Autophosphorylation of the Catalytic Subunit of the DNA-Dependent Protein Kinase Is Required for Efficient End Processing during DNA Double-Strand Break Repair

Qi Ding,1 Yeturu V. R. Reddy,2 Wei Wang,1 Timothy Woods,1 Pauline Douglas,3 Dale A. Ramsden,2,4 Susan P. Lees-Miller,3 and Katheryn Meek1*

College of Veterinary Medicine and Department of Pathobiology and Diagnostic Investigation, Michigan State University, East Lansing, Michigan 48824; Departments of Biochemistry and Molecular Biology and Biological Sciences, University of Calgary, Calgary, Alberta, Canada T2N 1N4; and Lineberger Comprehensive Cancer Center2 and Department of Biochemistry and Biophysics,4 University of North Carolina at Chapel Hill, Chapel Hill, North Carolina

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The DNA-dependent protein kinase (DNA-PK) plays an essential role in nonhomologous DNA end joining (NHEJ) by initially recognizing and binding to DNA breaks. We have shown that in vitro, purified DNA-PK undergoes autophosphorylation, resulting in loss of activity and disassembly of the kinase complex. Thus, we have suggested that autophosphorylation of the DNA-PK catalytic subunit (DNA-PKcs) may be critical for subsequent steps in DNA repair. Recently, we defined seven autophosphorylation sites within DNA-PKcs. Six of these are tightly clustered within 38 residues of the 4,127-residue protein. Here, we show that while phosphorylation at any single site within the major cluster is not critical for DNA-PK's function in vivo, mutation of several sites abolishes the ability of DNA-PK to function in NHEJ. This is not due to general defects in DNA-PK activity, as studies of the mutant protein indicate that its kinase activity and ability to form a complex with DNA-bound Ku remain largely unchanged. However, analysis of rare coding joints and ends demonstrates that nucleolytic end processing is dramatically reduced in joints mediated by the mutant DNA-PKcs. We therefore suggest that autophosphorylation within the major cluster mediates a conformational change in the DNA-PK complex that is critical for DNA end processing. However, autophosphorylation at these sites may not be sufficient for kinase disassembly.

Although DNA is the genetic blueprint for all living organisms, it is extremely sensitive to various forms of damage, including oxidation, hydrolysis, and methylation. Thus, efficient DNA repair systems are essential for the maintenance of chromosomal integrity. DNA double-strand breaks (DSBs) are perhaps the most lethal form of DNA damage. In eukaryotes, primarily two pathways repair DSBs: homologous recombination and nonhomologous DNA end joining (NHEJ). In higher eukaryotes, NHEJ is thought to be the major pathway that repairs these breaks (6, 29, 37, 38). Because NHEJ also functions in developing lymphocytes to repair the DSBs introduced during antigen receptor gene rearrangement, defects in this pathway result in a block in lymphocyte development and the disease known as severe combined immunodeficiency (SCID) (reviewed in references 14 and 29).

In the past decade, an intensive research effort has focused on NHEJ, resulting in a reasonable understanding of how DSBs are resolved. There are six known factors which unequivocally function in the NHEJ pathway. Three of these comprise the DNA-dependent protein kinase (DNA-PK) (reviewed in references 19, 20, and 25): the two subunits of the DNA end binding heterodimer Ku and the catalytic subunit of DNA-PK (DNA-PKcs) (18). Two other factors, XRCC4 and DNA ligase IV, form a stable DNA ligase complex (15, 28, 34). The sixth factor, Artemis, was described in 2001 (35); recent data indicate that it may play an important role in DNA end processing during NHEJ (31).

DNA-PK plays a central role in NHEJ because it initially recognizes and binds to damaged DNA and then targets other repair activities to the site of damage. The first step in NHEJ is binding of the Ku70/86 heterodimer to DNA ends at the site of DNA damage. When bound, Ku encircles the DNA. The Ku binding site covers approximately two turns of the DNA helix, but only the central 3 to 4 bp are completely encircled by the polypeptide chain. The DNA terminus is cupped in an accessible binding pocket (46), theoretically allowing DNA-PKcs to interact directly with the DNA end. Recruitment of DNA-PKcs results in translocation of Ku to internal sites of the linear DNA (51). When assembled onto DNA ends, DNA-PK is activated; it can then phosphorylate a variety of substrates including both Ku subunits, DNA-PKcs, XRCC4, and Artemis (4, 5, 8, 24, 31). Recent data from our laboratory and others demonstrate that the protein kinase activity of DNA-PK is essential during NHEJ (1, 21, 22), suggesting that phosphorylation by DNA-PK activates a downstream factor in the pathway.

In assay systems utilizing purified proteins, autophosphorylation of DNA-PK results in the release of DNA-PKcs from DNA-bound Ku and a decrease in the measurable protein kinase activity (4, 5, 8, 33). We suggest a model whereby autophosphorylation of DNA-PKcs and the resulting remodeling and/or disassembly of the DNA-PK complex could facilitate subsequent steps in the DNA repair process. To test this model, we have generated point mutations of the six clustered

* Corresponding author. Mailing address: Michigan State University, 350 FST, East Lansing, MI 48824. Phone: (517) 432-9505. Fax: (517) 353-9004. E-mail: kmeek@msu.edu.
autophosphorylation sites previously mapped within DNA-PKcs (9). We find that phosphorylation of any single site is not critical to DNA-PK's function; however, mutation of several phosphorylation sites abolishes the ability of DNA-PK to function in NHEJ. Finally, both in vivo and in vitro studies of the mutant protein suggest that autophosphorylation at the major cluster specifically facilitates DNA end processing, whereas further autophosphorylation is required to promote kinase disassembly.

