

Rad62 Protein Functionally and Physically Associates with the Smc5/Smc6 Protein Complex and Is Required for Chromosome Integrity and Recombination Repair in Fission Yeast

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Smc5 and Smc6 proteins form a heterodimeric SMC (structural maintenance of chromosome) protein complex like SMC1-SMC3 cohesin and SMC2-SMC4 condensin, and they associate with non-SMC proteins Nse1 and Nse2 stably and Rad60 transiently. This multiprotein complex plays an essential role in maintaining chromosome integrity and repairing DNA double strand breaks (DSBs). This study characterizes a *Schizosaccharomyces pombe* mutant *rad62-1*, which is hypersensitive to methyl methanesulfonate (MMS) and synthetically lethal with *rad2* (a feature of recombination mutants). *rad62-1* is hypersensitive to UV and gamma rays, epistatic with *rhp51*, and defective in repair of DSBs. *rad62* is essential for viability and genetically interacts with *rad60*, *smc6*, and *brc1*. Rad62 protein physically associates with the Smc5-6 complex. *rad62-1* is synthetically lethal with mutations in the genes promoting recovery from stalled replication, such as *rqh1*, *srs2*, and *mus81*, and those involved in nucleotide excision repair like *rad13* and *rad16*. These results suggest that Rad62, like Rad60, in conjunction with the Smc5-6 complex, plays an essential role in maintaining chromosome integrity and recovery from stalled replication by recombination.

Mutants in *Schizosaccharomyces pombe rad2*, *Saccharomyces cerevisiae RAD27*, and *Escherichia coli polA*, all of which are defective in processing Okazaki fragments, are synthetically lethal with mutations in recombination repair genes (8, 11, 23, 24, 35, 37, 40, 43, 44). In these mutants, double strand breaks (DSBs) are thought to be produced when replication forks encounter the single strand gaps or nicks remaining unsealed due to the inefficient processing of Okazaki fragments, so they require a highly efficient capacity to repair DSBs for survival. In an effort to identify novel genes involved in recombination repair, we isolated mutants that were hypersensitive to methyl methanesulfonate (MMS) and synthetically lethal with *rad2Δ* and cloned the genes by complementation of the MMS sensitivity. Genes identified in this way include *rhp57* (44), *rad60* (35), *rad32*, *nbs1* (45), and *fdh1*, which encodes an F-box DNA helicase (our unpublished data). A recent genome-wide search for mutations synthetically lethal with *rad27* in *S. cerevisiae* identified mutations in all of the well-studied recombination repair genes including *RAD52*, *RAD50*, *MRE11*, *XRS2*, *RAD51*, *RAD55*, *RAD57*, *RAD54*, *SGS1*, *MUS81*, and *MMS4* (43). The screen also identified genes involved in checkpoint regulation (*RAD9*, *RAD17*, *RAD24*, and *DDC1*) and those related to

regulation of chromatin structure (*CAC2*, *ESC2*, *HST1*, and *HST3*).

We report here a novel gene, *rad62*, which was identified by isolating mutants that were hypersensitive to MMS and synthetically lethal with *rad2* mutation and cloning the gene that complemented the MMS sensitivity of such a mutant. *rad62* mutants show very similar phenotypes to those of *rad60* mutants (7, 35) and *rad18* mutants (30, 47). They are hypersensitive to UV, MMS, and gamma rays, epistatic with *rhp51* with respect to the damage sensitivity, required for chromosome integrity during replication, and essential for growth. The *spr18* and *rad18* genes encode proteins whose structures are characteristic of the SMC family of proteins involved in chromosome condensation (Smc2-Smc4) and sister chromatid cohesion (Smc1-Smc3) (13, 21, 41). Since *RAD18* is a different gene in budding yeast, and the human homologs of *rad18* and *spr18* are named *hSMC6* and *hSMC5*, respectively (41), to avoid confusion we have used the names *smc6* and *smc5*, respectively.

SMC proteins share a common structure in that they contain two ATPase motifs composed of Walker A motifs at the N termini and Walker B motifs at the C termini, which are separated by long coiled-coil domains with a hinge region in the middle. The two coiled-coil domains interact and bring two ATPase motifs into close contact to form a globular structure (21). Eukaryotic SMC proteins including Smc5 and Smc6 form tight heterodimers (13). The condensin complex plays an essential role in folding extended chromosomes in an ATP-dependent manner during mitosis. The cohesin complex forms a ring structure which holds sister chromatids in close proximity after replication until they are separated at anaphase in mitosis

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TABLE 1. *S. pombe* strains used in this study

Strain	Genotype	Source or reference
MMPX1	<i>mat1PΔ17::LEU2 leu1-32 ura4-D18</i>	Laboratory stock
MMPX2	<i>mat1PΔ17::LEU2 rad62-1 leu1-32 ura4-D18</i>	This study
MMPX3	<i>smt-0 rhp51::ura4 leu1-32 ura4-D18</i>	Laboratory stock
MMPX4	<i>mat1PΔ17::LEU2 rad62-1 rhp51::ura4 leu1-32 ura4-D18</i>	This study
MMPX5	<i>smt-0 rhp57::his3 ura4-D18 leu1-32 his3-D1arg3-D1</i>	Laboratory stock
MMPX6	<i>smt-0 rad62-1 leu1-32 ura4-D18</i>	This study
MMPX7	<i>smt-0 rad62-1 leu1-32 ura 4-D18 his3-D1</i>	This study
MMPX8	<i>mat1PΔ17::LEU2 swi5::his3 leu1-32 ura4-D18</i>	Laboratory stock
MMPX9	<i>smt-0 rad62-1 swi5::his3 leu1-32 ura4-D18 his3-D1</i>	This study
MMP1	<i>h⁺ leu1-32 ura4-D18</i>	Laboratory stock
MMP2	<i>h⁺ rad62-1 leu1-32 ura4-D18</i>	This study
MMP5	<i>h⁻ rad2::ura4 leu1-32 ura4-D18 ade6-M210</i>	Laboratory stock
MMP6	<i>rad62-1 rad2::ura4 leu1-32 ura4-D18</i>	This study
MMP7	<i>h⁺ rad62-1 leu1-32 ura4-D18 his3-D1</i>	This study
MMP8	<i>rad62::ura4 leu1-32 ura4-D18 ade6-his7-366 (carrying pGFP62)</i>	This study
MMP9	<i>h⁻ rad13::ura4 leu1-32 ura4-D18 his3-D1</i>	This study
MMP10	<i>h⁺ rhp18::ura4 leu1-32 ura4-D18 his3-D1</i>	Laboratory stock
MMP11	<i>h⁻ rad62-1 rad13::ura4 leu1-32 ura4-D18 his3-D1</i>	This study
MMP12	<i>h⁺ rad62-1 rhp18::ura4 leu1-32 ura4-D18 his3-D1</i>	This study
MMP13	<i>h⁻ rad16::ura4 leu1-32 ura4-D18 his3-D1</i>	This study
MMP15	<i>rad62-1 rad50::LEU2 leu1-32 ura4-D18</i>	This study
MMP16	<i>h⁻ srs2::Kan^r leu1-32 ura4-D18 his3-D1</i>	This study
MMP17	<i>rad62-1 mus81::kanMX</i>	This study
MMP18	<i>smt-0 rad62-1 uvel::his3 leu1-32 ura4-D18 his3-D1</i>	This study
MMP20	<i>h⁻ leu1-32 ura 4-D18 rad62-FLAG-His smc5-Myc Kan^r</i>	This study
MMP21	<i>h⁻ leu1-32 ura 4-D18 rad62-FLAG-His Kan^r</i>	This study
MMP22	<i>h⁺ leu1-32 ura 4-D18 smc5-Myc Kan^r</i>	I. Miyabe
MMP23	<i>h⁻ leu1-32 ura4-D18 his3-D1 uvel1::his3</i>	This study
MMP24	<i>leu1-32 ura4-D18 his3-D1 rhp57::his3 rad62-1</i>	This study
MMP30	<i>h⁻ mus81::kanMX</i>	This study
PS3	<i>ade6-M375 int::pUC8/ura4/ade6-L469</i>	17
MPS401	<i>rad62-1 ade6-M375 int::pUC8/ura4/ade6-L469</i>	This study
MPS501	<i>rhp51::his3 ade6-M375 int::pUC8/ura4/ade6-L469</i>	This study
298	<i>h⁻ rad50::LEU2 leu1-32 ura4-D18</i>	42
TE767	<i>rgh1::ura4</i>	27
NB2554	<i>mus81::kanMX</i>	6
NCYC1979	<i>h⁻ rad18-X</i>	NCYC ^a
MMPD1	<i>h⁺/h⁻ leu1-32/leu1-32 ura4-D18/ura4-D18 his7-366/+ ade6-M210/ade6-M216 rad62::ura4/+</i>	This study

^a NCYC, National Collection of Yeast Cultures.

