmentalization	–
Spermatogonia	mitotic	–
Primary spermatocytes	M I, recombination, and synapsis	+
Secondary spermatocytes	M II	–
Spermatids	early sperm cell	– ^a
Spermatozoa	sperm cell	–

^aImmunoblot analysis shows Atr in round spermatid fraction. This may reflect expression of Atr in early stages of spermiogenesis or contamination of this fraction with zygotene spermatocytes.

tion of this fraction with zygotene spermatocytes. Regardless, these data demonstrate that Atr expression in testes was restricted to cells that were in meiotic prophase. Atr expression in meiotic cells is consistent with the elevated levels of *MEC1* transcript reported in budding yeast meiosis (Kato and Ogawa 1994), the requirement for Mec1p in establishing a meiotic checkpoint, and the elevated expression of Mei-41 in *Drosophila* ovaries (Hari et al. 1995).

To more precisely define the cell types that express Atr, sections of mouse testes were examined by immunohistochemical staining. Testes contain seminiferous tubules that are mainly composed of the seminiferous epithelium (a complex stratified epithelium containing the spermatogenic cells) and structural Sertoli cells. The spermatogenic cells are at various stages of meiotic differentiation. The most immature cells are peripherally localized nearest to the basement membrane, and as cells undergo spermatogenesis, they move toward the lumen of the tubule. Testicular sperm are released from the seminiferous epithelium into the lumen of the tubule. There are several morphologically distinct stages in spermatogenesis. Spermatogonia are mitotically dividing stem cells that line the basal lamina. Cells undergoing meiosis are the primary spermatocytes (meiosis I) and secondary spermatocytes (meiosis II); those undergoing condensation and differentiation into spermatozoa are early and late spermatids (Fig. 4A). Consistent with the immunoblot analysis, Atr protein was expressed highly in the primary spermatocytes (Fig. 4; Table 1) and is not present in round spermatids. This analysis also enabled us to determine the subcellular localization of Atr. Nuclear Fast Red, a nuclear counterstain, colocalized with the immunogold stain, establishing the presence of Atr in the nuclei of the cells in meiosis I (Fig. 4B). To our knowledge this is the first localization of a mammalian checkpoint homolog to the nuclei of early meiotic cells.

These results suggest that Atr may be involved in establishing checkpoints in meiotic cells.

Atr and Atm have associated protein kinase activity

While the structure of the kinase domain in the Rad3 subgroup of proteins is related to the lipid kinases, proteins with this domain function as protein kinases. The best-characterized example of this is DNA-PK_{CS}, which phosphorylates a wide range of protein substrates *in vitro* (Anderson and Lees-Miller 1992) but, to date, has not been shown to phosphorylate lipids (Hartley et al. 1995). The lipid kinases, Vps34p (Stack and Emr 1994), the p110 subunit of PI3-kinase (Carpenter et al. 1993; Dhand et al. 1994), and Frap (Brown et al. 1995) have protein kinase activity. Therefore, we tested Atr for protein kinase activity. Immunoprecipitation of an adult mouse testis extract with anti-Atr antisera followed by a

protein kinase assay demonstrated clearly the presence of several phosphorylated proteins (Fig. 5A). The major phosphorylated protein in the reaction comigrated with Atr and was blocked by the fusion protein to which the antisera were made. Other phosphorylated species are likely to represent phosphorylation of proteins present in the immune complex. Phosphoamino acid analysis revealed serine phosphorylation (data not shown). These results demonstrate that Atr, like *S. pombe* Rad3 (Bentley et al. 1996), has associated protein kinase activity.

Because we wanted to compare and contrast the functions of Atr with Atm, we generated an Atm-specific antibody. This antisera, 6076, was reactive with a ~350-kD protein in immunoprecipitates of [³⁵S]methionine-labeled MRC-5 fibroblasts, and this species was blocked with the immunizing peptide P45 (Fig. 5B). In comparisons, the Atm protein migrates ~50 kD larger than Atr from ³⁵S-labeled lysates (E. Christenson, unpubl.), con-

