

Drosophila melanogaster MNK/Chk2 and p53 Regulate Multiple DNA Repair and Apoptotic Pathways following DNA Damage

Michael H. Brodsky,^{1,2*} Brian T. Weinert,² Garson Tsang,^{2,3} Yikang S. Rong,⁴ Nadine M. McGinnis,¹ Kent G. Golic,⁵ Donald C. Rio,² and Gerald M. Rubin^{2,3}

Program in Gene Function and Expression and Program in Molecular Medicine, University of Massachusetts Medical School, Worcester, Massachusetts 01605¹; Department of Molecular and Cell Biology, University of California, Berkeley, California 94720-3200²; Howard Hughes Medical Institute³ and Department of Biology,⁵ University of Utah, Salt Lake City, Utah 84112; and Laboratory of Molecular Cell Biology, National Cancer Institute, Bethesda, Maryland 20892⁴

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We have used genetic and microarray analysis to determine how ionizing radiation (IR) induces p53-dependent transcription and apoptosis in *Drosophila melanogaster*. IR induces MNK/Chk2-dependent phosphorylation of p53 without changing p53 protein levels, indicating that p53 activity can be regulated without an Mdm2-like activity. In a genome-wide analysis of IR-induced transcription in wild-type and mutant embryos, all IR-induced increases in transcript levels required both p53 and the *Drosophila* Chk2 homolog MNK. Proapoptotic targets of p53 include *hid*, *reaper*, *sickle*, and the tumor necrosis factor family member *Eiger*. Overexpression of *Eiger* is sufficient to induce apoptosis, but mutations in *Eiger* do not block IR-induced apoptosis. Animals heterozygous for deletions that span the *reaper*, *sickle*, and *hid* genes exhibited reduced IR-dependent apoptosis, indicating that this gene complex is haploinsufficient for induction of apoptosis. Among the genes in this region, *hid* plays a central, dosage-sensitive role in IR-induced apoptosis. p53 and MNK/Chk2 also regulate DNA repair genes, including two components of the nonhomologous end-joining repair pathway, *Ku70* and *Ku80*. Our results indicate that MNK/Chk2-dependent modification of *Drosophila* p53 activates a global transcriptional response to DNA damage that induces error-prone DNA repair as well as intrinsic and extrinsic apoptosis pathways.

In organisms ranging from bacteria to humans, DNA damage induces transcription of genes that promote survival and limit mutations following genotoxic stress (22). In *Escherichia coli*, the SOS response to DNA damage induces the coordinated expression of multiple DNA repair systems, including an error-prone polymerase that leads to increased mutation rates (63). In yeast, the DNA damage checkpoint pathway activates both a general stress response as well as a DNA damage-specific response that upregulates genes involved in homologous recombination and subunits of ribonucleotide reductase (23). In mammalian cells, the p53 transcription factor regulates DNA damage-induced transcription (4, 19, 20, 68). p53 target genes include p21, which regulates cell cycle progression, RNR2, GADD45, which may enhance DNA repair, and Mdm2, which acts in a negative feedback pathway to down-regulate p53 activity. At least 10 p53 target genes encode potential proapoptotic proteins, including regulators of the intrinsic apoptotic pathway, such as APAF1 and several BH3-containing proteins, as well as tumor necrosis factor receptor family members that regulate the extrinsic apoptotic pathway. Knock out of the mouse BAX or Bid genes decreases damage-induced apoptosis in some cell types (41, 55). Why so many proapoptotic genes are regulated by p53 and the relative contributions of these genes to damage-induced apoptosis are unclear.

Although p53 is not conserved in yeast, five mammalian proteins related to yeast DNA damage-responsive kinases may directly or indirectly regulate p53 activity. The ATM, ATR, and DNA-PKcs kinases are members of the phosphatidylinositol 3-kinase superfamily and have sequence similarity to budding and fission yeast checkpoint kinases (2, 32, 53). ATM mutant mice and cells are defective in p53 activation following ionizing radiation (IR) (5, 36, 76). Analysis of dominant-negative forms of ATR has led to conflicting reports of ATR-dependent activation of p53 (45, 65). Two downstream checkpoint kinases, Chk1 and Chk2, can phosphorylate p53 in vitro. Analysis of Chk2 mutant mice and dominant-negative constructs supports a central role for this kinase in p53 activation following DNA damage (13, 29, 30, 50).

These DNA damage-responsive kinases regulate p53 activity through multiple mechanisms (67, 69). In unstressed cells, the MDM2 protein mediates p53 ubiquitination and degradation. Following damage, phosphorylation of p53 and MDM2 blocks MDM2-mediated turnover of p53, leading to a rapid accumulation of nuclear p53. Overexpression of p53 can induce target gene expression, indicating that p53 accumulation is sufficient for at least some p53 functions. Direct phosphorylation and acetylation may also contribute to p53 activation. In vivo, p53 activation is generally accompanied by both modification and accumulation. The relative contributions of these activation mechanisms are likely to reflect factors including cell type and environmental signals.

Model organisms provide an opportunity to examine the most highly conserved aspects of a signaling pathway in the context of normal development. *Drosophila melanogaster* and

* Corresponding author. Mailing address: Program in Gene Function and Expression and Program in Molecular Medicine, 623, LRB, University of Massachusetts Medical School, Worcester, MA 01605. Phone: (508) 856-1640. Fax: (508) 856-5460. E-mail: michael.brodsky@umassmed.edu.

Caenorhabditis elegans have a single p53 homolog required for DNA damage-induced apoptosis (11, 18, 48, 56). The proapoptotic gene *reaper* is a transcriptional target of *Drosophila* p53 and is part of a gene complex required for damage-induced apoptosis (11, 51, 61, 71, 72). Some damage-induced apoptosis can be induced in the absence of *reaper*, suggesting that *Drosophila* p53 activates additional proapoptotic genes. Regulation of other DNA damage responses by *Drosophila* p53 has not been described.

The mechanism of damage-induced activation of p53 is also unclear. The *Drosophila* genome contains homologs of the conserved checkpoint kinases, but it does not reveal an obvious MDM2 homolog (57); this observation indicates that either the homolog of MDM2 has too little sequence similarity to be identified by simple sequence searches or that *Drosophila* does not utilize protein turnover to regulate p53 activity.

In this study, we have characterized the regulation and function of *Drosophila* p53 following DNA damage. A null mutation of p53 (52) blocks damage-induced apoptosis but is not required for viability, fertility, or damage-induced cell cycle arrest. After IR, p53 protein exhibits a phosphatase-sensitive change in gel mobility, but p53 levels do not change. MNK, the *Drosophila* homolog of the Chk2 kinase (47, 75), is required for IR-induced modification of p53. These results suggest that posttranslational modification is sufficient to activate *Drosophila* p53. To identify cellular pathways regulated by p53, we have performed a genome-wide analysis of irradiation-induced gene expression in wild-type and mutant *Drosophila* embryos. IR-induced genes include regulators of apoptosis, cell-cell signaling, and DNA repair, but not cell cycle progression. Both *mnk/Chk2* and *p53* are required for all IR-induced increases in gene expression. Two targets of p53, *Ku70* and *Ku80*, are involved in nonhomologous end joining, suggesting that error-prone DNA break repair may be an important feature of p53 signaling. Another target of p53, the *Drosophila* tumor necrosis factor (TNF) homolog *Eiger* (31, 43), can induce apoptosis when overexpressed but is not required for IR-induced apoptosis. We also demonstrate that three known regulators of apoptosis, *rpr*, *skl* (14, 62, 73), and *hid* (26), are targets of p53. We find that animals heterozygous for deficiencies spanning all three genes exhibit impaired IR induction of apoptosis and that *hid* in particular is haploinsufficient for this DNA damage response. Combined with previous observations that *hid* function is regulated by Ras activity (6, 7, 37) and micro-RNA expression (10), our results suggest that *hid* plays a central role in integrating signals from diverse signaling pathways to determine the apoptotic response to p53 activation.

MATERIALS AND METHODS

Genetics and transgenes. All experiments were performed at 25°C unless otherwise indicated. The following alleles were used for analysis of damage-induced apoptosis and cell cycle arrest: *mei-41^{29D}* (38), *mei-41^{RT2}* (27), *mus304^{D1}*, *mus304^{D2}* (12), and *grps^{ts1}* (60). Stocks were obtained from Hermann Steller, Kristin White, Scott Hawley, and the Bloomington *Drosophila* Stock Center.

The *mnk⁶* allele was generated by transposase-mediated mobilization of a P[lacW] P-element insertion in the *barren* gene (8) followed by PCR to identify lines with insertions in the *mnk* coding region and not in *barren*. This line complemented the lethality associated with the original *barren* insertion. The insertion was in nucleotide position 465 of the long form of the *mnk* coding region, which corresponds to the second intron of the short form of *mnk* (47). A deletion associated with this insertion removed 218 nucleotides of *mnk* genomic

sequence and 823 nucleotides of the 3' end of the P[lacW] DNA. The sequence junction of this deletion was as follows: *mnk* genomic, GTGCTGGAGT/TCT TGAAGTG, P[lacW] DNA. A rescue construct for *mnk* was generated by PCR amplification. The oligonucleotide sequences used were as follows: 523 bases 5' to the start of transcription, GGCCTCTAGAAACGACGCCGCAATTTAG GGC; 72 bases 3' to the end of transcription, GGCCGCGCCCGTGTAGCAA TTTGCCCGCCTCCG. The underlined sequences correspond to *XbaI* and *NorI* sites that were used to clone the genomic fragment into the pCaSpeR-2 transposon vector.

