Repair of a DNA-Protein Crosslink by Replication-Coupled Proteolysis

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SUMMARY

DNA-protein crosslinks (DPCs) are caused by environmental, endogenous, and chemotherapeutic agents and pose a severe threat to genome stability. We use Xenopus egg extracts to recapitulate DPC repair in vitro and show that this process is coupled to DNA replication. A DPC on the leading strand template arrests the replisome by stalling the CMG helicase. The DPC is then degraded on DNA, yielding a peptide-DNA adduct that is bypassed by CMG. The leading strand subsequently resumes synthesis, stalls again at the adduct, and then progresses past the adduct using DNA polymerase ζ. A DPC on the lagging strand template only transiently stalls the replisome, but it too is degraded, allowing Okazaki fragment bypass. Our experiments describe a versatile, proteolysis-based mechanism of S phase DPC repair that avoids replication fork collapse.

INTRODUCTION

Chromosomes contain myriad structural and regulatory proteins that ensure the stability, expression, and duplication of the genome. These proteins sometimes form covalent DNA-protein crosslinks (DPCs) through the action of ionizing radiation (IR), UV light, endogenous and exogenous reactive aldehydes, and chemotherapeutics such as nitrogen mustards, cisplatin, and 5-aza-2’-deoxycytidine (azaC) (reviewed in Barker et al., 2005; Ide et al., 2011). Because of their bulky nature, DPCs are predicted to inhibit DNA replication and transcription and thereby interfere with genome integrity. Despite its relevance to human health, DPC repair is poorly understood.

In bacteria, current evidence suggests that small DPCs (less than 11 kDa) are repaired via nucleotide excision repair (NER), whereas larger DPCs are repaired by homologous recombination (HR) (Ide et al., 2011; Nakano et al., 2007). As seen in bacteria, the eukaryotic NER machinery only incises DPCs smaller than 11 kDa (Baker et al., 2007; Nakano et al., 2009; Novakova et al., 2003; Reardon and Sancar, 2006). To account for the repair of larger DPCs, the proteasome is proposed to reduce the protein to a small peptide that is removed by NER (Baker et al., 2003; Reardon and Sancar, 2006). However, other reports concluded that the proteasome and NER are not involved in DPC removal (Nakano et al., 2009; Zecevic et al., 2010). In a distinct model, eukaryotic DPCs are processed via HR, presumably during replication (Ide et al., 2011; Nakano et al., 2009). This idea is based on findings that chicken and mammalian cells deficient in the Fanconi anemia (FA) pathway and HR are sensitive to DPC-inducing agents (Nakano et al., 2009; Orta et al., 2013; Ridpath et al., 2007), although the involvement of HR has been challenged (Rosado et al., 2011). In summary, there is currently no clear consensus on how DPCs are repaired, especially in vertebrates.

The effect of DPCs on DNA replication has been investigated in bacteria and using purified DNA polymerases. In vitro, DPCs block synthesis by DNA polymerases (Chvalová et al., 2007; Novakova et al., 2003; Yeo et al., 2014). In E. coli, DNA methyltransferase-based DPCs induce replication fork stalling (Kuo et al., 2007), presumably by inhibiting the replicative DnaB helicase. We previously showed that the eukaryotic replicative helicase, CMG (a complex of Cdc45, MCM2-7 and GINS; Ives et al., 2010), readily bypasses a biotin-streptavidin roadblock on the lagging strand template but not on the leading strand template (Fu et al., 2011). This result indicates that CMG encircles and translocates on the leading strand template in the 3’ to 5’ direction. Similarly, only a DPC on the leading strand template is predicted to act as a helicase barrier (Ide et al., 2011). However, the functional interplay between DNA replication and DPC repair has not been examined in any organism.

Fanconi anemia is a bone marrow failure syndrome caused by defects in 16 “FANC” proteins best known for their roles in DNA interstrand crosslink (ICL) repair (Kottmann and Smogorzewska, 2013). Importantly, FANC-deficient cells are hypersensitive to formaldehyde (Ridpath et al., 2007; Rosado et al., 2011). Moreover, mice deficient in both FANCD2 and the aldehyde-detoxifying enzyme, ALDH2, phenocopy the symptoms of FA (Garaycoechea et al., 2012; Langevin et al., 2011). These results suggest that endogenous aldehydes cause FA, raising the possibility that the FA pathway helps repair DPCs.

