

Two Distinct Modes of ATR Activation Orchestrated by Rad17 and Nbs1

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<http://dx.doi.org/10.1016/j.celrep.2013.04.018>

SUMMARY

The ATM- and Rad3-related (ATR) kinase is a master regulator of the DNA damage response, yet how ATR is activated toward different substrates is still poorly understood. Here, we show that ATR phosphorylates Chk1 and RPA32 through distinct mechanisms at replication-associated DNA double-stranded breaks (DSBs). In contrast to the rapid phosphorylation of Chk1, RPA32 is progressively phosphorylated by ATR at Ser33 during DSB resection prior to the phosphorylation of Ser4/Ser8 by DNA-PKcs. Surprisingly, despite its reliance on ATR and TopBP1, substantial RPA32 Ser33 phosphorylation occurs in a Rad17-independent but Nbs1-dependent manner *in vivo* and *in vitro*. Importantly, the role of Nbs1 in RPA32 phosphorylation can be separated from ATM activation and DSB resection, and it is dependent upon the interaction of Nbs1 with RPA. An Nbs1 mutant that is unable to bind RPA fails to support proper recovery of collapsed replication forks, suggesting that the Nbs1-mediated mode of ATR activation is important for the repair of replication-associated DSBs.

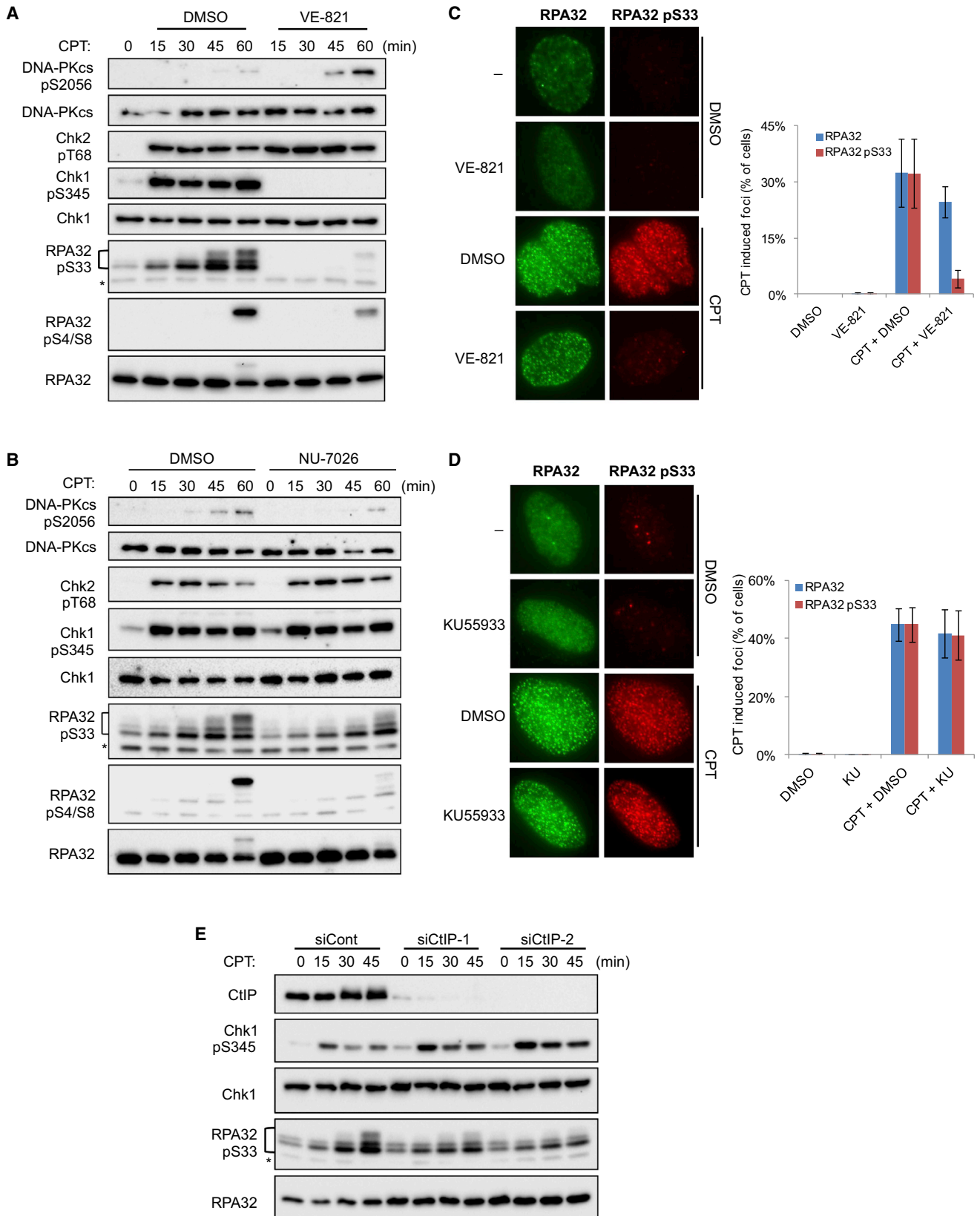
INTRODUCTION

The ability of cells to sense and signal DNA damage in their genomes is crucial for genomic stability. In human cells, the *ataxia telangiectasia* mutated (ATM) and the ATM- and Rad3-related (ATR) kinases are two master regulators of DNA damage signaling (Ciccia and Elledge, 2010). ATM, ATR, and their related DNA-dependent protein kinase catalytic subunit (DNA-PKcs) belong to the PI3K-like kinase (PIKK) family. Whereas ATM and DNA-PKcs are primarily activated by DNA double-stranded breaks (DSBs), ATR responds to a broad spectrum of DNA damage (Cimprich and Cortez, 2008; Flynn and Zou, 2011). Unlike ATM and DNA-PKcs, ATR is essential for cell survival even in the absence of extrinsic DNA damage, underscoring the critical function of ATR in coping with intrinsic genomic stress (Barlow et al., 2013; Brown and Baltimore, 2000; Murga et al., 2009; Toledo et al., 2011). Although the DNA-damage specificities

and functions of ATM, ATR, and DNA-PKcs are clearly distinct, how they distinguish different types of DNA damage and execute their unique functions is still poorly understood. In particular, it is largely unknown how ATR is activated by different types of DNA damage and replication stress.

Studies in different organisms have revealed some of the key principles of ATR activation. In response to DNA damage and replication stress, the complex of ATR and its functional partner, ATR-interacting protein (ATRIP), is recruited to sites of DNA damage and stalled replication forks by replication protein A (RPA)-coated single-stranded DNA (RPA-ssDNA) (Byun et al., 2005; Costanzo et al., 2003; Zou and Elledge, 2003). The activation of ATR-ATRIP requires additional regulators, such as the Rad17-RFC complex, the Rad9-Rad1-Hus1 (9-1-1) complex, and TopBP1 (Kumagai et al., 2006; Lin et al., 2012; Navadgi-Patil and Burgers, 2009; Zou et al., 2002). Independently of the recruitment of ATR-ATRIP to RPA-ssDNA, the Rad17-RFC complex recognizes the junctions of ssDNA and double-stranded DNA (dsDNA) and loads 9-1-1 complexes onto dsDNA (Ellison and Stillman, 2003; Zou et al., 2003). Through a process that is still not fully understood, TopBP1 is recruited to damaged DNA and interacts with Rad17, 9-1-1, and autophosphorylated ATR (Cotta-Ramusino et al., 2011; Delacroix et al., 2007; Lee and Dunphy, 2010; Lee et al., 2007; Liu et al., 2011; Wang et al., 2011; Yan and Michael, 2009). The engagement of TopBP1 with ATR-ATRIP allows TopBP1 to stimulate the kinase activity of ATR and to facilitate ATR to recognize its substrates (Kumagai et al., 2006; Liu et al., 2011; Mordes et al., 2008). In this model of ATR activation, ATR is activated by Rad17 and TopBP1 around ssDNA/dsDNA junctions. Indeed, checkpoint kinase 1 (Chk1), an effector kinase of ATR that is critical for the replication stress response and cell-cycle arrest, is phosphorylated by ATR in a Rad17-, TopBP1-, and ssDNA/dsDNA-junction-dependent manner (Liu et al., 2006; MacDougall et al., 2007; Van et al., 2010; Yamane et al., 2003; Zou et al., 2002). However, it is important to note that although Chk1 phosphorylation has been widely used as a surrogate for ATR activation, it remains unclear whether Chk1 phosphorylation accurately evinces the active mode of ATR in all situations.