Figure 5. Atr and Atm have associated kinase activity. (A) Atr has associated kinase activity. An immunoprecipitation of a mouse testis cell lysate with either a nonimmune rabbit IgG (RbIgG) or 4779 was carried out followed by a protein kinase assay. The phosphorylated protein indicated by the arrow was shown by immunoblot analysis to comigrate with Atr. (B) The anti-Atm peptide antibody recognizes a ~350-kD molecular weight protein in MRC-5 cells. Immunoprecipitation of [³⁵S]methionine-labeled human diploid fibroblasts was performed using rabbit IgG (as a negative control), the Atm antisera, 6076, or 6076 preincubated with 0.04 mg of the peptide against which it was made (P45). (C) Atm has an associated protein kinase activity. Immunoprecipitation with rabbit IgG, 6076, or 6076 preincubated with the peptide against which the antisera was made (P45), washed and incubated with ³²P-γATP in the presence of kinase buffer. The numbers at the left signify the approximate molecular weights of protein standards (in kD). (D) His-tagged Atr expressed in *S. cerevisiae* shows protein kinase activity. His-tagged Atr expression under the control of the *GAL1* promoter was induced by the addition of galactose or repressed by growth in glucose. Either vector alone or vector containing the full-length, his-tagged Atr were grown in the presence or absence of glucose or galactose as indicated by (-) or (+). Atr was purified on Ni⁺-affinity resin and assayed for protein kinase activity (right). The presence of Atr was followed by immunoblot analysis (left). The lower molecular weight proteins are likely to represent degradation products. The number on the left indicates the approximate molecular weight of the standards (in kD).

firming the size difference of Atm and Atr predicted from their open reading frames and the specificity of both Atm and Atr antisera. Furthermore, the anti-Atm sera immunoprecipitates truncated Atm proteins from cell lines derived from A-T patients that contain termination codon mutations (E. Christenson, unpubl.). Immunoprecipitation using anti-Atm antisera followed by a protein kinase assay (Fig. 5C) showed ^{32}P incorporation into the same ~350-kD protein that was immunoprecipitated in [^{35}S]methionine-labeled cells, demonstrating that Atm also has an associated protein kinase activity. This protein kinase activity is missing in A-T cell line AT7LA (Wright et al. 1996), which contains a termination codon mutation amino terminal to the Atm protein kinase domain (E. Christenson, unpubl.).

The immunoprecipitation and protein kinase assays for Atr and Atm reveal that both proteins have an associated protein kinase activity. Both DNA-PK_{CS} and FRAP have been shown to autophosphorylate in the absence of any associated cofactors (Lees-Miller et al. 1990; Dvir et al. 1993; Brown et al. 1995). To extend our analysis of the protein kinase activity of Atr, we expressed this protein in *S. cerevisiae* as a His-tagged protein and purified the protein using Ni^{2+} -affinity resin. As observed for immunoprecipitations from cell extracts, Ni^{2+} -His-affinity purified Atr contains protein kinase activity (Fig. 5D). While these results do not rule out the presence of associated proteins, affinity purification of the His-tagged Atr from yeast demonstrates that protein expressed in a heterologous system also shows kinase activity. Thus we have demonstrated that Atr that has been partially purified by two independent means show an associated kinase activity.

DNA-PK_{CS} has been shown to phosphorylate a wide variety of proteins, including p53, RPA, and casein (Anderson and Lees-Miller 1992). Because there is structural similarity between Atr, Atm, and DNA-PK_{CS}, we tested whether there was any overlap in substrate specificity. Neither immunoprecipitated Atr nor Atm phosphorylated purified RPA, and Atr did not phosphorylate a peptide from p53 that contained the DNA-PK_{CS} phosphorylation site. However, Atr showed a three- to five-fold increase in casein phosphorylation (data not shown), suggesting that there may be some overlap in substrate specificity. These observations suggest that the proposed role for Atm and Rad3-like proteins such as Atr in p53 and RPA effector phosphorylation may occur through an intermediate protein kinase like Rad53p (Carr 1996).

