

c-Jun N-terminal kinase-mediated Rad18 phosphorylation facilitates Pol η recruitment to stalled replication forks

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ABSTRACT The E3 ubiquitin ligase Rad18 chaperones DNA polymerase η (Pol η) to sites of UV-induced DNA damage and monoubiquitinates proliferating cell nuclear antigen (PCNA), facilitating engagement of Pol η with stalled replication forks and promoting translesion synthesis (TLS). It is unclear how Rad18 activities are coordinated with other elements of the DNA damage response. We show here that Ser-409 residing in the Pol η -binding motif of Rad18 is phosphorylated in a checkpoint kinase 1-dependent manner in genotoxin-treated cells. Recombinant Rad18 was phosphorylated specifically at S409 by c-Jun N-terminal kinase (JNK) *in vitro*. In UV-treated cells, Rad18 S409 phosphorylation was inhibited by a pharmacological JNK inhibitor. Conversely, ectopic expression of JNK and its upstream kinase mitogen-activated protein kinase kinase 4 led to DNA damage-independent Rad18 S409 phosphorylation. These results identify Rad18 as a novel JNK substrate. A Rad18 mutant harboring a Ser \rightarrow Ala substitution at S409 was compromised for Pol η association and did not redistribute Pol η to nuclear foci or promote Pol η -PCNA interaction efficiently relative to wild-type Rad18. Rad18 S409A also failed to fully complement the UV sensitivity of Rad18-depleted cells. Taken together, these results show that Rad18 phosphorylation by JNK represents a novel mechanism for promoting TLS and DNA damage tolerance.

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Abbreviations used: ATR, ATM- and Rad3-related; B[a]P, benzo[a]pyrene; BPDE, (+)-7,8-dihydroxy-t-9,10-epoxy-7,8,9,10-tetrahydrobenzo[a]pyrene; Cdc7, cell division cycle 7; Chk1, checkpoint kinase 1; CPD, cyclobutane pyrimidine dimer; Dbf4, Dumbbell former 4; DDK, Dbf4-dependent kinase; DDR, DNA damage response; DSB, double-stranded break; GFP, green fluorescent protein; GST, glutathione S-transferase; JNK, c-Jun N-terminal kinase; PCNA, proliferating cellular nuclear antigen; PIP, PCNA-interacting peptide; Pol η , DNA polymerase eta; Pol κ , DNA polymerase kappa; Rad6, radiation sensitive 6; Rad18, radiation sensitive 18; RPA, replication protein A; ssDNA, single-stranded DNA; TLS, translesion synthesis; UV, UV radiation; XPV, xeroderma pigmentosum variant; YFP, yellow fluorescent protein.

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INTRODUCTION

Various environmental agents induce DNA lesions termed "adducts" that impede the progression of the DNA replication machinery. For example the chemical carcinogen benzo[a]pyrene generates genotoxic metabolites, including (+)-7,8-dihydroxy-t-9,10-epoxy-7,8,9,10-tetrahydrobenzo[a]pyrene (BPDE), which reacts primarily with the N² amino group of guanine to form a BPDE-DNA (BPDE-N²-dG) adduct (Thakker et al., 1976; Conney, 1982; Dipple, 1995). Similar helix-distorting DNA lesions result from solar UV radiation, which generates cyclobutane pyrimidine dimers (CPDs; formed by linkage of two adjacent pyrimidines) and to a lesser extent pyrimidine (6-4) pyrimidone photoproducts (Pfeifer et al., 2005). BPDE adducts, CPDs, and many other DNA lesions block the progression of replicative DNA polymerases and cause stalling of active replication forks. Failure to preserve stalled replication forks can lead to fork collapse and acquisition of DNA double strand breaks (DSBs). Moreover, error-prone replicative bypass of damaged DNA can lead to

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mutations and cancer. Therefore appropriate replication and repair of damaged DNA is necessary to prevent genomic instability.

Cell cycle checkpoints and translesion synthesis are two ubiquitous responses to DNA damage and replication fork stalling. Both processes are hypothesized to play important roles in maintaining genome stability and preventing cancer. Checkpoints are DNA damage-induced signal transduction pathways that delay cell cycle phase transitions and promote repair mechanisms, thereby integrating cell cycle progression with DNA repair (Zhou and Elledge, 2000; Sancar *et al.*, 2004; Cimprich and Cortez, 2008). Many cell cycle checkpoint mechanisms have been described, and these respond to different forms of DNA damage (Zhou and Elledge, 2000; Sancar *et al.*, 2004; Cimprich and Cortez, 2008). The S-phase checkpoint induced by bulky lesions such as BPDE or CPD is mediated via the ATR and Chk1 protein kinases (Guo *et al.*, 2002; Heffernan *et al.*, 2002; Liu *et al.*, 2006; Barkley *et al.*, 2007; Heffernan *et al.*, 2007). ATR/Chk1-mediated S-phase checkpoint signaling is triggered by uncoupling of replicative DNA helicase (comprising Cdc45, Mcm2-7, and the GINS complex) and DNA polymerase activities (Byun *et al.*, 2005; Cimprich and Cortez, 2008). When replicative DNA polymerases are blocked by DNA lesions (CPD and BPDE–DNA adducts), helicase activity continues to unwind DNA ahead of the stalled polymerase (Byun *et al.*, 2005; Cimprich and Cortez, 2008). Replicative helicase–polymerase uncoupling results in accumulation of single-stranded DNA (ssDNA), which is coated by replication protein A (RPA; Zou and Elledge, 2003; Zou *et al.*, 2003). RPA-coated ssDNA helps to recruit and activate the ATR kinase (Burrows and Elledge, 2008; Mordes and Cortez, 2008; Mordes *et al.*, 2008), leading to phosphorylation and activation of the checkpoint kinase Chk1 (a key effector of the ATR pathway). Chk1 signaling inhibits DNA synthesis at unfired origins, stabilizes stalled replication forks, and inhibits entry into mitosis in the presence of unreplicated DNA (Sancar *et al.*, 2004; Cimprich and Cortez, 2008). The precise mechanisms by which stalled forks are stabilized and the mechanisms by which late origins are inhibited by Chk1 in mammalian cells are not well understood. Nevertheless, fork stabilization and repression of origin firing by Chk1 are believed to play key roles in cell cycle progression and maintenance of genomic stability even in cells that do not acquire DNA damage from exogenous sources (Sorensen *et al.*, 2004).

Concomitant with activation of ATR/Chk1 signaling, translesion synthesis (TLS) DNA polymerases are recruited to stalled replication forks (Bi *et al.*, 2005, 2006; Bomgarden *et al.*, 2006). In contrast to the DNA polymerases that perform bulk replicative DNA synthesis during S phase, “Y-family” TLS polymerases have low fidelity and processivity on undamaged DNA (Ohmori *et al.*, 2001; Prakash *et al.*, 2005). However, because of active sites that can accommodate helix-distorting lesions, TLS enzymes can bypass various forms of DNA damage with relative accuracy compared with replicative polymerases (Prakash *et al.*, 2005). In eukaryotes, the main TLS polymerases are DNA polymerase η (Pol η), DNA polymerase κ (Pol κ), DNA polymerase ι (Pol ι), REV1, and DNA polymerase ζ (Pol ζ). Collectively, these enzymes are responsible for translesion synthesis of various DNA adducts.

In a process termed polymerase switching, replicative DNA polymerases that encounter DNA lesions are replaced by TLS enzymes. The molecular basis of the DNA polymerase switch is best understood for *Saccharomyces cerevisiae* Rad30 and its mammalian homologue Pol η (Kannouche and Lehmann, 2004; Ulrich, 2004). When replication forks stall at sites of DNA damage, proliferating cell nuclear antigen (PCNA) is monoubiquitinated on lysine 164 (Ohmori *et al.*, 2001; Prakash *et al.*, 2005). PCNA K164 monoubiquitination is

mediated mainly by Rad18 (an E3 ubiquitin ligase) and Rad6 (an E2 ubiquitin–conjugating enzyme). PCNA ubiquitination results in part from DNA damage–induced recruitment of Rad18 to stalled replication forks. The Ulrich and Tateishi laboratories proposed that Rad18 associates with stalled replication forks via direct interactions with RPA-coated ssDNA (Davies *et al.*, 2008; Huttner and Ulrich, 2008; Tsuji *et al.*, 2008), and the p95/NBS protein may facilitate RPA-mediated Rad18 recruitment (Yanagihara *et al.*, 2011). DNA damage–induced degradation of the PCNA deubiquitinating enzyme USP1 may also contribute to accumulation of ubiquitinated PCNA in genotoxin-treated cells (Huang *et al.*, 2006). Monoubiquitinated PCNA is considered to promote recruitment of TLS polymerases to the replication fork. Specialized ubiquitin (Ub)-binding domains (termed UBZ or UBM motifs) facilitate interactions between TLS polymerases and monoubiquitinated PCNA (Bienko *et al.*, 2005). After lesion bypass, another switch is presumed to restore the replicative DNA polymerase to the fork.