The *p53¹* mutation was generated by homologous recombination (52). The p53 cDNA transgene (GUS-p53) has been described previously (11). This construct moderately overexpresses p53 in the developing eye at a level insufficient to generate a rough eye phenotype. Much higher levels of expression are generated by coexpression of GMR-Gal4, resulting in the rough eye phenotype seen below in Fig. 4. Rescue of p53-dependent apoptosis in the developing eye was accomplished using GUS-p53 in the absence of GMR-Gal4.

The *Eiger¹* and *Eiger²* mutations were generated by transposase-mediated mobilization of the KG02299 P-element insertion immediately upstream of the *Eiger* transcript. CyO-Δ2-3 was used as a source of transposase. PCR was used to screen chromosomes that had lost the *white* marker on the KG02299 element.

Overexpression constructs for *mnk* and *grps* were generated by PCR amplification of the coding region of each gene flanked by gateway recombination sites (oligonucleotide sequences available on request). Amplification oligonucleotides introduced a translation start sequence immediately 5' of the start codon and included the stop codon for each gene. Gateway recombination reactions transferred these sequences into pGUSgw, a gateway-modified pGUS plasmid (M. H. Brodsky, unpublished data). A predicted kinase-dead form of MNK was generated by using PCR to introduce an Asp (GAC) to Ala (GCC) mutation at amino acid position 303 in the *mnk* coding region. The *mnk* coding sequence with this mutation was introduced into pGUSgw.

Detection of p53 and MNK/Chk2. p53 antibodies were generated using C-terminal and N-terminal fragments of p53 (11). Two mouse monoclonal lines raised against the C-terminal fragment of p53, C7A10 and B9A10, were capable of recognizing p53 overexpressed in *Drosophila* S2 tissue culture cells by immunofluorescence, immunoblotting, and immunoprecipitation. Guinea pig polyclonal antibodies were raised against glutathione S-transferase fusions to N-terminal and C-terminal fragments of p53. These antibodies were affinity purified using maltose binding protein fusions to the same fragment. These antibodies also detect overexpressed p53 in *Drosophila* tissue culture cells.

To detect damage-induced modification of p53, protein lysates were isolated from staged (3- to 15-h) embryos treated with irradiation or mock treated and incubated for 30 min at room temperature. Embryos were dechorionated and frozen in liquid nitrogen. Frozen embryos were lysed in radioimmunoprecipitation assay (RIPA) buffer (50 mM Tris [pH 8.0], 300 mM NaCl, 0.5% sodium deoxycholate, 1% NP-40, 0.1% sodium dodecyl sulfate [SDS]) containing protease inhibitors (1 mM phenylmethylsulfonyl fluoride, and 0.5 μg each of aprotinin, pepstatin, leupeptin, chymotrypsin, and antipain/μl) and phosphatase inhibitors (10 mM sodium fluoride, 0.4 mM sodium *meta*- and *ortho*-vanadate [each], 0.1 mM β-glycerophosphate, 10 mM sodium pyrophosphate, 1 μg of phosphitin/ml, 0.5 μM microcystin-LR, and 0.5 μM okadaic acid) using a Wheaton glass homogenizer (A pestle). Homogenized lysates were centrifuged at 35,000 rpm in a Beckman Ti70 rotor at 4°C for 30 min. Total protein was determined by Bradford assay, and 50 mg was used in each immune precipitation (IP). IPs were performed for 20 h at 4°C, with 20 μg of an equal mixture of two p53 monoclonal antibodies covalently coupled to protein A/G-agarose (Oncogene Research Products). Precipitates were washed five times in RIPA buffer, resuspended in 50 μl of 2× Laemmli sample buffer, and heated to 100°C for 5 min. For phosphatase treatment, samples were resuspended to 50 μl in RIPA buffer, 2 mM MnCl₂, and 400 U of Lambda phosphatase (New England Biolabs) and incubated for 30 min at room temperature. Samples were analyzed by SDS-polyacrylamide gel electrophoresis (SDS-PAGE) with 9% gels, transferred to nitrocellulose, and detected with a 1:5,000 dilution of affinity-purified, p53 N-terminal-specific polyclonal guinea pig antiserum.

To detect MNK, clarified lysates from the p53 IPs were mixed with Laemmli sample buffer and heated to 100°C for 5 min. Protein samples (approximately 10 to 20 μg of total protein/sample) were separated by SDS-9% PAGE and probed with a 1:1,000 dilution of affinity-purified rabbit polyclonal anti-MNK antiserum (47).

Irradiation-induced phenotypes. Irradiation-induced apoptosis and cell cycle arrest were performed as previously described (11, 12).

Gene expression analysis was performed using embryos collected 3 to 5 h after egg laying. Each embryo collection plate was cut in half; one half was X-irradiated for 15 min with 4,000 rads, and the other half was used as the untreated

TABLE 1. DNA damage response homologs in mammals, flies, and yeast

Mammalian protein ^a	Fly protein	Yeast proteins (budding, fission)
ATR (upstream kinase)	MEI-41	Mec1, Rad3
ATRIP (ATR binding protein)	MUS304	Ddc2, Rad26
ATM (upstream kinase)	ATM	Tel1, Tel1
DNA-PKcs (upstream kinase)		
Chk1 (downstream kinase)	GRPS	Chk1, Chk1
Chk2 (downstream kinase)	MNK	Rad53, Cds1
p53 (transcription factor)	p53	
p63 (transcription factor)		
p73 (transcription factor)		

^a A general description of the activity of each human protein is indicated in parentheses.

control. Embryos were incubated at room temperature and frozen in liquid nitrogen. Embryos treated at the 15-min time point were irradiated for 12 min with 3,200 rads. Time points indicate the time following the initiation of irradiation. Each experimental condition was performed in triplicate. Fifty to 100 μ l of packed embryos was isolated for each sample. Total RNA was isolated by using RNAwiz (Ambion). Five micrograms of total RNA was used to generate biotin-labeled cRNA following a standard protocol provided by Affymetrix. Washes and scanning were performed according to the Affymetrix protocol. Image analysis was performed using Microarray Suite version 4. Expression values were determined using PM/MM models in dCHIP (39). Analysis of variance by least squares using the SAS proc glm program (SAS Institute, Cary, N.C.) was performed with the 30-, 60-, and 120-min time points from each genotype. *P* values for treatment effects and treatment by genotype effects were determined, and genes were selected with a false discovery rate of less than or equal to 5%. Only genes that increased or decreased at least 1.7-fold are included below in Table 2.

Real-time PCR was performed using the same total RNA samples used for microarray analysis. mRNA was isolated from 50 μ g of total RNA using Oligotex purification resin (Qiagen). Predesigned Taqman oligonucleotides and probes for most genes were purchased from Applied Biosystems (ABI). Oligonucleotides and probes for *ku80* and *mre11* were designed using Primer Express software (sequences available on request). mRNA was isolated from 380 ng of total RNA using Oligotex beads (Qiagen). Real-time PCR was performed on an ABI 5700 apparatus using reverse transcription and PCR reagents from ABI. Relative levels in untreated and treated samples were determined using a standard curve for each probe. Glycerinaldehyde-3-phosphate dehydrogenase probes were used to normalize for differences in total mRNA levels in each experiment.

***mei-41* and *ATM* cDNAs.** A putative full-length clone for *mei-41* was isolated from the LD embryonic cDNA library prepared by Ling Hong (Berkeley Drosophila Genome Project [www.fruitfly.org]). This clone is approximately 300 bp longer than the previously described cDNA sequence for *mei-41* and contains an open reading frame with 100 additional amino acids. A putative full-length clone for *ATM* was isolated from the LP adult head cDNA library prepared by Ling Hong. This clone contains a 5-bp deletion when compared to a shorter head cDNA clone (GenBank accession number AY395749) or to the genome sequence (www.fruitfly.org). Addition of the missing 5 bp created a large open reading frame that extended across the entire cDNA and showed sequence similarity to ATM throughout the predicted amino acid sequence.

Nucleotide sequence accession numbers. The full-length cDNA sequences obtained for *mei-41* and *CG6535/ATM* have been deposited in GenBank with the accession numbers AY282465 and AY395748, respectively.

RESULTS

Regulation of DNA damage-induced apoptosis by p53 and *mnk*. To identify genes required for IR-induced apoptosis in *Drosophila*, we examined several homologs of yeast and mammalian damage response genes (Table 1). BLAST searches and ClustalW alignments with these sequences confirmed earlier conclusions (57) that MEI-41 is most similar to mammalian ATR and CG6535 is most similar to ATM (S. Oikemus and M. H. Brodsky, unpublished data). Based on these observa-

tions, we refer to CG6535 as *Drosophila* ATM. No clear homolog of the related kinase DNA-PKcs was present in the *Drosophila* genome. Homologs of Chk1 (*GRPS*), Chk2 (*MNK*), and p53 have been previously described (11, 21, 47, 48). MUS304 is a DNA damage response gene with nearly identical mutant phenotypes to MEI-41 (12). ATR-IP appears to be the mammalian homolog of MUS304 and the yeast checkpoint proteins Rad 26 and Ddc2 (16). In support of this conclusion, we found that MUS304 interacted with MEI-41 in a yeast two-hybrid assay (M. H. Brodsky, G. Tsang, and G. M. Rubin, unpublished data).