In this study, we asked whether ATR is always activated by the Rad17-TopBP1 circuitry after DNA damage occurs. In particular, we wondered whether ATR is activated by Rad17 and TopBP1 at extensively resected DSBs, such as those generated in S phase



(legend on next page)

at collapsed replication forks. When long ssDNA is generated at DSBs by resection, a fraction of ATR could be recruited to the RPA-ssDNA distal to ssDNA/dsDNA junctions, raising the question as to whether and how this fraction of ATR is activated on RPA-ssDNA. To address this question, we analyzed the activation of ATR by camptothecin (CPT), which induces replication-associated DSBs that undergo rapid and efficient resection (Avemann et al., 1988; Sartori et al., 2007). We found that ATR is activated in two distinct modes toward Chk1 and RPA32. In one mode, ATR phosphorylates Chk1 rapidly, whereas in the other mode, ATR phosphorylates RPA32 Ser33 progressively during resection. The activation of ATR toward RPA32 is driven by resection and requires TopBP1. Surprisingly, Nbs1, a component of the Mre1-Rad50-Nbs1 (MRN) complex (Carney et al., 1998; Costanzo et al., 2001; Difilippantonio et al., 2005; Stracker and Petrini, 2011), plays a more important role than Rad17 in the phosphorylation of RPA32. The function of Nbs1 in RPA32 phosphorylation can be separated from ATM activation and DSB resection, and is dependent upon a direct interaction between Nbs1 and RPA. An Nbs1 mutant that is unable to bind RPA is compromised in its ability to support the recovery of collapsed replication forks. Together, these results suggest that Nbs1 mediates a TopBP1-dependent but Rad17-independent mode of ATR activation on RPA-ssDNA, allowing ATR to phosphorylate substrates such as RPA32 independently of ssDNA/dsDNA junctions and promote repair of replication-associated DSBs.

RESULTS

ATR Phosphorylates Chk1 and RPA32 Ser33 via Distinct Mechanisms

ATR is known to be activated by the replication-associated DSBs induced by CPT (Avemann et al., 1988; Sartori et al., 2007). Using VE-821, a specific ATR inhibitor (Reaper et al., 2011), we confirmed that a number of ATR substrates, including Chk1, Rad17, RPA32, and ATR itself, were phosphorylated in an ATR-dependent manner after CPT treatment (Figure S1A). In contrast to these ATR substrates, ATM and DNA-PKcs underwent efficient autophosphorylation in the presence of VE-821 (Figure S1B). Furthermore, the phosphorylation of several ATM and/or DNA-PKcs substrates, including Chk2, p53, and H2AX, was not affected by VE-821 (Figure S1B). These results confirm that VE-821 specifically inhibits ATR, but not ATM and DNA-PKcs, in CPT-treated cells.

In response to CPT, RPA32 is phosphorylated at multiple sites, including Ser4/Ser8, Thr21, and Ser33 (Anantha et al., 2007;

Block et al., 2004; Sartori et al., 2007). Using phosphospecific antibodies, we found that RPA32 Ser33 was phosphorylated within 15 min after CPT treatment, whereas RPA32 Ser4/Ser8 was phosphorylated at ~60 min (Figure 1A). Furthermore, the phosphorylation of RPA32 Ser33 was abolished by VE-821, whereas RPA32 Ser4/Ser8 phosphorylation was only partially reduced (Figure 1A). The phosphorylation of RPA32 Ser4/Ser8 was shown to be DNA-PKcs dependent (Anantha et al., 2007; Liaw et al., 2011; Liu et al., 2012). Indeed, the CPT-induced phosphorylation of RPA32 Ser4/Ser8 was abolished by the DNA-PKcs inhibitor NU7026 (Figure 1B). In contrast, RPA32 Ser33 phosphorylation was not affected by NU7026 (Figure 1B). Thus, compared with the phosphorylation of RPA32 Ser4/Ser8, the phosphorylation of RPA32 Ser33 is a more direct and reliable marker for ATR activation.

The phosphorylation of RPA32 has been linked to DSB resection (Sartori et al., 2007). A recent study suggested that ATR promotes resection in *Xenopus* extracts (Peterson et al., 2013). Consistent with this finding, at 1 hr after CPT treatment, the formation of RPA32 foci was reduced by VE-821 (Figure 1C). Nevertheless, a significant fraction of VE-821-treated cells displayed RPA32 foci, suggesting that ATR is not essential for resection (Figure 1C). Importantly, the majority of VE-821-treated cells that displayed RPA32 foci were negative for phospho-Ser33 foci, suggesting that ATR is required for RPA32 Ser33 phosphorylation after resection (Figure 1C). In contrast to VE-821, the ATM inhibitor KU55933 drastically diminished the foci of both RPA32 and phospho-Ser33 1 hr after CPT treatment (Figure S1C). At 2 hr, KU55933-treated cells no longer displayed a significant reduction in RPA32 foci and phospho-Ser33 foci (Figure 1D), showing that inhibition of ATM delays resection transiently. In KU55933-treated cells, RPA32 foci were always colocalized with phospho-Ser33 foci (Figure 1D), suggesting that ATM does not have a direct role in RPA32 Ser33 phosphorylation post resection. Consistent with RPA32 Ser33 phosphorylation being a specific marker of ATR activation, this phosphorylation event was clearly detected in the ATM-deficient AT cells and the DNA-PKcs-deficient M059J cells, even when they were treated with NU7026 and KU55933, respectively (Figures S1D and S1E).

Having established RPA32 Ser33 phosphorylation as a marker for CPT-induced ATR activation, we asked whether this phosphorylation event is regulated in the same way as Chk1 phosphorylation. In response to CPT, Chk1 was rapidly phosphorylated at Ser345 in an ATR-dependent manner (Figure 1A). In contrast to the phosphorylation of RPA32 Ser33, which

Figure 1. ATR-Mediated RPA32 Ser33 Phosphorylation in Response to CPT-Induced DSBs

(A) U2OS cells were treated with 1 μ M CPT in the presence or absence of 10 μ M VE-821 and collected at the indicated times. The levels of the indicated proteins and the CPT-induced phosphorylation events were analyzed by western blot; *a protein cross-reacting to the phospho-Ser33 antibody.

(B) Cells pretreated with 10 μ M NU-7026 or DMSO for 1 hr were exposed to 1 μ M CPT in the presence or absence of 10 μ M NU-7026. Extracts were analyzed as in (A).

(C) Cells were treated with 1 μ M CPT for 1 hr, or mock treated, in the absence or presence of VE-821. Immunofluorescence analysis of RPA32 and phospho-Ser33 was performed with specific antibodies. Representative images of cells are shown in the left panel. The fractions of cells that displayed RPA32 or phospho-Ser33 foci are shown in the right panel. Error bars: SDs from three independent experiments (n = 3).

(D) Cells were treated with 1 μ M CPT for 2 hr, or mock treated, in the absence or presence of KU55933. Immunofluorescence analysis was performed and quantified as in (C). Error bars: SDs from three independent experiments (n = 3).