TLS can be error-free or error-prone. Whereas error-free TLS suppresses mutagenesis and promotes DNA damage tolerance, error-prone TLS is mutagenic. TLS polymerases display specificity for bypass of different lesions *in vitro*. For example, Pol η correctly inserts A–A across T–T dimers, whereas other TLS polymerases (likely Pol ι and Pol κ) perform error-prone insertion across CPD (Ziv *et al.*, 2009). Therefore Pol η is crucial for error-free TLS, maintenance of ongoing DNA replication, and DNA damage tolerance in UV-treated cells. Indeed, UV-sensitivity and mutagenesis are hallmarks of xeroderma pigmentosum variant cells that lack Pol η (Maher *et al.*, 1976; Johnson *et al.*, 1999; Masutani *et al.*, 1999). Clearly, selection of the appropriate TLS polymerase for a particular lesion is crucial for maintaining genomic stability. However, the molecular mechanism for recruiting appropriate TLS polymerase(s) and ensuring error-free lesion bypass is not fully understood. Pol η is a versatile enzyme and may be the preferred TLS polymerase and the default enzyme selected for bypass of most lesions. Rad18 interacts directly with and helps to guide Pol η to sites of replication fork stalling (Watanabe *et al.*, 2004; Day *et al.*, 2010), leading to stable engagement of Pol η with PCNA. In part, the preferential selection of Pol η over other TLS polymerases might result from specific and direct associations between Pol η and Rad18.

Although S-phase checkpoint signaling and TLS are initiated independently via a common damage-induced DNA replication intermediate (RPA-coated ssDNA; Chang *et al.*, 2006a), there are some indications that TLS and checkpoint signaling are coupled in a manner that is necessary for appropriate responses to fork-stalling DNA lesions. For example, defective TLS results in elevated ATR/Chk1 signaling, most likely due to persistence of the postreplicative ssDNA gaps generated by polymerase/helicase uncoupling (Bi *et al.*, 2005, 2006; Bomgarden *et al.*, 2006). Conversely, certain aspects of TLS may be facilitated by checkpoint signaling. For example, we showed that PCNA monoubiquitylation and its association with Pol κ are partially dependent on ATR/Chk1 signaling (Bi *et al.*, 2006). Zou and colleagues also showed that DNA damage–inducible PCNA ubiquitination is ablated in Chk1-depleted cells (Yang *et al.*, 2008; Yang and Zou, 2009). Lehmann and colleagues found that Pol η is an ATR substrate whose genotoxin-inducible phosphorylation modulates checkpoint signaling (Gohler *et al.*, 2011). Nevertheless, the mechanisms that integrate checkpoint signaling with TLS are incompletely understood.

Because checkpoints are mediated by protein kinases, we hypothesized that Chk1-dependent phosphorylation of TLS proteins might provide a potential mechanism for linking S-phase checkpoint signaling with TLS, thereby coordinating distinct branches of DNA

damage response. Rad18 plays a central role in TLS and additional DNA repair pathways (Huang *et al.*, 2009; Watanabe *et al.*, 2009). Therefore we hypothesized that Rad18 represented a likely target of Chk1 signaling. Consistent with a hypothetical link between Chk1 and Rad18, we show here that Rad18 is phosphorylated inducibly (and in a Chk1-dependent manner) in response to DNA damage. We identified Ser-409 as a DNA damage-induced phosphorylation site that resides within the Pol η -binding motif of Rad18. We demonstrate that Rad18 phosphorylation at S409 specifically promotes Rad18–Pol η associations, facilitates efficient recruitment of Pol η to the replication machinery, and confers DNA damage tolerance. Surprisingly, however, we show that Rad18 S409 is not phosphorylated directly by checkpoint kinases. Instead we identify c-Jun N-terminal kinase (JNK; a member of the stress-activated protein kinase [SAPK] family) as the Rad18 S409 kinase. It is well established that JNK and other SAPKs mediate transcriptional events that modulate survival and apoptotic signaling in response to various cellular stresses, including UV radiation (Davis, 2000). There are recent indications that SAPKs may also participate in checkpoint control when the genome is compromised (Reinhardt *et al.*, 2007; Reinhardt and Yaffe, 2009). However, a relationship between SAPK signaling and TLS had not previously been described. Therefore our results demonstrate novel and unanticipated integration of checkpoint kinase signaling, SAPK cascades, and TLS, three major effector branches of the cellular response to UV irradiation.

RESULTS

Rad18 is phosphorylated in response to DNA damage

During immunoblot analyses we noticed that Rad18 in extracts from genotoxin-treated cells migrated with retarded electrophoretic mobility relative to Rad18 in control (untreated) cell extracts (Figure 1A). Moreover, the mobility shift of Rad18 was reversed by incubation with λ phosphatase (Figure 1A), indicating that Rad18 was phosphorylated inducibly in response to DNA damage. In immunoblotting experiments anti-phosphoserine/phosphothreonine antibodies preferentially detected Rad18 that was immunoprecipitated from genotoxin-treated cells (Figure 1B). Moreover, the DNA damage-induced phosphorylation of Rad18 was ablated in Chk1-depleted cells (Figure 1B), further consistent with DNA damage-inducible Rad18 phosphorylation. The genotoxin-inducible electrophoretic mobility shifts and anti-phosphoserine/threonine immunoreactivity were evident in both soluble and chromatin-associated pools of Rad18 (Figure 1C). In metabolic labeling experiments with radiolabeled orthophosphate, immunoprecipitated Rad18 from genotoxin-treated cells contained basal levels of incorporated ^{32}P , which were further increased following acquisition of DNA damage (Figure 1D). We conclude that Rad18 is phosphorylated basally and in a DNA damage-inducible manner.

JNK phosphorylates Rad18 at S409

In a global screen for phosphoproteins, Nousiainen *et al.* (2006) detected phosphorylation of Rad18 at serine 409, although they did not explore the mechanism or significance of this phosphorylation event. We considered the possibility that S409 phosphorylation might contribute to the DNA damage-inducible changes in total Rad18 phosphorylation detected in Figure 1. Therefore we raised phosphospecific antiserum against the phosphopeptide CFSQSKLD[pS]PEEL, corresponding to residues 398–413 of hRad18 (see *Materials and Methods*). As shown in Figure 2B, the anti-Rad18 pS409 antibody detected a UV-inducible species in anti-Rad18 immunoprecipitates. The S409-reactive protein was not detected in Rad18 immunoprecipitates from Chk1-depleted

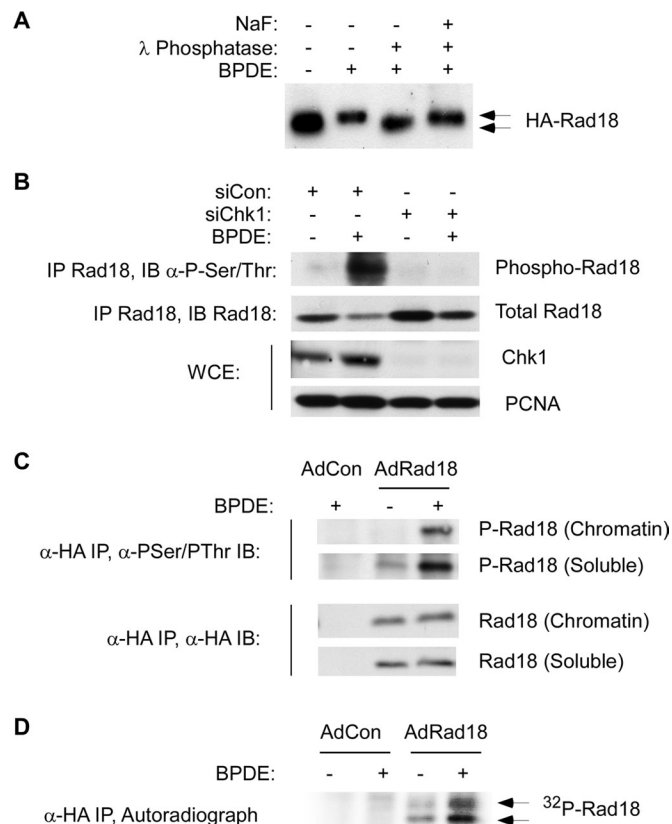


FIGURE 1: Rad18 phosphorylation is genotoxin inducible and requires Chk1. (A) HA-Rad18-expressing H1299 cells were treated with 600 nM BPDE for 4 h or were left untreated for controls. CSK extracts from the cells were incubated with λ phosphatase (1 U/ μ l) in the presence or absence of 50 mM NaF at 37 C for 30 min. The resulting extracts were analyzed by immunoblotting with anti-HA antibodies. BPDE-induced electrophoretic mobility shifts are indicated by the arrows. (B) Control or Chk1-depleted HA-Rad18-expressing H1299 cells were treated with 600 nM BPDE for 4 h or were left untreated for controls. Anti-HA immune complexes were analyzed by SDS-PAGE and immunoblotting sequentially with anti-phosphoserine/phosphothreonine and anti-HA antibodies. Whole-cell extracts were analyzed for Chk1 expression, and PCNA was used as a loading control. (C) AdCon or AdHA-Rad18-infected H1299 cells were treated with 600 nM BPDE for 4 h or were left untreated for controls. Chromatin and soluble fractions from the cells were immunoprecipitated with anti-HA antibodies, and the resulting immunoprecipitates were analyzed by SDS-PAGE and immunoblotting with anti-phosphoserine/phosphothreonine and anti-HA antibodies. (D) AdCon or AdHA-Rad18-infected H1299 cells were transferred to phosphate-free DMEM and incubated for 4 h in the presence of 0.2 mCi/ml of ^{32}P -orthophosphate. The ^{32}P -orthophosphate-labeled cells were treated with 600 nM BPDE for 4 h or were left untreated for controls. Whole-cell extracts from the cells were immunoprecipitated with anti-HA antibodies, and the resulting immunoprecipitates were resolved by SDS-PAGE. The resulting gels were washed extensively in 40% methanol/10% acetic acid. The fixed gel was dried, and radioactive proteins were detected by autoradiography.

cells (Figure 2B). Of importance, the anti-Rad18 pS409 antibody did not react with a Rad18 mutant harboring an S \rightarrow A substitution at residue 409 (see later discussion of Figure 4), demonstrating phosphospecificity. We conclude that S409 is a DNA damage-inducible Rad18 phosphorylation site.