(16). The Smc5-Smc6 (Smc5-6) complex is proposed to play a role in repairing DSBs by recombination by holding broken DNA together (41). In common with the other eukaryotic SMC complexes, the Smc5-6 heterodimer is associated with non-SMC protein subunits, Nse1 and Nse2, and interacts with another essential protein, Rad60 (7, 19, 33, 35). Like *smc5* and *smc6*, *rad60*, *nse1*, and *nse2* are essential for growth, and the hypomorphic mutants are sensitive to DNA damaging agents.

Analysis of the genetic interactions between *rad60* and *smc6* mutants suggests that the products of these genes play a role in a common pathway for repairing DSBs and maintaining chromosome integrity (35). A recent study demonstrated that substoichiometric amounts of Rad60 were associated with Smc5-6 (7). Chromosomes were fragmented when a temperature-sensitive *rad60-1* mutant was grown at nonpermissive temperature, suggesting that the Rad60 protein is required to prevent chromosome breakage during replication (35). Like *smc6* and *rad60* mutants (35, 47), the *rad62-1* mutant is defective in joining DSBs produced by ionizing radiation and *rad62* is essential for growth. From the phenotypic similarities and analysis of genetic and physical interactions, we propose that

Rad62 plays a role as a non-SMC component of the Smc5-6 complex in maintaining chromosome integrity and repairing DSBs.

MATERIALS AND METHODS

***S. pombe* media and methods.** Yeast cells were grown in complete medium (YES), minimal medium (EMM2), or sporulation medium (ME), and standard yeast genetic and molecular procedures were as described previously (34). Genetic interactions were assessed through standard mating techniques, followed by tetrad analysis on YES plates as described previously (35).

Strains and plasmids. The strains and plasmids used in this study are shown in Tables 1 and 2. pUR19 is a multicopy plasmid which contains the *arsI* and *ura4* genes (3). pREP41 is a plasmid that contains a weak *nmt* promoter, which is inducible by thiamine depletion (39).

Gamma ray, MMS, and UV sensitivity tests. Cells were grown to mid-log phase in YES medium, resuspended in water, and then irradiated with gamma rays from a ⁶⁰Co source. After irradiation, appropriately diluted samples were plated on YES plates and incubated at 30°C for 4 to 7 days before colony counting. For test of UV sensitivity, appropriately diluted samples were plated on YES plates, irradiated with the indicated dose of UV, and incubated at 30°C for 4 to 7 days. For the test of MMS sensitivity, 3 μl of sequential 10-fold dilutions of stationary-phase cells were spotted on EMM plates with the indicated MMS concentration.

TABLE 2. Plasmids used in this study

Plasmid	Insertion ^a	Plasmid used for derivation	Source or reference
pSLR4	<i>rad62</i> genomic	pUC118	This study
pSLR4D	<i>rad62::ura4</i>	pSLR4	This study
pREP41	<i>ars1 LEU2 nmt1*</i>		31
pREP41X	<i>ars1 LEU2 nmt1*</i>		12
pBRC1	<i>ars1 LEU2 nmt1* brc1</i> cDNA	pREP41X	M. O'Connell
pRAD62	<i>ars1 LEU2 nmt1* rad62</i> cDNA	pREP41	This study
pRAD60	<i>ars1 LEU2 nmt1* rad60</i> cDNA	pREP41	This study
pREP41EGFP-N	<i>ars1 LEU2 nmt1*</i> EGFP	pREP41X	9
pGFP62	<i>ars1 LEU2 nmt1*</i> EGFP- <i>rad62</i> cDNA	pREP41EGFP-N	This study

^a *, nmt medium-strength promoter.

Cloning of the *rad62* gene. *rad62-1* mutant cells were transformed with an *S. pombe* genomic DNA library (3), and MMS-resistant transformants were selected on plates containing 0.004% MMS (44). The plasmids that conferred MMS resistance to the mutant were isolated, and the region responsible for the recovery of the repair activity was identified (Fig. 1A) and sequenced.

Determination of the *rad62* coding region. The *rad62* cDNA was amplified by PCR with an *S. pombe* cDNA library constructed on the pGAD GH vector (Clontech) as a template and analyzed as follows. Analysis of the 5' region with primers GAD5-1 (5'-TATTCGATGATGAAGATACC-3'), corresponding to an upstream flanking sequence of the vector, and V1R1 (5'-ATAGTAGGTTGAG TTAGCAC-3'), corresponding to a middle region of a putative *rad62* exon, revealed three introns, and analysis of the 3' region with primers V1F2 (5'-GT CCTTATCATTTTCGCAAG-3'), corresponding to a region in the same putative *rad62* exon, and T7 (5'-AATACGACTCACTATAG-3'), corresponding to a downstream flanking sequence of the vector, revealed an additional three introns, as shown in Fig. 1B. A 5' rapid amplification of cDNA ends (5' RACE) experiment to identify the transcription initiation site was carried out by using the FirstChoice RLM-RACE kit (Ambion) and the V1R1 primer. Nucleotide sequence analysis of the six clones obtained indicated that there were at least four initiation sites corresponding to the nucleotide numbers 162, 197, 216, and 217 (Fig. 1B). A 3' RACE experiment was carried out by using the LA PCR kit, version 2 (TAKARA). DNA sequence analysis of the two clones indicated the poly(A) sites at nucleotide numbers 1592 and 1596, respectively.

Based on the above results, the *rad62* open reading frame encoding 300 amino acids was identified (Fig. 1B). The cDNA for the *rad62* coding region was amplified with primers S4NB (5'-GCGGATCCCATATGTCCTCCATTGATA AAC-3'), corresponding to the region for translational initiation, and S4CB (5'-GCGGATCCTCAGCCATACCAAGTATTAC-3'), corresponding to the region for translational termination by using the LA PCR kit, version 2. The resulting reverse transcription-PCR product was digested with BamHI and ligated to the BamHI site of pUC118. Nucleotide sequence analysis of the cDNA confirmed the existence of six introns.

Construction and characterization of *rad62* deletion strain. A *rad62* deletion mutant was constructed essentially as described previously (15). pSLR4, a derivative of pUC118 carrying the 2.7-kb BamHI-BamHI region containing the *rad62* gene (Fig. 1A), was used to generate a plasmid construct for gene disruption. The 0.9-kb HpaI-XhoI region of the *rad62* gene on the plasmid was removed, and the staggered ends were converted to blunt ends by filling them in with *E. coli* DNA polymerase I. The ends of the 1.8-kb HindIII fragment containing the *ura4* gene from pREP2 (31) were converted to blunt ends, and the fragment was ligated with the plasmid to generate pSLR4D (Fig. 1A). pSLR4D was linearized by BamHI digestion and transformed into the diploid *S. pombe* strain MPD1. The Ura⁺ transformed colonies were isolated and checked for the *rad62* deletion by genomic Southern analysis. They were sporulated and subjected to tetrad analysis. For cytological observation, the diploid cells were incubated on ME at 26°C for 2 days for sporulation. The spores were treated with 0.5% glusulase, washed in water three times, and then spread onto an EMM plate lacking uracil. After 2 to 3 days, germinated spores of the *rad62::ura4*⁺ strain were collected and washed in water three times. The cells were then fixed in 70% ethanol and stained with 1.5 µg of 4, 6-diamidino-2-phenylindole (DAPI)/ml. The stained cells were observed under an epifluorescent microscope.

PFGE. To examine the ability of the *rad62-1* mutant to repair DSBs, the cells were irradiated with gamma rays at 600 Gy and the repair of DSBs was followed by pulsed-field gel electrophoresis (PFGE) as described previously (35), with

slight modifications in the PFGE conditions: voltage gradient, 2 V/cm; pulse time, 30 min; angle, 120°; time, 48 h; temperature, 16°C.

Measurement of mitotic homologous recombination frequency. The mitotic homologous recombination of *rad62-1* was measured by the efficiency of integration of a linear DNA carrying the *leu1*⁺ gene into the *leu1-32* locus on the chromosome as described previously (44).

Spontaneous mitotic intrachromosomal recombination frequencies of *rad62-1*, *rhp51Δ*, and wild-type cells were measured by using the strains PS3, MPS401, and MPS501, containing a nontandem direct repeat of *ade6*⁻ heteroalleles flanking *ura4*⁺ as described previously (17, 38).