MATERIALS AND METHODS

Cell lines and culture conditions. The DNA-PKcs-deficient DSb repair mutant CHO cell line V3 (47) was the generous gift of Martin Gellert. Cells were maintained in α-MEM with 10% fetal calf serum (Gibco BRL, Gaithersburg, Md.). Stable transfecants were maintained with 400 μg of G418/ml.  

Oligonucleotides. Oligonucleotides used in this study are as follows (5’ to 3’):

<table>
<thead>
<tr>
<th>Oligonucleotide</th>
<th>Sequence</th>
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<tr>
<td>KAM97 (mutant A)</td>
<td>TTGTTGGAGGGCTTAGGCTTCA (complementary to wild-type DNA-PKcs)</td>
</tr>
<tr>
<td>KAM101 (mutant B)</td>
<td>GTGGTCAAGGAGGCTGTTCTCTCA</td>
</tr>
<tr>
<td>KAM103 (mutant C)</td>
<td>CGGCGCTAGCAGCATGACTTCACACTG</td>
</tr>
<tr>
<td>KAM105 (mutant D)</td>
<td>TTTGTGGAGCAGGAGTCGT</td>
</tr>
<tr>
<td>KAM146 (mutant E)</td>
<td>CAGGCCAAGGGCAGCTGCT</td>
</tr>
<tr>
<td>KAM103 (mutant CD)</td>
<td>GTGGTCAAGGAGGCTGTTCTCTCA</td>
</tr>
<tr>
<td>KAM105 (mutant D)</td>
<td>TTTGTGGAGCAGGAGTCGT</td>
</tr>
<tr>
<td>KAM146 (mutant E)</td>
<td>CAGGCCAAGGGCAGCTGCT</td>
</tr>
</tbody>
</table>

The two oligonucleotides flanking the phosphorylation site cluster are KAM97 (GATGTCGGTCAAGGAGGTGTTG) and KAM98 (Stratagene, La Jolla, Calif.) was utilized to introduce mutations into the subfragment of the human DNA-PKcs cDNA. It was subsequently subcloned into the wild-type human DNA-PKcs expression vector.  

Individual colonies were screened for DNA-PKcs expression by immunoblot analysis and were further cultured. Construction of the human DNA-PKcs cDNA. It was subsequently subcloned into the wild-type human DNA-PKcs expression vector has been described previously (11). Briefly, wild-type human DNA-PKcs expression vector was digested with BamHI and HindIII, ligated to the HindIII linker, and cloned into the HindIII site of the pCD8 expression vector (Promega Corp., Madison, Wis.). bam ends were fixed with ethanol and stained with crystal violet, and colony numbers were assessed.

Immunoblot analysis. The two oligonucleotides flanking the phosphorylation site cluster are KAM97 (GATGTCGGTCAAGGAGGTGTTG) and KAM98 (Stratagene, La Jolla, Calif.) was utilized to introduce mutations into the subfragment of the human DNA-PKcs cDNA. It was subsequently subcloned into the wild-type human DNA-PKcs expression vector.  

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and then washed three times in buffer A. DNA-cellulose fractions were incubated in buffer A with [γ-32P]ATP for 15 min at room temperature and then analyzed by SDS-PAGE and autoradiography.

**Purification of recombinant DNA-PKcs.** The purified recombinant human Ku heterodimer was generated as previously described (36). Purified human DNA-PKcs was obtained from 5 liters of V3 cells stably transformed with either wild-type or ABCDE mutant DNA-PKcs. The washed cell pellet was extracted by homogenization in 50 ml of a buffer containing 50 mM Tris-HCl (pH 7.5), 600 mM KCl, 10% glycerol, 1 mM DTT, 5 mM EDTA, and 0.5 ml of a 30% slurry of polyethyleneimine-cellulose (Sigma). Polyethyleneimine-cellulose with absorbed nucleic acid was removed by filtration. The filtrate was then dialyzed to buffer A (25 mM HEPES-KOH [pH 7.5], 10% glycerol, 1 mM DTT, 0.1 mM EDTA) plus 50 mM KCl (see Fig. 3A, lane 1) and loaded onto an affinity column (80 peptide) generated by coupling 5 mg of a peptide from the C terminus of Ku80 (KSGGEEGGD) to a 1-ml, 5% N-hydroxysuccinimide-activated HiTrap column (Amersham Biosciences). Bound proteins were eluted by a gradient to buffer A plus 1 M KCl. DNA-PKcs-containing fractions were identified by SDS-PAGE analysis, pooled (Fig. 3A, lane 3), and dialyzed against buffer A plus 100 mM KCl. The dialyze was then loaded onto a 100-μl Mono S ion-exchange column (Amersham Biosciences) and eluted with a 2-ml gradient to buffer A plus 350 mM KCl (Fig. 3A, lane 5). Ion-exchange chromatography both removed the remaining major contaminant (~180-kDa band in Fig. 3A, lanes 3 and 4) and concentrated the protein ~25-fold. Fractions were flash-frozen in liquid nitrogen in small aliquots.