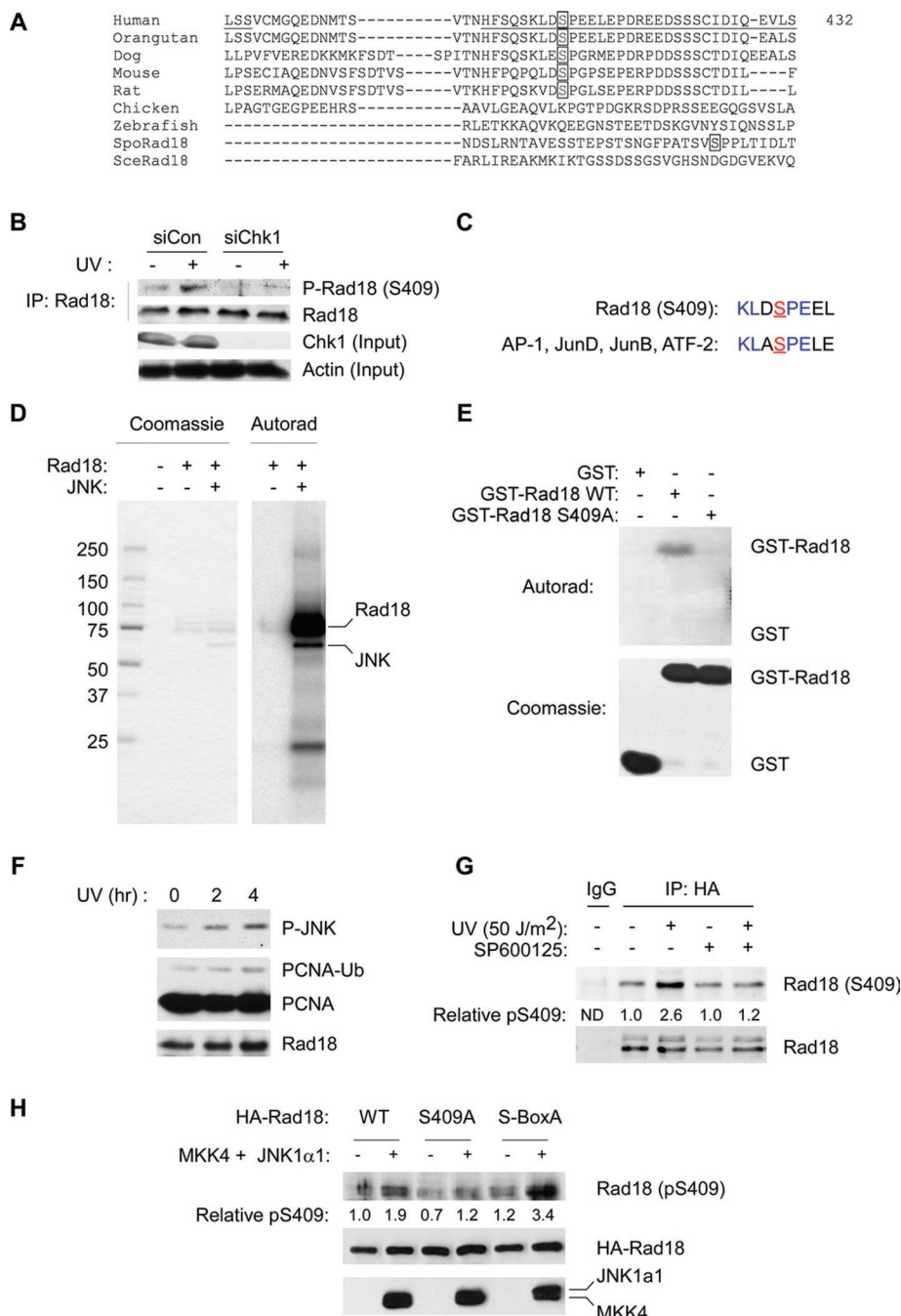


FIGURE 2: Rad18 S409 is phosphorylated by JNK. (A) Amino acid sequence alignment showing conservation of S409 among different species. (B) Control or Chk1-depleted HA-Rad18-expressing H1299 cells were treated with UV (20 J/m²) or left untreated for controls. After 4 h, anti-HA immune complexes were prepared from chromatin fractions and analyzed by SDS-PAGE and immunoblotting with anti-phospho-Ser-409 and anti-HA antibodies. Soluble extracts were analyzed for Chk1 expression to confirm efficient siRNA-mediated knockdown. Immunoblotting with an anti-actin antibody was used to verify equivalent protein loading between lanes. (C) Comparison of sequences flanking Rad18 S409 (underlined) with JNK1 consensus phosphorylation sites. (D) Recombinant Rad18–Rad6 complex was subject to in vitro phosphorylation by JNK. The reaction contained 0.4 μg (4 pmol) of Rad18–Rad6, and half (0.2 μg; 2 pmol) was resolved on SDS-PAGE. The phosphorylated species contained 4000 cpm (determined by scintillation counting of the excised gel slice), which is equivalent to 2 pmol of incorporated ATP incorporation. (E) Recombinant GST-Rad18 398–495 (WT), GST-Rad18 S409A, and GST (control) were tested as JNK substrates using in vitro kinase assays. Reaction products were resolved on SDS-PAGE. The resulting gel was fixed, stained, and dried. Top, an autoradiogram; bottom, the Coomassie-stained proteins. (F) Exponentially growing H1299 cells were irradiated with UVC (20 J/m²) or sham irradiated for controls. Cells were harvested 2 or 4 h

Because Rad18 S409 phosphorylation was Chk1 dependent, we considered the possibility that Chk1 phosphorylated this site directly. Therefore we performed in vitro kinase assays in which we tested recombinant baculovirus–encoded Rad18–Rad6 as well as bacterial glutathione S-transferase (GST)–Rad18 as potential Chk1 substrates. In those experiments, Rad18 was phosphorylated very inefficiently compared with Cdc25C, a bona fide Chk1 substrate. The modest Chk1-mediated in vitro phosphorylation of recombinant Rad18 was unaffected by S409A mutation (unpublished data). Furthermore, the conserved sequences flanking Rad18 S409 do not correspond to the preferred substrate motifs for Chk1 (O'Neill *et al.*, 2002). Therefore we conclude that Rad18 S409 is not a direct Chk1 target.

Because S409 precedes a proline residue, we hypothesized that cyclin-dependent kinases (CDKs) and/or SAPKs, representing the two major families of proline-directed protein kinases (Miller *et al.*, 2008), were likely candidate kinases responsible for Rad18 S409 phosphorylation. SAPKs (but not CDKs) are activated by DNA-damaging agents such as UV. Therefore we considered it most likely that SAPK member(s) mediate Rad18 S409 phosphorylation. Using in vitro kinase assays with recombinant CDK1 and CDK2, we were unable to detect Rad18 S409 phosphorylation, and we conclude that CDKs are most likely

after UVC treatment and lysates were analyzed for pJNK and PCNA by SDS-PAGE and immunoblotting. (G) Exponentially growing H1299 cells expressing HA-Rad18 were treated for 1 h with 1 μM SP600125 or left untreated for controls. Control and SP600125-treated cells were treated with UVC (20 J/m²) or sham irradiated. After 2 h, Rad18 was immunoprecipitated from cell extracts. Immune complexes were resolved by SDS-PAGE and probed sequentially with antibodies against Rad18 pS409 and HA. (H) H1299 cells were cotransfected with expression vectors encoding HA-Rad18 (WT), HA-Rad18 (S409A), FLAG-MKK4, and FLAG-JNK1α1 or with “empty” vector control plasmid as indicated. At 48 h posttransfection, cells were lysed and normalized for protein content. Lysates were immunoprecipitated with anti-HA, and the resulting immune complexes were resolved by SDS-PAGE, transferred to nitrocellulose, and probed sequentially with anti-Rad18 pS409 and anti-HA antibodies. Total cell lysates were also resolved and transferred to nitrocellulose and then probed with anti-FLAG to validate MKK4 and JNK expression.

not relevant protein kinases for this site. We noticed that the residues surrounding Rad18 S409 were very similar to the consensus target sites for c-Jun N-terminal kinase (Figure 2C). Indeed, full-length Rad18–Rad6 complex was phosphorylated very efficiently by JNK1 α 1 in vitro (Figure 2D). Under our standard experimental conditions ~1 mol of ATP was incorporated per mole of Rad18, indicating that Rad18 phosphorylation by JNK is very efficient.