Here, we report that Xenopus egg extracts support efficient repair of a chemically-defined DPC and that this repair is strictly coupled to DNA replication. Collision of the replisome with the DPC on the leading strand template stalls the CMG helicase and triggers DPC proteolysis, reducing the DPC to a short
peptide. The peptide adduct is then bypassed, first by the CMG helicase and then by the nascent leading strand. A lagging strand DPC is readily bypassed by CMG but must be destroyed for the completion of lagging strand synthesis. Extension of nascent strands past the peptide adduct requires DNA pol. Our results describe a versatile mechanism of DPC repair that avoids replisome disassembly and double-strand DNA break formation, two major sources of genome instability.

RESULTS

Replication of a Plasmid Containing a Site-Specific DPC

To generate a plasmid containing a site-specific DPC (pDPC), the DNA methyltransferase HpaII (M.HpaII, 45 kD) was covalently linked to its recognition site, CCGG, via 5-fluoro-2-deoxycytosine (Chen et al., 1991) (Figure 1A). M.HpaII was crosslinked to the top or bottom strands of the plasmid, generating pDPCTop and pDPCBot (Figure 1A and Figure S1A available online). pCTRL (the fluorinated plasmid lacking M.HpaII) and pDPCTop were replicated in nucleus-free Xenopus egg extract (Walter et al., 1998). In this system, a single, complete round of plasmid DNA replication can be monitored via incorporation of [α-32P]dATP. Replication of pCTRL quickly yielded supercoiled daughter molecules (Figure 1B, lanes 1–6) (Walter and Newport, 2000). In contrast, replicated pDPCTop first accumulated as a 50:50 ratio of open circular and supercoiled molecules (Figure 1B, lane 9) before gradual conversion into the supercoiled form (Figure 1B, lanes 10–12). Because M.HpaII is linked to one DNA strand, we postulated that replication of the undamaged strand quickly yielded supercoiled products, while the damaged strand yielded gapped molecules containing the DPC (Figure 1D). Consistent with this interpretation, when replicated DNA was not treated with proteinase K (ProtK) before electrophoresis, the gapped molecules were selectively retarded in the gel, migrating as a smear (Figure 1B, compare lanes 20–22 to 8–10). This smear was converted to supercoiled products by 60 min (Figure 1B,

Figure 1. Replication of a Plasmid Containing a Site-Specific DPC

(A) Schematic of pDPCTop.

(B) pCTRL and pDPCTop were replicated in egg extract in the presence of [α-32P]dATP. Samples were treated with Proteinase K (ProtK), as indicated, and analyzed by agarose gel electrophoresis. OC, open circular; SC, supercoiled.

(C) pCTRL and pDPCTop were replicated, digested with FspI, and analyzed by 2D gel electrophoresis. The lower cartoon illustrates relevant DNA intermediates. Arrows, see main text.

(D) Model for how replication of pDPC initially yields 50% open circular (OC) and 50% supercoiled (SC) products.
lanes 23 and 24), suggesting that the DPC was removed from DNA by the time the gap was filled in (Figure 1D). Indeed, we detected the presence of a gap surrounding the DPC (Figures S1A and S1B). Interestingly, the gap first increased in size due to 5’ to 3’ resection of nascent lagging strands (Räschi et al., 2008), but disappeared by 60 min, when the DPC had also been removed (Figures S1A and S1B). These data show that DPCs are efficiently removed from DNA during replication in Xenopus egg extracts.

We next examined whether a DPC stalls DNA replication forks. Replicating pCTRL and pDPCTop were analyzed by two-dimensional (2D) gel electrophoresis after linearization with Fsp1 (Figure 1C, left cartoon). After 7 min, replicated pCTRL migrated mostly as linear molecules (Figure 1C, red arrow). In contrast, most of the pDPCTop migrated at the apex of a double-Y arc, consistent with fork convergence and stalling at the lesion (Figure 1C, blue arrow). By 15 min, when most of the gapped plasmids still retained the DPC (Figure 1B, lane 21), virtually all replication intermediates migrated as linear products (Figure 1C, green arrow). We conclude that converging forks transiently stall at the DPC; after resolution of the two daughter molecules, the DPC is removed and the gap is filled in (Figure 1D).