(E) U2OS cells transfected with control or CtIP siRNAs were treated with 1 μ M CPT and collected at the indicated time points. Extracts were analyzed as in (A). See also Figure S1.

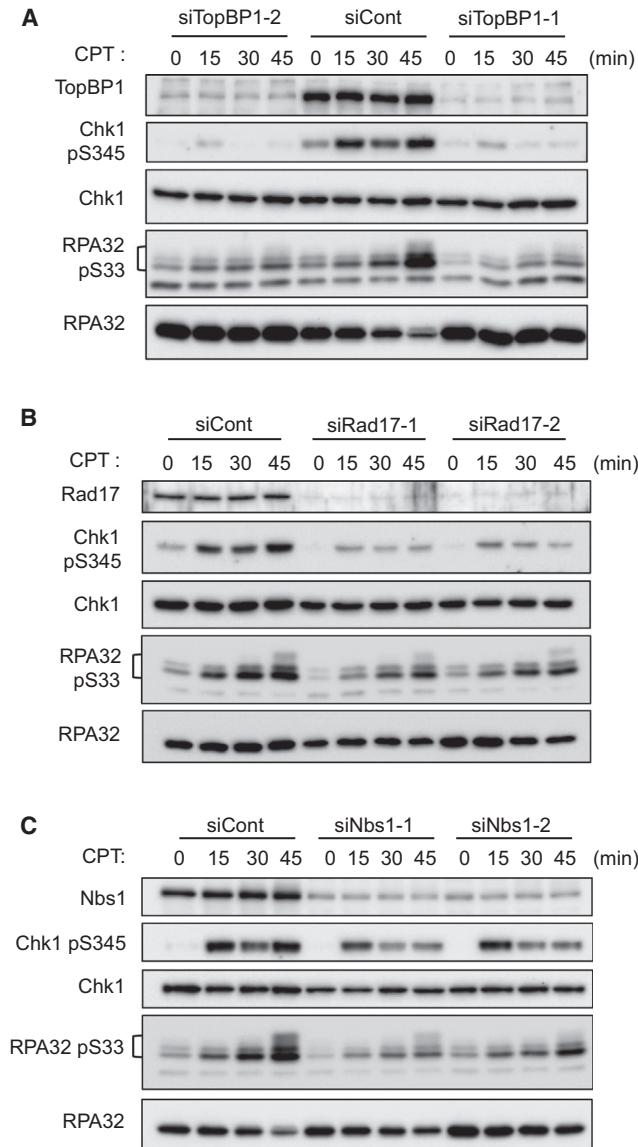


Figure 2. RPA32 and Chk1 Are Phosphorylated by ATR through Distinct Mechanisms

(A–C) U2OS cells transfected with control, TopBP1 (A), Rad17 (B), or Nbs1 (C) siRNAs were treated with 1 μ M CPT and collected at the indicated time points. The levels of the indicated proteins and the CPT-induced phosphorylation events were analyzed by western blot.

See also Figure S2.

gradually accumulated during the first 60 min after CPT treatment, Chk1 phosphorylation plateaued within 15 min (Figure 1A). Thus, Chk1 and RPA32 Ser33 are phosphorylated by ATR with distinct kinetics. Consistent with the sequential phosphorylation of RPA32 by ATR and DNA-PKcs (Anantha et al., 2007), the phosphorylation of RPA32 Ser4/Ser8 by DNA-PKcs occurred significantly later than Ser33 phosphorylation and Chk1 phosphorylation (Figure 1A; Kousholt et al., 2012).

The phosphorylation of RPA32 Ser4/Ser8 by DNA-PKcs requires CtIP, which is critical for resection (Sartori et al., 2007).

Prior to the abrupt phosphorylation of Ser4/Ser8, Ser33 was gradually phosphorylated by ATR (Figure 1A), suggesting that Ser33 phosphorylation may be the initial phosphorylation event on RPA32 that is driven by resection. Indeed, knockdown of CtIP with two independent small interfering RNAs (siRNAs) drastically reduced the phosphorylation of RPA32 Ser33 during the first 45 min after CPT treatment (Figure 1E). On the other hand, consistent with a recent report (Kousholt et al., 2012), depletion of CtIP did not significantly affect the phosphorylation of Chk1 during this early phase of the CPT response. Thus, in contrast to the rapid phosphorylation of Chk1, the phosphorylation of RPA32 Ser33 by ATR occurs progressively during resection.

Distinct Roles of Rad17 and Nbs1 in RPA Phosphorylation

The distinct kinetics and resection dependency of Chk1 and RPA32 phosphorylation prompted us to investigate whether ATR is activated toward these two substrates in different ways. To dissect the genetic requirements for RPA32 Ser33 phosphorylation, we used siRNAs to knock down the upstream regulators of the ATR pathway. Consistent with the effects of VE-821, knockdown of ATR abolished the phosphorylation of both Chk1 and RPA32 Ser33 (Figure S2A). Two independent siRNAs targeting TopBP1, the key activator of ATR, also drastically reduced the phosphorylation of Chk1 and RPA32 Ser33 (Figure 2A). Therefore, similar to Chk1 phosphorylation, RPA32 Ser33 phosphorylation requires both ATR and TopBP1.

TopBP1 and Rad17 function together in Chk1 phosphorylation (Delacroix et al., 2007; Lee and Dunphy, 2010). As expected, knockdown of Rad17 with two independent siRNAs significantly reduced Chk1 phosphorylation (Figure 2B). Surprisingly, however, substantial RPA32 Ser33 phosphorylation was detected in Rad17 knockdown cells (Figure 2B), suggesting the existence of a Rad17-independent mechanism of RPA32 Ser33 phosphorylation. The MRN complex has been implicated in ATR regulation and RPA phosphorylation (Jazayeri et al., 2006; Manthey et al., 2007; Myers and Cortez, 2006; Olson et al., 2007a; Stiff et al., 2005; Yoo et al., 2009; Zhong et al., 2005). Knockdown of Nbs1 with two independent siRNAs clearly reduced RPA32 Ser33 phosphorylation without altering cell-cycle distribution (Figures 2C and S2B). Consistent with Nbs1 knockdown, the Nbs1-deficient NBN-ILB1 cells displayed much lower RPA32 Ser33 phosphorylation than the Nbs1-complemented cells (Figure S2C). Compared with Rad17 knockdown, Nbs1 knockdown reduced RPA32 Ser33 phosphorylation to a greater extent. When both Rad17 and Nbs1 were knocked down, RPA32 Ser33 phosphorylation was further reduced (Figure S2D), suggesting that Rad17 and Nbs1 function in parallel to activate ATR toward RPA32 Ser33. Interestingly, although Nbs1 knockdown clearly reduced RPA32 Ser33 phosphorylation, its effects on Chk1 phosphorylation were modest (Figure 2C). Thus, in contrast to Rad17, Nbs1 plays a more important role in RPA phosphorylation than in Chk1 phosphorylation.