To identify the Rad18 region(s) phosphorylated by JNK, we expressed a series of partially overlapping GST fusion fragments of Rad18 spanning the entire length of the Rad18 protein (Day *et al.*, 2010). The various GST-Rad18 fragments (comprising residues 1–98, 1–121, 110–173, 165–251, 247–295, 267–402, and 398–495) were tested as possible in vitro substrates for JNK. Of the Rad18 fragments, only GST-Rad18 398–495 (comprising the C-terminal 103 amino acids of Rad18) was phosphorylated by JNK (Figure 2E). A mutant form of GST-Rad18 398–495 harboring a serine \rightarrow alanine substitution at S409 (GST-Rad18 S409A) was not phosphorylated by JNK (Figure 2E). Therefore, although Rad18 harbors multiple SP sites (including S158 and S164), JNK phosphorylates Rad18 very specifically at S409. In similar experiments, p38 (another SAPK family member) did not phosphorylate Rad18 in vitro (unpublished data). We conclude that Rad18 phosphorylation by JNK is relatively specific (although we have not yet performed an exhaustive analysis of Rad18 phosphorylation by other SAPKs, and it is possible that additional JNK-related protein kinases contribute to Rad18 phosphorylation at S409).

Because the results of Figure 2 identified a JNK phosphorylation site in very close proximity to previously described Dbf4-dependent kinase (DDK) phosphorylation sites (including S421–434; Day *et al.*, 2010), we asked whether Rad18 S409 phosphorylation by JNK influenced its phosphorylation by DDK (or vice versa). Therefore we performed kinase assays in which we measured phosphorylation of recombinant Rad18–Rad6 complex by JNK and DDK individually or in combination. In those experiments, no synergistic enhancement or inhibition of phosphorylation was observed (for either substrate) when we coinubated DDK and JNK (unpublished data). Similar results were obtained when we performed these kinase assays using a synthetic peptide substrate corresponding to Rad18 residues 393–437 (unpublished data). We conclude that there is no phosphoprimering or hierarchical sequence of Rad18 phosphorylation by JNK and DDK and that S409 and S-box phosphorylations are independent events.

We performed experiments to determine whether JNK contributes to Rad18 S409 phosphorylation in intact cells. The kinetics of JNK phosphorylation and PCNA ubiquitination in UV-treated cells were similar, indicating a temporal correlation between JNK signaling and TLS (Figure 2F). Therefore we determined the effect of JNK inhibition on Rad18 S409 phosphorylation. As shown in Figure 2G, acute (1 h) pretreatment with the pharmacological JNK inhibitor SP600125 ablated UV-induced phosphorylation of Rad18 at S409. The apparent SP600125-resistant basal S409 phosphorylation may represent unmodified Rad18 (detected due to the incomplete discrimination between unphosphorylated and phosphorylated epitopes observed for many commonly used commercial antibodies). Alternatively, there are many SAPKs, and it is already known that there is considerable overlap in substrates among different SAPK members. For example, in their recent study Cook and colleagues found that both JNK and p38 contributed to phosphorylation of Cdt1 (Chandrasekaran *et al.*, 2011). Nevertheless, the reduced UV-dependent Rad18 S409 phosphorylation in SP600125-treated cells indicates that JNK contributes significantly to Rad18 phosphorylation at this site. We cannot exclude the possibility that

other protein kinases also target S409 for phosphorylation either basally or following genotoxin treatment.

JNK may be activated in a DNA damage and stress-independent manner by ectopic expression of active forms of its upstream kinase mitogen-activated protein kinase kinase 4 (MKK4). Therefore we determined the effects of coexpressing MKK4 and JNK on Rad18 S409 phosphorylation. As shown in Figure 2H, coexpression of MKK4 and JNK led to a 1.9-fold increase in Rad18 S409 phosphorylation, even in the absence of UV irradiation. As expected, MKK4/JNK-inducible phosphorylation of a Rad18 S409A mutant was not observed. We noticed that a DDK phosphorylation-resistant Rad18 mutant (designated Rad18 S-box-A) was phosphorylated at S409 more efficiently than Rad18 wild type (WT). These data may indicate that failure to phosphorylate the Rad18 S-box is associated with compensatory phosphorylation at S409. Taken together, the results of Figure 2 indicate that JNK is a Rad18-directed kinase that contributes to UV-inducible S409 phosphorylation.

Rad18 S409 phosphorylation specifically promotes Rad18–Pol η interactions

To determine the significance of Rad18 S409 phosphorylation, we compared the various activities of WT and S409A-mutant forms of Rad18. First we asked whether S409A substitution affected its association with chromatin. In cell fractionation experiments, there was no difference in association of ectopically expressed hemagglutinin (HA)-Rad18 (WT) and HA-Rad18 S409A with the detergent-insoluble chromatin (Supplemental Figure S1).

Rad18 contributes to TLS in part via its DNA damage-inducible interaction with Pol η , an association that helps chaperone the TLS polymerase to sites of replication stalling (Watanabe *et al.*, 2004; Day *et al.*, 2010). Because S409 resides in the Pol η -binding domain of Rad18, we considered the possibility that phosphorylation of this residue might influence associations with Pol η (Watanabe *et al.*, 2004).

To test this hypothesis, we expressed wild-type HA-Rad18 or HA-Rad18 S409A in RAD18^{-/-} HCT116 cells. Control and Rad18-expressing cells were treated with UV to induce DNA damage, and then Rad18 immunoprecipitates from solubilized chromatin extracts were analyzed by immunoblotting.

As expected, S409A substitution largely ablated the basal and DNA damage-induced phosphorylation of Rad18 and also inhibited the UV-inducible association between Rad18 and Pol η (Figure 3A). Phosphomimetic substitutions are sometimes conferred when serine or threonine sites are mutated to acidic residues. Therefore we generated an adenovirus vector encoding Rad18 harboring an S409 \rightarrow E substitution. However, basal and UV-inducible associations with endogenous Pol η were only very modestly increased in the Rad18 S409E mutant relative to WT Rad18 (Figure 3B), most likely because the glutamate residue is an imperfect chemical mimic of a phosphate moiety (unpublished data). Nevertheless, those data indicate that the inhibitory effect of the Rad18 S409A mutation on Pol η association is relatively specific. We conclude that Rad18 phosphorylation at S409 promotes formation of complexes containing Pol η .

To test whether S409 phosphorylation specifically affected Rad18–Pol η association, we determined the effect of the S409A mutation on additional Rad18 activities, including Rad6 association, PCNA ubiquitination, and ssDNA binding. Rad6 is an E2 ubiquitin-conjugating enzyme that binds and cooperates with Rad18 to ubiquitinate PCNA and other Rad18 substrates. In coimmunoprecipitation experiments, Rad18 S409A formed complexes with Rad6 as efficiently as WT Rad18 (Figure 4A). In subcellular fractionation studies, the binding of Rad18 S409A to chromatin was not impaired

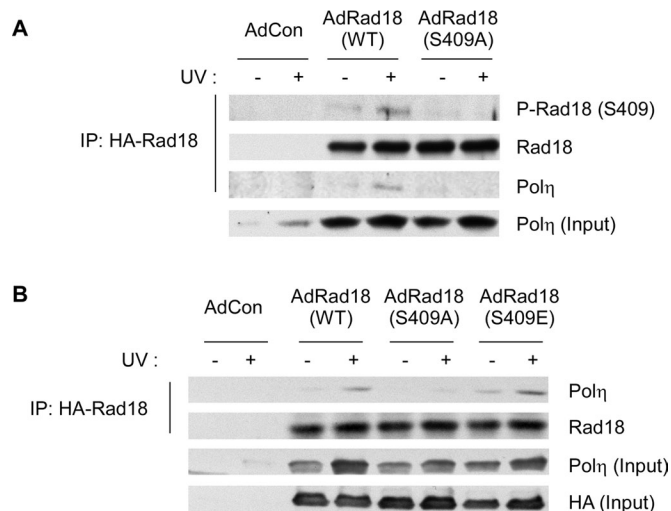


FIGURE 3: Rad18 S409 phosphorylation promotes association with Pol η . (A) *RAD18*^{-/-} cells were coinfectd with adenovirus vectors encoding HA-Rad18 WT or HA-Rad18 S409A or with an “empty” adenovirus vector (AdCon). Cells were treated with UV (20 J/m²) and harvested after 6 h. Chromatin fractions from the cells were solubilized and immunoprecipitated with anti-HA antibodies. The resulting immune complexes were resolved by SDS–PAGE and analyzed by immunoblotting with antibodies against Rad18 S409, HA, and Pol η . Total chromatin input fractions were also analyzed using anti-Pol η . (B) *RAD18*^{-/-} cells were infected with adenovirus vectors encoding HA-Rad18 WT, HA-Rad18 S409A, HA-Rad18 S409E, or with an “empty” adenovirus vector (AdCon). Cells were UV irradiated, and chromatin fractions were prepared and analyzed by SDS–PAGE and immunoblotting using antibodies against Rad18 and Pol η as described in A.