The Adducted Strand Is Replicated after the Unadducted Strand

Our data suggest that replication of the adducted strand is delayed compared to the unadducted strand (Figure 1D). To confirm this, we examined the fate of both nascent strands during replication, pCTRL, pDPCTop, and pDPCBot were replicated, and at different times, nascent strands were digested with NcoI and AatII and separated on a denaturing polyacrylamide gel. The overhangs created by NcoI and AatII digestion yield top strand extension products (Top, 412 nt) that are 8 nt shorter than bottom strand extension products (Bot, 420 nt), allowing us to differentiate between replication of the damaged and undamaged strands (Figures 2A and 2B). During replication of pCTRL, Top and Bot extension products accumulated simultaneously, as expected (Figure 2C, upper panel, lanes 1–7). In contrast, with pDPCTop, replication of the damaged (Top) strand was delayed by ~20 min relative to the undamaged (Bot) strand (Figure 2C, upper panel, lanes 8–14). The same result was obtained with pDPCBot, only now the damaged (Bot) strand was replicated after the undamaged (Top) strand (Figure 2C, upper panel, lanes 15–21). In conclusion, the undamaged strand is replicated well before the DPC-containing strand.

Multistep Bypass of a DPC

We next addressed why replication of the DPC-containing strand was delayed relative to the unadducted strand. When two replication forks converge on a DPC (Figures 1C and 1D), one fork encounters the DPC on the leading strand template, whereas the converging fork encounters the same lesion on the lagging strand template (Figure 2A). Given that CMG translocates on the leading strand template (Fu et al., 2011), we suspected that the fork that encounters the DPC on the leading strand template will stall. In contrast, the fork that encounters the DPC on the lagging strand template might quickly bypass the lesion, allowing rapid replication of the unadducted strand.

To test these predictions, we monitored the leading strand of the rightward fork as it encountered a DPC on the leading (Figure 2B) or lagging strand templates (Figure 2A). Nascent strands of pCTRL, pDPCTop, and pDPCBot were digested with NcoI, which cleaves ~150 nt to the left of the DPC (Figures 2A and 2B), and products were separated on a denaturing gel. Replication of pCTRL yielded no ~150 nt products, reflecting the absence of any fork stalling (Figure 2C, lower panel, lanes 1–7). In contrast, when the fork encountered a DPC on the leading strand template in pDPCBot, nascent leading strands stalled ~29 to 42 nt from the DPC and persisted there for up to 30 min (Figure 2C, lanes 15–19, red bracket). We previously determined that when forks encounter an ICL, the leading strand stalls on average 30 nt from the lesion due to steric hindrance by CMG (Fu et al., 2011; Räschi et al., 2008). The slightly increased distance observed here (~36 nt) is consistent with the helicase being obstructed by a DPC, which is bulkier than an ICL. We conclude that the DPC causes prolonged CMG stalling. After this initial arrest, nascent leading strands were further extended, stalling again 15 to 24 nt from the DPC (Figure 2C, green bracket; see Discussion for the possible source of these intermediates).

Next, the leading strands reached the crosslinked DNA base, where they stalled at the 0, +1, and +2 positions (Figure 2C, blue bracket; Figures S2D and S2E show that the 0 position corresponds to the site of the damaged base). Finally, by 60 min, the leading strands bypassed the lesion and accumulated as extension products (Figure 2C, upper panel, lanes 20–21, Bot extension product).

When the rightward fork encountered the DPC on the lagging strand template (pDPCTop, Figure 2A), the outcome was different. Nascent leading strands stalled ~34–47 nt from the DPC, but much more transiently (Figure 2C, lanes 8–9, orange bracket), before being extended past the lesion (Figure 2C, upper panel, lanes 9 and 10, Bot extension product). Thus, a DPC on the lagging strand template appears to only briefly delay movement of the CMG helicase. When leftward and rightward forks converging on the same DPC were compared by cutting pDPCTop on the right of the DPC with AatII (Figure S2A) or on the left of the DPC with NcoI (Figure 2A), we observed the same result (Figures S2C and 2C). We conclude that the fork whose helicase travels on the undamaged strand stalls transiently (~5 min) before moving past the lesion and completing replication of the undamaged template (Figure S2F, rightward fork). The helicase traveling on the damaged strand stalls for ~20–30 min (~29 to ~42” species), accounting for delayed replication of the damaged strand (Figure S2F, leftward fork).