Atr and Atm proteins associate with meiotic chromosomes

If Atm and Atr are involved in establishing the meiosis I checkpoints, it may be possible to elucidate their functions by studying their localization. The size of the mammalian genome, the degree of compaction of the genetic material at meiosis, and the relatively slow rate of progression of spermatocytes through meiosis I allows the spatial and temporal localization of proteins and can indicate the sites and timing of their activity (Ashley et

al. 1995). We have used immunostaining techniques on surface spreads of mouse spermatocytes to localize Atm and Atr to meiotic chromosomes (Figs. 6–8). Rabbit antibodies recognizing Atr and Atm were colocalized with mouse antibodies against Cor1 to assist us in timing and substaging meiotic events. Cor1 is a component of axial/lateral elements of synapsing chromosomes (Dobson et al. 1994). Cor1 chromosomal staining appears when the axial elements begin to form between the sister chromatids of each homolog in leptotema of meiotic prophase, prior to the initiation of synapsis (Dobson et al. 1994). As homologous bivalents synapse, the axial elements from the two homologs align and a central element forms between them, completing the structure called the synaptonemal complex (SC).

When short stretches of Cor1 begin to appear prior to any evidence of synapsis, neither Atr nor Atm is detectable. As homologs start synapsis, both proteins are seen at pairing forks; however, the location and behavior of the two proteins differ markedly. In normal zygotene nuclei, the stage during which homologs synapse, Atr is present in small amounts and transiently at discrete foci along the asynapsed (unpaired) axes. As homologs synapse, Atr disappears from these locations. However, at regions delayed in synapsis, often seen near the proximal ends of autosomal bivalents, there was an accumulation of Atr foci along the unsynapsed axes. Atr is detected at similar locations on the two axial elements. In nuclei where an entire autosome fails to find its homologous pairing partner, Atr foci were detected along the entire lengths of these asynapsed axes. In males, where the X chromosome has no homolog, Atr foci were localized along the unpaired axis (Fig. 8).

Atm is also visualized as foci and is first detected during zygonema as homologs synapse (Fig. 7). The Atm localization is different than Atr (Fig. 8). Atm is first observed along synapsed axes when homologous autosomal axial elements come into contact. However, during mid-pachynema, after autosomal synapsis has been completed, Atm foci appear on the X chromosome axis (data not shown). Atm localization persists on fully synapsed bivalents into pachynema, a substage that lasts 3 days in mouse oocytes and 6 days in mouse spermatocytes. During pachynema, the number of foci drops gradually, stabilizing briefly in mid-pachynema before eventually disappearing mid- to late pachynema. These data indicate important and complementary roles for the Atr and Atm protein kinases at distinct stages in meiosis I.

Discussion

Atr is a mammalian counterpart of Rad3, Mec1p, and Mei-41

We have characterized a mammalian Rad3, Mec1p, Mei-41 and Atm-related protein, Atr. This protein is more homologous to the Rad3, Mec1p, and mei-41 checkpoint proteins than it is to the Atm protein (Bentley et al. 1996; Cimprich et al. 1996), whereas Tel1p is more re-

Figure 6. Atr localization on meiotic chromatin. Immunofluorescence localization of ATR in meiotic prophase of spermatocytes of mouse. *A* and *C* show the localization of Cor1, a component of the axial elements of the synaptonemal complex (SC), and *B* and *D* show the merged Cor1 (white) and Atr (red) images. ATR localizes to sites along unsynapsed axes in zygonema (*B*) and pachynema (*D*). The unpaired axes of a partially synapsed autosome in early pachynema (*D*) have Atr foci, as does the unpaired X axis shown at higher magnification in Fig. 8.

lated to Atm (Bentley et al. 1996). *rad3* and *MEC1* control the G₂-M DNA damage checkpoints in response to UV and ionizing radiation and the S-phase checkpoint following inhibition of DNA replication (Al-Khodairy and Carr 1992; Weinert et al. 1994; Paulovich and Hartwell 1995). In contrast, cells derived from A-T patients are sensitive to a subset of the DNA-damaging agents in comparison to yeast cells that lack Rad3 function (Jimenez et al. 1992; Beamish and Lavin 1994; Bentley et al. 1996). The Atm protein controls the checkpoints that respond to a narrow range of DNA damaging agents, all of which produce DNA strand breaks as a consequence of free-radical attack. The response of A-T cells to UV and certain chemical carcinogens is normal as is the re-

sponse to inhibition of DNA synthesis. In addition, we report that Atr and Atm, like *MEC1* (Kato and Ogawa 1994) and *mei-41* (Baker and Carpenter 1972; Baker et al. 1976; Hari et al. 1995), appear to play a role in meiosis. These observations suggest that Atr is a mammalian checkpoint protein with analogous function to the Rad3 proteins and shares complementary functions with Atm in human cells. The definition of precisely which checkpoints are dependent on Atr function in mammalian cells awaits the availability of Atr-deficient cells.