Nbs1 Mediates RPA32 Phosphorylation Independently of Rad17 and ssDNA/dsDNA Junctions

The MRN complex regulates ATM activation and DSB resection (Symington and Gautier, 2011; Uziel et al., 2003). Furthermore,

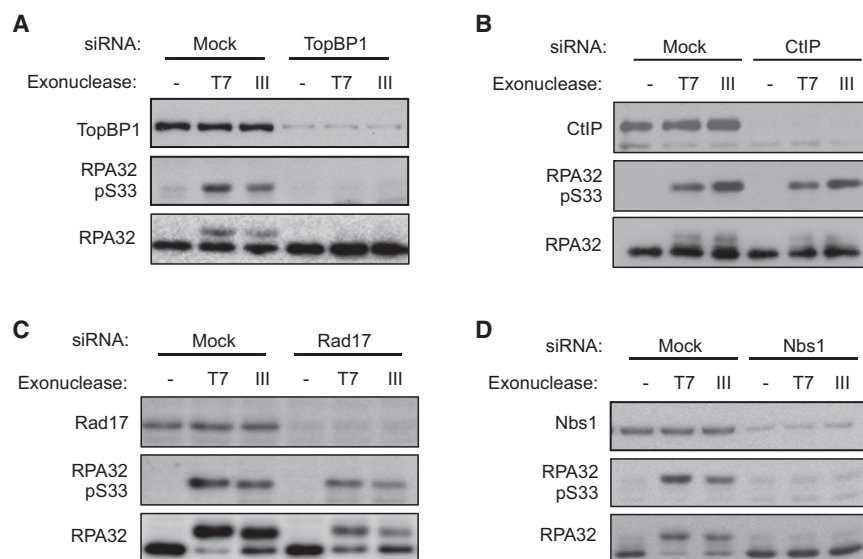


Figure 3. Nbs1 Regulates RPA32 Phosphorylation Independently of ATM Activation and DSB Resection

(A) Nuclear extracts were prepared from HeLa cells transfected with TopBP1 siRNA or mock transfected. A 6 kb dsDNA fragment was resected with T7 exonuclease or exonuclease III as described in the [Experimental Procedures](#). Resected or unresected dsDNA was added to cell extracts, and the resected-dsDNA-induced phosphorylation of RPA32 Ser33 was analyzed with phosphospecific antibody.

(B) The resected-dsDNA-induced phosphorylation of RPA32 Ser33 was analyzed in extracts of CtIP knockdown cells and control cells as in (A).

(C) The resected-dsDNA-induced phosphorylation of RPA32 Ser33 was analyzed in extracts of Rad17 knockdown cells and control cells as in (A).

(D) The resected-dsDNA-induced phosphorylation of RPA32 Ser33 was analyzed in extracts of Nbs1 knockdown cells and control cells as in (A).

the MRN complex is implicated in the removal of Top1 from CPT-induced DSBs ([Sacho and Maizels, 2011](#)). To determine whether Nbs1 regulates RPA32 phosphorylation directly or indirectly, we analyzed the function of Nbs1 in RPA32 phosphorylation using an *in vitro* assay that we recently developed ([Shiotani and Zou, 2009](#)). In this assay, linear dsDNA is resected by T7 exonuclease or exonuclease III to generate ssDNA overhangs, and subsequently is added to HeLa cell nuclear extracts to activate ATR ([Shiotani and Zou, 2011](#)). Using this assay, we have shown that RPA32 Ser33 is phosphorylated in an ATR-dependent manner. Since both ATM and DNA-PKcs are inhibited in extracts, and dsDNA is already resected before it is added to extracts, this assay provides a unique opportunity to test whether Nbs1 has an ATM-, resection-, and end-processing-independent function in RPA32 phosphorylation.

Consistent with its dependency on TopBP1 in cells, RPA32 Ser33 phosphorylation was not induced by resected dsDNA in extracts derived from TopBP1 knockdown cells ([Figure 3A](#)). In contrast, RPA32 Ser33 phosphorylation occurred efficiently in extracts from CtIP knockdown cells, confirming that the role of MRN-CtIP in DSB resection is bypassed by the use of pre-resected dsDNA in this assay ([Figure 3B](#)). Also consistent with the observation in cells, the phosphorylation of RPA32 Ser33 in extracts was only modestly reduced by Rad17 depletion ([Figure 3C](#)). Strikingly, even in this resection-independent *in vitro* assay, Nbs1 was clearly required for RPA32 Ser33 phosphorylation ([Figure 3D](#)). These results reveal a postresection function of Nbs1 in RPA32 phosphorylation.

Since substantial RPA32 Ser33 phosphorylation occurred independently of Rad17 *in vivo* and *in vitro*, we asked if this phosphorylation event is absolutely dependent upon ssDNA/dsDNA junctions. In nuclear extracts, ssDNA alone was sufficient to induce RPA32 Ser33 phosphorylation in a length-dependent manner ([Figure 4A](#)). We noted that unlike resected dsDNA, ssDNA did not induce a mobility shift of RPA32 in protein gels. The reason for this difference is currently unknown. Importantly,

the ssDNA-induced RPA32 Ser33 phosphorylation was dependent upon ATR, TopBP1, and Nbs1 ([Figures 4B–4D](#)), suggesting that a process driven by ssDNA lengthening and mediated by Nbs1 and TopBP1 is able to activate ATR toward RPA independently of ssDNA/dsDNA junctions.

Nbs1 Targets the MRN Complex to RPA-ssDNA

We next asked how Nbs1 executes its postresection function in ATR activation. The MRN complex interacts with RPA ([Oakley et al., 2009; Olson et al., 2007b](#)). Consistent with previous studies, we found that in extracts, Mre11, Rad50, and Nbs1 all bound to ssDNA in an RPA-dependent manner ([Figure 5A](#)). The RPA complex containing the RPA70 t-11 mutant, whose N-terminal oligonucleotide/oligosaccharide-binding (OB) fold was disrupted by point mutations, was unable to recruit MRN to ssDNA ([Figure 5B](#)). Furthermore, purified MRN complex bound to ssDNA in an RPA-dependent manner, showing that MRN associates with RPA-ssDNA directly ([Figure 5C](#)). Even in the absence of Mre11 and Rad50, purified Nbs1 bound to ssDNA in an RPA-dependent manner, suggesting a direct interaction between Nbs1 and RPA-ssDNA ([Figure 5D](#)).

To determine whether the recruitment of MRN to RPA-ssDNA is dependent on Nbs1, we generated extracts from Nbs1 knockdown cells. Consistent with previous studies, loss of Nbs1 did not affect the levels of Mre11 and Rad50 ([Figure 5E; Stewart et al., 1999](#)). However, in extracts of Nbs1 knockdown cells, neither Mre11 nor Rad50 bound to RPA-ssDNA efficiently ([Figure 5E](#)). In contrast to Mre11 and Rad50, ATRIP still associated with RPA-ssDNA efficiently in the absence of Nbs1 ([Figure 5E](#)). Together, these results suggest that Nbs1 targets the MRN complex to RPA-ssDNA.

Nbs1 Binds RPA-ssDNA via a Previously Uncharacterized Domain

To determine whether the interaction between Nbs1 and RPA-ssDNA is important for the phosphorylation of RPA32 Ser33 by

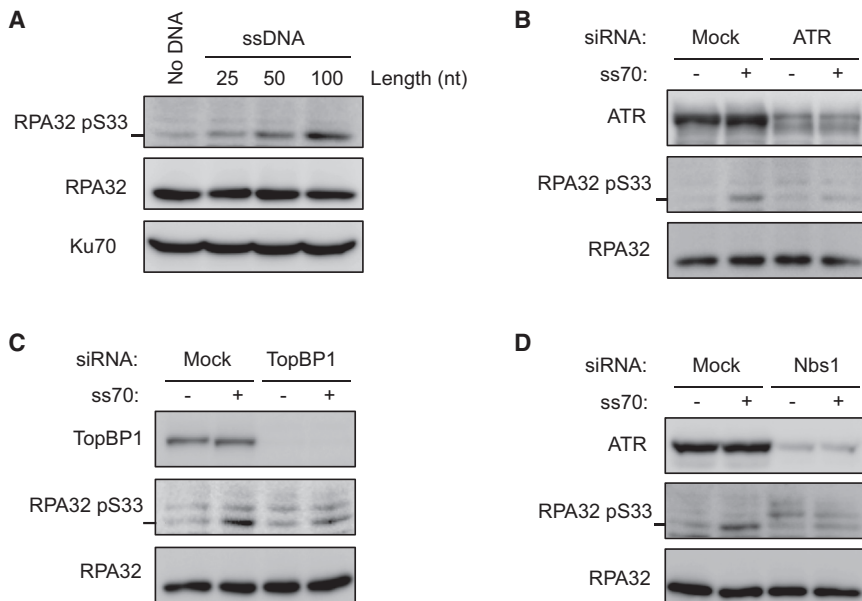


Figure 4. ssDNA Is Sufficient to Induce RPA32 Phosphorylation

(A) ssDNA of the indicated lengths was added to nuclear extracts, and the ssDNA-induced phosphorylation of RPA32 Ser33 was analyzed with phosphospecific antibody. (B) The ssDNA-induced phosphorylation of RPA32 Ser33 was analyzed in extracts of ATR knockdown cells and control cells as in (A). (C) The ssDNA-induced phosphorylation of RPA32 Ser33 was analyzed in extracts of TopBP1 knockdown cells and control cells as in (A). (D) The ssDNA-induced phosphorylation of RPA32 Ser33 was analyzed in extracts of Nbs1 knockdown cells and control cells as in (A).