The DPC Is Processed into a Short Peptide Adduct

The fact that the DPC-containing strand is ultimately fully replicated suggested that the DPC is degraded or excised. Importantly, we did not detect any incision on the parental DNA in the vicinity of the DPC (Figure S3A, lanes 6–11). We therefore postulated that the DPC might be degraded on DNA (Figure S2F). If this hypothesis is correct, a peptide adduct might remain on the parental DNA after completion of replication. To address this possibility, we replicated pCTRL or pDPCTop in egg extract for 120 min in the absence of radiolabeled nucleotides.
The DNA was then digested to excise a 165 nt fragment encompassing the DPC (Figure 3A). The DNA was separated on a denaturing gel, and the top strand was visualized by strand-specific Southern blotting. As expected, pCTRL produced a single, 165 nt band (Figure 3B, lanes 1 and 2). In contrast, replicated pDPCTop generated the same 165 nt band and a ladder of more slowly migrating bands, indicating that adducts of different sizes remained attached to DNA (Figure 3B, lane 4). When the same samples were treated with ProtK prior to electrophoresis, the slower migrating species collapsed into a single, new band that migrated just above the unadducted strand (Figure 3B, compare lanes 4 and 10). Importantly, ProtK is predicted to leave a 4-residue peptide of M.HpaII crosslinked to DNA. We infer that the ladder of bands observed in lane 4 corresponds to short peptide adducts of different lengths. These results demonstrate that DPC repair involves reducing the DPC to a series of short DNA-peptide adducts.

To examine the kinetics of DPC removal, we used a C-terminally biotinylated M.HpaII to generate pDPCTop, which we replicated in egg extract containing [α-32P]dATP. At different times, biotinylated M.HpaII was precipitated using streptavidin beads, and the recovery of radiolabeled DNA was monitored on an agarose gel (Figure 3C). Once the DPC is degraded, DNA can no longer coprecipitate with M.HpaII. As shown in Figure 3D, DNA recovery drastically declined by 30 min, indicating completion of DPC processing (top panel, lane 4). Notably, only replication intermediates and gapped, monomeric molecules were coprecipitated, consistent with our earlier conclusion that

Figure 2. Multistep Bypass of a DPC
(A and B) Depiction of nascent leading strands generated after Ncol digestion of DPCTop (A) and pDPCBot (B). Extension products were monitored with Ncol and AatII digestion. Green hexamer, rightward CMG helicase; N, sequencing primer.
(C) pCTRL, pDPCTop, and pDPCBot replication intermediates were digested with Ncol and AatII (upper panel) or Ncol (lower panel), and separated on a denaturing polyacrylamide gel alongside a sequencing ladder. Nascent strands generated by the rightward fork are indicated in brackets. For lagging strand identification, see legend for Figure S2C. Red and gray arrows, extension products of the bottom (Bot) and top (Top) strands (upper panel). Black arrows, location of the DPC in pDPCTop and pDPCBot (bottom panel).

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gapped but not supercoiled molecules contain the DPC (Figure 3D, lanes 1–4). Whereas most DPC processing occurred before 30 min (Figure 3E, blue graph), nearly all lesion bypass was completed after this time (Figure 3E, red graph). Strikingly, when DNA replication initiation was blocked with Geminin (Wohlschlegel et al., 2000), the DPC persisted, demonstrating that DPC processing requires DNA replication (Figure S3B). Together, these experiments show that replication forks activate DPC proteolysis, followed by replication bypass.

**Bypass of the Peptide Adduct Requires DNA pol ζ**

Mutations in Rev1 and Rev7 (a subunit of DNA pol ζ) sensitize chicken cells to formaldehyde (Ridpath et al., 2007), suggesting a role for these TLS polymerases in DPC repair. In support of this idea, immunodepletion of Rev7 from egg extract (Figure 4A) inhibited the conversion of gapped to supercoiled molecules during replication of pDPCBot (Figure 4B, lanes 7–12). Rev7 depletion did not affect DPC destruction (data not shown). Importantly, in the absence of Rev7, leading strands accumulated at the 0, +1, and +2 positions, and extension products of the damaged (Bot) strand were greatly diminished (Figure 4C, lanes 7–12). Due to the difficulty of expressing vertebrate DNA pol ζ, we did not attempt to rescue the effect of Rev7 depletion. Instead, we examined the effect of depleting Rev1, which has a close physical and functional interaction with DNA pol ζ (Guo et al., 2003), including in Xenopus egg extracts (M. Budzowska and J.C.W., unpublished data). Rev1 depletion from egg extract precisely mimicked the effect of Rev7 depletion (data not shown). We conclude that following DPC destruction, a complex containing DNA pol ζ and Rev1 extends the leading strand past the peptide adduct.