Previous genetic analysis demonstrated that *mei-41*, *mus304*, and *gps* are required for DNA damage and DNA replication checkpoints (12, 27, 59, 60). We examined X-ray-induced apoptosis in developing wings (wing disks) from larvae mutant for each of these genes (Fig. 1). Very low levels of apoptosis were observed in the absence of irradiation (Fig. 1A). Following irradiation, extensive induction of apoptosis was induced in wild-type wing disks (Fig. 1B). Damage-induced apoptosis was still observed in *mei-41*, *mus304*, and *gps* disks (Fig. 1C to H). Thus, each of these genes is essential for IR-induced cell cycle arrest, but not for apoptosis.

To test if other damage response homologs are required for IR-induced apoptosis, we examined loss-of-function mutations in the remaining genes shown in Table 1. Initial characterization of *ATM* mutant animals indicated that this gene is required for viability (S. Oikemus, unpublished results), preventing a straightforward analysis of its role in IR-induced apoptosis. Mutations in *mnk* and *p53* (52) were created by transposon-mediated mutagenesis and homologous recombination, respectively. p53 protein was detectable in wild-type and *mnk* mutant embryos, but not in *p53* mutant embryos (Fig. 2B, upper panel). These antibodies recognize a C-terminal portion of p53, indicating that translation of p53 does not reinitiate following the stop codon. Anti-MNK antibodies detected a protein in wild-type and *p53* mutant embryos, but not in *mnk* mutants (Fig. 2B, lower panel). These results indicate that we have isolated null mutations in *mnk* and *p53*.

We examined IR-induced apoptosis in *p53* and *mnk* mutant larvae. In the absence of DNA damage, both mutants exhibited the low level of spontaneous apoptosis seen in wild-type animals (Fig. 1I and K). However, no increase in apoptosis was observed following irradiation of either *mnk* (Fig. 1J) or *p53* (Fig. 1L) mutant larvae. IR-induced apoptosis could be rescued in *mnk* mutant disks by a transgene containing *mnk* (see Materials and Methods) (Fig. 1M and N). Similarly, IR-induced apoptosis could be restored in the third-instar eye imaginal disk of *p53* mutant animals by a transgene expressing the *p53* cDNA under the control of a *Glass*-dependent promoter (11). In these animals, IR-induced apoptosis occurred only in the posterior region of the disk, where the *Glass*-responsive promoter directed expression of the *p53* transgene (Fig. 1O and P).

We also examined the ability of the *mnk* and *p53* mutants to undergo IR-induced cell cycle arrest. Following irradiation, wild-type imaginal disks exhibited a G₂ arrest (Fig. 3A and B) that could be assayed with a phospho-specific antibody against histone H3, specific for mitotic cells (12, 28). IR-induced cell cycle arrest was normal in *p53* mutant larvae (Fig. 3C), while *mnk* mutant larvae showed a very mild defect in IR-induced

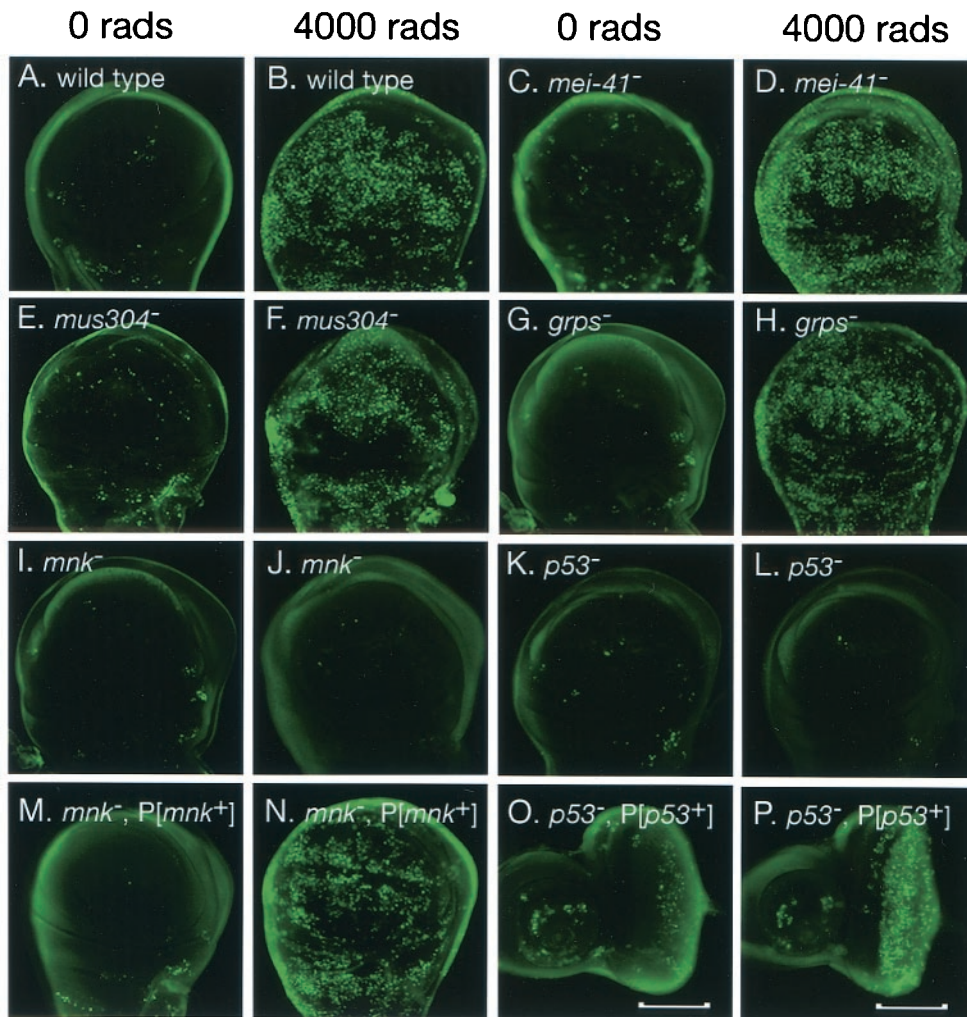


FIG. 1. *Drosophila* *p53* and *mnk/Chk2* are required for IR-induced apoptosis. Imaginal wing or eye disks were dissected from untreated (A, C, E, G, I, K, M, and O) or irradiated (B, D, F, H, J, L, N, and P) third instar larvae and stained with the vital dye acridine orange. IR-induced apoptosis was observed 4 h following irradiation in wild-type (B), *mei-41* (D), *mus304* (F), and *grps* (H) mutant larvae. No induction of apoptosis was observed in *mnk* (J) or *p53* (L) mutant larvae. Damage-induced apoptosis was restored to *mnk* mutant disks by a transgene containing the *mnk* promoter and coding sequence (M and N). Damage-induced apoptosis was restored to the posterior (shown by brackets) of *p53* mutant eye disks by a transgene driving expression of a *p53* cDNA under control of a *Glass*-responsive promoter (described in Materials and Methods).

cell cycle arrest (Fig. 3D). In contrast, *grps* mutant larvae demonstrated a reduction in IR-induced arrest (Fig. 3E) (12). Larvae with mutations in both *mnk* and *grps* were completely defective in IR-induced cell cycle arrest (Fig. 3F). Our results demonstrate that while *mnk* and *p53* function are absolutely required for IR-induced apoptosis, these genes do not play a major role in cell cycle arrest.

p53-deficient mice generally develop normally and die from early-onset cancers. Mice deficient for the *p53* homologs *p63* and *p73* exhibit developmental defects resulting in embryonic or neonatal lethality (42, 77, 78). *Drosophila p53* and *mnk* mutant flies are viable, fertile, and show no striking defects in external morphology (data not shown).

mnk/Chk2 is required for IR-induced modification of *p53*.

To examine whether DNA damage induced changes in *Drosophila p53* modification or abundance, we used IP and Western blotting to analyze *p53* from untreated and X-irradiated

embryos (Fig. 4A). *p53*-dependent changes in gene expression could be seen within 15 to 30 min of IR (see below). Thirty minutes following IR, *p53* exhibited reduced mobility, indicating that *p53* underwent a posttranslational modification (Fig. 4A, lanes 1 and 2). Total levels of *p53* protein were not substantially altered following IR. Similar results were seen at 15 and 60 min (data not shown). Following phosphatase treatment, there was no difference in the mobility of *p53* from untreated or irradiated embryos (Fig. 4A, lanes 3 and 4), indicating that the IR-induced shift was due to phosphorylation. In *mnk* mutant embryos, *p53* mobility was not altered following irradiation (Fig. 4A, lanes 5 and 6). The absence of an IR-induced shift in *p53* mobility indicated that *mnk* is specifically required for IR-induced phosphorylation of *p53*.