**Assays using purified proteins.** The kinase activity of purified DNA-PKcs was assessed in a buffer containing 50 mM Tris-HCl (pH 7.5), 150 mM NaCl, 0.1 mM EDTA, and 0.5 mM KCl, 10% glycerol, 1 mM DTT, 5 mM EDTA, and 0.2 mM MgCl2. The peptide substrate used was biotinyl-EPLSQEAFADLWK. Phosphorylation of the peptide was assessed as described previously (7). For EMSA analysis, we used a 60-bp double-stranded DNA substrate made by annealing the 5′-32P-labeled oligonucleotide DAR166 (defined above) to DAR167. One hundred nanomolar DNA-PKcs stocks were inactivated by treatment with 10 μM pervanadate or mock treated with an equivalent amount of a solvent (dimethyl sulfoxide) for 10 min on ice prior to dilution into the reaction mixture. Samples were prepared for EMSA analysis in two steps. In the first step (complex assembly and autophosphorylation), DNA-protein complexes were formed with 10 nM each protein and 100 nM duplex DNA by using a buffer containing 25 mM Tris-HCl (pH 8.0) supplemented with 5 mM MgCl2 and 200 μM ATP. Reaction mixtures were incubated for 10 min at 30°C before autophosphorylation was arrested by addition of 5 mM EDTA. In the second step (probing the stability of complexes), reaction products were adjusted with added NaCl to the indicated salt concentrations and were incubated for a further 10 min at room temperature. Stable DNA-protein complexes were then filtered by cross-linking using 0.25% glutaraldehyde and were resolved by electrophoresis on a 3.5% native PAGE gel in Tris-borate-EDTA at 18 V/cm. The composition of each indicated species was confirmed by antibody supershift analysis (data not shown).

**Construction of an Artemis-expressing baculovirus and coimmunoprecipitation.** A cDNA encoding full-length human Artemis was assembled from overlapping reverse transcription-PCR fragments. Although numerous splice variants were isolated, we assembled a full-length coding sequence analogous to that shown previously to complement the V(DJ) recombination deficiency in Artemis-deficient cell lines (35). The termination codon was not included, and an in-frame V5 His tag was added at the C terminus. The assembled fusion protein construct was cloned into the pAHC-T-C transfer vector (Pharmingen, San Diego, Calif.). A recombinant baculovirus encoding human Artemis was obtained by using Baculogold-linearized viral DNA (Pharmingen) and standard techniques. For the pulldown experiments for which results are shown in Fig. 5, whole-cell extracts of V3 transfectants (1 mg) and whole-cell extracts from either Artemis-infected or control S9 cells were coinubated for 30 min. Subsequently, 50 µl of Ni+ agarose was added, and the extracts were absorbed in buffer A with 25 mM imidazole. After 1 h, the Ni+ agarose was washed three times with buffer A containing 50 mM imidazole. Proteins were eluted with SDS-PAGE buffer and analyzed by immunoblotting.

**Analyses of V(DJ) recombination intermediates.** V3 cells were transfected either with the pH290 recombination substrate alone, with the substrate and the RAG expression plasmids, or with the substrate and the RAG and DNA-PKcs expression plasmids as described above. Hirt supernatants were prepared from transfected cells as follows. Forty-eight hours after transfection, cells were harvested, resuspended in 400 µl of Hirt buffer 1 (10 mM Tris [pH 8.0], 1 mM EDTA, 0.6% SDS), and incubated for 15 min at room temperature. One hundred microliters of Hirt buffer 2 (10 mM Tris [pH 8.0], 1 mM EDTA, 5 mM NaCl) was added, and the samples were incubated overnight at 4°C. After being spun for 10 min at 10,000 × g at 4°C, supernatants were extracted with phenol-chloroform, ethanol precipitated, and then resuspended in 10 µl of double-distilled water. Five microliters of each Hirt supernatant was ligated to 500 pmol of annealed oligonucleotides KAM150–KAM200 or KAM201–KAM200 at 16°C overnight. Ligated Hirt supernatants were ethanol precipitated and used in nested PCR amplifications. Signal ends were amplified first with the primer combination KAM214–KAM217 and then with KAM216–KAM215. Coding ends were amplified first with the primer combination KAM214–KAM218 and then with KAM206–KAM219. Forty cycles of amplification were performed using the following conditions: for primary PCRs, 94°C for 30 s, 58°C for 1 min, and 68°C for 1 min; for secondary PCRs, 94°C for 30 s, 56°C for 1 min, and 68°C for 1 min. Amplification products were analyzed by Southern hybridization using oligonucleotide KAM88 as a hybridization probe.

**RESULTS**

Six of seven autophosphorylation sites are located within a 38-amino-acid cluster within human DNA-PKcs. Recently, the locations of seven in vitro autophosphorylation sites within DNA-PKcs were reported (9). As diagrammed in Fig. 1A, six of the seven phosphorylation sites are tightly clustered in a short region of primary sequence in the central region of the molecule (amino acids 2609 to 2647). Four of the six clustered autophosphorylation sites (T2609, S2612, T2638, and T2647) are also phosphorylated in vivo in okadaic acid-treated cells. All four of these in vivo sites are completely conserved among the six sequenced vertebrate DNA-PKcs genes (human, horse, dog, mouse, chicken, and frog).

To begin to assess the functional relevance of autophosphorylation of DNA-PKcs, several point mutations of the DNA-PKcs cDNA were generated. Initially, five mutant constructs, designated A, B, C, D, and E, were generated, as indicated in Fig. 1A. Constructs A, C, D, and E each contain a single mutation of serine or threonine to alanine, as indicated. Mutation B includes two mutations of serine or threonine to alanine.