**FANCI-FANCD2 Is Not Essential for DPC Bypass**

Because Fanconi anemia has been linked to reactive aldehydes (Langevin et al., 2011; Rosado et al., 2011), we explored its connection to DPC repair. As shown in Figure S1C, replication of pDPCTop stimulated ubiquitylation of FANCD2. However, immunodepletion of the FANCI-FANCD2 complex from egg extract had no effect on DPC bypass (Figures S4A and S4B), even though it inhibited ICL repair (Figures S4C and S4D; Knipscheer et al., 2009). We conclude that FANCI-FANCD2 does not participate in replication-coupled DPC repair.

**Single Forks Efficiently Bypass Leading and Lagging Strand DPCs**

During replication of pDPC, two replication forks rapidly converge on the DPC. By contrast, in vivo, where the interorigin distance is large, one fork is generally expected to collide with a DPC well before a second fork arrives. We therefore wanted to know whether two forks are required to repair a DPC or whether one fork suffices. To address this issue, we created pDPC-LTop and pDPC-LBot in which an array of 48 lac operator (lacO) sites is placed ~260 nt to the left of top or bottom strand DPCs (Figures 5A and 5B). When bound by the lac repressor (Lacl), the lacO array blocked fork progression for up to 4 hr (Figure 5A, 346–357, October 9, 2014 ©2014 Elsevier Inc.)
Correlation of CMG Bypass with DPC Proteolysis

To examine how a single fork bypasses a leading strand DPC, replicating pDPC-LBot was digested with AatII alone (Figure 5A). Upon encounter of the fork with the DPC, the leading strands stalled, approached, and bypassed the lesion (Figure 5C, lanes 22–27, red, green and blue brackets) as seen during fork convergence (Figures 5C, lanes 4–9 and S2C), albeit with slightly slower kinetics. Strikingly, even when there was no converging fork, a significant portion of lagging strands had fully matured between 20 and 30 min, while nearly none of the leading strands had extended past the DPC (Figure 5C, upper panel, lanes 23–24).

DPC Processing Is Essential for Efficient Lesion Bypass

Our model predicts that failure to destroy a DPC on the leading strand template should inhibit CMG bypass. In contrast, blocking the destruction of a DPC on the lagging strand template should not significantly delay CMG bypass, but rather block completion of the lagging strand. To test these predictions, we sought to inhibit the destruction of M.HpaII by perturbing the ubiquitin system. Two different drugs that inhibited the destruction of M.HpaII had no evident effect on DPC repair (Figures S6B and S6D). As an alternative approach, we added ubiquitin-vinyl-sulfone (Ub-VS), a
deubiquitylating enzyme (DUB) inhibitor that blocks ubiquitin recycling and thereby leads to the depletion of free ubiquitin (Figure S6E) (Dimova et al., 2012). pDPC-LTop and pDPC-LBot were replicated in the presence of LacI in egg extracts that were preincubated with buffer, Ub-VS, or Ub-VS and free ubiquitin. Although Ub-VS treatment did not inhibit DNA replication, it dramatically inhibited M.HpaII destruction on both leading and lagging strand templates (Figure 6A, lanes 7–12 and 25–30). Addition of free ubiquitin restored M.HpaII destruction (Figure 6A, lanes 13–18 and 31–36). Therefore, M.HpaII destruction requires the presence of free ubiquitin but apparently not DUB or proteasome activity.

To address how lesion bypass is affected when the DPC is not destroyed, nascent strands from each reaction were analyzed following AatII digestion (as depicted in Figures 5A and 5B). When a leading strand DPC was stabilized with Ub-VS, the disappearance of the −29 to −42 nt leading strand products was delayed for up to 90 min, indicating prolonged CMG stalling at the DPC (Figures 6B, compare lanes 1–6 and 7–12, red bracket, and 6C). This effect was largely reversed by coaddition of free ubiquitin (Figures 6B, lanes 13–18, red bracket and 6C). Thus, efficient CMG bypass of a leading strand template DPC requires the proteolytic processing of the DPC into a peptide adduct. Although CMG bypass appeared to be severely inhibited, leading strands still approached the lesion over time (Figure 6B, lanes 7–12, blue bracket). By 3 hr, virtually all of the nascent leading strands had advanced from the −29 to −42 position to the crosslinked base, accumulating at the 0 position to the crosslinked base, accumulating at the 0 position.
even though the DPC was still intact (Figure 6A, lane 12). This approach of the leading strands to the lesion correlated with full maturation of lagging strands past the DPC (Figure 6B, upper panel, lanes 7–12). This surprising observation suggests that a replication fork can bypass an intact DPC on the leading strand template, albeit with slower kinetics than when the DPC is degraded into a peptide. We ruled out the possibility that lagging strand maturation was due to origin firing between the replication fork barrier and the DPC (Figures S6F and S6H). In the Discussion, we propose models that can explain replication fork bypass of an intact DPC.