Treatment of unirradiated samples with phosphatase increased the gel mobility of *p53* (Fig. 4A, lanes 1 and 3), indicating that *p53* is constitutively phosphorylated in the absence

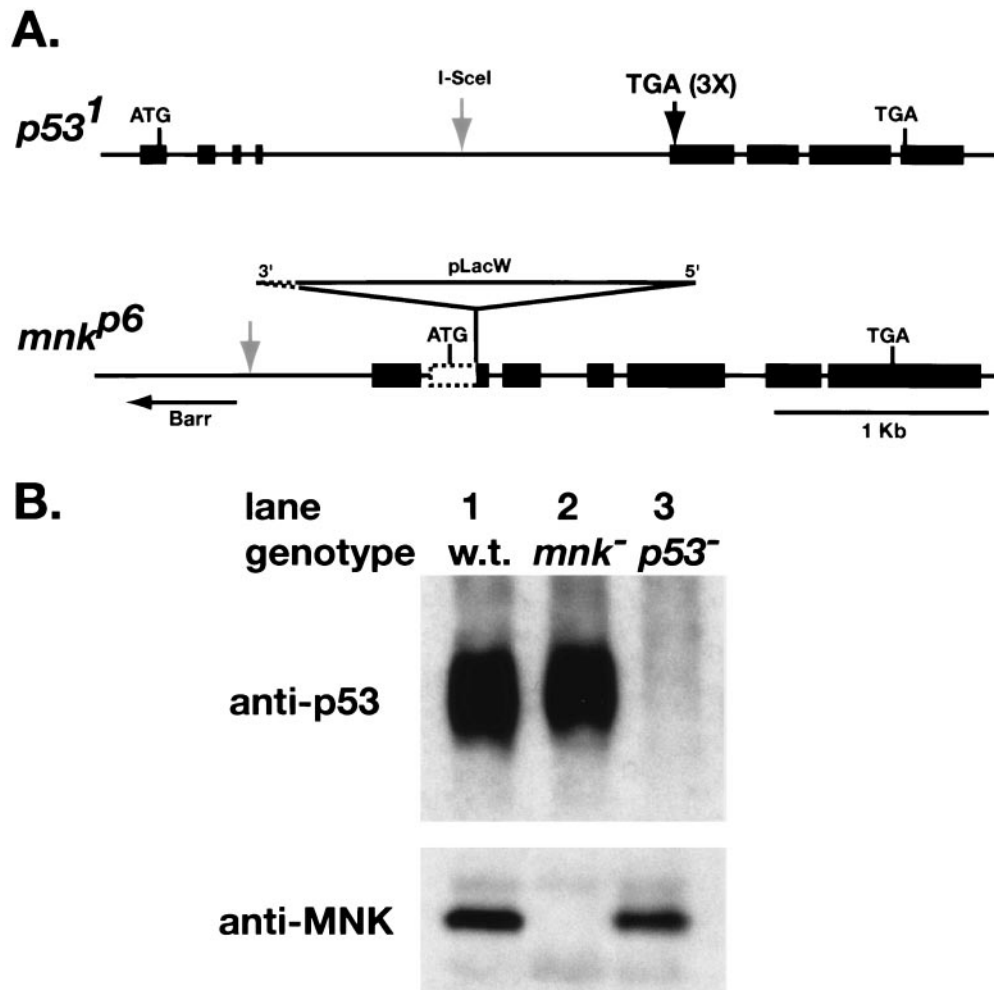


FIG. 2. Loss-of-function mutations in *Drosophila* MNK/Chk2 and p53. (A) Maps of *p53* and *mnk* loci. Exons are depicted as rectangles, and introns are lines. Homologous recombination was used to introduce stop codons in three reading frames at the beginning of the second exon of p53, eliminating the DNA binding domain and the C-terminal tetramerization and regulatory domains. The I-SceI site used for double-strand break-mediated recombination (arrow) was present in the targeting construct but not the final mutant. The *mnk*^{p6} allele contains a P-element insertion at amino acid position 52 and an associated deletion that removes 218 nucleotides of genomic sequence, including the *mnk* start codon. (B) p53 and MNK protein in lysates of 2- to 16-h embryos. p53 was immunoprecipitated with mouse anti-p53 monoclonal antibody, run on an SDS-PAGE gel, and detected with guinea pig anti-p53 polyclonal antibody. Signal was detected in wild-type (lane 1) and *mnk* mutant (lane 2) embryos, but not in *p53* mutant embryos (lane 3). These antibodies recognized a C-terminal portion of p53, indicating that translation of p53 does not reinitiate following the stop codon. MNK was detected with a polyclonal rabbit antibody (47). Signal was detected in wild-type (lane 1) and *p53* mutant (lane 3) embryos, but not in *mnk* mutant embryos (lane 2).

of exogenous DNA damage. This constitutive phosphorylation of p53 did not require *mnk* (lanes 1 and 5).

We also examined the gel mobility of MNK following IR. In unirradiated embryos, MNK migrated as a single band on SDS-PAGE (Fig. 4B, lane 1). Following IR, a second band with slower mobility was observed (Fig. 4B, lane 2). Phosphatase treatment of embryo lysates did not affect the mobility of MNK in untreated embryos (Fig. 4B, lane 3), but it eliminated the appearance of a second band in IR-treated embryos (Fig. 4B, lane 4). These results were consistent with IR-induced phosphorylation of MNK. The IR-induced shift in MNK mobility was present in both wild-type and *p53* mutant embryos (lanes 5 and 6), suggesting that *p53* acts downstream of *mnk*.

Although MNK was required for p53 phosphorylation, initial attempts to detect a stable association between MNK and

p53 by coimmunoprecipitation from untreated embryos or interaction in a yeast two-hybrid assay were unsuccessful (B. T. Weinert and G. Tsang, unpublished results). For an in vivo assay, we examined genetic interactions between *mnk* and *p53* transgenes expressed during fly eye development. Overexpression of *Drosophila p53* during eye development produced a reduced, rough eye phenotype (11, 48) (Fig. 4C and D). We have tested the effect of coexpressing wild-type *gpps*, *mnk*, and a kinase-dead form of *mnk*, *mnk(kd)*. Each of these transgenes had little or no effect on eye development on their own (data not shown). Coexpression of *gpps* did not alter the *p53*-dependent rough eye phenotype (Fig. 4E). Coexpression of *mnk(kd)* suppressed the *p53*-dependent rough eye phenotype (Fig. 4F). When raised at 25°C, animals coexpressing *p53* and wild-type *mnk* died during pupation (data not shown), probably reflect-

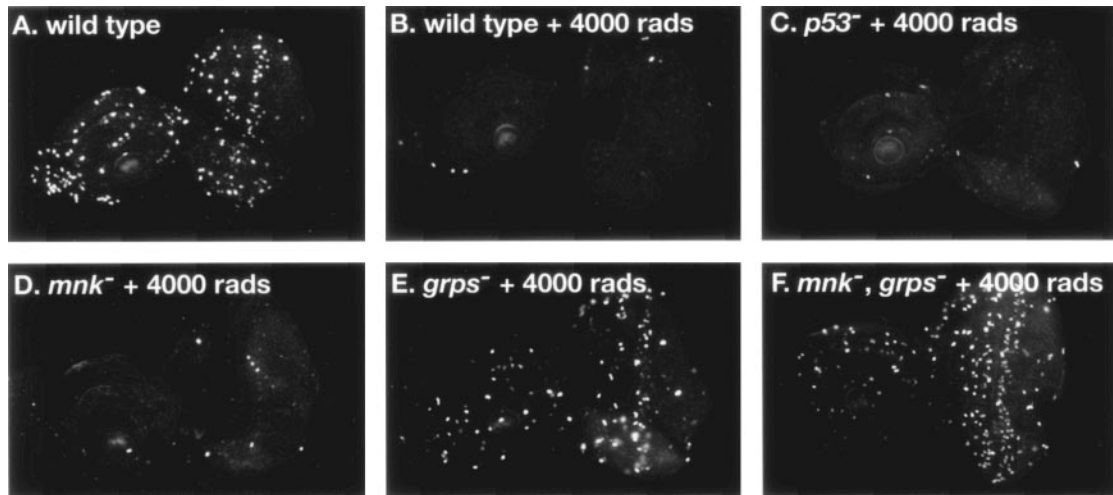


FIG. 3. Cell cycle regulation in *p53* and *mnk/Chk2* mutant animals. Imaginal eye disks were dissected from untreated (A) or irradiated (B to F) larvae, fixed, and stained with a phospho-specific antibody, anti-phosphorylated histone H3, which specifically recognizes mitotic cells in eukaryotes. Cell cycle arrest was assayed by the absence of mitotic cells. IR-induced arrest was observed in wild-type (A and B), *p53* (C), and *mnk* (D) mutant larvae. Partial arrest was observed in *grps* mutant larvae (E). No arrest was observed in *mnk grps* double mutant larvae (F).

ing enhanced activity of *p53* in tissues other than the eye. When raised at 18°C, animals expressing *p53* alone or *p53* and *mnk* could be recovered. Under these conditions, coexpression of *mnk* enhanced the *p53*-dependent rough eye phenotype (Fig. 4G and H). Similar genetic interactions have been reported previously (50). These results are consistent with a direct interaction between MNK and *p53*.

Genome-wide analysis of IR-induced transcription by *p53* and *MNK/Chk2*. Previously, we and others provided evidence that the proapoptotic genes *reaper* and *sickle* are targets of *p53* following IR (11, 61). To determine which *Drosophila* mRNAs are induced by IR, we used Affymetrix oligonucleotide microarrays to probe expression of greater than 13,000 genes in both untreated and irradiated embryos (Table 2). We examined 3- to 5-h embryos, a stage when high levels of apoptosis can be induced by DNA damage but little developmentally regulated apoptosis is observed (3, 46). RNA was prepared from wild-type, *p53* mutant, or *mnk* mutant embryos at five time points following irradiation. To limit gene expression variation due to minor age differences among different embryo collections, each collection was split and one half was irradiated while the other half was the unirradiated control. As a second assay for transcript levels, the relative expression level of a subset of genes was assayed at the 1- or 4-h time points using real-time quantitative PCR (Q-PCR) (Table 2). In each example tested, mRNA changes observed by microarray analysis were confirmed by Q-PCR.