(Figure 4B), and ectopic expression of Rad18 S409A in *RAD18*^{-/-} cells induced DNA damage-independent PCNA ubiquitylation at least as efficiently as WT Rad18 (Figure 4B). It was suggested that the recruitment of Rad18 to stalled replication forks is mediated via its associations with RPA-coated ssDNA (Davies *et al.*, 2008; Tsuji *et al.*, 2008). Indeed, as shown in Figure 4C, WT HA-Rad18 (derived from *RAD18*^{-/-} cell extracts after complementation with Ad HA-Rad18) was recovered by streptavidin beads in an ssDNA- and RPA-dependent manner. Using *in vitro* pull-down assays, we detected no difference in RPA-ssDNA binding of HA-Rad18 S409A compared with WT HA-Rad18. Taken together, the results of Figure 4 indicate that S409 phosphorylation of Rad18 does not generally affect Rad18 activities. We conclude that the effect of S409 phosphorylation on associations between Rad18 and Pol η is relatively specific.

Rad18 S409 phosphorylation facilitates recruitment of Pol η to PCNA

We and others have shown that Rad18–Pol η interactions help guide the TLS polymerase to sites of replication fork stalling (Watanabe *et al.*, 2004; Day *et al.*, 2010). Because the results of Figure 3 showed that Rad18 S409A associates with Pol η inefficiently (compared with WT Rad18), we asked whether Rad18-induced formation of nuclear Pol η foci was affected by the S409A mutation. Therefore *RAD18*^{-/-} cells were complemented with WT Rad18 or Rad18 S409A.

We examined formation of yellow fluorescent protein (YFP)–Pol η foci in the Rad18-complemented cells using fluorescence microscopy. As expected, YFP–Pol η was distributed broadly and diffusely in the nuclei of Rad18-depleted H1299 cells, and this pattern was insensitive to UV treatment in the absence of Rad18 (Figure 5A). Com-

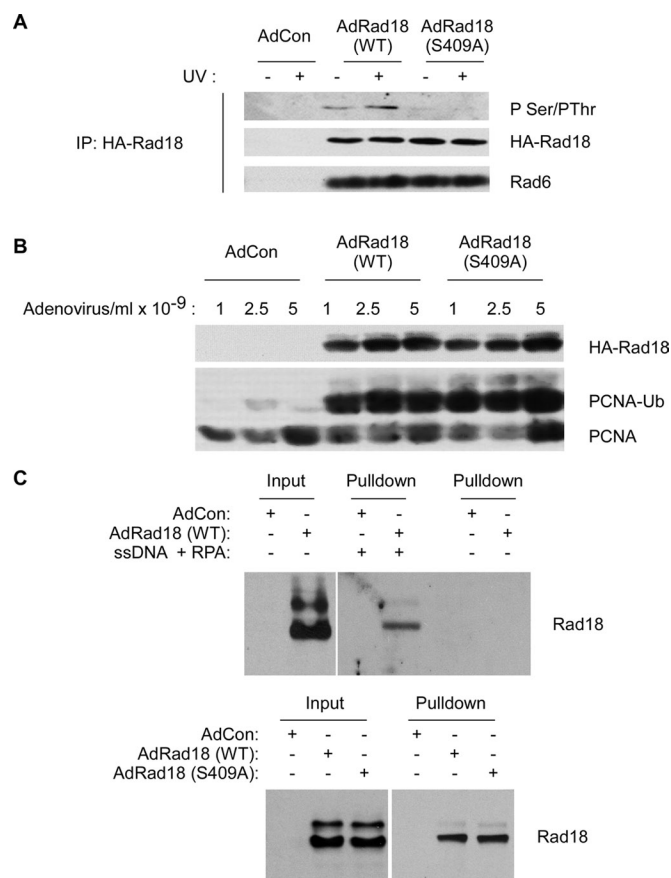
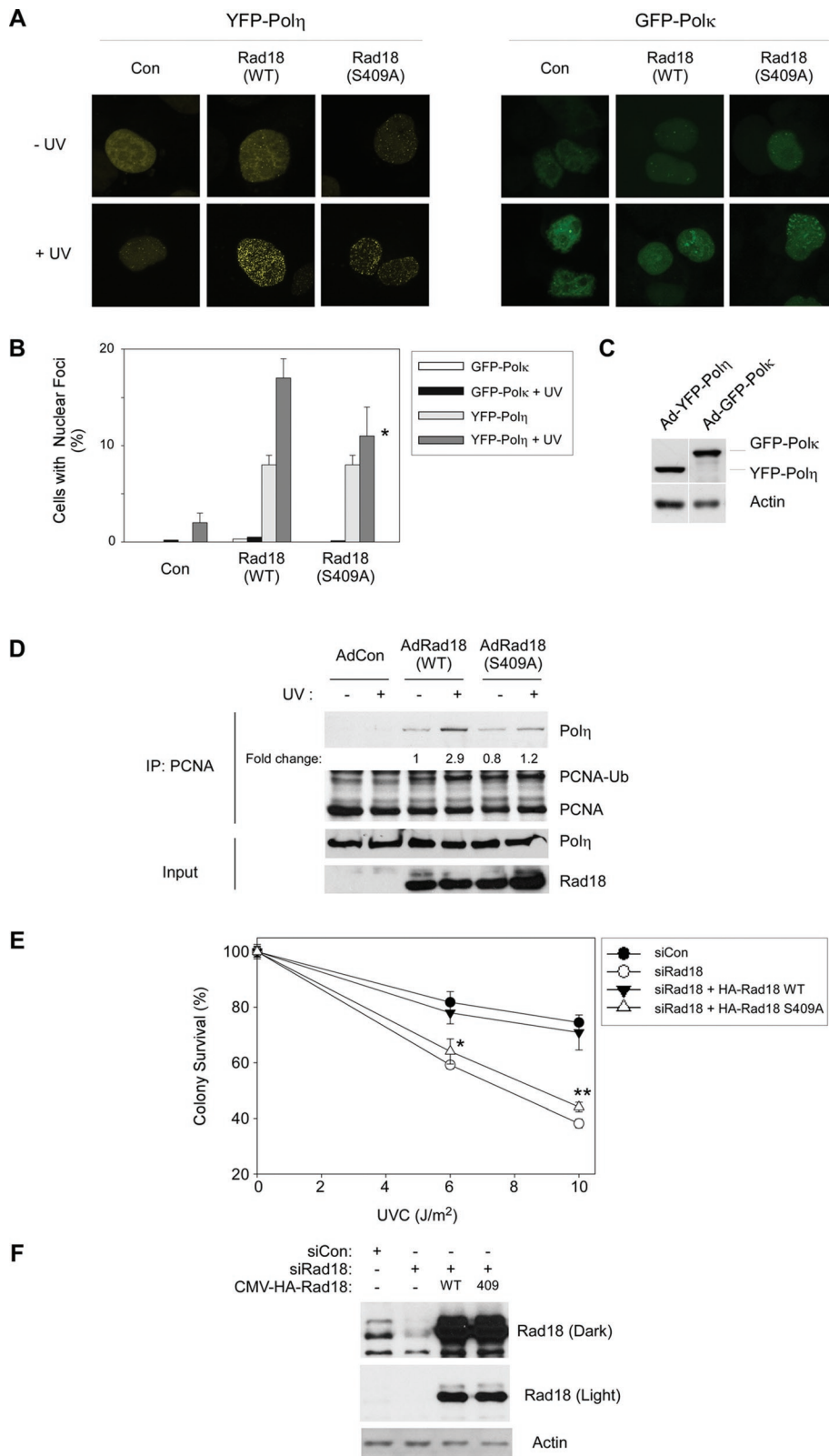


FIGURE 4: Rad18 S409 is dispensable for Rad18–Rad6 interactions, PCNA-directed E3 ligase activity, and binding to RPA-coated ssDNA. (A) HA-Rad18-expressing H1299 cells were treated with UV (20 J/m²) or left untreated for controls. Anti-HA immune complexes were prepared from chromatin fractions and analyzed by SDS–PAGE and immunoblotting with anti-HA and anti-Rad6 antibodies. (B) H1299 cells were infected with different doses of AdCon, AdRad18WT, and AdRad18 S409A. Twenty-four hours later, chromatin and soluble fractions were prepared and analyzed by SDS–PAGE and immunoblotting with HA and PCNA antibodies. (C) *RAD18*^{-/-} cells were infected with AdCon, AdRad18 WT, or Rad18 S409A. After 24 h, soluble extracts were prepared from the resulting cells. RPA/ssDNA-interacting proteins were captured as described in *Materials and Methods*. Proteins bound to the ssDNA-coated beads were analyzed by SDS–PAGE and immunoblotting. Top, the RPA/ssDNA-dependency of Rad18 binding to streptavidin-coated beads. Bottom, the comparison of WT Rad18 and Rad18 S409A binding to RPA/ssDNA-beads (and ssDNA and RPA were present in all pull-down samples).

plementation of Rad18-depleted cells with small interfering RNA (siRNA)–resistant WT Rad18 resulted in redistribution of YFP–Pol η and conferred a basal level of YFP–Pol η focus formation (Figure 5A).