In further agreement with our predictions, stabilizing the lagging strand DPC did not significantly prolong CMG stalling at the lesion (Figures 6B, lanes 19–30, orange bracket and 6D). This result implies that once the replication fork has moved past the lesion, a growing Okazaki fragment collides with the intact DPC. To examine the fate of this maturing Okazaki fragment, we digested replication samples to the left of the DPC with BssHII (Figure S6G). As shown in Figure S6H, nascent lagging strands approached the 0 position with similar kinetics in the presence and absence of Ub-VS (compare lanes 25–30 with lanes 19–24), consistent with unhindered bypass of the intact DPC by CMG. Interestingly, in the presence of Ub-VS, lagging

**Figure 6. DPC Processing Is Essential for Efficient Replicative DPC Bypass**

(A) pDPC-LTop and pDPC-LBot were replicated in the presence of LacI and buffer (+Buffer), 13 $\mu$M Ub-VS (+Ub-VS) or 13 $\mu$M Ub-VS and 80 $\mu$M ubiquitin (+Ub-VS +Ub.). Samples were analyzed as in Figure 3D.

(B) Samples from (A) were analyzed as in Figure 5C.

(C) Quantification of the −29 to −42 species of pDPC-LTop from (B). Values were normalized to the maximum peak value of the buffer control.

(D) −34 to −47 species of pDPC-LBot from (B) were quantified as in (C).
strands persisted at the 0 position for up to 3 hr and never advanced to the +1 and +2 positions (Figure S6H, lanes 25–30, 0 position). The same phenomenon was observed when leading strands collided with a stable DPC (Figure 6B, lanes 7–12). Together, these data suggest that a DNA polymerase is able to approach the base to which the intact DPC is attached, and insert a nucleotide across from it. However, advance to the +1 and +2 positions might be impossible due to steric hindrance by the DPC. Consistent with this view, TLS polymerases can bypass short peptide DNA-adducts in vitro, but not larger DPCs (Yamanaka et al., 2010; Yeo et al., 2014). Alternatively, or in addition, ubiquitin-depletion might inhibit recruitment of TLS polymerases (Goodman and Woodgate, 2013). We conclude that DPC degradation is dependent on ubiquitin and that a persistent DPC on the leading but not the lagging strand inhibits progression of the replication fork.

DISCUSSION

Here, we show that Xenopus egg extracts support DNA-protein crosslink (DPC) repair. Our system involves a chemically defined and sequence-specific DPC, and repair is measured directly, by conversion of the DPC to a peptide adduct and generation of supercoiled DNA products. Using this approach, we show that DPC repair requires DNA replication, and we elucidate how the two processes are coupled. A leading strand DPC must be degraded to a peptide to allow efficient bypass of the CMG helicase, consistent with CMG translocation on the leading strand template (Figure 7A, i–iv). Following CMG bypass of a leading strand DPC, the growing leading strand is extended to the adducted base, where it stalls (Figure 7A, iv). Concurrently, the lagging strand continues growing past the DPC, providing a clear instance of replicative uncoupling in vertebrates (Figure 7A, iv and v). Finally, DNA pol ζ allows the leading strand to bypass the peptide adduct (Figure 7A, v). In contrast, a DPC on the lagging strand template only transiently stalls the replisome (Figure 7B, i–iii), but it too is degraded to a peptide, allowing Okazaki fragment bypass (Figure 7B, iii–v). Although egg extracts do not support removal of the peptide adduct, this reaction might occur in vivo via excision repair (Figures 7A and 7B, vi). Our results describe a comprehensive model of replication-coupled DPC repair, implications of which are discussed below.