Of over 13,000 genes represented on these arrays, we identified 17 induced and 18 repressed genes with at least a 1.7-fold change in expression levels (Table 2). The upregulation of all 17 induced genes was partially or entirely dependent on both *p53* and *mnk*. The rapid (15- to 30-min) induction of many of these targets suggested that they may be direct targets of *p53*. In contrast, the decreased expression of only 1 of 18 repressed genes required *p53*, while all 18 required *mnk* for downregulation. These results suggest that *mnk* and *p53* act in a linear pathway to activate gene expression following IR, but that a

p53-independent mechanism can lead to decreased gene expression.

The annotations of the induced genes indicated that *Drosophila* *p53* regulates both intrinsic and extrinsic apoptosis pathways following irradiation (Table 2). Three of the five most highly induced genes, *rpr*, *skl*, and *hid*, are part of a group of genetically linked, proapoptotic genes at cytological position 75C (14, 26, 62, 72, 73). Like the mammalian proapoptotic genes SMAC/Diablo and HTRA2/Omi, each of these genes can induce apoptosis by blocking the caspase-inhibiting activity of IAP proteins (25, 40, 70). Overexpression of one of these genes is sufficient to induce apoptosis, while deletions that remove these genes reduce or eliminate developmentally programmed and stress-induced apoptosis (51, 71). Previous experiments have demonstrated that the *Drosophila* Apaf1 homolog *Ark* is induced following UV irradiation (79, 80). Following IR, *Ark* and other known components of the core apoptosis pathway in *Drosophila* did not show altered expression. In mammals, members of the TNF and TNF receptor families act in the cell-cell signaling, or extrinsic, apoptosis pathway. *Drosophila* has one member each in the TNF ligand and TNF receptor families (31, 35, 43). The *Drosophila* TNF ligand homolog *Eiger* is induced following DNA damage (Table 2), while the TNF receptor homolog is not (data not shown). *Eiger*, *rpr*, *skl*, and *hid* RNAs were all induced within 30 min of IR treatment.

One hour following IR, several DNA repair genes are induced. Two genes, *Mre11* and *Rad50*, encode proteins that are part of a heterotrimeric complex that also includes NBS1 (17). This repair complex has been implicated in multiple aspects of DNA repair, including homologous recombination, nonhomologous end joining, and delay of cell cycle progression. Two other induced genes, *Ku70* and *Ku80*, have more specific functions in DNA break repair as part of the nonhomologous end-joining pathway. The coordinate upregulation of two components of each repair complex indicated that these changes are functionally important.

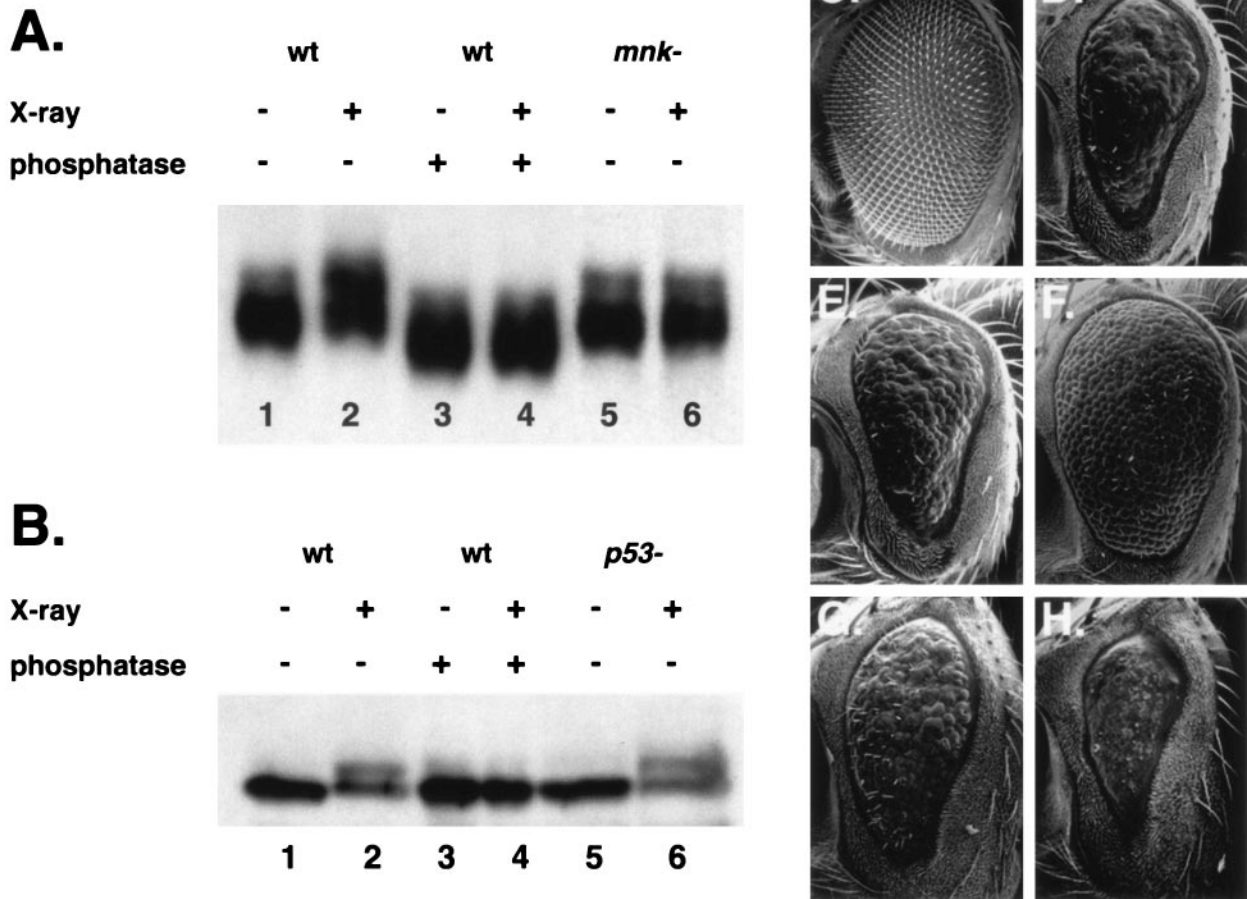


FIG. 4. *MNK/Chk2* is required for IR-induced modification of p53. (A) p53 protein was detected in lysates of untreated (lanes 1, 3, and 5) or irradiated (lanes 2, 4, and 6) 2- to 16-h-old embryos. p53 was immunoprecipitated with mouse anti-p53 monoclonal antibody, run on an SDS-PAGE gel, and detected with guinea pig anti-p53 polyclonal antibody. p53 mobility was reduced following irradiation of wild-type (lanes 1 and 2) but not *mnk* mutant (lanes 5 and 6) embryos. Phosphatase treatment (lanes 3 and 4) increased the mobility of p53 from both unirradiated (lane 3 versus lane 1) and irradiated (lane 4 versus lane 2) embryos. (B) MNK protein was detected in lysates of untreated (lanes 1, 3, and 5) or irradiated (lanes 2, 4, and 6) 2- to 16-h-old embryos. Lysates were run on an SDS-PAGE gel, and MNK was detected with rabbit anti-MNK polyclonal antibody (47). An anti-MNK signal with reduced mobility was detected in wild-type (lanes 1 and 2) and *p53* mutant (lanes 5 and 6) embryos. Phosphatase treatment (lanes 3 and 4) eliminated the signal, with reduced mobility from irradiated embryos (lane 4 versus lane 2). (C to H) Adult eyes from transgenic female animals were dehydrated and viewed by scanning electron microscopy. All animals carried a transgene expressing the Gal4 transcription factor under the control of the eye-specific promoter GMR. Expression of Gal4 alone did not affect eye development and is shown here as the wild-type control. Animals were raised at either 25°C (C to F) or 18°C (G and H). (C) GMR-Gal4/+; wild-type eye morphology. (D) GMR-Gal4 GUS-p53/+; p53-dependent rough eye phenotype at 25°C. (E) GMR-Gal4 GUS-p53/GUS-GRPS; the p53-dependent rough eye phenotype was unaffected by GUS-GRPS. (F) GMR-Gal4 GUS-p53/GUS-MNK(kd); the p53-dependent rough eye phenotype was suppressed by the kinase-dead form of MNK. (G) GMR-Gal4 GUS-p53/+; p53-dependent rough eye phenotype at 18°C. (H) GMR-Gal4 GUS-p53/GUS-MNK; the p53-dependent rough eye phenotype was enhanced by wild-type MNK.

Other targets of mammalian p53 include cell cycle regulators, including the cyclin-dependent kinase inhibitor p21, which mediates p53-dependent G₁-S arrest following DNA damage. In contrast, *dacapo*, the only member of the p21/p27 family in *Drosophila*, is repressed following IR (Table 2). RNA levels of other *Drosophila* cell cycle regulators are not altered following IR (data not shown). These results are consistent with our observation that *Drosophila* p53 is not required for IR-induced cell cycle arrest.

Besides proapoptotic genes and DNA repair genes, the remaining IR-induced targets do not resemble genes previously associated with mammalian p53 function. Two targets (CG17836 and CG5202) have motifs associated with known

transcriptional regulators. Other targets include a putative transmembrane protein (CG15658) and a member of the cytochrome P450 family.

Among the IR-repressed genes identified in our experiments, many are known or likely regulators of embryonic patterning or cell fate determination. It is possible that the decreased expression of these genes was a secondary effect of an *mnk*-dependent checkpoint.