ATR, we sought to map the RPA-ssDNA-interacting domain of Nbs1. We expressed a set of myc-tagged Nbs1 fragments in cells and tested them for binding to biotinylated ssDNA coated with RPA (Figures 6A, S3A, and S3B). An Nbs1 fragment containing the N-terminal FHA and BRCT domains (Np38) was unable to interact with RPA-ssDNA (Figure S3A). On the other hand, two Nbs1 fragments lacking the FHA and BRCT domains (Cp49 and Cp38) retained the ability to interact with RPA-ssDNA (Figure S3B). Moreover, an Nbs1 fragment lacking the C-terminal ATM and Mre11 binding domains (Np75) still bound to RPA-ssDNA (Figure S3A). Thus, the RPA-ssDNA-interacting domain of Nbs1 is distinct from its known functional domains.

Through testing the binding of additional Nbs1 fragments to RPA-ssDNA (Figures S3A and S3B), we found that all of the Nbs1 fragments that were able to bind RPA-ssDNA encompassed amino acids 536–567 (Figure 6A). Alignment of the sequences of Nbs1 from different species revealed that a stretch of charged amino acids in this region is conserved in higher vertebrates (Figure 6B). To pinpoint the residues that are critical for RPA-ssDNA binding, we mutated the conserved residues KKR, DD, and EDE to EEG, AA, and AAA in full-length Nbs1, respectively (Figure 6B; Nbs1^{KKR}, Nbs1^{DD}, and Nbs1^{EDE}). Only the Nbs1^{EDE} mutant, and not the Nbs1^{KKR} and Nbs1^{DD} mutants, lost the ability to bind RPA-ssDNA (Figure 6C). The Nbs1^{EDE} mutant still bound to Mre11 efficiently (Figure 6D), suggesting that loss of the EDE motif of Nbs1 specifically disrupts its binding to RPA-ssDNA.

The Recognition of RPA-ssDNA by Nbs1 Is Important for RPA32 Ser33 Phosphorylation and Recovery of Collapsed Forks

To determine whether the binding of Nbs1 to RPA-ssDNA is important for RPA32 Ser33 phosphorylation, we first tested the Nbs1^{EDE} mutant in vitro. We expressed siRNA-resistant wild-type myc-Nbs1 (myc-Nbs1^{WT}) and myc-Nbs1^{EDE} in cells that

were depleted of endogenous Nbs1 by siRNA and prepared extracts from these cells. In the myc-Nbs1^{WT}-containing extracts, resected dsDNA induced RPA32 Ser33 phosphorylation efficiently (Figure 7A). In contrast, RPA32 Ser33 phosphorylation was poorly induced by resected dsDNA in extracts containing myc-Nbs1^{EDE} (Figure 7A), suggesting that the ability of Nbs1 to recognize RPA-ssDNA is important for its postresection function in RPA32 Ser33 phosphorylation in vitro.

Next, we tested whether the Nbs1^{EDE} mutant is able to support RPA32 Ser33 phosphorylation in cells. We generated stable cell lines in which siRNA-resistant myc-Nbs1^{WT} or myc-Nbs1^{EDE} could be inducibly expressed. Upon induction of myc-Nbs1^{WT} in Nbs1 knockdown cells, RPA32 Ser33 phosphorylation was clearly induced by CPT (Figure 7B, lanes 3 and 4). In contrast, the levels of RPA32 Ser33 phosphorylation were not increased by CPT in cells expressing the myc-Nbs1^{EDE} mutant (Figure 7B, lanes 7 and 8). Thus, consistent with our in vitro experiments, the Nbs1^{EDE} mutant also fails to support RPA32 Ser33 phosphorylation in cells.

Finally, we sought to determine whether the recognition of RPA-ssDNA by Nbs1 is important for the DNA damage response. Because CPT induces replication-associated DSBs, we focused on the recovery of collapsed replication forks. We synchronized cells in S phase with thymidine and released them into CPT-containing media. As expected, γ H2AX foci were induced in cells regardless of whether endogenous Nbs1 was replaced by myc-Nbs1^{WT} or myc-Nbs1^{EDE} (Figure 7C). After the myc-Nbs1^{WT}-expressing cells were released from CPT, γ H2AX foci were largely gone in 24 hr (Figure 7C), indicating the recovery of collapsed forks. In contrast, a significant fraction of the myc-Nbs1^{EDE}-expressing cells retained γ H2AX foci 24 hr after the release from CPT (Figure 7C), suggesting that the recovery of collapsed forks was compromised. These results establish a functional link between Nbs1-mediated ATR activation and the repair of replication-associated DSBs.

DISCUSSION

Many investigators have studied the activation of ATR using Chk1 phosphorylation as a readout. Both in vivo and in vitro