In WT Rad18-expressing cells, the numbers of Pol η foci increased approximately twofold in response to DNA damage, as we and others previously reported (Kannouche *et al.*, 2004; Bi *et al.*, 2006). Of interest, however, Rad18 S409A failed to fully correct the UV-induced formation of Pol η foci in cells depleted of endogenous Rad18 (Figure 5B). In the experiment shown in Figure 5, A and B, the UV-inducible component of YFP–Pol η focus formation was reduced by ~66% ($p < 0.05$, $n = 3$) in Rad18 S409A-expressing cells compared with Rad18 WT-expressing cultures. For the purpose of comparison, in the same experiment we also determined the effects of Rad18 on



each treatment, number of cells positive for nuclear foci was expressed as a percentage of YFP/GFP-polymerase-positive cells (B). For GFP-Polκ- and YFP-Polη-expressing cells we performed analysis of variation (ANOVA) followed by Tukey's test to correct for experiment-wise error rates between multiple comparisons. For UV-irradiated cells coexpressing YFP-Polη and Rad18 WT or YFP-Polη and Rad18 S409A, the difference in number of foci was significant ($p < 0.05$). For GFP-Polκ-expressing cells under these experimental conditions there were no statistically significant differences between groups ($p > 0.05$). (C) We performed immunoblotting (with anti-GFP antibodies that recognize both GFP and YFP) to confirm that expression levels of GFP-Polκ and YFP-Polη were similar under these experimental conditions. (D) H1299 cells were infected with AdCon, AdRad18 WT, or AdRad18 S409A and treated with UV (20 J/m²) or left untreated for controls. PCNA was immunoprecipitated from the resulting cells, and immunoprecipitates (as well as appropriate input fractions) were analyzed by SDS-PAGE and immunoblotting. Relative levels of PCNA-associated Polη in the adjacent lanes were calculated after densitometric analysis of the Polη immunoblot. (E) H1299 cells were transfected with siRNA directed against the 3' untranslated region of endogenous Rad18 mRNA or with a nontargeting control siRNA, siCon. The resulting cells were transfected with CMV-driven mammalian expression plasmids encoding HA-Rad18 (WT), HA-Rad18 S409A, or "empty" pAC.CMV vector for control. Control and HA-Rad18-complemented cells were trypsinized and replated in replicate (five replicates per experimental condition) and then treated with varying doses of UV and analyzed for clonogenic survival as described in *Materials and Methods*. For each siRNA/complementation, the number of surviving colonies from UV-treated cultures was expressed as a percentage of colony number from unirradiated cells. On the survival curves, each data point represents the mean of five replicate determinations, and error bars represent the range. For each dose of UV, we performed ANOVA between groups followed by Tukey's multiple comparison of means test. For cells that received 6 or 10 J/m² of UV, in ANOVA the p value was < 0.0001 , which is significant. Results of the Tukey test were as follows: siCon vs. siRad18, $p < 0.001$ (indicating reduced UV tolerance); siCon vs. siRad18 + WT Rad18, $p > 0.05$ (indicating no significant difference and therefore full complementation by WT Rad18); siRad18 + Rad18 WT vs. siRad18 + Rad18 S409A, $p < 0.001$ (indicating significant difference in phenotype between Rad18 WT and Rad18 S409A, and indicated by asterisk and double asterisk for 6 and 10 J/m², respectively).

FIGURE 5: Rad18 S409A does not promote efficient recruitment of Polη to PCNA and fails to complement the UV sensitivity of Rad18-depleted cells. (A) Triplicate cultures of H1299 cells were depleted of endogenous Rad18 and complemented with siRNA-resistant HA-Rad18 (WT) and HA-Rad18 S409A. The Rad18 dependence of UV-induced YFP-Polη and GFP-Polκ redistribution to nuclear foci was determined as described in *Materials and Methods*. For each experimental condition (performed in triplicate) 60 cells were scored as nuclei containing > 20 TLS polymerase foci. Representative images of cells displaying YFP-Polη foci are shown in A. For

subcellular distribution of another Y-family TLS polymerase, namely Pol κ . In contrast with YFP-Pol η , the subcellular distribution of GFP-Pol κ (expressed at similar levels to that of YFP-Pol η as determined by immunoblotting experiments shown in Figure 5C) was largely unaffected by UV treatment, regardless of Rad18 expression. The relative insensitivity of GFP-Pol κ to UV irradiation is consistent with previous studies indicating that Pol κ does not participate significantly in bypass of CPD (Zhang *et al.*, 2000). We conclude that Rad18 phosphorylation at S409 specifically facilitates efficient recruitment of Pol η to sites of UV-induced replication fork stalling.

It was of interest to determine whether recruitment of endogenous Pol η to replication and repair foci was similarly dependent on Rad18 S409 phosphorylation. Unfortunately, however, we have been unable to detect endogenous levels of Pol η by immunofluorescence microscopy using available antibodies against Pol η . Therefore, in an alternative approach for studying recruitment of endogenous Pol η to replication forks, we investigated the Rad18-dependent association of endogenous Pol η with PCNA. In the experiment presented in Figure 5D, PCNA was immunoprecipitated from Ad HA-Rad18- or Ad HA-Rad18 S409A-infected cells. As shown in Figure 5B, UV treatment induced a 2.9-fold increase in the association of Pol η with PCNA in HA-Rad18 WT-expressing cells. In cells expressing HA-Rad18 S409A, UV only induced a 1.5-fold increase in the association of Pol η with PCNA. Taken together, the results of Figure 5, A and B, suggest that Rad18 phosphorylation at S409 promotes efficient Rad18-dependent recruitment of Pol η to sites of stalled replication.

Pol η recruitment to sites of DNA damage is considered important for DNA damage tolerance. Therefore we performed clonogenic survival assays to test whether impaired (Rad18 S409-dependent) recruitment of Pol η to replication forks is associated with reduced viability after UV treatment. For reasons that are not understood, the HCT116 RAD18^{-/-} cells used elsewhere in this study do not show UV sensitivity when compared with parental HCT116 RAD18^{+/+} cells (Shiomi *et al.*, 2007). Therefore we analyzed the role of Rad18 in UV tolerance in an alternative cell line. H1299 cells were selected for these studies because we found that Pol η is necessary for UV tolerance in this line. Moreover, Rad18 depletion fully phenocopies the UV sensitivity of Pol η -depleted H1299 cells (Supplemental Figure S2). We depleted H1299 cells of endogenous Rad18 (using siRNA) and then complemented with siRNA-resistant WT or mutant forms of Rad18. As expected, Rad18-depleted cells were moderately sensitive to UV, and UV tolerance was fully corrected by ectopically-expressed HA-Rad18 WT (Figure 5E). Of interest, HA-Rad18 S409A failed to correct the UV sensitivity of Rad18-depleted cells, thereby indicating a role for JNK-mediated Rad18 S409 phosphorylation in DNA damage tolerance via TLS (Figure 5E).

The UV sensitivity of the Rad18 S409A-complemented cells resembles the phenotype we previously reported for the Rad18 S-box-A mutant (which harbors DDK phosphorylation-site mutations at S421–423 and S434; Day *et al.*, 2010). It was of interest therefore to determine whether the DNA damage tolerance defects conferred by Rad18 S409 and S-box \rightarrow A mutations were additive or nonadditive. Therefore we performed UV survival assays in cells expressing Rad18 mutants harboring individual or combinatorial S \rightarrow A substitutions in the S-box serines and S409. As shown in Supplemental Figure S3, the survival curves for S409A, S-box \rightarrow A and S409, S-box \rightarrow A mutants were not significantly different. Therefore we conclude that independent phosphorylation events at S409 and S-box serine residues (mediated by JNK and DDK, respectively) are both necessary for Rad18 TLS functions and UV tolerance.

DISCUSSION

The interaction of Rad18 with Pol η represents one of several mechanisms that facilitate recruitment of Pol η to the replication machinery. It is generally believed that Rad18 guides Pol η to sites of stalled replication (Watanabe *et al.*, 2004) and that Rad18-mediated Pol η chaperone activity promotes stable engagement of Pol η with PCNA via PCNA-interacting protein (PIP) box and UBZ-mediated interactions (Bienko *et al.*, 2005). In this study we identified a novel DNA damage-inducible Rad18 phosphorylation site at the conserved residue Ser-409 that resides in the Pol η -binding domain of Rad18. We demonstrate that S409 phosphorylation is important for Rad18–Pol η association and confers more efficient chaperoning of Pol η to PCNA at sites of replication stalling. We previously identified a distinct cluster of DDK-dependent phosphorylation sites (serines 421–423 and 434 in the Pol η -binding domain) that similarly promote Pol η association and DNA damage tolerance (Day *et al.*, 2010). However, S409 is not phosphorylated by DDK (Day *et al.*, 2010), thereby distinguishing the new JNK-mediated pathway from the DDK-dependent mechanism that we previously described. Our finding that S409 and S-box phosphorylation-site mutations in Rad18