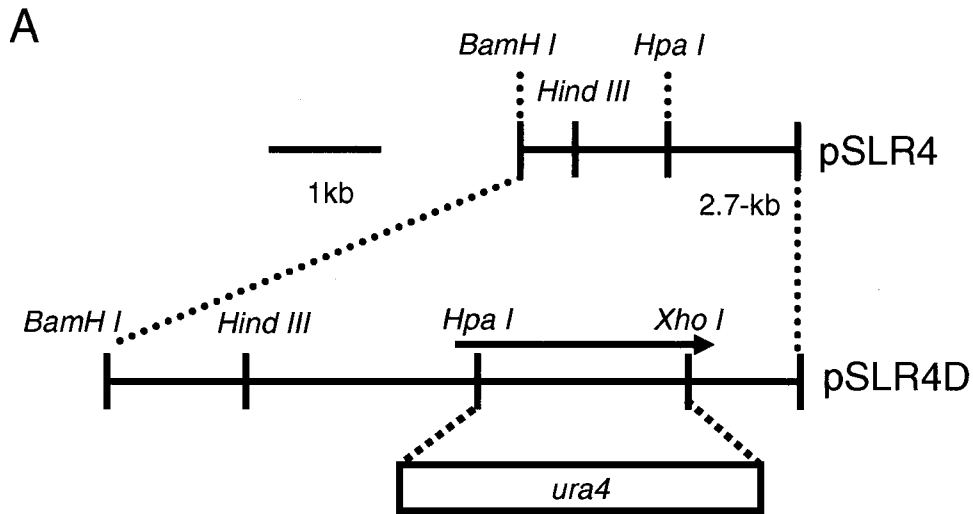
Construction of epitope-tagged genes. For immunoprecipitation experiments and protein complex purification, an epitope-tagged *rad62* gene generated as described previously (27a) by the PCR-based method to place a FLAG-His epitope at the C terminus of the protein and mark the allele with the *kanMx6* gene (27a). The *rad62-1* strain containing the tagged *rad62* gene was not sensitive to MMS and grew as normally as the wild-type strain, indicating that the tagged gene is functional.

Immunoprecipitation. The cells carrying the epitope-tagged gene were grown in YES medium. Mid-log-phase cells from a 50-ml culture were collected, resuspended in 500 µl of buffer A (10 mM HEPES [pH 8.0], 0.1% NP-40, 25 mM KCl, 1 mM phenylmethylsulfonyl fluoride) containing protease inhibitor cocktail, and lysed by acid-washed glass beads. The lysate was clarified by centrifugation (12,000 × g, 15 min). Ten microliters of protein G-Sepharose 4FF (Amersham Pharmacia Biotech) prewashed in buffer A was added to adsorb nonspecific binding proteins to protein G. The suspension was incubated for 1 h at 4°C, and the supernatant was collected by centrifugation. Anti-Myc antibody (PL14; MBL) with protein G-Sepharose 4FF or anti-FLAG M2 affinity gel (Sigma) was added to the supernatant, and the suspensions were rocked for 1 h at 4°C. Immunoprecipitates were washed three times with buffer A, resuspended in 40 µl of 5% sodium dodecyl sulfate (SDS), and incubated for 10 min at 37°C. After centrifugation, the supernatants were mixed with 10 µl of SDS-polyacrylamide gel electrophoresis (PAGE) sample buffer, boiled for 10 min, separated by SDS-PAGE, and analyzed by Western blotting with the ECL Advance Western blotting detection kit (Amersham Bioscience).

Nucleotide sequence accession number. Nucleotide sequence data, including the exon and intron information, have been deposited with the DDBJ under accession number AB158548.

RESULTS

Isolation of *rad62-1* mutant defective in repairing DSBs and cloning of the gene that complemented the repair deficiency. *rad62-1* is one of the seven mutants previously isolated as sensitive to MMS and synthetically lethal with the *rad2* mutation (44). The original mutant was repeatedly backcrossed with the wild-type strain, and the clones sensitive to MMS were selected. A genomic DNA fragment that conferred the MMS resistance to the mutant was isolated from an *S. pombe* genomic DNA library. The original plasmid carried a 4.1-kb genomic DNA insert, whose physical map is shown in Fig. 1A. The region that complemented the MMS sensitivity was sub-



B

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1  ttatagggttttatgeaattaatagaaaaaagtaoaaaggctococataaataattott  60
61  tgaaatgatccasatatactaataatgtttotgcoogtatatacoagttataatttgostatg  120
121  otogttgaoaaatattttotgogtttagoagcaactgtgttatttagoatcaactaac  180
181  gattaagotaacactttsatggcaacattotttctcaacgttttaottttttaatttttaa  240
241  aattattaatotaasaagooaatotttggagatttggottaaaggttgaagatttgotog  300
301  cgatcaaatcoctcatgaggaaotgggcaaatttttgtaoctcaaaatotaatttcgtoa  360
361  totagottatatttgaataaaaotatacaaaaaaooocagatatactcaatcattgata  420
421  acgggcatotttggaaacgcatatgkgaacttaataagcaaaagttcaagtcocatstaoo  480
      R D L R K R Y R N L I N K V Q E
481  tttaecttataaatacaacttttagoaaagatortttgaaactgctgagggaa  540
      S R L E L L V D E E
541  aacnataactttatatgaactattactactgcaatgattttttttttcagatgaatat  600
      N N N L Y E T I T T A N D L F S S V
601  caagtgtatotttgcactgttaacaataaataatgctgctgctgctgcaagcaat  660
      D A P T E A T L
661  takatgatttactgcttactaaacgcttactgctgctgctgctgctgctgctgctgctg  720
      D A L L L T K T V D L A S I K A R Q E L H
721  ttattgaaagcccaacttttaattttgcaatattcaagaaatattgaaacttctctca  780
      I G R P K F N I E L F T K N I K Q F L N
781  actatgctacttccactgcaatgcttactgcaatccaaagattgcaacacatgctcaa  840
      Y P T S H S N V T R I Q E I D T A W S R
841  gcatagkcaattgcaactaattgcaaaagcaagccgctctctctgcaacttctctatg  900
      L G K L A S N C E K Q P A S L N L M
901  taaatagttttocaaatgcttatotaaootttttagggctgctgcttctgctgctg  960
      V G P L S F R
961  agagaaagaaagaaagaaagaaagaaagaaagaaagaaagaaagaaagaaagaaagaa  1020
      K K E R N I Q R R E R L G K A P N V L T
1021  caagctactatgataatttttttttttttttttttttttttttttttttttttttttt  1080
      Q P T M L N E R
1081  ctaacattactaccgaagaaatgaagaaatgaagaaatgatttgaagaaatgatttga  1140
      N I T T Q E N N T T K N V L H I S R L L
1141  ttgaaagaaagaaagaaagaaagaaagaaagaaagaaagaaagaaagaaagaaagaa  1200
      Q A H Q P V N F L K F I T N P Q S Y P Q
1201  aaagaaagaaagaaagaaagaaagaaagaaagaaagaaagaaagaaagaaagaaag  1260
      T V E N L F Y V S F L F K E G K A A L V
1261  tgaagaaagaaagaaagaaagaaagaaagaaagaaagaaagaaagaaagaaagaa  1320
      E N E S G I L N L E
1321  ttattaaactgttagaaagaaagaaagaaagaaagaaagaaagaaagaaagaaagaa  1380
      F R I P P T D D Q V V A G E I
1381  tgaagaaagaaagaaagaaagaaagaaagaaagaaagaaagaaagaaagaaagaa  1440
      R N I D L V L D M I M D L Y E
1441  tgtttaaaatttttttaeocotocgaatgcaagaaatattgaagaaatcaaatgca  1500
      N I I K E Y N I K
1501  agaaagaaagaaagaaagaaagaaagaaagaaagaaagaaagaaagaaagaaagaa  1560
      E S I T P T R A P V E Y S T I N S N I W V
1561  tgcctgctactaatacaacaatttttaatttcaagtttaattaccatatttgacataac  1620
      G
1621  gottogtaattaagataaaaaaatatataatagatgataaaaaatagagcttcoata  1680
1681  oocooaaacttttttaacaaatcaagtcocatttcttttttttaacaaagaaagatog  1740
1741  toaataaaataaataaataaatttttagaagaaagaaatattttttgcatcaaaact  1800
1801  ttaggcttaagaaatgataaacaataaataaataaagaaagaaagaaagaaagaa  1860
1861  gtaggactaaagtatttttagaagaaatatttttttaacaaattgtaagaaatgata  1920
1921  gcaagaaagaaagaaagaaagaaagaaagaaagaaagaaagaaagaaagaaagaa  1980
1981  ototaacggctgatcttgt  2000
    
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FIG. 1. Physical map of the *rad62* genomic region and construction of a *rad62* deletion mutant. (A) A restriction map of the *rad62* region is shown at the top. The plasmid containing the *rad62* deletion was constructed by replacing the HpaI-XhoI fragment in *rad62* with the *ura4* gene, as shown at the bottom. (B) Nucleotide sequence of the *rad62* gene and predicted amino acid sequence of the Rad62 protein. The exons are underlined. Arrows show the transcription initiation sites. The poly(A) sites are shown by vertical lines. The amino acid alteration from Lys to Glu at position 223 caused by a mutation at position 1239 in *rad62-1* mutant is shown.