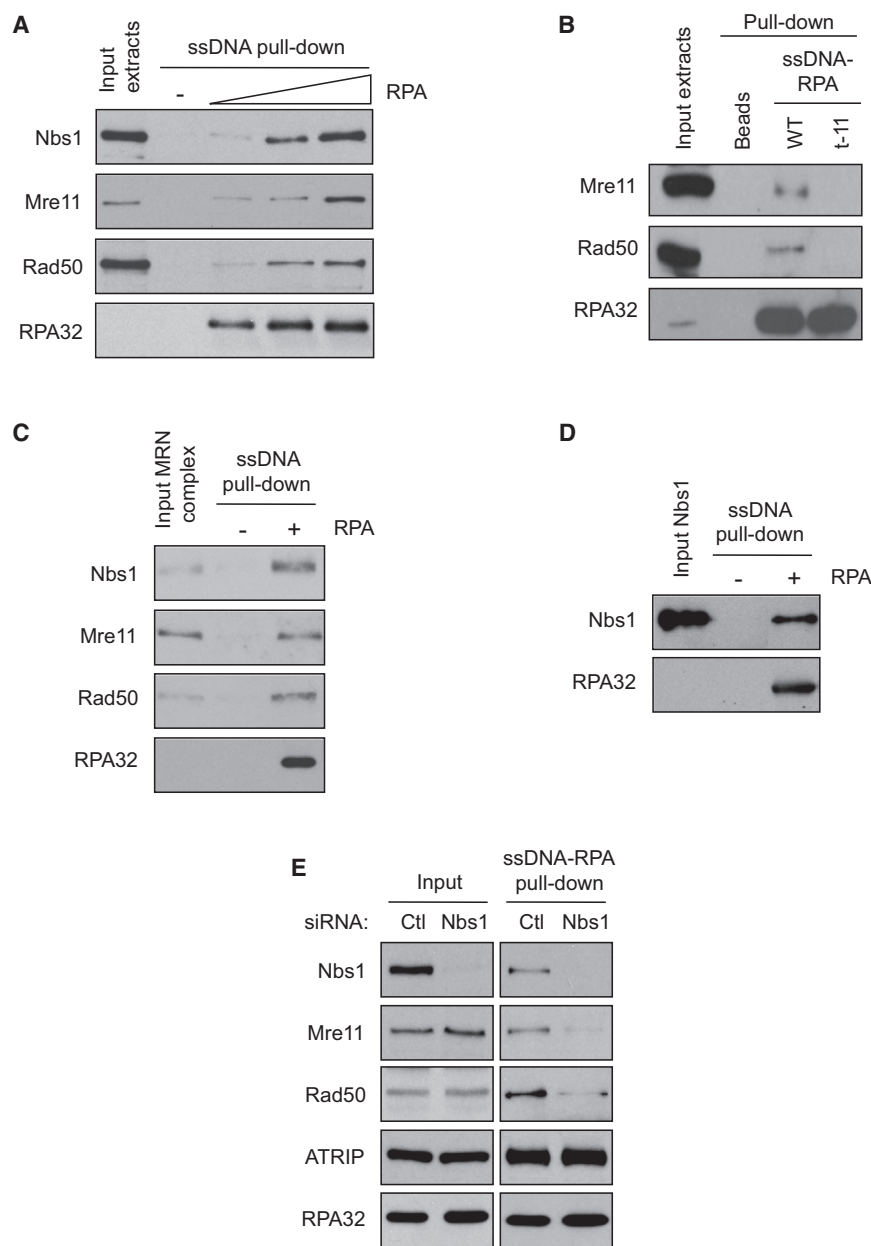


Figure 5. Nbs1 Targets the MRN Complex to RPA-ssDNA

(A) Biotinylated 70-nucleotide ssDNA (5 pmole) was coated with increasing amounts of purified RPA (0, 5, 10, and 20 pmole) and then incubated in cell extracts. The proteins bound to RPA-ssDNA were retrieved by streptavidin-coated beads and analyzed by western blot using the indicated antibodies. All components of the Mre11-Rad50-Nbs1 complex bound to ssDNA in an RPA-dependent manner.

(B) Biotinylated ssDNA was coated with either WT RPA or the RPA t-11 mutant complex purified from *E. coli*. The binding of Rad50 and Mre11 to RPA-ssDNA was tested as in (A).

(C) Biotinylated ssDNA coated with RPA was incubated with purified MRN complex. The MRN components bound to RPA-ssDNA were analyzed by western blot.

(D) Binding of purified Nbs1 to RPA-ssDNA was tested as in (C).

(E) Extracts were prepared from cells transfected with control or Nbs1-2 siRNA. Biotinylated ssDNA coated with RPA was incubated in the extracts, and the proteins bound to RPA-ssDNA were analyzed by western blot using the indicated antibodies.

the fact that ssDNA is not sufficient to trigger checkpoint signaling through Chk1 (Guo and Dunphy, 2000; MacDougall et al., 2007).

Although ssDNA and ssDNA/dsDNA junctions are commonly induced by DNA damage and replication stress, the relative abundance of these two DNA structures may vary in different contexts. For example, at replication forks stalled by aphidicolin, an increased number of primers and ssDNA/dsDNA junctions may be generated, which would enhance the activation of ATR toward Chk1 (Van et al., 2010). On the other hand, when DSBs undergo extensive resection in S phase, long stretches of ssDNA may be generated without increasing the number

of ssDNA/dsDNA junctions. We found that RPA32 Ser33 is progressively phosphorylated by ATR during the resection of CPT-induced DSBs. Although it is an ATR-mediated event, RPA32 Ser33 phosphorylation is regulated differently from Chk1 activation. In addition to its delayed kinetics and greater dependency on resection, RPA32 Ser33 phosphorylation is more dependent on Nbs1 but less dependent on Rad17 compared with Chk1 phosphorylation. Furthermore, unlike Chk1 phosphorylation, RPA32 Ser33 phosphorylation can occur on ssDNA in a length-dependent manner independently of ssDNA/dsDNA junctions. These findings suggest that, in a resection-driven manner, Nbs1 mediates a distinct mode of ATR activation toward RPA32.

experiments have suggested that Rad17 and TopBP1 are two key regulators of the activation of ATR toward Chk1 (Cimprich and Cortez, 2008; Flynn and Zou, 2011). In the current model of ATR activation, both RPA-ssDNA and ssDNA/dsDNA junctions are critical determinants of Chk1 phosphorylation (MacDougall et al., 2007). Whereas RPA-ssDNA presents a platform for ATR-ATRIP recruitment, the Rad17-RFC complex and 9-1-1 complexes are localized to ssDNA/dsDNA junctions. The colocalization of ATR-ATRIP, Rad17, 9-1-1, and TopBP1 around ssDNA/dsDNA junctions may create a protein-DNA assembly that allows TopBP1 to activate the ATR-ATRIP kinase complex and enable it to recognize its downstream substrates. This mechanism of ATR activation toward Chk1 is consistent with

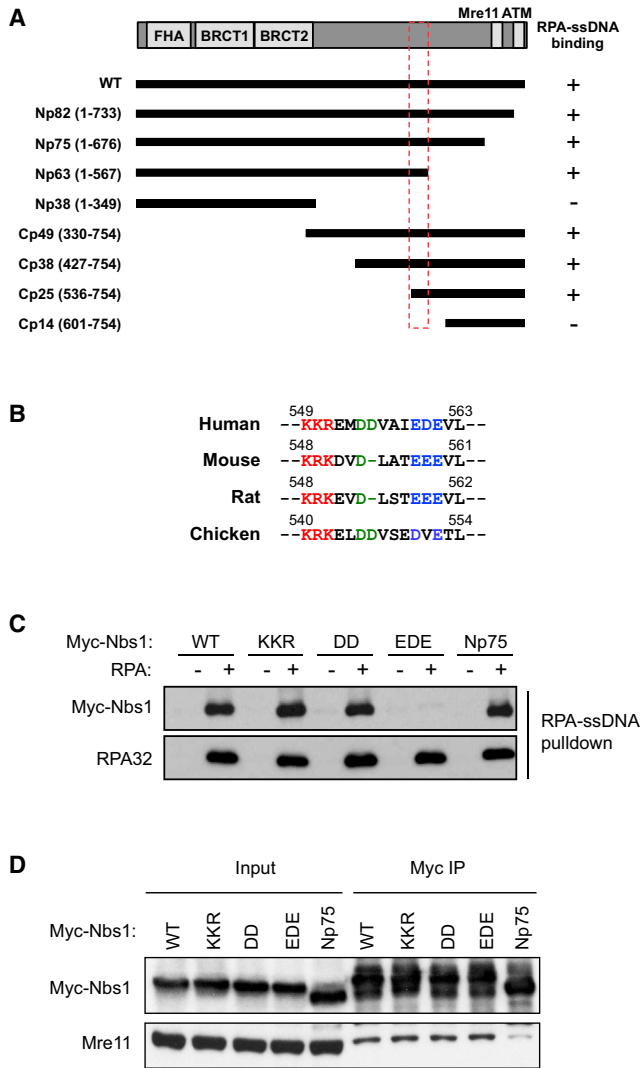


Figure 6. Mapping the RPA-ssDNA-Interacting Domain of Nbs1

(A) Schematic of the Nbs1 fragments used in this study and their ability to bind RPA-ssDNA.

(B) The RPA-ssDNA-binding domain of human Nbs1 is aligned to the corresponding regions in the Nbs1 of other higher vertebrates. The conserved charged amino acids are highlighted in colors.

(C) The conserved charged amino acids in the RPA-ssDNA-binding domain of Nbs1 were mutated in myc-tagged, full-length Nbs1 (KKR to EEG, DD to AA, and EDE to AAA). These Nbs1 mutants were transiently expressed in 293T cells, and their ability to bind RPA-ssDNA was tested in cell extracts using biotinylated ssDNA coated with RPA. WT Nbs1 and an Nbs1 mutant lacking the Mre11- and ATM-binding domains (Np75) were also tested as controls.