![FIG. 1. Single phosphorylation site mutants complement the radiosensitivity of V3 cells, whereas multiple phosphorylation site mutants do not. (A) Diagrammatic representation of seven autophosphorylation sites within DNA-PKcs (asterisks). Mutations are as follows: A, T2609A; B, T2609A and S2624A; C, T2638A; D, T2647A; E, S2612A. The A, B, C, D, and E mutations were introduced either alone or in combinations as indicated. (B) Immunoblot analyses of whole-cell extracts from V3 transfectants expressing either wild-type DNA-PKcs (lane 1), vector alone (lane 2), mutant A (lane 3), mutant B (lane 4), mutant C (lane 5), mutant D (lane 6), or mutant E (lane 7). (C) The radiation resistance of V3 transfectants expressing wild-type DNA-PKcs, vector alone, or each of the single autophosphorylation site mutants was assessed as described in Materials and Methods. Data are presented as percent survival relative to that of nonirradiated controls (set at 100%). Error bars, standard errors of the means of three separate experiments. (D) Immunoblot analyses of whole-cell extracts from V3 transfectants expressing either full-length DNA-PKcs (lane 1), vector alone (lane 2), mutant AB (lane 3), mutant AC (lane 4), mutant AD (lane 5), mutant ABCD (lane 6), or mutant ABCDE (lane 7). (E) The radiation resistance of V3 transfectants expressing either wild-type DNA-PKcs, vector alone, or mutant AB, AC, AD, ABC, or ABCDE was assessed. Data are presented as percent survival relative to that of nonirradiated controls (set at 100%). Error bars, standard errors of the means of three separate experiments.](http://mcas.asm.org/index.php?article=5839)
tion in V3 cells, as assessed by the plasmid substrate pJH290, which detects materials and Methods. RAG expression from plasmid vectors initiates recombination.

Transient V(D)J recombination assays were performed as described in Ma-

numbers from at least three separate experiments are presented.


recombination (%)*

<table>
<thead>
<tr>
<th>Transfected plasmid</th>
<th>No. of Amp&lt;sup&gt;+&lt;/sup&gt; Cm&lt;sup&gt;R&lt;/sup&gt; colonies/</th>
<th>Recombination (%)*</th>
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</thead>
<tbody>
<tr>
<td>pJH290 only</td>
<td>0/46,060</td>
<td>0</td>
</tr>
<tr>
<td>pJH290 + RAGS</td>
<td>1/30,968</td>
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<td>275/25,520</td>
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<tr>
<td>pJH290 + RAGS + mut A</td>
<td>283/31,130</td>
<td>0.91</td>
</tr>
<tr>
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<td>141/22,540</td>
<td>0.63</td>
</tr>
<tr>
<td>pJH290 + RAGS + mut C</td>
<td>209/22,990</td>
<td>0.91</td>
</tr>
<tr>
<td>pJH290 + RAGS + mut D</td>
<td>107/45,080</td>
<td>0.24</td>
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<tr>
<td>pJH290 + RAGS + mut E</td>
<td>203/300</td>
<td>0.61</td>
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<td>0</td>
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<td>0</td>
</tr>
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<td>pJH201 + RAGS + wild type</td>
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<td>0.5</td>
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<tr>
<td>pJH201 + RAGS + mut ABCD</td>
<td>6/1,200</td>
<td>0.5</td>
</tr>
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</table>

* Transient V(D)J recombination assays were performed as described in Materials and Methods. RAG expression from plasmid vectors initiates recombination in V3 cells, as assessed by the plasmid substrate pJH290, which detects coding joints, or the pJH201 substrate, which detects signal joints, as indicated.

* Numbers from at least three separate experiments are presented.

* Calculated as (number of chloramphenicol-resistant colonies/number of ampicillin-resistant colonies) × 100.

Single phosphorylation site mutant constructs complement V3’s radioresistance similarly to constructs encoding wild-type DNA-PKcs. To assess the function of each phosphorylation site mutant, expression vectors carrying each coding sequence were transfected into V3 cells (Fig. 1B). The V3 cell line is DNA-PKcs deficient and thus is defective in DNA DSB repair (47). Clones with similar levels of DNA-PKcs expression were selected for each mutant. We next performed cell irradiation assays for this panel of DNA-PKcs mutants and compared the radioresistance to that of clones expressing wild-type human DNA-PKcs as well as to that of an empty vector control clone. As seen in Fig. 1C, each of the single phosphorylation site mutant constructs complemented the radioresistance of V3 cells similarly to the construct encoding wild-type DNA-PKcs. In some experiments (four of six) (data not shown), the transfectant expressing mutant C was slightly less radioresistant than the wild-type transfectant, suggesting that phosphorylation of Thr<sup>2638</sup> is functionally relevant. Indeed, it was found previously that Thr<sup>2638</sup> was a major in vitro phosphorylation site in DNA-PKcs (9). Recently, Chan et al. (3) reported that an alanine mutant of Thr<sup>2638</sup> (analogous to mutant A) could only partially complement the radioresistance of V3 cells. In our studies, the V3 T2609A clone generated displayed a radiosensitivity similar to that of transfectants expressing wild-type DNA-PKcs. Two additional clones expressing mutant A were also studied, and these clones also displayed radiosensitivity similar to that of clones expressing wild-type DNA-PKcs (data not shown). In the report of Chan et al., the single T2609A mutant V3 transfectant studied showed 10% survival at a dose of only 2.4 Gy, whereas the wild-type-expressing V3 transfectant showed 10% survival at a dose of 5 Gy. In our studies, the wild-type transfectant showed 10% survival at an average dose of 5.28 Gy (range, 4.28 to 6.24), similar to the dose for the wild-type transfectant studied by Chan et al. The average dose for 10% survival for the mutant A V3 transfectant was 4.47 Gy (range, 3.28 to 4.67). The discrepancy in these results is most likely due to individual clonal variation. In any case, our data demonstrate that the radioresistance of transfectants expressing single phosphorylation site substitutions is not dramatically different from that of wild-type transfectants. Thus, we conclude that phosphorylation of any single site within the cluster is not essential to DNA-PKcs function in NHEJ.