Atr and Atm have associated protein kinase activity

The protein kinase activity associated with Atm and Atr

Figure 7. Atm localization on meiotic chromatin. Immunofluorescence localization of ATM in meiotic prophase of mouse oocytes (A,B) or spermatocytes (C,D). A and C show the localization of Cor1, a component of the axial elements of the synaptonemal complex (SC), and B and D show the merged Cor1 (white) and Atm (yellow-green) images. Atm localizes to synapsed axes only. By late zygonema in oocytes (A and B) pairing is proceeding. Atm foci are found along the entire lengths of the synapsed axes, but not the asynapsed axes (small arrow, B). In pachynema in spermatocytes (C and D), Atm foci are present along the synapsed autosomal axes and the paired portion of the XY (small arrow, D). In addition, two small foci are regularly observed on the unpaired axis of the X chromosome at this substage of pachynema.

may result from an intrinsic protein kinase activity of these molecules; however, we are unable to rule out the presence of an associated protein kinase in our immunoprecipitates. Possession of intrinsic protein kinase activity is consistent with the model of a checkpoint signal transduction cascade (Carr 1996). The presence of protein kinase domains in Atm and Atr make these molecules ideal candidates as sensors of aberrations in DNA structure. Detection of DNA damage or an incorrect DNA structure by Atm or Atr could stimulate the protein kinase activity of these proteins, thereby initiating a cascade leading to cell-cycle arrest and then, ultimately,

DNA repair, apoptosis, or cellular senescence in mitotic cells or genetic recombination, chromosome pairing, and segregation in meiotic cells. While the structure of the kinase domains of these two proteins is related most highly to lipid kinase domains, it is known that proteins with this domain are able to act as protein kinases and that DNA-PK_{CS} and FRAP are able to autophosphorylate independent of associated regulating subunits (Dvir et al. 1993; Brown et al. 1995). DNA-PK_{CS} phosphorylates a wide range of proteins *in vitro*; however, it has not been shown to phosphorylate lipids (Hartley et al. 1995). Rad3 has associated protein kinase activity and mutants

Figure 8. Atr and Atm show complementary localization along unpaired (Atr) and paired (Atm) axes. The Cor1 images are shown at *left*; the merged Cor1 plus Atr (red) at *upper right* and Cor1 plus Atm (yellow-green) at *lower right*. The Atr enlargement shows a completely synapsed autosomal bivalent on the *left*, the univalent X in the *center*, and a partially synapsed autosomal bivalent on the *right*. The synapsed portions (small arrows) of the autosomes have no Atr foci, while the asynapsed portions of the autosomal axes (large arrow) and the X axis have foci along their lengths. In the lower set of images there is a completely synapsed autosomal bivalent (*left*) and a partially synapsed autosomal bivalent (*right*). Atm foci are found only on the synapsed axes (small arrows), not on the asynapsed axes (large arrows).

in the protein kinase domain that render the kinase inactive are defective for checkpoint function (Bentley et al. 1996). In addition, Frap and Vps34 autophosphorylate (Stack and Emr 1994; Brown et al. 1995).

We have been unable to observe phosphorylation of proteins known to be substrates for DNA-PK_{CS} such as RPA or a p53 peptide containing the DNA-PK site. However, there are other proteins that may be potential substrates of Atr or Atm and these substrates have been suggested to be protein kinases. In fission yeast, phosphorylation of the Chk1 protein kinase, which is required for the mitotic arrest after DNA damage, has been shown to depend on a functional *rad3* gene (Walworth and Bernards 1996). The *S. cerevisiae* RAD53 gene, which encodes another protein kinase, is a multicopy suppressor of *mec1* lethality (Sanchez et al. 1996), and *mec1* mutants lack Rad53p hyperphosphorylation. Further, overexpression of the ATM homolog, TEL1 in a *mec1* strain restores Rad53p phosphorylation (Sanchez et al. 1996). Identification of the mammalian homologs

of these downstream effectors for Rad3 and Mec1p will allow us to determine whether they act as substrates for Atm and Atr. These will be key to understanding the pathways leading from DNA damage and DNA structural aberration to cell-cycle arrest and to the meiotic function of these proteins.