(D) Myc-tagged, full-length Nbs1 (WT) and its mutant derivatives (KKR, DD, EDE, and Np75) were transiently expressed in 293T cells. The myc-tagged Nbs1 proteins were immunoprecipitated with myc antibody, and the coprecipitated Mre11 was analyzed by western blot using Mre11 antibody.

See also Figure S3.

How do Rad17 and Nbs1 regulate the activation of ATR during resection? As resection is initiated at DSBs and ssDNA/dsDNA junctions are generated, the Rad17-RFC complex recognizes these junctions and loads 9-1-1 complexes onto dsDNA (Fig-

ure 7D). This Rad17- and ssDNA/dsDNA-junction-mediated process promotes ATR activation and Chk1 phosphorylation around the ssDNA/dsDNA junctions. In addition, a fraction of RPA adjacent to ssDNA/dsDNA junctions is phosphorylated by ATR in a Rad17-dependent manner during this early phase of resection. As resection continues, the ssDNA is gradually lengthened, and increasing amounts of RPA and ATR-ATRIP are placed on the ssDNA distal to ssDNA/dsDNA junctions (Figure 7D). In the late phase of resection, a fraction of RPA and ATR-ATRIP on ssDNA becomes “out of reach” for the Rad17 and 9-1-1 complexes at ssDNA/dsDNA junctions. In this situation, the MRN complex may directly recognize RPA-ssDNA via Nbs1, and activate ATR-ATRIP by recruiting TopBP1 (Yoo et al., 2009). Consistent with this model, substantial RPA32 Ser33 phosphorylation is driven by resection and is dependent upon both Nbs1 and TopBP1 in vivo and in vitro. Furthermore, the direct interaction between Nbs1 and RPA-ssDNA is needed for efficient RPA32 Ser33 phosphorylation both in vivo and in vitro. Since the Nbs1-mediated mode of ATR activation is independent of Rad17, which interacts with Claspin to promote Chk1 activation (Wang et al., 2006), this mode of ATR activation is specifically directed to RPA but not Chk1.

When and where does Nbs1 activate ATR in cells? In laser microirradiated cells, a fraction of Nbs1 precisely colocalizes with RPA in the ssDNA subcompartments at DSBs (Bekker-Jensen et al., 2006). In the context of DNA replication, Mre11 partially colocalizes with PCNA during S phase (Maser et al., 2001). Although RPA32 Ser33 phosphorylation is rapidly induced by CPT, it does not occur robustly in hydroxyurea (HU)-treated cells until DSBs become detectable in these cells (Figures 1A and S1F; Sakasai et al., 2006). This observation, together with the dependency of RPA32 Ser33 phosphorylation on resection, suggests that RPA32 Ser33 phosphorylation occurs primarily at replication-associated DSBs. Indeed, in previous studies, the phosphorylation of RPA32 was implicated in the progression and repair of stressed replication forks (Liu et al., 2012; Shi et al., 2010; Vassin et al., 2009). We found that when the interaction between Nbs1 and RPA-ssDNA was disrupted, the recovery of collapsed replication forks was compromised (Figure 7C). These findings suggest that the Nbs1-mediated mode of ATR activation may be particularly important for the repair of replication-associated DSBs. Consistent with this idea, the MRN complex was shown to colocalize with RPA at collapsed forks (Manthey et al., 2007). Furthermore, the MRN complex is required for preventing accumulation of DSBs during replication (Costanzo et al., 2001). In addition to the mechanism that we propose, recent studies suggested that Mre11 degrades stalled replication forks in the absence of BRCA1/2 and Fanconi anemia proteins, and promotes ATR activation via a nuclease-dependent mechanism at stalled forks (Lee and Dunphy, 2013; Schlacher et al., 2011, 2012). Another recent study showed that the *Xenopus* MRN complex recognizes the ssDNA/dsDNA junctions in synthetic DNA templates and recruits TopBP1 to DNA (Duursma et al., 2013). We note that the EDE motif of human Nbs1 is not conserved in *Xenopus* Nbs1, suggesting that the role of MRN in ATR activation may have expanded during evolution. The MRN complex may regulate the ATR response and protect replication forks through multiple mechanisms.

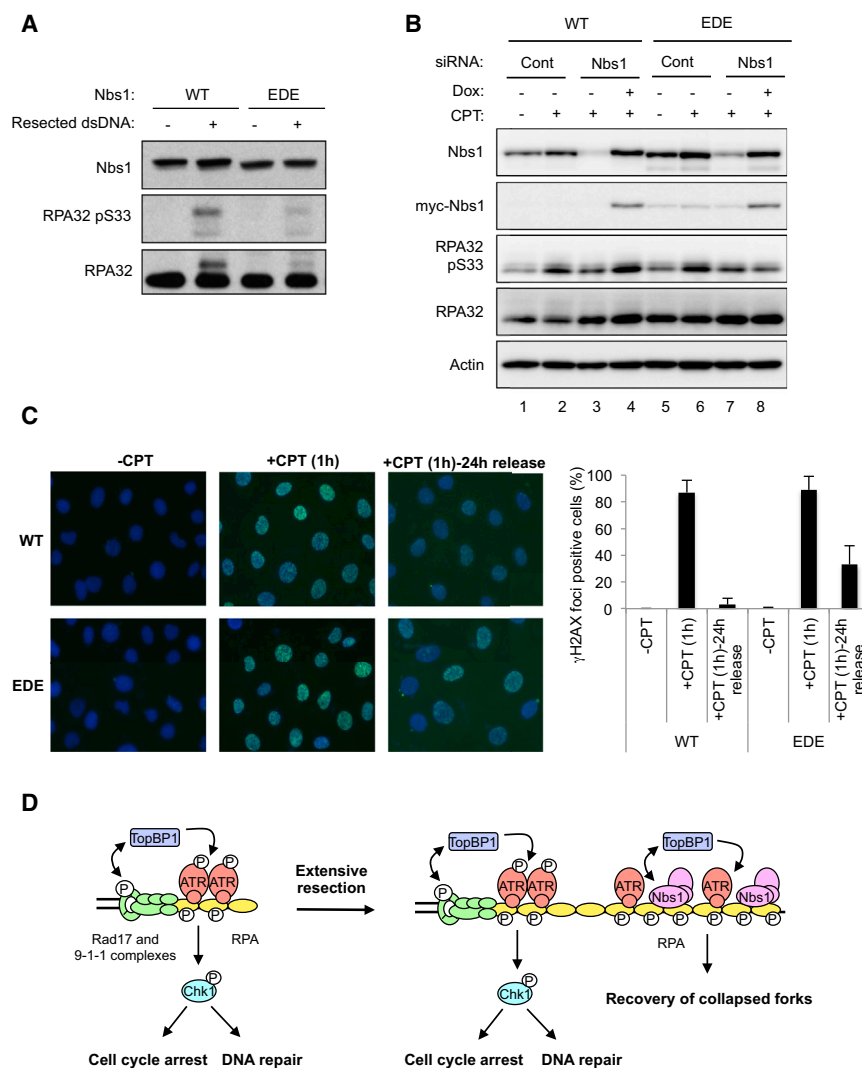


Figure 7. The Interaction of Nbs1 and RPA Is Required for Nbs1-Mediated RPA32 Phosphorylation and Recovery of Collapsed Replication Forks

(A) U2OS-derivative cell lines were established to express myc-tagged Nbs1^{WT} and Nbs1^{EDE} conditionally. As indicated, cells were treated with 1 μ g/ml doxycycline (Dox) to induce myc-tagged Nbs1 proteins, and transfected with siNbs1-2 to knock down endogenous Nbs1. Nuclear extracts were prepared from cells, and the ability of resected dsDNA to induce RPA32 Ser33 phosphorylation was analyzed with phosphospecific antibody.