V3 transfectants expressing DNA-PKcs with multiple phosphorylation site mutations are extremely radiosensitive. The fact that six of seven autophosphorylation sites are tightly clustered within approximately 1% of the total amino acid sequence of DNA-PKcs suggested that concurrent phosphorylation of several sites within the cluster might be functionally important. Thus, we next generated expression constructs with two, three, five, or six mutations of serine or threonine to alanine. These are designated mutants AB, AC, AD, ABCD, and ABCDE, and they combine the mutations described previously. Stable cell lines with similar DNA-PKcs expression levels were selected for each mutant (Fig. 1D), and their radiosensitivities were assessed (Fig. 1E). Mutants AB, AC, and AD only partially complemented the radiosensitive phenotype of the V3 cells. Significantly, clones expressing DNA-PKcs with five or six phosphorylation site substitutions (mutants ABCD...
and ABCDE) displayed radiosensitivities analogous to (or more severe than) those of the vector-only control clones. These data demonstrate that concurrent autophosphorylation of DNA-PKcs on at least two of these clustered sites is a critical step in DNA damage repair, and they define DNA-PKcs as a physiologically relevant target of its own protein kinase activity.

Transiently expressed DNA-PKcs with single but not multiple phosphorylation site mutations support wild-type levels of coding end joining in V3 cells. We next assessed the ability of the mutant DNA-PKcs to support V(D)J recombination (Table 1). Wild-type DNA-PKcs substantially complemented the coding end joining defect of V3 cells (assessed with substrate pJH290). Similarly, all the mutants, with the exception of AC, ABCD, and ABCDE, supported near-wild-type levels of coding joint formation. Mutant AC supported reduced levels of coding end joining, whereas the abilities of the multiple phosphorylation mutants ABCD and ABCDE to support V(D)J recombination were severely impaired. These data are consistent with reports that minimal function of the NHEJ pathway can suffice to support V(D)J recombination (21, 39, 49). Thus, only transfectants that are severely impaired in NHEJ show reduced levels of V(D)J recombination.

Signal end joining is variably depressed in different DNA-PKcs-deficient cell lines and in different animal models of DNA-PKcs deficiency (10, 32, 43, 52). The V3 cell line displays a fairly significant defect in signal end joining (22). This is evidenced by the fact that wild-type DNA-PKcs increased the level of signal joints retrieved in transient assays approximately 30-fold (Table 1, pJH201 transfections). In contrast, signal joining was still impaired in cells transfected with the multisite mutant ABCDE. We conclude that the multiple phosphorylation site mutant cannot support normal levels of either signal or coding end joining during V(D)J recombination.

**V3 transfectants expressing DNA-PKcs with multiple phosphorylation site mutations display wild-type levels of DNA-PK activity.** DNA-PK activity in extracts derived from V3 transfectants was next assayed by using a previously described DNA-cellulose pulldown assay (11) and a peptide substrate (derived from the N terminus of p53). Transfectants expressing mutant ABCD or ABCDE displayed DNA-PK levels similar to those of transfectants expressing wild-type DNA-PKcs. Though DNA-PK activity was consistently somewhat higher in the transfectant expressing mutant ABCD, this increase might be explained by the slightly higher DNA-PKcs expression in the clone analyzed (Fig. 1D). We also utilized recombinant XRCC4 and Artemis as substrates in pulldown assays; the ABCDE mutant phosphorylated XRCC4 and Artemis similarly to

![FIG. 2. Transfectants expressing multiple phosphorylation site substitutions express normal levels of DNA-PK activity.](http://mcb.asm.org/)

as indicated. Each cell extract was tested in duplicate, and three independent extracts were tested for each cell line. Error bars, standard deviations. (B) Whole-cell extracts (250 µg) from V3 cells transfected with vector alone, wild-type DNA-PKcs, mutant ABCD, or mutant ABCDE were assayed for DNA-PK activity in extracts derived from V3 transfectants was next assayed by using a previously described DNA-cellulose pulldown assay (11) and a peptide substrate (derived from the N terminus of p53). Transfectants expressing mutant ABCD or ABCDE displayed DNA-PK levels similar to those of transfectants expressing wild-type DNA-PKcs. Though DNA-PK activity was consistently somewhat higher in the transfectant expressing mutant ABCD, this increase might be explained by the slightly higher DNA-PKcs expression in the clone analyzed (Fig. 1D). We also utilized recombinant XRCC4 and Artemis as substrates in pulldown assays; the ABCDE mutant phosphorylated XRCC4 and Artemis similarly to
FIG. 3. Autophosphorylation within the 2609-to-2647 cluster is not required for autophosphorylation-induced kinase dissociation. (A) Wild-type and ABCDE mutant DNA-PKcs’s were purified as described in Materials and Methods. Coomassie-stained SDS-PAGE gel shows analysis of fractions from a purification of the recombinant ABCDE mutant of DNA-PKcs. Lane 1, 10 μg of clarified extract from V3 cells expressing the ABCDE mutant; lane 2, 10 μg of flowthrough after the extract was loaded onto an affinity column coupled with a peptide from the C terminus of Ku80 (80peptide); lane 3, 2 μg of pooled fractions containing DNA-PKcs from 80peptide eluate; lane 4, 2 μg of flowthrough of Mono S column; lane 5, 2 μg of pooled DNA-PKcs-containing fractions from Mono S; lane 6, 2 μg of purified DNA-PKcs from placenta. Asterisk indicates location of major contaminant in eluate from 80peptide column. Similar results were obtained by purifying recombinant wild-type DNA-PKcs. (B) Kinase activities of purified wild-type and ABCDE mutant DNA-PKcs were assessed as described in Materials and Methods. DNA-protein complexes were preformed by incubation (at 30°C for 10 min) of 1 nM purified recombinant Ku, 1 nM either purified wild-type (wt) or purified ABCDE mutant DNA-PKcs, and 10 μg of sheared cell thymus DNA/ml. Reactions were started by transfer to 37°C and addition of 400 μM peptide substrate, 200 μM [γ-32P]ATP (0.5 Ci/mmol), and 5 mM MgCl2. The impact of prior autophosphorylation (pre-autophos.) was assessed in parallel reactions by including the same concentrations of ATP and MgCl2 in the preincubation step as well. (C) The stabilities of DNA-PK complexes containing wild-type or mutant DNA-PKcs were assessed by EMSA as described in Materials and Methods. DNA-protein complexes were formed with a 60-bp radiolabeled duplex, Ku, and wt or ABCDE mutant (m) DNA-PKcs as indicated. Complexes were preincubated at 75 mM salt by using either mock-treated DNA-PKcs, active kinase (+ Auto-phos.), or wortmannin-inactivated kinase (– Auto-phos.). Autophosphorylation was arrested by addition of 5 mM EDTA. Reactions were then adjusted to the indicated salt concentrations and incubated for a further 10 min at room temperature before surviving DNA-protein complexes were fixed by cross-linking with 0.25% glutaraldehyde. DNA-protein complexes were resolved by electrophoresis on a 3.5% native polyacrylamide gel.