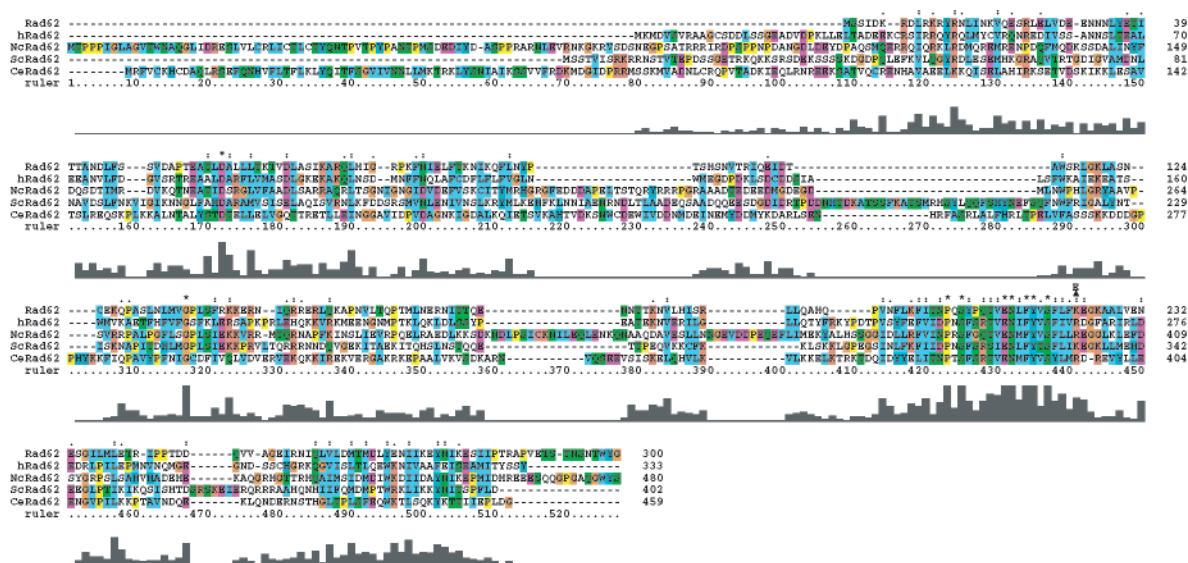


FIG. 2. Sequence alignment of Rad62 proteins from various organisms. Multiple-sequence alignment was performed with CLUSTAL X, version 1.81 (<http://ftp-igbmc.u-strasbg.fr/pub/ClustalX/>). *hRad62*, human Rad62 (accession number AAH27612.1); *NcRad62*, *Neurospora crassa* Rad62 (accession number EAA36561.1); *ScRad62*, *S. cerevisiae* Rad62 (Qri2) (accession number CAA55925.1); *CeRad62*, *Caenorhabditis elegans* Rad62 (accession number CAB09113.1). The arrow shows the position of the *rad62-1* mutation. The levels of sequence homology are shown by the heights of the blocks at the bottom.

cloned on a shuttle vector, and the DNA sequence of the 2.8-kb BamHI region was determined. The sequenced region matches a region contained on the cosmid c20F10 (EMBL accession number AL021747).

The *rad62* coding region was analyzed by the 5' and 3' RACE and reverse transcription-PCR methods, and the existence of seven exons in the *rad62* gene was shown by sequence analysis of the cDNA (Fig. 1B). The gene is unusual in having at least as many as four transcription initiation sites and such a long untranslated upstream exon. The open reading frame of *rad62* was determined from these analyses and is shown in Fig. 1B. It was different from the one described in the DNA database as SPBC20F10.04c. The gene encodes a protein with 300 amino acids with no apparent functional sequence motifs, as analyzed by using databases (<http://motif.genome.ad.jp/>). Database search revealed that the Rad62 homologs exist ubiquitously throughout eukaryotes, as shown in Fig. 2. The nucleotide sequence covering the *rad62* region of the *rad62-1* allele and its flanking genomic regions were amplified by PCR, and the nucleotide sequence was analyzed directly. A single transition mutation from A to G was found at the first base of codon 223, altering the AAA codon for lysine to a GAA codon for glutamic acid. Codon 223 is likely to be important for function, as the corresponding sites are conserved as positive residues, including Lys and Arg, and the mutation changing Lys to an acidic residue, Glu, caused the functional defect in the *rad62-1* mutant.

The *rad62-1* mutant is defective in repairing DNA damage and epistatic with *rhp51*. The sensitivities of the *rad62-1* mutant to DNA-damaging agents, UV and gamma rays, were examined by counting survival fractions after irradiation at various doses (Fig. 3). The mutant showed two phases of response to UV irradiation, a highly sensitive phase at low doses up to 25 J/m² and a relatively resistant phase at higher doses

than that. The *rhp51* gene, which is a homolog of *S. cerevisiae* *RAD51* and *E. coli* *recA*, plays a major role in repairing DSBs by homologous recombination and is required for the UVER repair pathway, which is unique to fission yeast and some fungi (29, 32). The *rad62-1* mutant was more sensitive to low doses (up to 50 J) of UV than the *rhp51Δ* mutant, but it was less sensitive at higher doses. *rhp51Δ* partially suppresses the UV sensitivity of *rad62-1* at the lower doses, and the *rad62-1 rhp51Δ* double mutant showed a similar UV survival curve to that of the *rhp51Δ* single mutant. This result suggests that, at low doses of UV, the Rhp51 protein converts DNA damage to a toxic recombination intermediate in the *rad62-1* mutant. The *rad62-1* mutant is also hypersensitive to gamma rays, but it is less sensitive than the *rhp51Δ* mutant at all doses tested. The double mutant showed the same sensitivity as the *rhp51Δ* single mutant. These results suggest that *rhp51* is epistatic to *rad62* with respect to the DNA repair function. *rhp57Δ* is also epistatic to *rad62-1* (data not shown). *rhp51* is epistatic to *rhp57*, and both of them are involved in recombination repair (25, 44). *rad60-1* and *smc6-X* (*rad18-X*) mutants are hypersensitive to UV and gamma rays and show a similar epistasis with *rhp51* (30, 35).

Since *rad62* belongs to the same epistasis group as the recombination repair genes *rhp51* and *rhp57* and the *rad62-1* mutant showed phenotypes very similar to those of the *rad60-1* and *smc6-74* (*rad18-74*) mutants (35, 47), we examined whether the mutant was also defective in repairing DSBs. DSBs were induced by irradiation at 500 Gy, and DNA isolated at various times after irradiation was analyzed by PFGE. As shown in Fig. 3B, fragmented chromosomes were mostly re-joined in wild-type cells by incubation for 3 h while they remained unjoined fragments in the *rad62-1* cells even after incubation for 6 h. The results strongly support the hypothesis that Rad62, Rad60, Smc5, and Smc6 function in the same