Atr is a nuclear protein expressed in early meiotic cells

We have demonstrated that Atr is expressed in cells in early meiotic prophase. Many of the checkpoint proteins identified in yeast as being required for the mitotic arrest after DNA damage or S-phase arrest (for review, see Carr and Hoekstra 1995) are also required for the checkpoints that function during meiosis. The checkpoint genes *MEC1* (Weber and Byers 1992; Kato and Ogawa 1994) and *mei-41* (Baker and Carpenter 1972; Baker et al. 1976; Hari et al. 1995) and DNA repair genes such as *MLH1* (Baker et al. 1996; Edelmann et al. 1996) and the *RAD50–RAD57* series (Petes et al. 1991; Friedberg et al. 1995) have been shown to play roles in meiotic processes. For yeast, it has been suggested that meiotic recombination is initiated by DSBs (for review, see Roeder 1990; Kleckner 1996) and involves strand invasion and repair synthesis. This potential role for DSBs as initiating lesions for meiosis I recombination would predict a requirement for checkpoint and repair proteins in signaling DSBs and/or repair synthesis during meiotic prophase. An alternative model of synaptic initiation in multicellular eukaryotes suggests that a subfraction of the eukaryotic genome is not replicated in the premeiotic S phase, but is delayed until zygonema (for review, see Stern and Hotta 1984), thus the function of checkpoint proteins during zygonema could be to monitor the completion of DNA replication before further meiotic progression. In this model nicks or breaks in DNA may not occur until a check for homology is completed.

Complementary roles for Atr and Atm in meiosis

We have demonstrated that both Atr and Atm form foci at discrete sites along meiotic chromosomal axes as homologs synapse. The involvement of Atr appears to be transient and restricted to an early event in the synaptic process. In fact, in zygotene nuclei, the interaction appears to be sufficiently transient that Atr detection at many pairing forks is low. One interpretation is that a small amount of Atr is present, and then only briefly. Alternatively, Atr localization to chromosomes may be absent when chromosome synapsis is proceeding normally. In support of this second explanation, Atr appears to accumulate at autosomal pairing sites where synapsis has been delayed. Since the X chromosome has no homolog in spermatocytes, it can never undergo completely homologous synapsis, although it undergoes a small amount of homologous synapsis and extensive nonhomologous synapsis with the Y chromosome. We find that Atr remains on the asynapsed X axis into

pachynema (Fig. 8). These results suggest that Atr may play a role in monitoring asynapsis of chromosomes.

While the role for Atr in early meiotic prophase is transient, the role for Atm is more prolonged. Atm associates with axes as they synapse, producing multiple foci along these axes. It appears likely that those Atm foci seen in zygonema and early pachynema may reveal a participation in events associated with synapsis. However, some Atm foci are present in mid-pachynema, the stage at which reciprocal recombination is thought to occur, suggesting Atm may be involved in both synaptic and recombination events.

The time of appearance of Atr and Atm in zygotene nuclei, along with the localization of Atr at sites along unpaired chromosomal axes, and Atm along paired axes, suggest they are both involved in the synaptic process. Although there is evidence that the meiotic search for homology is initiated by DSBs and accomplished by gene conversion events in yeast (Sun et al. 1989; Cao et al. 1990; Bishop 1994), this may not be the sequence of events in multicellular eukaryotes (Plug et al. 1996). The check for homology in mouse may precede breakage and gene conversion events (Plug et al. 1996), and our current observations are consistent with this suggestion. When Xu et al. (this issue) disrupted the Atm gene they found fragmentation of synaptonemal complex axes, but not until later in meiotic prophase, suggesting that unrepaired breaks could be a secondary, not primary consequence of Atm disruption. In this model, Atr would be predicted to be monitoring breaks associated with a search for homology. An unsuccessful search (with unrepaired breaks) should result in fragmentation of univalent axes. While univalent axes accumulate Atr (Figs. 6 and 8), they do not fragment, suggesting that breaks may not have yet occurred.