wild-type DNA-PKcs (Fig. 2B). Finally, by using SDS-PAGE with a low percentage of acrylamide, a marked reduction in mobility was observed for highly phosphorylated wild-type DNA-PKcs. (The indicated phosphoproteins were confirmed to be DNA-PKcs by immunoblotting [data not shown].) As can be seen (Fig. 2C), although the ABCDE mutant protein can still be phosphorylated (possibly at position 3205 outside the major phosphorylation site cluster or perhaps at another, as yet unidentified site) hyperphosphorylation of the mutant protein (as evidenced by a gel shift) was not observed. In summary, DNA-PKcs containing multiple autophosphorylation site mutations is active toward exogenous substrates such as a synthetic peptide, XRCC4, and Artemis; therefore, the radiosensitive phenotype of cells expressing mutant ABCDE is not due to a lack of protein kinase activity per se.

DNA-PK complexes containing wild-type or ABCDE mutant DNA-PKcs display similar stabilities. To begin to assess the biochemical basis of defective NHEJ in cells expressing the ABCDE mutant, we purified the wild-type and ABCDE mutant proteins from the V3 transfectants (see Materials and Methods) (Fig. 3A). We have shown previously using purified proteins that autophosphorylation of DNA-PKcs results in dissociation of DNA-PKcs from Ku bound to DNA and loss of kinase activity. Thus, we tested whether preautophosphorylation of the mutant protein could affect kinase activity. Although the mutant protein is slightly more resistant to preautophosphorylation, preautophosphorylated DNA-PK complexes containing either wild-type or mutant DNA-PKcs have dramatically lower protein kinase activity than nonphosphorylated DNA-PK (Fig. 3B). The modest difference in resistance to autophosphorylation likely explains the variance in protein kinase activity observed over time with the mutant versus wild-type proteins without preautophosphorylation. Still, these data suggest that phosphorylation at additional sites facilitates kinase disassembly.

To more formally assess complex stability after autophosphorylation, a modified EMSA was also performed. Samples were prepared for analysis in two steps. First, complexes were allowed to assemble onto the DNA probe by using either wortmannin-inactivated DNA-PKcs or non-wortmannin-treated DNA-PKcs in the presence of ATP to allow autophosphorylation. After kinase activity was stopped (by the addition of EDTA), the relative stabilities of the complexes were probed by a second incubation in increasing NaCl concentrations. As can be seen (Fig. 3C), under more stringent conditions, neither autophosphorylated wild-type nor autophosphorylated ABCDE mutant DNA-PKcs forms stable complexes with Ku-
bound DNA, although complexes containing the mutant protein are consistently slightly more stable than wild-type complexes. These data suggest that further autophosphorylation is required for kinase disassembly.

An ABCDE mutant with aspartic acid substitutions is partially competent in NHEJ. The data presented above suggest that autophosphorylation outside of the major cluster might be important to signal kinase inactivation, at least in vitro. In an attempt to mimic constitutively phosphorylated DNA-PKcs in living cells, we next constructed an expression vector with aspartic acid substitutions for each of these six phosphorylation sites within the major cluster. We considered two possibilities. If phosphorylation within the major cluster induces release of DNA-PKcs bound to DNA, although complexes containing the mutant protein are consistently slightly more stable than wild-type complexes. These data suggest that further autophosphorylation is required for kinase disassembly.

Another possibility is that autophosphorylation within the major cluster is important for another reason, perhaps by inducing some conformational change in the repair complex that facilitates subsequent steps in the repair process. In this case, the mutant protein should not complement the NHEJ defect in V3 cells, and we would expect to observe reduced kinase activity in extracts from these cells. Another possibility is that autophosphorylation within the major cluster is important for another reason, perhaps by inducing some conformational change in the repair complex that facilitates subsequent steps in the repair process. In this case, the mutant protein should at least partially complement the NHEJ defect in V3 cells, and we would expect the mutant protein to be an active protein kinase.