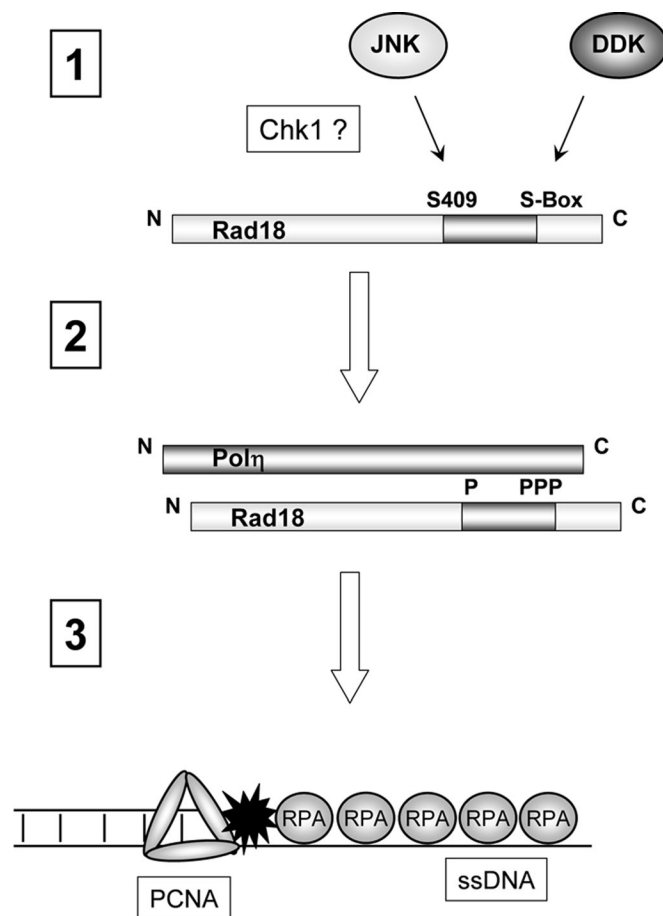


FIGURE 6: Hypothetical model showing roles of DDK and JNK in UV-inducible Rad18 phosphorylation and DNA damage tolerance. UV induces JNK-mediated phosphorylation of S409 and DDK-mediated phosphorylation of the S-box serine cluster. UV-induced phosphorylation of both S409 and the S-box is Chk1 dependent (step 1). S409 and the S-box reside in the Pol η -binding region of Rad18 (residues 401–445). S409 and S-box phosphorylations promote Rad18–Pol η interactions (step 2), thereby facilitating Rad18-dependent chaperoning of Pol η to sites of replication stalling, where the Rad18–Pol η complex associates with RPA-coated ssDNA (step 3).

confer similar and nonadditive phenotypes shows that efficient Rad18-mediated DNA damage tolerance requires independent JNK- and Cdc7-mediated phosphorylation of Rad18 (Figure 6).

To our knowledge, this study represents the first demonstration of a relationship between JNK signaling and TLS. The activation of JNK in response to replication stress-inducing agents such as UV radiation (Weston and Davis, 2007) and BPDE (Li *et al.*, 2004) has been studied extensively, yet few studies have explored the integration of SAPK and checkpoint signaling in the DNA damage response (Wang *et al.*, 2009). Our finding that JNK regulates TLS in a manner that is Chk1 dependent further demonstrates the intricate coordination of the various effector branches of the DNA damage response, including checkpoint signaling, SAPK cascades, and TLS.

JNK is widely viewed as a mediator of apoptotic responses, and it is perhaps counterintuitive that JNK may play a protective role in Rad18–Pol η recruitment and fork recovery. However, not all stimuli that activate JNK lead to apoptosis, perhaps because of active survival signaling pathways that prevent the apoptotic response (Xia *et al.*, 1995). In addition, the kinetics of JNK activation may determine whether or not there is an apoptotic outcome. For example, transient JNK activation (e.g., in response to cytokines) may mediate survival, whereas prolonged activation can mediate apoptosis (Ventura *et al.*, 2006). Although SAPK signaling has not previously been implicated in TLS or postreplication repair, recent studies by other labs have implicated JNK in control of DNA replication. Miotto and Struhl (2011) reported that JNK phosphorylates Cdt1 (a replication licensing factor) on T29, leading to dissociation of HBO1 from replication origins and blocking DNA replication licensing. In an independent study, Cook and colleagues identified multiple JNK and p38 target sites on Cdt1 the phosphorylation of which serves to rapidly inactivate Cdt1 and inhibit origin licensing (Chandrasekaran *et al.*, 2011). Therefore JNK helps to ensure appropriate DNA replication and genome maintenance in genotoxin-treated cells via at least two mechanisms: inhibition of origin licensing (Chandrasekaran *et al.*, 2011; Miotto and Struhl, 2011) and stimulation of TLS at sites of ongoing DNA replication (the present study). It will be interesting to determine whether additional proteins involved in other stages of DNA synthesis (e.g., initiation) or other modes of DNA repair are similarly subject to regulation by SAPKs.

Potentially consistent with additional roles for SAPKs in DNA replication and repair, Lannigan and colleagues demonstrated that extracellular signal-regulated kinase 8 (ERK8; a SAPK family member) contains a conserved PIP box that mediates PCNA binding (and subsequent PCNA turnover via Mdm2; Groehler and Lannigan, 2010). PCNA represents a central hub for numerous DNA repair and replication processes. Therefore recruitment of ERK8 and perhaps additional PIP box-containing mitogen-activated protein kinase (MAPK) family members to PCNA may provide the basis for crosstalk between MAPK signaling and DNA replication and repair events. By analogy, JNK2 was recently detected at RPA-coated ssDNA (a replication intermediate generated by DNA damage that acts as a proximal trigger of both the TLS and ATR/Chk1 pathways; Chen *et al.*, 2010). Therefore, similar to ERK8, JNK2 may be poised to regulate DNA replication and repair events in the vicinity of stalled replication forks. Further work is needed to investigate potential regulators and targets of the SAPKs at sites of DNA replication and repair.

Increasingly it is apparent that TLS is coordinated with other elements of the DNA damage response. JNK- and DDK-dependent Rad18 phosphorylation (of S409 and the S-box serines, respectively) is clearly Chk1 dependent (Day *et al.*, 2010), demonstrating further links between S-phase checkpoint signaling and TLS. The Chk1 de-

pendence of these Rad18 phosphorylation events may indicate that both JNK and DDK activities are influenced by Chk1 or that Chk1 inhibits phosphatase(s) that target the S409- and S-box-phosphorylated forms of Rad18. Alternatively, the effect of Chk1 status on Rad18 phosphorylation might be indirect, possibly resulting from Chk1-dependent stabilization of stalled forks, which promotes a state that is permissive for regulated changes in Rad18 phosphorylation. In preliminary experiments, Chk1 inhibition did not influence phosphorylation of other JNK or DDK substrates. Therefore we favor the hypothesis that Chk1 influences putative Rad18-directed protein phosphatases or that Chk1 promotes formation or stabilization of replication fork structures that facilitate Rad18 phosphorylation by its upstream kinases, including JNK and DDK. Further studies are needed to distinguish between these (non-mutually exclusive) mechanisms of Chk1 action.

There is a strong precedent for phosphorylation-dependent regulation of E3 ubiquitin ligases. For example, sustained (but not transient) JNK signaling mediates tumor necrosis factor α -induced cell death via the E3 ligase Itch (which degrades the caspase inhibitor cFLIPL; Chang *et al.*, 2006b). Another E3 ligase, Siah2, is regulated by p38-dependent phosphorylation (Khurana *et al.*, 2006). Rad18 plays a central role in TLS, DSB repair (Huang *et al.*, 2009; Palle and Vaziri, 2011), and perhaps additional DNA repair pathways. Therefore Rad18 phosphorylation has the potential to regulate multiple effector pathways, perhaps by influencing associations with its substrates (e.g., PCNA, RFC2, 53BP) or regulatory binding partners (e.g., Pol η , Rad6, Rad5). Rad18 phosphorylation may serve as a molecular switch for determining various physiological outcomes including but not limited to TLS. Unpublished mass spectrometry data from our lab and global proteomic profiling data from other labs suggest that Rad18 is phosphorylated on multiple sites (Nousiainen *et al.*, 2006). Further work is underway to validate additional Rad18 phosphorylation sites, test the physiological significance of these putative phosphorylation events, and identify the relevant kinases.

We also consider it likely that many other TLS proteins will be subject to regulation by phosphorylation. Rad6 is also phosphorylated by CDKs (Sarcevic *et al.*, 2002), and this modification may provide a mechanism for coordinating Rad6-dependent ubiquitination events with specific stages of the cell cycle. A study by Cleaver and colleagues suggested that Pol η recruitment to stalled replication forks is regulated via its direct phosphorylation, most likely by protein kinase C family members (Chen *et al.*, 2008). Lehmann and colleagues also demonstrated that Pol η is an ATR substrate whose phosphorylation modulates S-phase checkpoint signaling (Gohler *et al.*, 2011). Taken together, our results in this report and a previous study (Day *et al.*, 2010), together with studies from other labs (Chen *et al.*, 2008; Gohler *et al.*, 2011), indicate that multiple phosphorylation-based mechanisms facilitate the efficient recruitment of Pol η to sites of replication fork stalling. Further studies are underway to determine how protein kinases regulate Rad18 and integrate TLS (and perhaps other Rad18-dependent DNA repair processes) with the cell cycle.