(B) Myc-tagged Nbs1^{WT} and Nbs1^{EDE} were used to replace endogenous Nbs1 as in (A). Cells were synchronized in S phase as described in [Experimental Procedures](#) and subsequently treated with CPT (1 μ M) for 45 min or mock treated. The levels of the indicated proteins and the phosphorylation of RPA32 Ser33 were analyzed by western blot.

(C) Myc-tagged Nbs1^{WT} and Nbs1^{EDE} were used to replace endogenous Nbs1 as in (A). Cells were synchronized in S phase, treated with CPT (1 μ M) for 1 hr or mock treated, and then released into CPT-free medium. After 24 hr, cells were immunostained with antibody to γ H2AX and the fractions of cells with γ H2AX foci were quantified. Error bars: SDs from three independent experiments (n = 3).

(D) A model that depicts the two distinct modes of ATR activation at resected DSBs.

EXPERIMENTAL PROCEDURES

Cell Culture and Drug Treatment

HeLa, U2OS, and 293T cells were cultured in Dulbecco's modified Eagle's medium (DMEM) supplemented with 10% fetal bovine serum (FBS). NBS-ILB1 cells were cultured in DMEM supplemented with 15% FBS. NBS1-ILB1 cells stably expressing the full-length Nbs1 transgene were maintained in the same medium containing

It is important to note that although the phosphorylation of RPA32 Ser33 is a marker for Nbs1-mediated ATR activation, ATR may have additional substrates when it is activated in this mode. Many of the known or putative substrates of ATR, including a number of DNA repair proteins, interact with RPA and may accumulate on RPA-ssDNA during resection ([Matsuoka et al., 2007](#)). The Nbs1-mediated mode of ATR activation may contribute to the phosphorylation of these ATR substrates and to DNA repair, particularly during the late phase of the DNA damage response. Together, Rad17 and Nbs1 may orchestrate the phosphorylation of two distinct sets of ATR substrates in two sequential phases of the DNA damage response. Whereas Rad17 may promote DNA damage signaling, cell-cycle arrest, and the early events of DNA repair through Chk1 phosphorylation, Nbs1 may promote the late events of DNA repair through progressive phosphorylation of ATR substrates on RPA-ssDNA. The coordination of these two distinct mechanisms of ATR activation may be important for the full function of ATR in the DNA damage response.

500 μ g/ml G418. For RNAi, cells were transfected with 40 nM siRNA using RNAi MAX transfection reagent (Invitrogen). The sequences of the siRNAs used in this study are listed in the [Extended Experimental Procedures](#). Plasmid transfection of 293T cells was performed with Lipofectamine 2000 (Invitrogen) and the transfected cells were analyzed after 48 hr. U2OS derivative cell lines stably expressing myc-Nbs1 or mutants were cultured in DMEM with 10% FBS and 100 μ g/ml Zeocin (Invitrogen). Expression of myc-Nbs1 was induced with 1 μ g/ml doxycycline (Sigma). To synchronize cells in S phase, cells were treated with 2.5 mM thymidine (Sigma) for 20 hr, washed thoroughly with PBS, and released into thymidine-free medium for 3 hr. To assess the recovery of collapsed replication forks, synchronized S phase cells were treated with CPT for 1 hr and then released into CPT-free medium. Kinase inhibitors VE-821, KU55933, and NU7026 were used at 10 μ M. CPT was used at 1 μ M.

Antibodies

The phosphospecific antibodies to ATR pT1989 were previously described ([Liu et al., 2011](#)). We obtained ATR, Nbs1, Mre11, Rad50, TopBP1, phospho-Rad17 (Ser645), and phospho-RPA32 (Ser33 and Ser4/Ser8) antibodies from Bethyl; Rad17 and Chk1 antibodies from Santa Cruz; phospho-Chk1 (Ser345), phospho-Chk2 (Thr68), and phospho-p53 (Ser15) antibodies from Cell Signaling; phospho-H2AX (Ser139) antibody from Millipore; phospho-ATM (Ser1981) antibody from Epitomics; RPA32 antibody from Thermo

Scientific; and Myc antibody from Medical & Biological Laboratories. Phospho-DNA-PKcs (Ser2056) and CtIP antibodies were kindly provided by Drs. B. Chen and R. Baer, respectively.

Extract-Based ATR Activation Assay

An extract-based ATR activation assay was performed as described previously (Shiotani and Zou, 2011). To analyze ssDNA-induced ATR activation, nuclear extracts were pretreated with 10 μ M of KU-55933 and NU7026 for 15 min on ice to inhibit ATM and DNA-PKcs, and supplemented with the reaction buffer (buffer R), which brought the final buffer compositions to 10 mM HEPES (pH 7.6), 50 mM KCl, 0.1 mM MgCl₂, 1 mM phenylmethanesulfonyl fluoride, 0.5 mM dithiothreitol, 1 mM ATP, 10 μ g/ml creatine kinase, and 5 mM phosphocreatine. ssDNA of various lengths was incubated in the extracts for 30 min at 37°C. The sequences of the DNA oligonucleotides used in this study are listed in the Extended Experimental Procedures.

RPA-ssDNA Binding Assay

Biotinylated ssDNA (5'TGCAGCTGGCACGACAGGTTTAAATGAATCGGCCA-ACGCGCGGGGAGAGGCGGTTTGGCGCT-biotin-3') was attached to streptavidin-coated magnetic beads in binding buffer (10 mM Tris-HCl [pH 7.5], 100 mM NaCl, 10% glycerol, 0.01% NP-40, and 10 μ g/ml bovine serum albumin) followed by incubation with or without purified RPA at room temperature for 30 min. To analyze the binding of various proteins to RPA-ssDNA, we incubated RPA-ssDNA with purified MRN, Nbs1, or cell lysates in the binding buffer. To map the RPA-ssDNA-interacting domain of Nbs1, we used lysates prepared from cells expressing various myc-Nbs1 mutants. After a 30 min incubation at room temperature, beads were retrieved and washed twice with the binding buffer. The proteins bound to beads were denatured in the SDS sample buffer, separated on SDS-PAGE, and analyzed by western blot. Recombinant RPA complex was purified from *Escherichia coli* as previously described (Henricksen et al., 1994). The MRN complex and Nbs1 were purified from insect cells infected with baculoviruses expressing Mre11, Rad50, and Nbs1 (Paull and Gellert, 1999).

SUPPLEMENTAL INFORMATION

Supplemental Information includes three figures and Extended Experimental Procedures and can be found with this article online at <http://dx.doi.org/10.1016/j.celrep.2013.04.018>.

LICENSING INFORMATION

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ACKNOWLEDGMENTS

We thank Dr. K. Cimprich for communicating previously unpublished results (Duursma et al., 2013); Drs. S. Elledge, A. Nussenzweig, T. Paull, X. Wu, B. Chen, R. Baer, P. Concannon, and K. Komatsu for reagents; and members of the Zou lab for helpful discussions. P.H. was supported by a postdoctoral fellowship from the Swedish Research Council. A.M. is supported by an FRSQ fellowship. L.Z. is a Jim and Ann Orr Massachusetts General Hospital Research Scholar. This work was supported by grants from the Japanese Ministry of Education, Culture, Sports, Science and Technology to B.S. (KAKENHI 24800044) and H.T. (KAKENHI 23300363), and grants from the NIH (GM076388), the ACS (RSG-08-297), and the Federal Share of MGH Proton Program to L.Z.

Received: October 16, 2012

Revised: April 5, 2013

Accepted: April 19, 2013

Published: May 16, 2013

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