Genetic analysis of *Eiger* and *Df(3L)Cat* function during IR-induced apoptosis. Although two TNF receptor family members, FAS and DR5/Killer, are targets of p53, neither is known to be required for damage-induced apoptosis. Previous analysis of the single *Drosophila* TNF ligand, *Eiger*, indicated

TABLE 2. Gene expression changes following IR^a

Gene	Fold change in expression										P (t)	P (t × g)	Domain and/or functional annotation		
	For wild type at min:					For <i>p53</i> mutants at min:								For <i>mkk</i> mutants at min:	
	015	030	060	120	240	030	060	120	030	060				120	
CG17836	4.9	6.2	10.1	9.1	6.2	1.2	0.7	1.4	1.2	0.6	1.6	1.1E-05	2.3E-07	AT-hook/HMG DNA-binding domain	
rpr	2.2	6.3	7.6	4.7	8.1	0.7	0.5	1.0	1.1	1.1	0.9	2.9E-08	3.8E-14	grh, apoptosis activator	
skl	1.2	4.0	7.6	7.0	9.1	0.5	0.7	0.9	1.1	0.9	1.0	8.4E-09	1.1E-15	grh, apoptosis activator	
Cyp307a1	1.2	1.6	5.0	5.5	*	1.0	0.8	1.7	1.2	1.3	1.2	1.7E-06	6.2E-05	cytochrome P450	
hid	1.1	2.1	3.2	2.2	1.8	0.6	0.6	0.6	0.7	1.0	0.8	3.4E-01	2.7E-07	grh, apoptosis activator	
CG13714	1.5	2.5	2.7	2.0	0.9	1.1	0.9	1.0	1.0	0.9	1.0	1.3E-06	6.0E-10	tumor necrosis factor family, apoptosis activator	
Eiger	1.0	1.6	2.3	1.9	2.0	0.9	1.0	0.9	1.0	1.0	1.2	1.1E-04	1.9E-07	DNA binding, DNA Repair	
Ku70	1.1	0.9	2.2	3.1	3.7	1.0	1.1	1.5	1.0	1.2	1.2	4.0E-05	4.3E-03	Leucine-rich repeat (LRR) transmembrane protein	
CG15658	1.3	1.5	2.0	1.6	0.9	1.0	1.1	1.0	1.1	1.3	1.0	1.0E-05	1.3E-04	DNA binding, DNA Repair	
CG10965	1.4	2.0	2.0	2.1	2.1	1.0	0.9	0.9	1.1	1.2	1.3	2.1E-06	1.7E-06	DNA binding, DNA Repair	
Ku80	1.1	0.8	1.9	2.3	2.5	1.1	0.9	1.2	1.0	1.1	1.1	1.7E-04	2.9E-03	WD40 domains, similar to Embryonic Ectoderm Development	
CG15479	1.2	1.3	1.8	2.6	2.3	1.0	1.0	1.2	1.1	1.1	1.1	2.2E-05	3.5E-05	coiled-coil domain, DNA repair	
CG16815	1.3	1.5	1.7	1.6	1.3	1.2	1.1	1.1	1.0	0.8	1.1	2.0E-05	2.1E-05	DNA Repair	
CG5202	1.0	1.0	1.4	2.1	2.4	0.9	1.0	1.0	1.0	1.1	1.3	6.6E-07	1.4E-05	WD40 domains, similar to Embryonic Ectoderm Development	
CG15480	1.4	1.6	1.4	1.7	3.4	1.1	1.0	1.3	1.1	1.1	1.1	6.7E-05	1.9E-02	coiled-coil domain, DNA repair	
Rad50	1.1	1.0	1.4	2.5	2.5	1.1	1.2	1.8	1.0	1.1	1.3	5.2E-09	4.5E-03	DNA Repair	
Mre11	1.1	0.8	1.2	2.0	3.1	1.1	1.0	1.2	1.1	1.1	1.3	1.9E-04	4.0E-01	DNA Repair	
Wnt8	1.0	0.6	0.4	1.1	*	1.0	1.1	1.5	1.0	1.2	1.4	4.8E-01	8.6E-06	secreted signaling protein	
inv	1.2	0.5	0.5	0.6	0.6	1.0	0.5	0.5	0.8	1.0	0.9	8.7E-06	1.9E-02	transcription factor, homeobox domain, development	
ac	1.0	0.4	0.6	0.6	1.8	0.8	0.7	1.0	1.0	0.9	1.0	1.3E-04	3.1E-03	transcription factor, Helix-loop-helix domain, neurodevelopment	
wun2	1.0	0.7	0.6	0.6	0.9	1.0	0.8	0.8	1.0	1.1	1.1	3.6E-05	6.4E-06	phosphatidic acid phosphatase a-2, cell migration	
ab	1.0	0.7	0.6	0.5	0.4	0.9	0.7	0.8	0.8	1.1	0.9	1.7E-04	5.5E-02	transcription factor, BTB and C2H2 type Zinc finger domains, development	
Oli	0.9	0.6	0.6	0.5	0.7	1.1	0.5	0.5	0.6	1.1	0.6	1.6E-04	5.4E-01	transcription factor, Helix-loop-helix domain	
l(l)sc	0.8	0.4	0.6	0.8	1.8	0.8	0.7	1.0	1.1	1.2	1.2	8.7E-04	4.5E-05	transcription factor, Helix-loop-helix domain, neurodevelopment	
nerfin-1	1.1	0.5	0.6	0.7	0.9	0.7	0.6	0.7	0.6	1.0	0.8	5.4E-07	3.4E-01	transcription factor, zinc finger domain, neurodevelopment	
CG8886	1.3	0.8	0.6	0.5	0.9	1.0	0.8	0.7	1.0	1.1	0.7	1.6E-04	4.2E-02	similar to p21/p27, cyclin-dependent protein kinase inhibitor	
CG8045	1.0	0.9	0.6	0.4	0.3	0.8	0.5	0.5	0.6	1.2	0.7	2.1E-06	2.3E-01	transmembrane receptor protein tyrosine kinase, development	
dap	1.1	0.6	0.6	0.4	0.6	0.9	0.7	0.8	0.9	1.0	0.9	1.7E-05	5.6E-03	transcription factor, ligand-dependent nuclear receptor, development	
drl	1.0	0.6	0.7	0.6	0.5	0.7	0.7	0.7	0.9	1.2	0.8	3.3E-07	2.6E-03	transcription factor, BTB and C2H2 type Zinc finger domains, development	
Dref	1.1	0.8	0.7	0.6	0.8	1.1	0.8	0.7	1.0	1.1	0.9	3.3E-05	2.9E-03	transcription factor, DNA Replication	
tlf	1.1	0.6	0.7	0.8	0.6	0.8	0.5	0.7	1.0	0.9	1.0	3.1E-06	1.4E-02	transcription factor, coiled-coil, proliferation control	
prod	1.2	0.7	0.7	0.6	0.8	1.0	0.7	0.7	0.9	1.0	0.9	1.9E-04	7.3E-02	chromatin associated, coiled-coil, transcription factor	
tkk	0.8	0.7	0.7	0.6	0.5	0.9	0.5	0.8	0.8	1.0	0.8	8.4E-05	2.1E-01	transcription factor, BTB and C2H2 type Zinc finger domains, development	
CG8147	1.0	0.8	0.8	0.6	0.5	1.0	0.8	0.6	1.0	1.1	0.9	1.6E-04	1.6E-02	alkaline phosphatase	
en	1.0	0.6	0.8	0.5	0.7	0.8	0.7	0.8	0.9	1.0	0.9	1.7E-04	4.2E-02	RNA polymerase II transcription factor, homeobox domain	

^a mRNA was prepared 15 to 240 min following treatment of wild-type, *p53* mutant, or *mkk* mutant embryos. Each value is the average of three experiments determining the increase or decrease of expression in irradiated and untreated samples. The O-PCR values indicate the increases following irradiation of wild-type embryos. O-PCR was performed using samples collected 60 min following irradiation, except for Ku80 and Mre11, which were analyzed at 240 min following irradiation. P values are given for treatment (t) and treatment by genotype (t × g). Red shading indicates increases greater than or equal to 1.5-fold. Blue shading indicates decreases less than or equal to 0.7-fold. Gene names are from www.flybase.org. Asterisks indicate that negative expression values resulted in negative (Cyp307a1) or very high (Wnt8) expression ratios.

that overexpression of *Eiger* can induce apoptosis by activating the Jnk pathway (31, 43). To examine the role of *Eiger* in IR-induced apoptosis, we generated gain- and loss-of-function alleles of *Eiger* (Fig. 5A). We confirmed that overexpression of *Eiger* in the developing wing was sufficient to induce apoptosis (Fig. 5B and C). Transposon-mediated mutagenesis was used to make deletions in the *Eiger* locus (Fig. 5A). Both deletions removed the predicted start codon. As previously reported (31), animals with these deletions were viable and fertile (data not shown). *Eiger* mutant animals had normal levels of IR-induced apoptosis at 4 and 8 h following irradiation with 4,000-rad X rays (Fig. 5D and E and data not shown). At lower X-ray doses, induction of apoptosis in wild-type animals was more variable. The levels observed in *Eiger* mutants overlapped with those in wild-type animals (data not shown). These results indicate that while *Eiger* overexpression is sufficient to induce apoptosis, it is not required for IR-induced apoptosis.