Replication-Dependent Proteolysis of a DPC

It has been postulated that replication fork collision with DPCs provides a sensing mechanism that triggers DPC repair (Kuo et al., 2007; Nakano et al., 2007; Reardon et al., 2006). Our experiments provide direct support for this model as they establish the existence of a replication-dependent protease that reduces DPCs to small DNA-peptide adducts. Notably, inhibition of proteasome activity in egg extracts had no significant effect on DPC repair, suggesting this process is proteasome-independent (although we cannot formally rule out a role for residual proteasome activity). In contrast, ubiquitin depletion inhibited DPC repair. The role of ubiquitin is presently unclear, but we speculate that DPCs might be ubiquitylated prior to destruction by the replication-dependent protease. A potential candidate for this protease is the p97 cofactor Spartan/DVC1, which has homology to yeast Wss1 (Mosbech et al., 2012). Wss1 was recently shown to remove trapped topoisomerase I complexes and to confer resistance to formaldehyde, probably by removing DPCs (Stingele et al., 2014). Intriguingly, Wss1 destroys a wide variety of different DNA binding proteins in a DNA-dependent manner. Analogously, the replication-dependent protease in egg extract appears to employ a general recognition strategy, given its ability to degrade a bacterial methyltransferase. These
parallels raise the possibility that Spartan/DVC1 functions in replication-dependent DPC repair. However, we have not been able to generate useful Spartan/DVC1 antibodies to test this idea.

Two Mechanisms of Leading Strand DPC Bypass?
Our data suggest that a DNA replication fork can bypass a DPC on the leading strand template by at least two mechanisms. In the primary mechanism, DPC proteolysis allows CMG to pass over the lesion (now a peptide) and continue unwinding DNA. In support of this model, the disappearance of CMG’s footprint (–29 to –42 arrest) was closely correlated with DPC degradation (Figure S5E). In addition, when DPC destruction was blocked by Ub-VS, the CMG footprint persisted for up to 2 hr (Figure 6C). Consistent with this mechanism, a purified MCM helicase can translocate past ~30 amino acid but not ~45 amino acid peptide adducts (Nakano et al., 2013). Interestingly, after the –29 to –42 CMG footprint disappears, a new population appears that is stalled at the –11 to –24 positions (Figures 6B, green bracket). We speculate that these species might reflect slow unwinding by CMG of the DNA containing the peptide adduct.

Strikingly, in the presence of Ub-VS, all of the leading strands ultimately approached the added base by 3 hr, and lagging strands fully matured past the lesion (Figure 6B). Therefore, even when DPC proteolysis is blocked, the fork can eventually bypass the lesion. We can imagine at least two mechanisms for this secondary, proteolysis-independent DPC bypass. First, CMG unwinds past the intact DPC by transiently opening its ring (Figure S7A), as proposed for the viral T antigen helicase (Yardimci et al., 2012). Alternatively, a 5’ to 3’ DNA helicase might load onto the lagging strand template and translocate past the DPC (Figure S7B). Interestingly, the –11 to –24 leading strand species normally seen during DPC bypass are absent in the presence of Ub-VS (Figure 6B, compare lanes 1–6 and 7–12, green bracket). One interpretation is that CMG ring-opening allows the helicase to pass over the DPC without stalling. Alternatively, the 5’ to 3’ helicase invoked above might displace CMG as it unwinds past the DPC (Figure S7B). Whatever the precise details, proteolysis-dependent and proteolysis-independent mechanisms of DPC bypass appear to be mechanistically distinct.

The Mechanism of TLS during DPC Repair
We show that during bypass of a leading or lagging strand DPC, the nascent strand does not arrest immediately before the added base (~1 position). Instead, 0, +1, and +2 products accumulate. This finding demonstrates that the insertion of a nucleotide across from the added base (by a replicative or TLS polymerase) is not rate-limiting. In our DPC substrate, the crosslinked cytosine is predicted to be flipped out of the double helix, generating an abasic-like site (Klimasauskas et al., 1994). Importantly, both pol ε and pol δ are able to insert nucleotides opposite abasic sites (Haracska et al., 2001; Sabouri and Johansson, 2009). Pol ε and pol δ might also be able to extend the leading strand to the +1 and +2 positions. Once the adduct blocks further synthesis, the nascent strand is extended past the lesion by a complex containing DNA pol ζ and Rev1. In yeast cells lacking Wss1, Rev3-dependent mutagenesis is reduced (Stingele et al., 2014), consistent with our observation that DPC destruction and TLS are coupled. Given the heterogeneity in the chemistry of DPCs, it is likely that different TLS polymerases will be employed to bypass different types of DPCs.