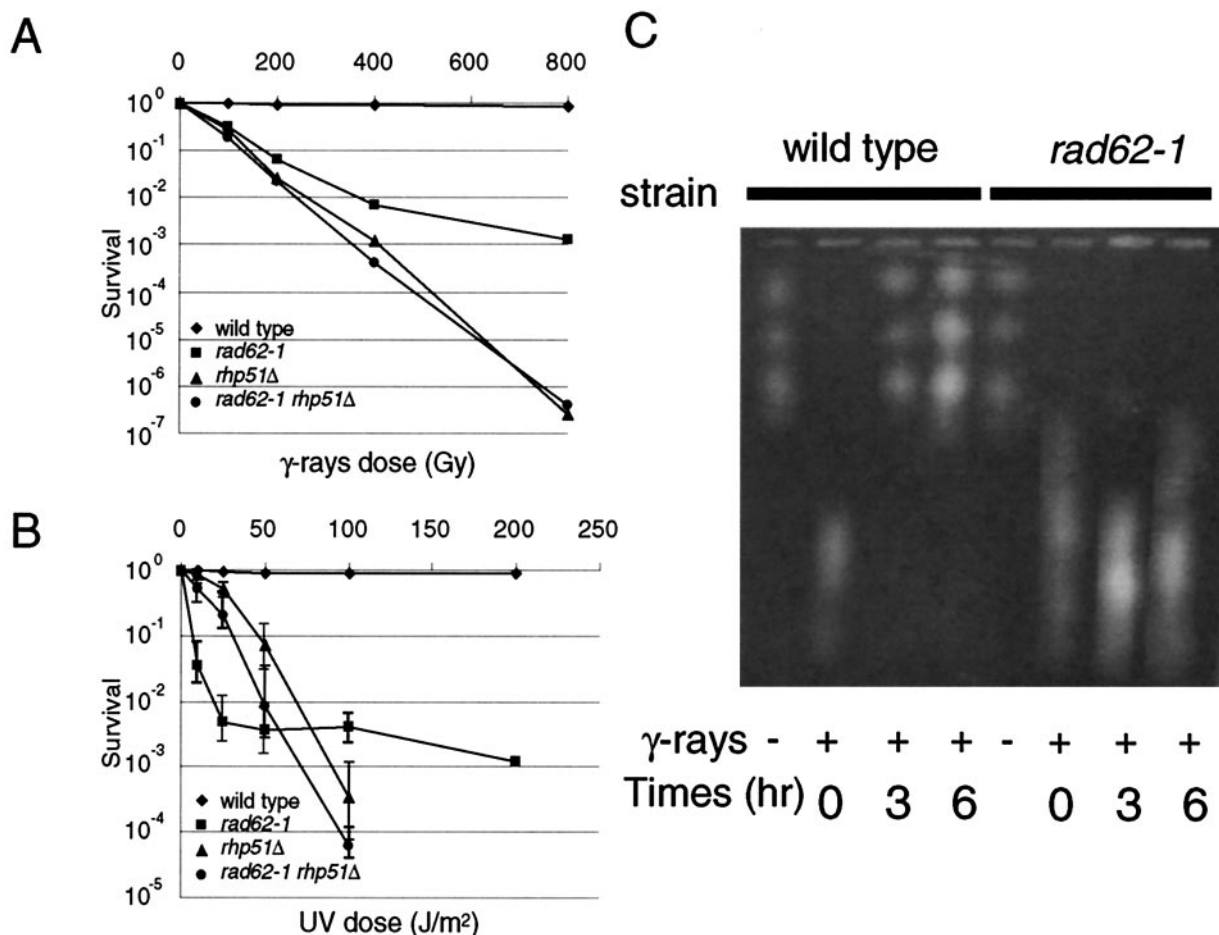


FIG. 3. The *rad62-1* mutant is defective in DNA repair. (A and B) Sensitivities of *rad62-1* mutant to UV and gamma rays. The cells at mid-logarithmic phase were irradiated with UV or gamma rays at the indicated doses, and the relative plating efficiencies were determined. Wild-type strain, MMP1 (◆); *rad62-1* strain, MMP2 (■); *rhp51 Δ strain, MMPX3 (▲); *rad62-1 rhp51 Δ strain, MMPX4 (●). The data represent the averages of the results from three experiments. (C) The *rad62-1* mutant is defective in repairing DNA DSBs. Genomic DNA of wild-type and *rad62-1* strains was analyzed by PFGE. Samples were taken at the indicated times (in hours) after irradiation by gamma rays at 500 Gy (+) or before irradiation (-).**

pathway with recombination proteins such as Rhp51 and Rhp57 for repairing damaged DNA.

The synthetic lethality of *rad62-1* with *rad2* Δ was examined by constructing a diploid strain by mating the two mutant strains *rad62-1* (MMP7) and *rad2* Δ (MMP5) and analyzing the resultant tetrads. The double mutants formed extremely small colonies, and when they were streaked on a YES plate, they showed little sign of growth, confirming that the two mutations are synthetically lethal (data not shown).

***rad62-1* is not defective in Rhp51-dependent genetic recombination.** Since the efficiency of homology-dependent genetic recombination is reduced in *rhp51* and *rhp57* mutants, we examined the ability of the *rad62-1* mutant to carry out the recombination. The efficiency of integrating the *leu1*⁺ gene into the *leu1-32* locus of the *rad62-1* mutant (MMPX6) was not reduced, whereas it was reduced threefold (34%) in the *rhp57* Δ mutant (MMPX5). This shows that the *rad62-1* mutant is not defective in recombination during mitosis, as examined by the integration of linear homologous DNA into the chromosome.

rad60-1 was also not defective in such recombination (our unpublished data).

Next we examined spontaneous mitotic intrachromosomal recombination frequencies in the *rad62-1* mutant containing a nontandem direct repeat of *ade6*⁻ heteroalleles flanking *ura4*⁺, and intrachromosomal mitotic recombination was assayed by the formation of Ade⁺ recombinants. Two main classes of Ade⁺ recombinants could be distinguished: Ade⁺ Ura⁻ deletion type recombinants and Ade⁺ Ura⁺ conversion type recombinants.

The frequency of conversion type of recombination was much reduced in the *rhp51* Δ mutant (0.16 ± 0.07 per viable 10^4 cells) compared to wild-type cells (1.58 ± 0.30), but it was not reduced in *rad62-1* cells (1.42 ± 0.72 per 10^4 viable cells). This result suggests that the *rad62-1* mutant is not defective in Rhp51-dependent spontaneous intrachromosomal recombination such as the gene conversion type of recombination. On the other hand, deletion type recombination was significantly increased in the *rhp51* Δ mutant (9.68 ± 1.44 per 10^4 viable cells)

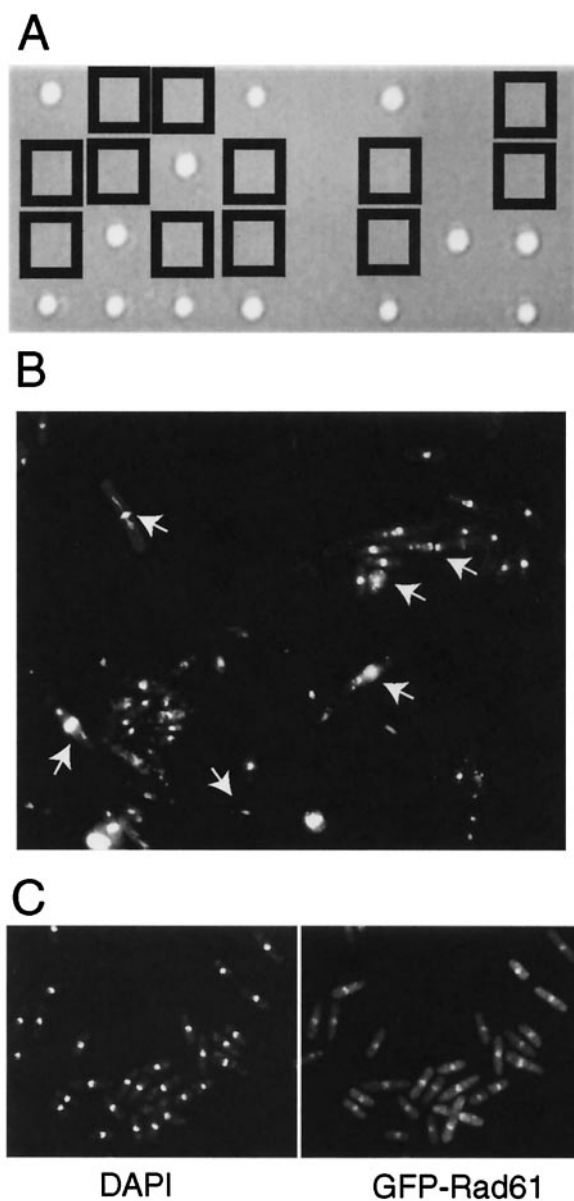
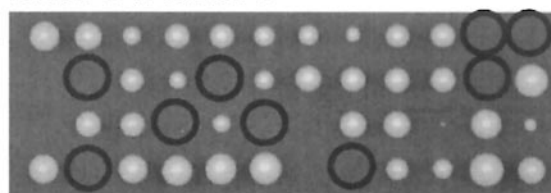


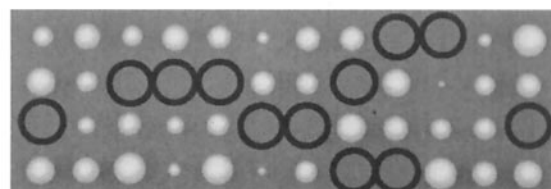
FIG. 4. Rad62 is an essential nuclear protein. (A) Tetrad analysis of *rad62* heterozygous diploids. The diploid strain heterozygous for the *rad62* deletion mutation was sporulated and subjected to tetrad analysis. The segregants from the dissected spores were grown on a YES plate at 30°C for 5 days. The *rad62*Δ segregants are boxed. (B) Terminal morphology of *rad62*Δ cells. The diploid strain MMPD1 was sporulated and germinated on an EMM2 plate lacking uracil. The *rad62*Δ cells grown for 2 days were fixed with 70% ethanol, stained with DAPI (1 μg/ml) and calcofluor white (20 μg/ml), and photographed. The abnormal nuclei are shown by arrows. (C) Nuclear localization of Rad62 protein. Cells of the *rad62*Δ (MMP8) carrying a plasmid for expression of GFP-Rad62 fusion protein, pGFP62, were fixed, stained with DAPI (1 μg/ml), and observed under an epifluorescence microscope. Fluorescence images of DAPI (left) and GFP-Rad62 (right) are shown.

compared to wild-type cells (1.29 ± 0.85 per 10^4 viable cells), and it was not affected in the *rad62-1* mutant (1.30 ± 0.59 per 10^4 viable cells). Therefore, the *rad62-1* mutant is similar to the wild-type strain and different from *rhp51* and *rhp57* in regard to the recombination phenotypes examined by these methods.

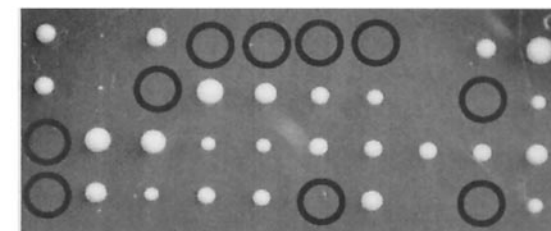
rad62-1 × *rad60-1*



rad62-1 × *smc6-X*



rad62-1 × *brc1Δ*



○ double mutant

FIG. 5. *rad62-1* is synthetically lethal with *rad60-1*, *smc6-X* and *brc1Δ*. *rad62-1* was crossed with *rad60-1*, *smc6-X* or *brc1Δ*, and spores were subjected to tetrad analysis at 26°C. The double mutants are circled. Some single mutants formed small colonies, but they showed normal growth when restreaked on plates.