MATERIALS AND METHODS

Adenovirus construction and infection

Adenovirus construction and infections were performed as described previously (Guo *et al.*, 2002; Bi *et al.*, 2006). In brief, cDNAs encoding HA-Rad18 WT, HA-Rad18 S409A, and HA-Rad18 S409E were subcloned into pAC-CMV. The resulting shuttle vectors were cotransfected into 293T cells with the pJM17 plasmid to generate recombinant adenovirus as described previously (Bi *et al.*, 2005). H1299 and HCT 116 cells were routinely infected with adenovirus at a multiplicity of infection of 20 and 50, respectively. To control for

adenoviral infection, cells received AdCon ("empty" adenovirus vector) or AdGFP.

Cell culture

Human lung carcinoma H1299 cells and *RAD18*^{-/-} HCT116 cells (Shiomi *et al.*, 2007) were cultured in DMEM supplemented with 10% fetal bovine serum, streptomycin sulfate (100 µg/ml), and penicillin (100 U/ml).

In vitro kinase assays

Recombinant Rad18–Rad6 complex from insect cells and GST–Rad18 from bacteria were expressed, purified, and tested as DDK and JNK substrates using in vitro phosphorylation reactions as described previously (Day *et al.*, 2010). Recombinant DDK was purified from insect cells as described previously (Day *et al.*, 2010), and recombinant JNK1α1 was purchased from Upstate (Millipore, Billerica, MA).

Genotoxin treatments

BPDE (National Cancer Institute Carcinogen Repository) was dissolved in anhydrous Me₂SO and added directly to the growth medium as a 1000× stock to give various final concentrations, as indicated in the figure legends. For UVC treatment, the growth medium was removed from the cells, reserved, and replaced with phosphate-buffered saline (PBS). The plates were transferred to a UV cross-linker (Stratagene, Santa Clara, CA) and then irradiated. The UVC dose delivered to the cells was confirmed with a UV radiometer (UVP BioImaging Systems, Upland, CA). The reserved medium from the cells was replaced, and cells were returned to the incubator.

RNA interference

Cells were plated into 60- or 100-mm culture dishes. Twenty-four hours later, when cells were 50–75% confluent, the cultures were placed in antibiotic-free medium (2 or 5 ml per plate). siRNA transfections were performed using Lipofectamine, using protocols recommended by the manufacturer (Invitrogen, Carlsbad, CA). After overnight transfections, the Lipofectamine-containing medium was removed and replaced with standard culture medium. All tubes, tips, and solutions used for RNA interference experiments were certified RNase free.

Clonogenic survival assays

H1299 cells were grown to ~30–50% confluence in 60-mm plates and transfected with 4 µg of pAC.CMV GFP vector (for controls) or with pAC.CMV expression vectors encoding wild-type or mutant Rad18 using Lipofectamine. Transfected cells were treated with UVC (as described) and then split into replicate 10-cm plates at a density of 1000 cells/plate. After 10–14 d, colonies on the plates were fixed in methanol, stained with crystal violet, and counted as described previously (Day *et al.*, 2010).

Fluorescence microscopy

H1299 cells at near-confluency were transfected with siRNA targeting the 3' untranslated region of Rad18. Ten hours later, the cells were split into glass-bottomed dishes (MatTek Corporation, Ashland, MA) to give a density of ~50% after 24 h. The cells were then infected with adenovirus encoding Rad18, YFP–Polη, or GFP–Polκ (or "empty" vector control). Six hours later, the virus-infected cells were transfected one more time with siRNA against Rad18. Twenty hours after the second siRNA transfection, the cells were sham or UV irradiated (10 J/m²). Two hours after UV treatment, the cells were rinsed with cold PBS, immersed briefly (<5 s) in PBS containing 0.1% Triton

X-100, rinsed again in ice-cold PBS, and then fixed in PBS containing 4% formaldehyde for 5 min. Fixed nuclei were 4',6-diamidino-2-phenylindole stained and mounted with Vectashield solution (Vector Laboratories, Burlingame, CA). Slides were imaged and analyzed using a Zeiss 710 Laser Scanning Microscope at 63× magnification with digital zoom (Zeiss, Jena, Germany). We collected 0.5 µm z-sections of representative cells from different experimental conditions and deconvolved them to generate the images presented.

Immunoblotting

Cells were separated into cytosolic and nucleosolic or chromatin fractions using CSK buffer as described previously (Bi *et al.*, 2005; Liu *et al.*, 2006). The resulting samples were normalized for protein content, then separated by SDS–PAGE, transferred to nitrocellulose, and analyzed by immunoblotting with the following antibodies: rabbit anti-Chk1 (Santa Cruz Biotechnology, Santa Cruz, CA), mouse monoclonal anti-PCNA (Santa Cruz Biotechnology), rabbit anti-phospho-Chk1 Ser-317 (Cell Signaling, Beverly, MA), polyclonal HA tag antibody (Santa Cruz Biotechnology), polyclonal PCNA antibody (Santa Cruz Biotechnology), polyclonal Polη (Bethyl Laboratories, Montgomery, TX), GFP (Invitrogen Molecular Probes, Eugene, OR), and Rad18 (Bethyl Laboratories). Phospho–serine/threonine antibodies were from BD Biosciences (San Diego, CA). Rad18 S409 phosphospecific antibodies were generated by immunizing rabbits with the peptide FSQSKLD[pS]PEELEPC. To ensure phosphospecificity, the resulting sera were depleted using unmodified immunogen and purified using the phosphopeptide. Peptide synthesis, immunizations, and antibody purification steps were performed by 21st Century Biochemicals (Marlboro, MA).

Coimmunoprecipitation of Polη with PCNA and Rad18

Cells were plated in 10-cm culture dishes and infected with adenovirus as described. Genotoxin treatments were performed at ~70% confluence. In some experiments, 2% formaldehyde was added for 5 min to cross-link chromatin-associated proteins before cell lysis. Cross-linking reactions were quenched by the addition of 1 M glycine for 5 min before PBS washes and CSK lysis. Samples containing chromatin fractions were generated as described and were incubated for 30 min at room temperature with 1 unit/µl RQ DNase (Promega, Madison, WI) and then sonicated (for three 15-s pulses at 30% maximum output). Pulses were separated by a 10-s interval on ice to prevent excessive heating. The sonicated samples were clarified by centrifugation at 10,000 × g for 5 min. Supernatants were removed and normalized for protein concentration (~600 µg of protein in 1 ml was used for each immunoprecipitation). PCNA or HA–Rad18 was immunoprecipitated overnight at 4°C using anti-PCNA or anti-HA monoclonal antibodies. Replicate immunoprecipitations were performed using immunoglobulin G to control for specificity of protein–protein associations. After antibody incubations, 25 µl pf protein A/G beads were added to each sample for 4 h. The beads were recovered by brief centrifugation and washed five times with 1 ml CSK (5–10 min per wash). The washed immune complexes were boiled in protein loading buffer for 10 min to release and denature immunoprecipitated proteins before separation on SDS–PAGE.

ssDNA-binding assays

The binding of Rad18 to ssDNA was detected as described by Tateishi and colleagues (Tsuji *et al.*, 2008). To prepare ssDNA-coated beads, 0.1 ml of TetraLink Tetrameric Avidin Resin (Promega) was washed twice with 1 ml of buffer A (50 mM Tris–HCl, pH 7.6, 1 mM dithiothreitol). Washed beads were suspended in 50 µl of buffer A. We mixed 10 µl of 10 µM 5'-biotinylated oligonucleotides, whose

nucleotide sequences are described by Tsuji *et al.* (2008), with the beads and incubated them for 15 min at room temperature. In parallel reactions beads were prepared without addition of biotinylated oligonucleotides, and these were subsequently used to control for the ssDNA dependence of binding reactions. We added 1 ml of a solution containing 3% skim milk, 20 mM Tris-HCl (pH 7.5), and 150 mM NaCl to the beads and incubated the mixture for 20 min at room temperature. The beads were then collected by centrifugation, washed once with 1 ml of buffer A, and resuspended in 0.3 ml of buffer A (DNA-beads solution). Bovine serum albumin was added to the DNA-beads solution to give a final concentration of 0.1 mg/ml. We removed 20- μ l aliquots of the resulting mixture and added them to fresh microfuge tubes. RPA, 0.5 μ M, was added to each 20- μ l aliquot of ssDNA-coated beads. We mixed 100- μ l aliquots of Rad18-containing CSK extracts (normalized to a protein concentration of 1 μ g/ μ l) with the RPA-coated beads and incubated them for 10 min at 37°C. The binding reactions were then diluted by addition of 1 ml of buffer A. The beads containing bound Rad18 were collected by centrifugation (1000 \times g, 5 min) and washed three times in buffer A. After the final wash the packed beads were resuspended in 50 μ l of buffer A containing 0.4 M NaCl and incubated for 5 min at room temperature. The samples were centrifuged for 5 min at 10,000 \times g. A 20- μ l amount of the supernatant was analyzed by SDS-PAGE and immunoblotting with anti-Rad18 antibodies.

Reproducibility

All data shown are representative of experiments that were repeated at least three times, with similar results on each occasion.

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