The expression vector with aspartic acid substitutions at ABCDE (designated S/T→D) was transfected into the V3 cell line, and two cell clones with similar DNA-PKcs expression levels were studied (Fig. 4A). DNA-PK activity was assessed using the DNA-cellulose pulldown assay and the p53 peptide substrate. Transfectants expressing the S/T→A mutant displayed DNA-PK levels similar to those of transfectants expressing wild-type DNA-PKcs or the ABCDE mutant (Fig. 4B). As can be seen in Fig. 4C, although cells expressing the S/T→D mutant are still very radiosensitive compared to cells expressing wild-type DNA-PKcs, these cells are slightly more radioreistant than the vector-only control and significantly more radioreistant than the ABCDE mutant. Similarly, coding end resolution was slightly better in transient transfections comparing the S/T→D mutant to the ABCDE mutant (Table 1). We attribute this weak partial phenotype to poor mimicking of phosphoserine or phosphothreonine by the aspartic acid substitutions, although other explanations are also possible. Still, these data are consistent with a model whereby phosphorylation within the major cluster induces a conformational change that is requisite for subsequent steps in DNA end joining.

Autophosphorylation within the 2609-to-2647 cluster is not required for the interaction of DNA-PKcs with Artemis or to facilitate opening of coding end hairpins. The recent elegant studies of Ma et al. demonstrated that DNA-PKcs interacts with Artemis to generate an active DNA end-processing complex (31). To assess whether autophosphorylation within the 2609-to-2647 cluster is required for the interaction of Artemis with DNA-PKcs, a baculovirus vector expressing V5-His-tagged Artemis was constructed. Although only a fraction of the baculovirus-expressed, V5-His-tagged Artemis was soluble (Fig. 5A), a significant portion of the soluble protein fractionated onto Ni\(^{2+}\) agarose beads. Extracts from V3 transfectants expressing either wild-type DNA-PKcs or the DNA-PKcs mutant ABCD were incubated with extracts from Artemis-infected Sf9 cells or extracts from control Sf9 cells and then absorbed onto Ni\(^{2+}\) agarose. Wild-type DNA-PKcs and the multiple phosphorylation site mutant copurified equivalently with Ni\(^{2+}\) agarose-immobilized Artemis (Fig. 5A, lanes 11 and 12), suggesting that autophosphorylation within the major phosphorylation site cluster is not required for the interaction of DNA-PKcs with Artemis.

Ma et al. also demonstrated that the DNA-PKcs–Artemis complex opens hairpinned DNA ends (like the ones that result from RAG-mediated cleavage) and that DNA-PK's kinase activity is requisite for this activity. However, it was not determined whether DNA-PK's phosphorylation of Artemis or phosphorylation of DNA-PKcs itself is important for activating the hairpin nuclease. Thus, we next assessed broken DNA intermediates generated during transient V(D)J assays by us-
ing an LMPCR assay (Fig. 5B). Signal ends were detected by using a blunt-ended oligonucleotide in the ligation step. As expected, blunt signal ends are readily detected from Hirt supernatants prepared from cells transfected with vector alone (lane 4), wild-type DNA-PKcs (lane 5), or mutant ABCD (lane 6); lanes 7 to 9, Ni⁺-agarose fractions of control Sf9 extracts coincubated with extracts from V3 cells transfected with vector alone (lane 7), wild-type DNA-PKcs (lane 8), or mutant ABCD (lane 9); lanes 10 to 12, Ni⁺-agarose fractions of extracts from Sf9 cells expressing Artemis coincubated with extracts from V3 cells transfected with vector alone (lane 10), wild-type DNA-PKcs (lane 11), or mutant ABCD (lane 12). (B) LMPCR was performed on Hirt supernatants prepared from V3 cells transiently transfected with substrate alone (lanes 1 and 6), substrate and RAG expression vectors (lanes 2 and 7), substrate plus RAG and wild-type DNA-PKcs expression vectors (lanes 3 and 8), or substrate plus RAG and mutant ABCDE expression vectors (lanes 4 and 9). Lanes 5 and 10 included no input DNA in the LMPCRs. In lanes 1 to 5, a blunt oligonucleotide was ligated to Hirt fractions, and PCR amplifications to detect signal ends were performed. In lanes 6 to 10, an oligonucleotide with a 4-bp overhang complementary to a potential opened coding end in substrate pH290 was ligated to Hirt fractions, and PCR amplifications to detect coding ends were performed.

Thus, we conclude that the ABCDE mutant is fully capable of activating the hairpin-opening activity of Artemis.

**Coding joints mediated by mutant ABCDE have minimal nucleotide loss from joined coding ends.** Structural analyses of signal and coding joints as well as recombination intermediates has provided significant insight into the mechanistic details of both V(D)J recombination and NHEJ. Thus, we next sequenced rare coding joints mediated by the ABCDE mutant isolated from transient V(D)J recombination assays and compared them to sequences obtained from transfections including wild-type DNA-PKcs. As expected, coding joints obtained from recombination assays including wild-type DNA-PKcs yielded a diverse collection of sequences (Fig. 6; Table 2). Nucleotide loss from each joint ranged from 0 to 14 nucleotides, with an average of 4.61 bp lost per joint. Of the joints with at least one complete coding end (37 of 122 [30%]), P segments were apparent in 10 of 37 (27%). Short sequence homologies of 1 to 2 bp could be observed in 44% of the joints.
FIG. 6. Coding joints mediated by mutant ABCDE have minimal nucleotide loss from joined ends. The sequences of coding ends as they occur in the pJH290 substrate are shown above the sequences of the recombinant junctions. The number of observations of each sequence is given in parentheses to the right of the sequence. All duplicate sequences were derived from separate transfections. Nucleotides in columns headed by "p" are palindromic nucleotides added to each junction. Nucleotides that cannot be unequivocally assigned to a particular coding end are underlined and listed in the 5′/3′-most location. Portions of the wild-type (WT) and RAG-only coding joints are from references 21 and 49.
Though only a few rare coding joints were sequenced from transfections with no DNA-PKcs, structurally these were consistent with classic SCID joints. These joints display excessive nucleotide loss and a high incidence of long P segments when a coding end is complete (12 of 13 [92%]).