Atr and Atm may play essential, but indirect, roles in meiosis. In mitotic cells, Atm is required for a DNA damage-dependent signal transduction cascade that activates multiple cell-cycle checkpoints (Meyn 1995; Shiloh 1995). In yeast and flies, Atr-like proteins are important for DNA damage and replication-like checkpoints. In spermatocytes, Atr and Atm may help coordinate the various events of meiotic prophase by performing similar checkpoint functions. Early meiotic prophase checkpoints would be predicted to signal breaks and monitor repair synthesis as well as coordinately monitor homologous synapsis and DNA replication before further meiotic progression. The Atr- and Atm-dependent signal transduction networks could be triggered by DSBs and/or related structures occurring in synapsing meiotic chromosomes. Alternatively, Atr may control an S-phase meiotic checkpoint that responds to single-stranded regions of DNA, whereas Atm may recognize direct DNA strand lesions during recombination processing events.

The focal pattern of Atr along the asynapsed axes, and of Atm along the synapsed axes, is similar to the focal pattern described for hRad51 (Ashley et al. 1995). Recent work has suggested that during early meiotic prophase Rad51 is a component of early recombination nodules (RNs), thought to be involved in the check for homology

and synaptic initiation (Bishop 1994; Ashley et al. 1995; Terasawa et al. 1995). The time of appearance of Rad51p in yeast appears to coincide with the time of appearance of both DSBs and SC precursors (Bishop 1994). Rad51 in mammals associates with chromosomes as early as premeiotic S phase, before the appearance of SC precursors (Plug et al. 1996). We find no evidence that either Atr or Atm are present on chromosomes at this stage. Atr and Atm foci appear as homologous chromosomes begin to synapse. The staining sites on synapsing chromosomes appear to correspond to similar sites of Rad51 distribution in mammalian spermatocytes and oocytes.

While the location of Atr and Atm along meiotic chromosomal axes is similar to the focal pattern of hRad51 along axes of synapsing and synapsed chromosomes, the timing and appearance and disappearance of each of these proteins differs, indicating a dynamic nature and suggesting multiple activities for these protein complexes. The pattern of association of Atr and Atm with meiotic chromatin is consistent with each playing separate roles in monitoring DNA structure or behavior during early synaptic and recombination events.

Materials and methods

Northern blot hybridization

A 1.3-kb PCR product was amplified in the presence of 32 P-dCTP using primers 279-3 (TGGATGATGACAGCTGTGTC) and 279-6 (TGTAAGTCGCTGCTCAATGTC). A nylon membrane containing 2 μ g of size-fractionated poly(A)⁺ RNA from a variety of human tissue sources (Clontech Laboratories) was probed with the labeled PCR fragment described above as recommended by the manufacturer except that the final wash was performed at 55°C to minimize the possibility of cross-hybridization to related sequences. A mouse cDNA clone was isolated by low-stringency hybridization. Oligonucleotides mrad3-1 (5'-CGAGATCCATTGTGCTGGAAGG-3') and mrad3-2 (5'-CTGCAGTAGAGCGGCAATATGC-3') were used to produce an ~300-bp probe as above that was used for hybridization of a Northern blot containing 2 μ g of poly(A)⁺ RNA from a variety of mouse tissues and from total embryos at days 7, 11, 15, and 17 as described above.

Cell lines and media

MRC-5 cells were obtained from the ATCC at passage 19 and maintained in Minimal Essential Medium (MEM) supplemented with 10% fetal bovine serum, 100 units/ml penicillin, and 100 μ g/ml streptomycin, and 100 μ M MEM nonessential amino acids. 293T were maintained in MEM with the supplements listed above. G361 were maintained in McCoy's 5A containing 10% fetal bovine serum and 100 units/ml penicillin and streptomycin. Media and media supplements were obtained through Gibco Life Technologies. Cell lines were maintained in a water-saturated 37°C incubator with 5% CO₂.

Antibody preparation

Atr-antibodies to two different glutathione-S-transferase (GST) fusion proteins were generated in rabbits. Antigen DH-2 (4779) contains Atr amino acid residues 2341 to 2644 fused to GST in pGEX1. Antigen DH-3 (4747) contains Atr residues 1892 to 2340 fused to GST in pGEX3 (Pharmacia). Antigens were gel-purified

by SDS-PAGE and electroelution and used to immunize rabbits. Antibodies were affinity purified with inclusion body preparations of antigen coupled to cyanogen bromide-activated Sepharose (Pharmacia) as per manufacturer's instructions. Both antisera immunoprecipitate an associated kinase activity that is blocked by the antigen against which the antibodies were generated.