Previous studies have demonstrated that deletions that span the region containing the *hid*, *grm*, *rpr*, and *skl* genes reduce or eliminate apoptosis in *Drosophila* (51, 71). In embryos homozygous for *Df(3L)H99*, developmentally regulated and DNA damage-induced apoptosis was absent. However, the proapoptotic gene *skl* is not included in this deficiency but is highly induced following DNA damage. Thus, IR-induced expression of *skl* is not sufficient to induce apoptosis in the absence of the other proapoptotic genes in this region (14, 61). One possible explanation for this result is that following DNA damage, induction of multiple proapoptotic genes, *hid*, *rpr*, and *skl*, is required to produce a proapoptotic signal that exceeds a threshold level required to irreversibly commit cells to the apoptotic program. In this model, reducing the expression of all three genes by 50% might prevent the proapoptotic signal from reaching the threshold. To test this possibility, we examined IR-induced apoptosis in animals heterozygous for chromosome deficiencies that remove all of the damage-induced transcripts in the 75C region. In *Df(3L)Cat/+* or *Df(3L)WR10/+* animals, induction of apoptosis 4 h following IR was greatly reduced (Fig. 5F and data not shown). In contrast, animals heterozygous for other deficiencies that do not remove these genes showed normal induction of apoptosis (data not shown). Although the deficiencies showing reduced apoptosis also removed one copy of additional genes, *rpr*, *skl*, and *hid* are the only IR-induced genes within the affected regions. These results demonstrate a dose-sensitive requirement for genes in the 75C region for IR-induced apoptosis and are consistent with a threshold model for IR-induced apoptosis.

To further delineate which genes contribute to haploinsufficiency, we examined IR-induced apoptosis in animals heterozygous for deficiencies or point mutations removing a subset of genes in this region (Fig. 5G to K). Surprisingly, animals heterozygous for a deficiency that removes *rpr* and *skl* still exhibited a robust induction of apoptosis (Fig. 5G). In contrast, animals heterozygous for deficiencies or point mutations that remove *hid* all showed a significant decrease in IR-induced apoptosis.

DISCUSSION

Roles of MNK/Chk2 and p53 in damage response pathways.

Previous studies have established that *Drosophila* p53 mediates

X-irradiation-induced apoptosis and expression of *rpr* and *skl* (11, 48, 61). Here, we characterized the pathway that transduces the DNA damage signal to the apoptosis and cell cycle machineries. Our results indicated that a number of genes in this pathway are largely specific to the cell cycle or apoptotic response. Both cellular assays and transcriptional profiling suggest that *Drosophila* p53 is required for IR-induced regulation of apoptosis but is not required for G₂ arrest. In contrast, *mei-41*, *mus304*, and *grps* were required for cell cycle arrest, but not induction of apoptosis. Our biochemical experiments suggested that *mnk*, which encodes a conserved damage-activated kinase, is required for phosphorylation of p53 following IR. All IR-induced transcription required both *mnk* and p53. The absence of genes that only required *mnk* or p53 was consistent with a linear signaling pathway of MNK activating p53, which acts as a global regulator of IR-induced transcription (Fig. 6).

Although *mnk* and p53 mutant animals have similar defects in IR-induced transcription, *mnk* also acts in p53-independent pathways. In animals with mutations in double-strand break repair enzymes, unrepaired breaks formed during meiotic recombination activate an *mnk*-dependent checkpoint signal that disrupts oocyte patterning and nuclear morphology (1, 24). Induction of the meiotic checkpoint differs from IR-induced transcription in at least two respects. First, activation of *mnk* during meiosis requires *mei-41*, the *Drosophila* homolog of *ATR*. Second, p53 is not required for this damage response pathway. In a different damage response pathway, *mnk*, but not p53, is required for damage-induced inactivation of centrosomes (64). In this study, we found that IR induced a p53-independent decrease in RNA levels of at least 17 genes, including many developmental regulators. Although this observation could indicate a transcriptional repressor that is regulated by *mnk*, we favor a model in which an *mnk*-dependent cell cycle delay following IR has a secondary effect on the developmental induction of these genes. Together, our results and previous studies indicate that *mnk* regulates multiple signaling pathways in addition to p53-dependent induction of gene expression.

In mammals, Chk2 and other checkpoint kinases block Mdm2-mediated turnover and inhibition of p53. Several lines of evidence suggest that this regulatory mechanism is not conserved in *Drosophila*. First, simple sequence searches have not revealed an obvious Mdm2 homolog in the *Drosophila* genome. Second, the *Drosophila* p53 protein sequence does not contain a conserved binding site for Mdm2. Finally, we found that p53 protein levels were not dramatically altered following IR. p53 did exhibit an IR-induced change in gel mobility due to *mnk*-dependent phosphorylation. Thus, our results provide a clear example of damage-induced activation of p53 without changes in p53 protein levels.

Phosphorylation of p53 by Chk2 may represent an important step in the evolution of DNA damage responses in multicellular animals. Checkpoint pathways regulating cell cycle control and DNA repair have been highly conserved in eukaryotes, including unicellular organisms such as yeast. In contrast, induction of apoptosis during development or in response to cellular stress is confined to multicellular organisms. We found that p53 phosphorylation by Chk2/MNK is a conserved molecular link between DNA damage detection and the core apoptotic machinery in metazoans. Mdm2 adds an additional layer

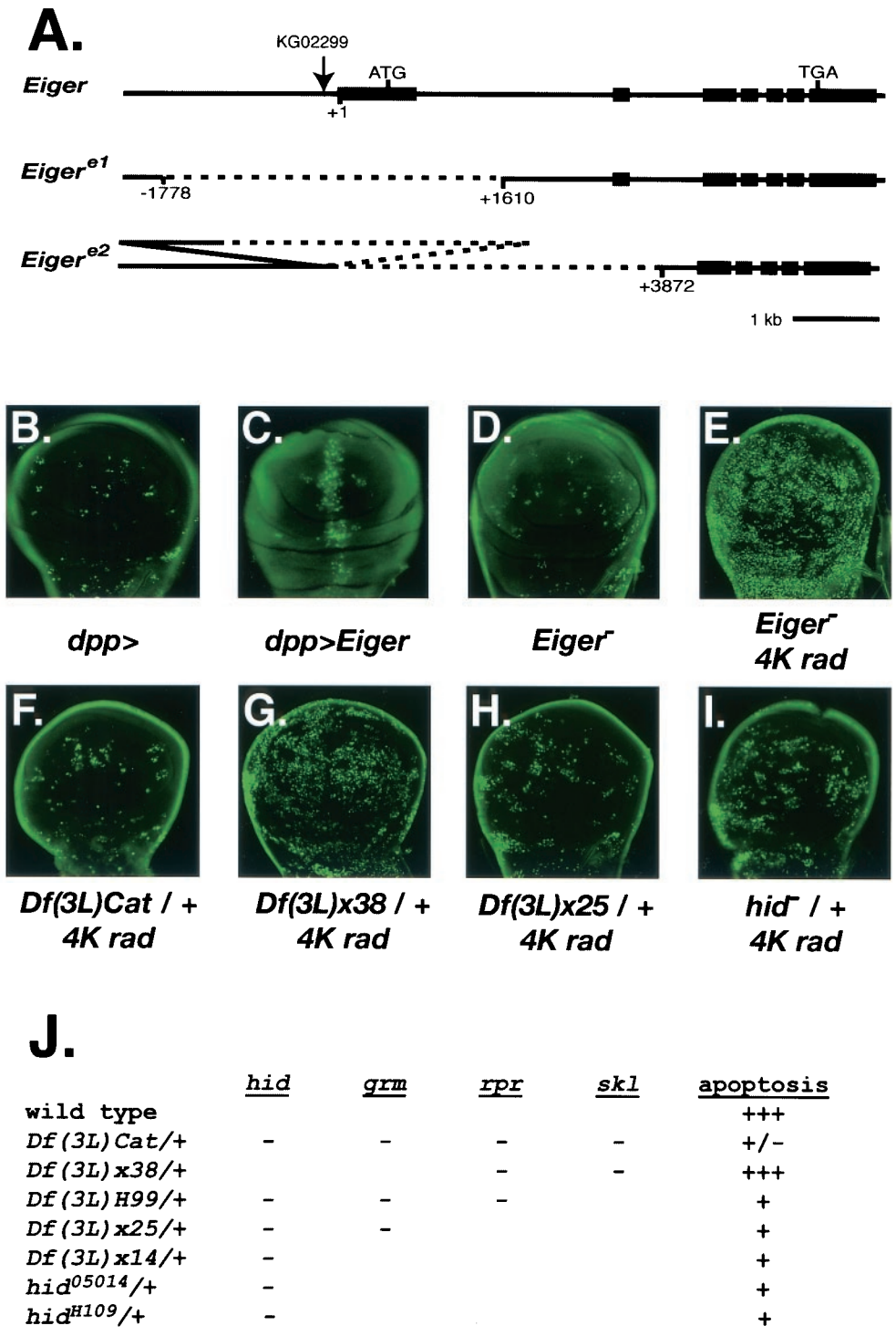


FIG. 5. Genetic analysis of p53 targets and DNA damage-induced apoptosis. (A) Maps of *Eiger* mutations. Exons are depicted as rectangles, while introns are lines. The P-element insertion KG02299 (arrow) was used to generate two deletions in the *Eiger* locus. *Eiger*^{e1} removes the first exon. *Eiger*^{e2} removes the first two exons and leaves part of the KG02299 insertion. (B to F) Imaginal wing disks were dissected from third instar larvae and stained with the vital dye acridine orange to detect apoptotic cells. (B) *dpp*-Gal4 does not induce apoptosis. (C) *dpp*-Gal4 driving UAS-*Eiger* induces apoptosis at the anterior-posterior boundary. (D) Normal spontaneous apoptosis in homozygous *Eiger*^{e1} disks. (E) Normal IR-induced apoptosis in homozygous *Eiger*^{e1} disks 4 h following 4,000-rad X rays. (F to I) Imaginal wing disks were dissected from third instar larvae 4 h following 4,000-rad X rays and stained with the vital dye acridine orange. IR-induced apoptosis was reduced in *Df(3L)Cat*/+ (F), *Df(3L)x25*/+ (H), and *hid*⁰⁵⁰¹⁴/+ (I) and normal in *Df(3L)x38*/+ (G). (J) Summary of apoptotic response in animals heterozygous for deficiencies or point mutations in the 75C region. Genes disrupted in a given deficiency or mutant are indicated by a minus sign.