***rad62* is essential for growth.** To study the phenotype of the *rad62* null mutant, 70% of the *rad62* coding region of one of the chromosomes in a diploid strain was replaced with the DNA fragment containing the *ura4* gene by homologous recombination. The heterozygous diploid cells (MMPD1) were sporulated and subjected to tetrad analysis. Thirty asci were dissected and allowed to germinate on YES medium at 30°C, and only two viable segregants arose from each of the 30 asci (Fig. 4A). All 60 segregants required uracil for growth, indicating that these segregants did not carry the *rad62::ura4*⁺ allele and that the *rad62*⁺ gene is essential for growth. Microcolonies consisting of a few 10s of elongated cells were formed from the *rad62::ura4*⁺ segregants. The terminal morphology of the mutant cells grown for 2 days on a minimal plate lacking uracil was examined under an epifluorescent microscope after staining with DAPI (Fig. 4B). They were elongated, and the nuclei were mostly abnormal, sometimes extended, shrunk, fragmented, bisected by a septum (cut phenotype), or biased in position. The terminal morphology was very similar to that of *rad60* and *smc5* (*spr18*) deletion cells (13, 35).

The localization of Rad62 protein in the cell was examined by expressing Rad62 protein tagged with enhanced green fluorescent protein (EGFP) at its N terminus in the *rad62*Δ strain

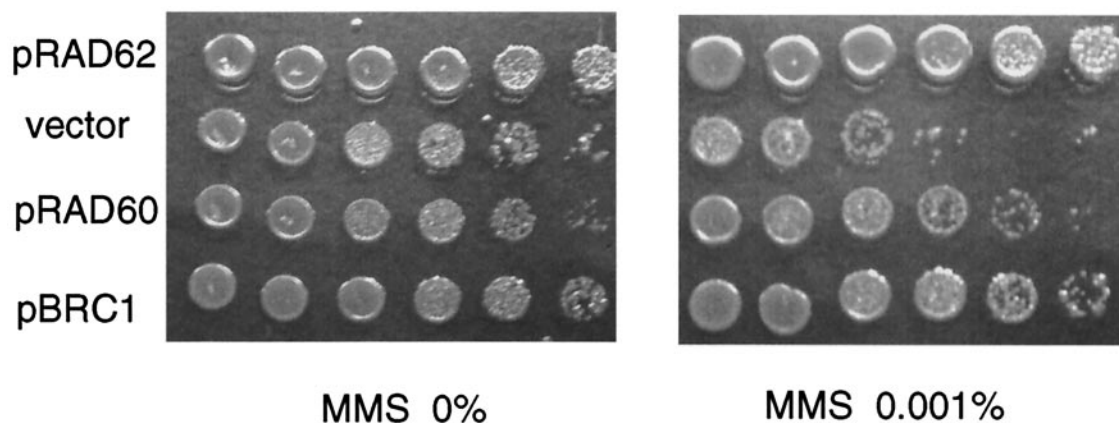


FIG. 6. Suppression of MMS sensitivity of *rad62-1* by overexpression of Rad60 or Brc1. *rad62-1* cells (MMP7) carrying the pRAD62 plasmid, a vector plasmid, pREP41, the pRAD60 plasmid, or the pBRC1 plasmid were grown to saturation on EMM with required supplements. Serial 10-fold dilutions were spotted on YES plates with or without MMS. The plates were incubated at 30°C for 3 days.

(MMP8). The cells were fixed, DNA stained with DAPI, and observed under an epifluorescent microscope (Fig. 4C). Rad62 colocalized with DNA, suggesting that it localizes to the nucleus.

***rad62* genetically interacts with *rad60*, *smc6*, and *brc1*.** Since the phenotypes of *rad62* mutants were very similar to those of *rad60* and *smc6* mutants, we suspected that these genes may be involved in a common pathway for maintaining chromosome integrity and repairing DNA damage by recombination. The *brc1* gene was isolated as an allele-specific multicopy suppressor of the *smc6-74* (*rad18-74*) mutation, and the deletion mutant, which is viable, was synthetically lethal with *smc6* mutants, indicating close functional interaction with *smc6* (47). The genetic interactions among the above genes were examined by constructing double mutants between *rad62-1*, *rad60-1*, *smc6-X* (*rad18-X*), and *brc1*. The mutants were crossed between each other, and the spores were subjected to tetrad analysis at 26°C. The double mutants combining any of the above mutations were inviable (Fig. 5).

The *rad60* gene overexpressed from a multicopy plasmid partially suppressed the MMS hypersensitivity of the *smc6-X* (*rad18-X*) mutant (35). We examined whether overexpression of the *rad60* or *brc1* gene suppressed the MMS hypersensitivity of *rad62-1*. The pBRC1 plasmid carrying *brc1*⁺ under the thiamine-repressible promoter suppressed the MMS and UV sensitivities of the *rad62-1* mutant either under repressed or derepressed conditions (Fig. 6). The pRAD60 plasmid carrying

rad60⁺ under the weaker *nmt1* promoter suppressed the MMS and UV sensitivities of the *rad62-1* mutant under derepressed conditions (Fig. 6). Furthermore, overexpression of Brc1 partly suppressed the MMS and UV sensitivity and raised the permissive temperature of the *rad60-1* mutant (data not shown). The results of the genetic interactions revealed here and by previous works (35, 47) are summarized in Tables 3 and 4. These results strongly suggest that Rad62 is functionally related to Rad60, Smc5, Smc6, and Brc1 and that these proteins collaborate in repairing DNA damage and maintaining chromosome integrity during replication.

Rad62 physically interacts with the Smc5-6 complex. To identify proteins associated with Rad62, we purified a complex containing FLAG-His-tagged Rad62. MMP20 cells carrying the FLAG-His-tagged *rad62* gene and the control cells were grown, and the extracts were prepared. The protein complex containing the tagged Rad62 was purified and analyzed by SDS-PAGE. In addition to a band corresponding to the tagged Rad62 protein (38 kDa), a 120-kDa protein band was specifically identified in the preparation from the MMP20 cells (data not shown). Since multiple genetic interactions of Rad62 with the Smc5-6 complex were observed in this work and the size of Smc5 and Smc6 proteins are about 120 kDa, we suspected that

TABLE 3. Synthetic lethality of double mutations

Gene	Lethality upon double mutation with:				
	<i>rad62-1</i>	<i>rad60-1</i>	<i>brc1Δ</i>	<i>smc6-74</i>	<i>smc6-X</i>
<i>rad62-1</i>		Lethal	Lethal	ND ^c	Lethal
<i>rad60-1</i>			Lethal	ND	Lethal ^b
<i>brc1Δ</i>				Lethal ^a	Lethal ^a
<i>smc6-74</i>					
<i>smc6-X</i>					

^a Verkade et al. (47).

^b Morishita et al. (35).

^c ND, not determined.

TABLE 4. Suppression of UV or MMS sensitivity by overexpression of Smc5-6 related proteins

Protein	Result with overexpression of:				
	<i>rad62-1</i>	<i>rad60-1</i>	<i>brc1Δ</i>	<i>smc6-74</i>	<i>smc6-X</i>
Rad62		No effect	ND ^{f,g}	ND	No effect ^e
Rad60	Suppressed ^c		ND ^f	ND	Suppressed ^{b,e}
Brc1	Suppressed ^c	Suppressed ^c		Suppressed ^{a,d}	Suppressed ^{a,d}
Smc6	No effect	No effect	ND ^f		

^a Verkade et al. (47).

^b Morishita et al. (35).

^c UV and MMS sensitivity.

^d UV sensitivity.

^e MMS sensitivity.

^f *brc1Δ* is not sensitive to UV.

^g ND, not determined.