Antipeptide antibodies were generated against the human Atm protein by coupling a 15-amino-acid peptide (residues 1359–1373) to Keyhole Limpet Hemocyanin (KLH) using 1-ethyl-3-[dimethylaminopropyl]-carbodiimide hydrochloride (EDC) as described by the manufacturer (Pierce), followed by injection of the coupled immunogen into rabbits. The antibodies were first precipitated from the serum (#6076) with an equal volume of saturated ammonium chloride followed by resuspension and dialysis against PBS. Affinity purification was carried out using a peptide column prepared by coupling the antigenic peptide to CNBr-activated Sepharose (Pharmacia) as described by the manufacturer. The antibodies were then bound to the peptide column and washed with 2 M KCl-PBS. Elution was carried out with 20 ml 5 M NaI (in 1 mM sodium thiosulfate), which was dialyzed immediately against PBS. Specificity for the Atm sera was examined by immunoblot analysis using recombinant Atm and by immunoprecipitation of ³⁵S-labeled lysates. Anti-Atm and anti-Atr recognize different sized proteins that migrate at the predicted molecular mass for Atm and Atr when compared from the ³⁵S-labeled lysates.

The ATM1 antibody, used for staining of surface spreads, was generated against the Atm protein by coupling a 14-amino-acid peptide (residues 1347–1360) to KLH, followed by injection of the coupled immunogen into New Zealand white rabbits. Antibodies were purified using protein A agarose and tested for anti-Atm reactivity by immunoblotting against extracts of Daudi and Hela cells (positive controls) and the A-T lymphoblast line ATF169 (negative control). ATM1 was provided by Dave Hill (Oncogene Research, Products/Calbiochem, San Diego, CA).

Immunoprecipitations and kinase assays

Cell extracts were prepared by lysing in lysis buffer [50 mM NaPO₄ at pH 7.2, 0.5% TritonX-100, 2 mM EDTA, 2 mM EGTA, 25 mM NaF, 25 mM 2-glycerophosphate, 1 mM phenylmethylsulfonyl fluoride (PMSF), 1 μg/ml leupeptin, 1 μg/ml pepstatin A, and 2 mM DTT]. Mouse testes extracts were prepared by dounce homogenization of decapsulated testes in lysis buffer followed by centrifugation at 10,000g. For Atr IPs, 500 μg of extract was incubated with either 10 μg of affinity-purified 4779 antibody or 10 μg purified rabbit IgG in 1 ml of lysis buffer for 1 hr on ice. Thirty microliters of protein A-agarose slurry (Pierce) was added and incubated for 30 min at 4°C on a rocking platform. The immune complexes were then washed 3× with lysis buffer, once with kinase buffer (25 μM HEPES at pH 7.7; 50 mM KCl; 10 mM MgCl₂; 0.1% NP-40; 2% glycerol; 1 mM DTT), and incubated in kinase buffer with 10 μM ATP (50 Ci/mmol) plus 10 μCi [γ -³²P]ATP for 20 min at 37°C. The reactions were stopped with 20 μl 2× SDS sample buffer prior to separation on 6% polyacrylamide gels. Kinase reactions were then transferred to Immobilon, exposed to film, and then subsequently probed with 4779 to confirm that the phosphorylated band comigrated with Atr.

MRC-5 cell extracts were prepared by lysis of a 10-cm plate of cells in 0.5 ml of the above lysis buffer on ice. Cells were scraped from plates using rubber spatulas and sonicated in a cup horn sonicator (Sonifier 250, Branson Ultrasonics Corp., Danbury, CT) at 100% output for 90 sec. Lysates were then clarified

in a 4°C microfuge for 2 min. Preclearing was done by adding 10 μg purified rabbit IgG and 30 μl protein A-agarose slurry followed by incubation at 4°C for 60 min while rocking. To the precleared lysates, 10 μg of affinity purified 6076 antisera (or 10 μg 6076 preblocked with 0.04 mg peptide for 30 min) was added and incubated on ice for 60 min. Immunoprecipitates were washed and a kinase reaction was performed as described above.