The most obvious difference between joints isolated from transfections with wild-type DNA-PKcs versus the ABCDE mutant is that there is remarkably less nucleotide loss at the site of joining in joints mediated by the mutant protein. Nucleotide loss from each joint ranged from 0 to 7 nucleotides, with an average of only 1.43 bp lost per joint. Furthermore, 70% of the coding ends (39 of 56) had no nucleotide loss versus only 30% (37 of 122) for joints mediated by wild-type DNA-PKcs. Of the joints with at least one complete coding end, P segments were apparent in 10 of 40 (25%). This is completely analogous to P segment incidence in joints mediated by wild-type DNA-PKcs and substantiates our conclusion that hairpin opening is normal in joints mediated by mutant ABCDE. Although four P segments were slightly longer than the normal 2 bp, slightly longer P segments were also occasionally observed in wild-type joints. Remarkably, short sequence homology at the junctures could not be observed in any of the 28 coding sequences, suggesting that this mechanism to facilitate end joining might be blocked in cells expressing the ABCDE mutant (although this may be partially attributed to the lower extent of base loss in these joints).

We also sequenced 25 coding joints mediated by the S/T→D mutant protein. These joints more closely resembled those isolated by wild-type DNA-PKcs: they averaged 3.08 bp lost per joint, and 19 of 50 coding ends (38%) were complete. Of the joints with at least one complete coding end, 5 of 19 (26%) had a P element, again analogous to P segment incidence in joints mediated by either wild-type DNA-PKcs or mutant ABCDE. Finally, short sequence homologies of 1 to 2 bp were apparent in 52% of the joints, also similar to the percentage in joints mediated by wild-type DNA-PKcs. In summary, these data suggest that end processing during V(D)J recombination is in some way impeded in the presence of the ABCDE mutant. We suggest that phosphorylation within the major cluster induces a conformational change that facilitates DNA end alignment and/or end processing prior to end joining.

### DISCUSSION

DNA-PK has been vigorously studied for more than a decade, and numerous potential substrates have been reported. In fact, more than 40 different proteins have been shown to be efficient in vitro targets of DNA-PK. To date, DNA-PKcs itself is the only unequivocally defined, functionally relevant target of its own kinase activity (3, 44).

Numerous spontaneous and targeted mutations of DNA-PKcs have been studied previously. These include three spontaneous germ line mutations (30, 32, 48), five spontaneous mutations in cultured cell lines (26, 52), four targeted germ line mutations (2, 12, 23, 45), and three reports of specific targeted mutations of DNA-PKcs expressed ectopically in cultured cells (21, 22, 49). In each of these, loss of DNA-PKcs function in vivo was associated with loss of DNA-PK activity. The multiphosphorylation site DNA-PKcs mutants are the first DNA-PKcs mutants that retain protein kinase activity but are incapable of supporting NHEJ. Thus, these mutants may be helpful in discerning other postulated functions (i.e., apart from NHEJ) of DNA-PKcs (16, 42).

We have explored several possible molecular mechanisms that could account for the radiosensitive phenotype of cells expressing the multisite autophosphorylation mutants of DNA-PKcs. Mutations in the autophosphorylation sites do not affect the in vitro protein kinase activity of DNA-PK, as judged by the ability of extracts from V3 cells containing DNA-PKcs with multiple autophosphorylation site mutations to phosphorylate synthetic peptide substrates, recombinant XRCC4, or recombinant Artemis. This suggests that the radiosensitive phenotype is due not to a lack of DNA-PK activity but rather is a consequence of the inability of DNA-PKcs to undergo autophosphorylation.

Similarly, in the autophosphorylation sites of DNA-PKcs do not affect the ability of DNA-PKcs to interact with or activate Artemis. In fact, if lack of Artemis activation were the only important deficit in cells expressing mutant ABCDE, one might expect complementation of V3’s signal joint deficit by mutant ABCDE, since Artemis is not required for signal end joining (35, 40). This is not the case. In sum, these data demonstrate that the function of DNA-PKcs in NHEJ goes beyond its role in activating the Artemis endonuclease. This is consistent with conclusions of a recent report directly comparing mice with targeted deletions of Artemis and DNA-PKcs (40).

The most revealing data as to the functional block of the multi-phosphorylation site mutants are the sequence analyses of coding joints mediated by the mutants. These joints have minimal nucleotide loss from rarely joined coding ends. Furthermore, there is no evidence of joining via short sequence homologies, a mechanism commonly observed in joints mediated by wild-type DNA-PKcs. These data imply that autophosphorylation of DNA-PKcs facilitates alignment via short se-
Several studies have shown that DNA-PKcs facilitates alignment or synapses of DNA-PK-bound DNA ends (7, 33, 50). Furthermore, this conformational change leads to remodeling of the DNA-PK complex. Nucleic Acids Res. 31:18992–18998.