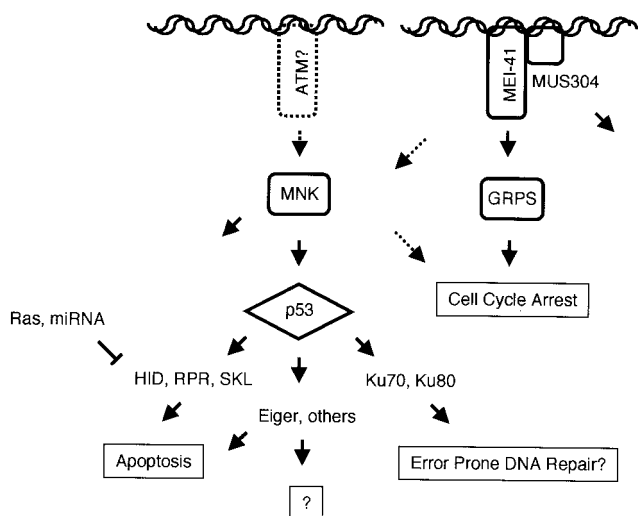


FIG. 6. A working model for the organization of DNA damage response pathways in *Drosophila*. See the text for a discussion.

of complexity to the regulation of mammalian p53 compared to *Drosophila* p53. Regulation of p53 turnover by Mdm2 may provide mammalian cells with greater control of the levels or timing of p53-dependent transcription.

DNA damage-induced targets of p53. We have used microarray analysis to perform a comprehensive analysis of p53 targets following exposure to IR. The number of genes identified in these experiments was substantially smaller than the number of p53 targets identified in mammals. In part, this observation may reflect underlying differences in the damage response pathway in flies and mammals. For example, induction of p21 by mammalian p53 mediates G₁ arrest following damage. To the best of our knowledge, IR-induced G₁ arrest has not been described in *Drosophila*, consistent with our observation that the *Drosophila* p21/p27 homolog *dacapo* is not induced by IR. However, the smaller number of targets identified in *Drosophila* also reflects experimental differences. We have examined expression changes induced by IR and during a defined window of embryonic development. In contrast, targets of mammalian p53 have been identified in many different cell types following different types of DNA damage or simply overexpression of p53. It is likely that additional targets of *Drosophila* p53 will be identified using other types of cellular stresses in different cell types or developmental stages. For example, UV irradiation of *Drosophila* embryos has been shown to induce Apaf1 through either E2F or *mei-41*, depending on the developmental stage (80).

The most prominent group of p53 targets identified in our study regulates two apoptotic pathways that are also targeted by mammalian p53. *hid*, *rpr*, and *skl* are part of a group of genes that induce apoptosis by blocking the caspase-inhibiting activity of IAP proteins (25, 40, 70). Recent experiments have confirmed that HTRA2, a functional homolog of these genes, is a target of mammalian p53 (34). The *Drosophila* p53 target *Eiger* is a member of the TNF ligand family and can induce apoptosis when overexpressed (31, 43). In mammals, FAS and DR5/Killer are p53 targets that can regulate apoptosis by acting as receptors for TNF ligand family members (19, 68). Thus,

we have identified two examples of mammalian and *Drosophila* p53 regulating common signaling pathways. Combined with the many other proapoptotic targets of mammalian p53, these results support the general hypothesis that multiple components of proapoptotic signaling pathways can be targets for transcriptional regulation following stresses such as DNA damage.

Although FAS and DR5/Killer are targets of mammalian p53 and act in the extrinsic apoptosis pathway, it is unclear what role they play in DNA damage-induced apoptosis (49, 58, 74). Our analysis of deletion mutations in the *Drosophila* p53 target *Eiger* indicated that this gene is not required to initiate IR-induced apoptosis. This negative result was not due to redundancy with a related molecule, since *Eiger* is the only TNF-related gene in the *Drosophila* genome sequence. It is possible that the conserved activation of the TNF pathway by p53 is required for the induction of apoptosis under specific conditions not tested in our experiments. Alternatively, induction of *Eiger* may activate other cellular responses to DNA damage. Further characterization of *Eiger* function should reveal how cell-cell signaling contributes to survival or genomic stability following DNA damage in multicellular organisms.

Our analysis of the remaining proapoptotic targets of p53 indicated that they are part of a dosage-sensitive mechanism that regulates IR-induced apoptosis. In contrast to *Eiger*, the proapoptotic genes in the genetic region containing *hid*, *rpr*, and *skl* are both sufficient and necessary for apoptosis. Animals with deletions that include genes in this region are defective in IR-induced apoptosis (51, 71). Because these proapoptotic genes act, at least in part, by inhibiting a common target (IAP1/Thread), it has been proposed that they contribute to a rheostat-like mechanism in which the added activity of all proapoptotic proteins present must pass a threshold before a cell undergoes the irreversible decision to undergo programmed cell death. Following our observation that three of these genes are induced following DNA damage, we tested the effect of lowering the dose of all proapoptotic genes in this region by half. We found that deletions in this region were haploinsufficient for IR-induced apoptosis. Dose sensitivity may represent an important feature of damage-induced apoptosis. Animals heterozygous for these deletions exhibit apparently normal morphology and fertility, suggesting that they are not haploinsufficient for developmentally regulated apoptosis. One possible interpretation of these results is that the apoptotic signal in many developmental contexts is well past the threshold required to commit to apoptosis, while the apoptotic signal following DNA damage is closer to that threshold. A lower apoptotic signal following DNA damage may allow cells to monitor DNA repair and block apoptosis if repair is successful. Haploinsufficiency of some tumor suppressor genes, including p53, has been proposed to contribute to cancer development (15, 66). If stress-induced apoptosis in mammals is sensitive to the dose of p53 target genes, haploinsufficiency of these genes may also contribute to suppression of apoptosis, particularly in cells with extensive aneuploidy.

Analysis of animals heterozygous for deletions that removed a subset of genes revealed that loss of one copy of *hid* was sufficient to reduce IR-induced apoptosis. A greater reduction was observed in larger deletions, indicating that additional genes in this region, likely *rpr* and *skl*, also contribute to IR-induced apoptosis. Previous analysis of animals heterozygous

for two overlapping deletions [*Df(3L)H99* and *Df(3L)xr38*] that remove both copies of *rpr* demonstrated reduced levels of IR-induced apoptosis (51). Our results indicate that part of that reduction is due to haploinsufficiency of *hid* and other genes in this region. Although the induction of *hid* RNA was lower than that observed for *rpr* and *skl*, *hid* may exhibit a greater absolute difference in RNA and protein levels following IR. Because null mutations in *hid* are embryonic lethal, we have not examined the effects of completely removing *hid* function. The dose-sensitive effects of *hid* suggest that total loss of *hid* would completely block IR-induced apoptosis. However, even in animals with normal levels of *hid*, increased levels of *rpr* and *skl* may be required to pass the proapoptotic signaling threshold required for a full DNA damage response. The Ras pathway (6, 7, 37, 54) and a micro-RNA in the *bantam* locus (10) regulate *hid* expression. These and other pathways regulating *hid* may help determine which cells in the developing wing are most sensitive to DNA damage.

The other class of p53 targets identified in our experiments includes components of the Ku and Mre11 DNA repair complexes. Both of these complexes participate in repair of double-strand DNA breaks by nonhomologous end joining (NHEJ) (33). Compared with homologous recombination, NHEJ is a potentially error-prone mechanism for DNA repair. Mutagenic DNA repair mechanisms are a prominent feature of the SOS response in bacteria that apparently promotes cell survival following DNA damage at the expense of genomic integrity (63). The ability of multicellular animals to eliminate damaged cells by apoptosis might suggest that low-fidelity mechanisms of DNA repair would not be favored following damage. However, the induction of NHEJ components by p53 suggests that mechanisms such as apoptosis or cell cycle arrest that are presumed to prevent mutations following DNA damage may compete with mechanisms that promote cell survival and prevent aneuploidy by error-prone DNA repair. The previous demonstration that an isoform of Ku86 is also a target of mammalian p53 (9, 44) suggests that this is an evolutionarily conserved response to DNA damage in metazoans that may modulate mutagenesis following DNA damage.

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ADDENDUM IN PROOF

Jassim et al. have shown that *hid* is also induced by UV irradiation but that p53 protects cells from UV-induced apoptosis, apparently through enhanced DNA repair (O. W. Jassim, J. L. Fink, and R. L. Cagan, *EMBO J.* **22**:5622–5632, 2003). Although the targets of p53 that mediate enhanced repair of UV damage are unknown, these results are consistent with our conclusion that *Drosophila* p53 regulates DNA repair.

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