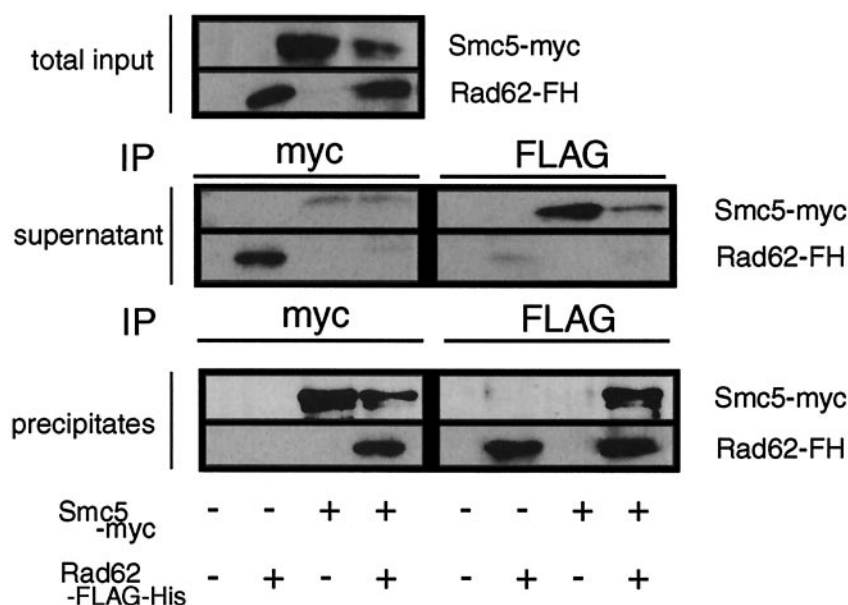


FIG. 7. Physical association of Rad62 with Smc5. The association of Rad62 and Smc5 in vivo was examined by coimmunoprecipitation of Rad62-FLAG-His (Rad62-FH) and Smc5-Myc. Cells expressing both tagged proteins (MMP20) or singly tagged protein (MMP21 or MMP22) were prepared and immunoprecipitated with anti-Myc or anti-FLAG. The immune complexes were separated by SDS-PAGE and immunoblotted with antibodies against Myc or FLAG. Sample volumes were adjusted to make a direct comparison of the relative amounts of Rad62-FH and Smc5-Myc in the extracts, supernatants, and precipitates possible.

the 120-kDa protein found in the Rad62 complex was either Smc5, Smc6, or the doublet of them.

To examine whether Rad62 was included in the Smc5-6 complex, we performed coimmunoprecipitation experiments with FLAG-His-tagged Rad62 and Myc-tagged Smc5 expressed from their cognate promoters. From the cells expressing FLAG-His-tagged Rad62 and Myc-tagged Smc5, the tagged Rad62 was coimmunoprecipitated with the tagged Smc5 by the anti-Myc antibody and the tagged Smc5 was coimmunoprecipitated with the tagged Rad62 by the anti-FLAG M2 antibody (Fig. 7). As a negative control experiment, Rad62 was not immunoprecipitated by the anti-Myc antibody from the cells expressing FLAG-His-tagged Rad62 and untagged Smc5. The tagged Smc5 was not immunoprecipitated by the anti-FLAG M2 antibody from the cells expressing Myc-tagged Smc5 and untagged Rad62. From the comparison of the amounts of Rad62 and Smc5 in the extracts, supernatants, and immunoprecipitates, we conclude that large majorities of Rad62 and Smc5 were physically associated with each other (Fig. 7). These results suggest that Rad62 forms a protein complex with Smc5 in vivo.

Relationship of *rad62-1* with genes involved in recovery from replication block. It occurred to us that one of the functions of *rad62*⁺ may be to provide a pathway for a resolution of the replication block by recombination. To test this hypothesis, we examined the synthetic lethality of the *rad62-1* mutation with other mutations in the genes involved in repairing DNA damage and resolving the replication fork block one way or another (Fig. 8). The mutant became lethal when combined with mutations in *rqh1*⁺, encoding a RecQ DNA helicase homolog involved in recombination repair (36); *srs2*⁺, whose budding yeast homolog encodes a DNA helicase proposed to remove

Rad51 recombination protein from toxic recombination intermediates (28, 46); or *mus81*, encoding a structure-specific endonuclease proposed to resolve recombination intermediates and rescue stalled replication forks (5, 6, 22). It showed a severe growth defect when combined with a mutation in *rhp18*⁺, which encodes a subunit of the ubiquitin ligase that regulates postreplication repair (48). It is also synthetically lethal or nearly so when combined with mutations in the genes involved in nucleotide excision repair, *rad16*⁺ or *rad13*⁺, encoding an excision endonuclease (29) (Fig. 8). Since these genes play roles in repairing various types of DNA damage, the only common feature caused by the mutations will be a block to replication, and together, these data suggest that the function defective in *rad62-1* becomes essential for viability when replication blocks are not resolved during replication.

DISCUSSION

In this study, we identified another novel essential gene, *rad62*, whose function is very similar to the one previously reported for *rad60* (35). Both of them are essential for viability, required to repair DSBs, and epistatic with recombination genes *rhp51* and *rhp57* with respect to their DNA repair function. The terminal morphologies of the deletion mutant cells were very similar; cells are elongated with abnormal nuclear structures, diffuse, shrunk, fragmented, or cut, where the nuclear material is bisected by a septum. These phenotypes are also shared by *smc6* (*rad18*) mutants (13, 30, 47) and the recently identified *nse1* and *nse2* mutants (19, 33). *S. cerevisiae* *nse1* mutants showed similar phenotypes (14). Smc6, like other SMC proteins, forms a heterodimer with its partner Smc5 (13), and they form a tight complex with Nse1 and Nse2 (33). Only

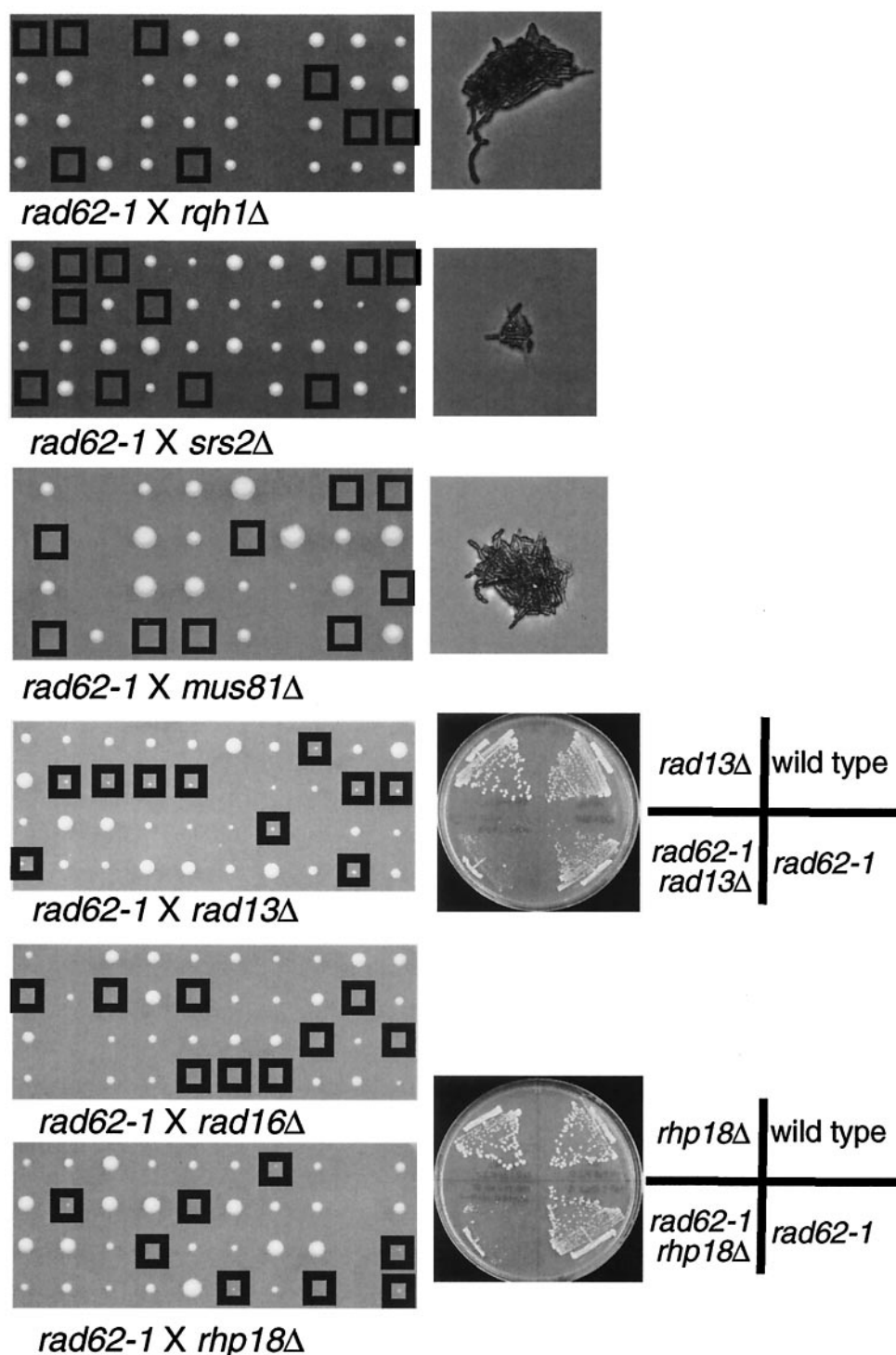


FIG. 8. Synthetic growth defect of *rad62-1* with *rqh1Δ*, *srs2Δ*, *mus81Δ*, *rad13Δ*, *rad16Δ*, and *rhp18Δ*. *rad62-1* was crossed with *rqh1Δ*, *srs2Δ*, *mus81Δ*, *rad13Δ*, *rad16Δ*, or *rhp18Δ*, and spores were subjected to tetrad analysis at 30°C. The double mutants are boxed. Small colonies were formed from the double mutants, *rad62-1 rad13Δ* and *rad62-1 rhp18Δ*, and they were tested for growth by streaking on plates. Some single mutants formed small colonies like the double mutants on the figure, but they showed normal growth when restreaked on plates.

a very small proportion of Rad60 (0.5 to 2.0%) was detected to be associated with Smc5-6 (7). We have demonstrated in this work that *rad62* shows very intimate genetic interactions with *rad60*, *smc6*, and *brc1*. These results strongly suggest that

Rad62 collaborates with the Smc5-6 complex for maintenance of chromosome integrity and repairing DSBs. Indeed, we demonstrated that the majority of Rad62 is associated with Smc5 in vivo, and thus, it is a subunit of the Smc5-6 complex (Fig. 7).