Expression and purification of recombinant Atr

An ATR *Bam*HI–*Sal*I fragment containing an amino-terminal -His tag was cloned into the *Bam*HI–*Xho*I site of the CEN-based galactose inducible yeast vector p416 GAL1 (gift from D. Morrow, Johns Hopkins University, Baltimore, MD). Plasmid DNA was transformed into an *esr1-1* diploid yeast strain (*MATa leu2-1 his4-4 can1 ura3 cyh2 ade6 ade2 esr1-1/MATa leu2-27 his4 trp1 met2 ade2 esr1-1*), and cells were grown to mid-log phase in either galactose or glucose containing medium. Cells were pelleted, washed, and all steps performed at 4°C. Cell pastes were resuspended in buffer (20 mM Tris at pH 8.0, 300 mM NaCl, 10% glycerol, 0.1 mM PMSF, 0.25 mg/ml pepstatin, leupeptin, and aprotinin) and lysed in a French Press or using glass beads. Lysis was verified by microscopy following a low-speed (10K) spin and a high-speed spin (100K), and the supernatant was loaded onto a 1.5-ml Ni-NTA agarose (Qiagen, Inc., Chatsworth, CA) column prewashed in 1× buffer. The column was washed with six column volumes of buffer. The column was eluted stepwise with 8 ml of 10 mM, 50 mM, 100 mM, and 250 mM imidazole in buffer. Fractions were collected and Western analysis was performed using 15 μl of each elution peak. For protein kinase assays, 2 μl of the 50 mM imidazole elution pool was added to kinase buffer. The reactions were incubated at 37°C for 20 min. Samples were analyzed by SDS-PAGE prior to autoradiography or Western analysis.

Spermatocyte preparation, immunoblot analysis, and immunocytochemistry

For tissue sections, testis tissue from Balb/c mice was sectioned, fixed in acetone, blocked with 30% rat serum, 5% goat serum, and 1% BSA before incubation with either affinity-purified 4747 or 4779 antisera, the equivalent preimmune sera, or anti-GST sera. Antibody was detected with biotinylated goat antirabbit antibody and streptavidin-gold conjugate. Sections were counterstained with nuclear fast red. Signal was detected with both 4747 and 4779 sera, but not detected with preimmune sera or antibodies to GST alone. For purified cell populations, decapsulated testes of adult CD-1 mice were dissociated sequentially with collagenase and trypsin-DNase I (Romrell et al. 1976). Dissociated cells were separated into populations by sedimentation in 2–4% BSA gradients (Bellve et al. 1977). Pachytene spermatocyte and round spermatid populations were at least 85% pure. The condensing spermatid population was 40–50% pure (contaminating material was primarily anucleated residual bodies and some round spermatids). Sperm were obtained from the cauda epididymides. Cell populations were dissolved directly in SDS sample buffer for immunoblot analysis.

Surface spreads were prepared using the method of Speed (1982) as modified by Antoine Peters (pers. comm.). The antibody detection procedure was as described previously (Ashley et al. 1995). The polyclonal Cor1 antibody, raised in mouse against the Syrian hamster Cor1 protein (Dobson et al. 1994) was a gift from Drs. Peter Moens and Barbara Spyropoulos (York University, Toronto, Canada). The production of the Cor1 antibody and its specificity for the axial/lateral element of the synaptonemal complex has been described previously (Dobson et al.

1994). Affinity-purified 4779 antibody was detected with goat antirabbit IgG conjugated with fluorescent isothiocyanate (FITC) and the mouse anti-Cor1 with goat antimouse IgG conjugated with Texas Red. Incubation of spermatocytes with 4779 preimmune serum produced only a nonspecific background reaction in all cell types. One hundred twenty-one spermatocyte nuclei were imaged from six males 3 to 12 weeks old for localization studies on Atr and 176 nuclei from four males 3 to 12 weeks old for localization with Atm. In addition 25 oocyte nuclei were imaged from two female fetuses 16 days postconception. Preimmune serum from 4779 and from Atm1 produced only a faint generalized nuclear staining pattern on spermatocyte preparations.

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