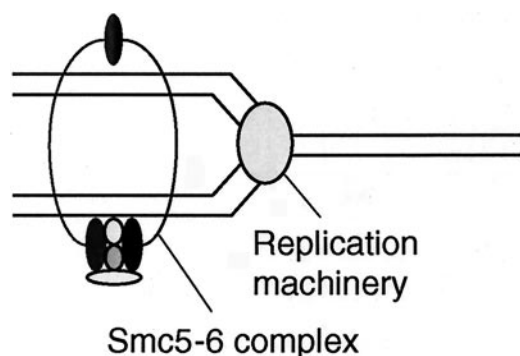


FIG. 9. Model of the Smc5-6 complex for maintenance of chromosome integrity and repair of DSBs at the replication fork. The Smc5-6 complex holds replicated chromatids in close proximity at the replication fork. Smc5 and Smc6 proteins form a heterodimeric tight complex similar to other SMC protein complexes like condensin and cohesin. Nse1, Nse2, and Rad62 associate tightly with the complex. Rad60 and Brc1 may transiently associate with the complex.

We could not detect the association of Rad60 with Smc6 by similar experiments (unpublished data). Therefore, Rad60 association with the Smc5-6 complex is different from the association of the other subunits, as previously reported (7). Although, Rad62 was shown to associate with Smc5, it remains to be shown whether the interaction is direct or through other subunits of the complex. According to a recent paper that identified proteins copurifying with Smc5 and Smc6 in *S. cerevisiae*, the Smc5-6 proteins form two distinct complexes, one with Qri2 (Rad62 homolog), Nse1, Mms21 (Nse2 homolog), and an uncharacterized protein, YDR288W, and another with two uncharacterized proteins, YML023C and Kre29 (20). The homologs of the components of the first complex have been characterized for *S. pombe*, and physical interactions among Smc5, Smc6, Nse1, and Nse2 have been demonstrated (13, 19, 33). The phenotypes of the mutants were very similar. We recently identified a homolog of YDR288W in *S. pombe* named *nse3* by a two-hybrid experiment with *rad62* as a bait, and the mutants show similar phenotypes with those of the mutants of the *smc5-6*-related genes (H. Morikawa, I. Miyabe, T. Morishita, and H. Shinagawa, unpublished data). Although a fraction of *S. pombe* Rad60 was found in association with the Smc5-6 complex (7), its homolog in *S. cerevisiae*, Esc2, was not found in either of the Smc5-6 complexes. The *ESC2* gene is not essential for viability and is involved in the silencing of genes in silent mating type loci and telomere regions (10). Since none of the non-SMC components of the latter complex have been studied, whether or not the two subcomplexes are involved in the same function is an open question.

We made an extensive analysis of the genetic interactions of *rad62* with other genes involved in resolving the replication fork block in various ways (Fig. 8 and Table 1). The *rad62-1* mutant was synthetically lethal with mutations in the *rqh1*, *srs2*, *mus81*, and *rhp18* genes, all of which are required for repairing DNA damage or bypassing blocks to replication (26). To our surprise, it was also synthetically lethal with mutations in *rad16* and nearly so with mutations in *rad13*, which encodes nucleases required for making incisions at the sites flanking large DNA adducts, such as pyrimidine dimers in nucleotide excision re-

TABLE 5. Effects of repair mutations on UV sensitivity of *rad62-1* mutant

Mutation ^a	UV sensitivity
<i>rhp51</i> Δ	Epistatic
<i>rhp57</i> Δ	Epistatic
<i>rad50</i> Δ	Additive
<i>swi5</i> Δ	Additive
<i>uve1</i> Δ	Additive

^a Strains used are shown in Table 1.

pair (29). These results are consistent with Rad62 being required for viability when replication is blocked by spontaneous damage in the absence of the functions that repair the damage or override the block by translesion DNA synthesis. The *rad62* gene seems to possess at least two discrete functions, one essential for growth and the other for repair. The latter function is epistatic with *rhp51*, which plays a dominant role in homologous recombination and recombination repair. The *rhp51* mutation could not suppress the lethality of a *rad62* deletion mutant. So lethality is not due to the accumulation of recombination intermediates. Both the *rad62-1* and *rad60-1* mutants were not defective in homology-dependent recombination, as examined by integration of homologous linear DNA into the chromosome and by formation of spontaneous intra-chromosomal conversion type recombinants. Preliminary results suggest that *rad60-1* is not defective in meiotic recombination (data not shown). However, *rad62* and *rad60* belong to the same epistasis group with *rhp51* and *rhp57* with respect to repair function. The apparent discrepancies could be resolved if we assume that they function for recombination repair only at the replication fork but are not required for recombination in nonreplicating regions of chromosome. However, we cannot rule out a role for *rad60* and *rad62* in genetic recombination, since they are essential genes and these hypomorphic alleles may not give a strong change in recombination rates.

The Smc5-Smc6-Nse1-Nse2-Rad62 complex together with Rad60 may be a specific SMC complex functioning only at the replication fork (Fig. 9). Human Smc6 localizes to the nuclear region during the interphase and is excluded from condensed mitotic chromosomes (41). The Smc5-6 complex may provide a scaffold for the recombination reaction by holding the newly replicated sister chromatids in close proximity so that DSBs produced by replication block can be efficiently repaired by homologous recombination. The essential role of this SMC complex may be to maintain proper structure of the chromosome at the fork during replication. Spontaneous fragmentation of chromosomes and the diffuse nuclear structure observed in *rad60-1* cells grown at the nonpermissive temperature may be caused by the defect of this function. The fact that the *rhp51*Δ mutation partially suppresses UV sensitivity at lower doses than the *rad62-1* mutant (Fig. 3B) may suggest that Rhp51 bound to the UV-damaged fork may be toxic rather than beneficial when the newly replicated chromatids are not properly held. Since Rad60 seems transiently associated with Smc5-6 in contrast to other non-SMC components, which are more tightly associated, and is phosphorylated in response to a replication block in a manner dependent on Cds1 kinase (7), Rad60 may regulate the function of the Smc5-6 complex in response to the replication block. Further experiments are

required to prove that the Smc5-6 complex functions at the replication fork.

We examined the epistatic relationship with other repair and recombination genes, and the results are summarized in Table 5. Interestingly the UV sensitivity of the *rad62-1* mutant was additive with *rad50Δ* and *swi5Δ*. *rad50* has been proposed to play a role in promoting recombination between sister chromatids in collaboration with the cohesin complex (18). *Swi5* has dual roles, one in mating type switching and the other in recombination repair in a novel pathway dependent on *rhp51* but independent of *rhp55-57* (1). Since the *swi5Δ rhp57Δ* double mutants are additive in UV sensitivity and equally as sensitive as *rhp51Δ*, the recombination repair pathway involving the Smc5-6 complex function is likely to be the same as the one involving the *rhp55-57* functions. The additive UV sensitivity in the *rad62-1 uve1Δ* double mutant indicates that *rad62* may function in a pathway different from the UV nucleotide excision repair pathway (32).

It should be noted that cohesion- and condensin-related genes like *rad21/scc1* (4, 21) and *cmd2* (2) are also involved in repairing DNA damage. The *cmd2-1* mutant is sensitive to hydroxyurea, and *cmd2* is required for activation of the checkpoint kinase Cds1 (2). Rad60 is phosphorylated in a manner dependent on the *cds1* function upon replication block (7), and both *rad60-1* and *rad62-1* are sensitive to hydroxyurea and defective in maintaining checkpoint arrest (our unpublished data) in a similar manner to that of *smc6-74* and *nse1* mutants (19, 47). Therefore, the integrity of the Smc5-6 complex seems to be essential for the maintenance of the checkpoint. The complex may be monitoring the completion of DNA repair, as suggested recently (19), and premature resumption of mitosis in these mutants may cause disintegration of the chromosomes, resulting in stretched or bisected nuclear structures. The essential role of the condensin complex in mitosis is well documented, but the involvement of the condensin complex in DNA repair in the interphase has been only recently demonstrated by the analysis of the phenotypes of the *cdn2-1* mutant (2). Conversely, defects in S-phase function in the Smc5-6 complex affect mitosis. *rad60*, *rad62*, and *smc6* mutations cause abnormal nuclear segregation. Recent studies on various SMC complexes have revealed interacting roles of the SMC complexes in the functions related to the DNA damage response, and the importance of the Smc5-6 complex in these processes is just beginning to be unveiled.

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