Implications for Fanconi Anemia
The synthetic sickness between FANC mutations and mutations affecting reactive aldehyde metabolism suggested that the biological function of the FA pathway might be to repair DPCs. We found that unlike ICL repair, DPC repair does not require FANCI-FANCD2. In hindsight, this observation is not unexpected. We showed previously that FANCI-FANCD2 promotes incision of replication forks that have stalled at an ICL, leading to ICL unhooking and lesion bypass (Klein Douwel et al., 2014; Knipscheer et al., 2009; Zhang and Walter, 2014) (Figure S4E). In contrast, the DPC repair pathway we describe here does not involve the incision of parental DNA strands, obviating the need for FANCI-FANCD2. A reasonable conclusion is that ICLs are in fact the DNA lesions that drive FA, as widely believed. Consistent with this view, formaldehyde is known to cause ICLs (Huang and Hopkins, 1993). However, we cannot rule out the possibility that the repair of some aldehyde-induced DPCs requires incisions and therefore involves the FA pathway.

Proteolytic versus HR Repair of DPCs
Some mammalian studies find a role for HR in DPC repair (Nakano et al., 2009; Ridpath et al., 2007). In yeast, Wss1 and Rad52 exhibit nonepistasis with regard to formaldehyde sensitivity, suggesting that DPC proteolysis and HR represent distinct means of removing DPCs in S phase (Stingele et al., 2014). Consistent with this view, our experiments show that replication-dependent DPC repair in vertebrates proceeds without the formation of double-strand DNA breaks, which pose a major threat to genome stability. We propose that replication-dependent proteolysis represents the preferred pathway of DPC repair and that HR participates in the bypass of rare DPCs that cannot be degraded.

EXPERIMENTAL PROCEDURES
Preparation of DNA Constructs
To generate pDPCTop and pDPCBot, we cloned 5’-CCTCAGCATCCCAGTACC TCAGC-3’ between the EcoRI and NdeI sites of pUC.HSO (Wold et al., 1987) to generate pHY10 (Yardimci et al., 2012). We then inserted a 5-fluoro-2’-deoxyctydine (C-flu)-modified oligonucleotide (Biosynthesis, Lewisville, TX) into either strand. pHY10 was either nicked with Nb.BbvCI and annealed/ligated to 5’-TCAGCATCCCAGTACC-3’ to modify the top strand, or nicked with Nb.BbvCI and annealed/ligated to 5’-TCAGGTACCCAGTACC-3’ to modify the bottom strand. Modified DNA was gel-purified and mixed with M.HpaII-His, or M.HpaII-biotin-His in reaction buffer (50 mM Tris-HCl, pH 7.5, 5 mM 2-mercaptoethanol, 10 mM EDTA) supplemented with 100 μM S-adenosylmethionine (NEB, Ipswich, MA) for 12 hr at 37°C. To make pCTRL, the plasmid containing the fluorinated cytosine on the top strand was used in an identical reaction without M.HpaII.

Protein Expression and Purification
To purify M.HpaII-biotin-His6 (biotinylated M.HpaII), pHpaII-Avitag-His6 and an identical reaction without M.HpaII. We showed previously that FANCI-FANCD2 promotes incision of replication forks that have stalled at an ICL, leading to ICL unhooking and lesion bypass (Klein Douwel et al., 2014; Knipscheer et al., 2009; Zhang and Walter, 2014) (Figure S4E). In contrast, the DPC repair pathway we describe here does not involve the incision of parental DNA strands, obviating the need for FANCI-FANCD2. A reasonable conclusion is that ICLs are in fact the DNA lesions that drive FA, as widely believed. Consistent with this view, formaldehyde is known to cause ICLs (Huang and Hopkins, 1993). However, we cannot rule out the possibility that the repair of some aldehyde-induced DPCs requires incisions and therefore involves the FA pathway.
was purified over Ni-NTA resin (QIAGEN, Valencia, CA). M.Hpall-His$_6$ (unbiotinylated M.Hpall) was purified as described above omitting the addition of biotin during induction.

**Xenopus Egg Extracts and Replication**

For DNA replication in Xenopus egg extracts (Lebofsky et al., 2009; Walter et al., 1998), plasmid DNA was incubated in a high-speed supernatant (HSS) of egg cytoplasm for 30 min (7.5 ng/et al., 1998), plasmid DNA was incubated in a high-speed supernatant (HSS) was quantified using ImageJ (NIH, USA).

Sequencing ladders were generated with the indicated primers using Cycle enzymes followed by addition of 0.5 volumes of nucleoplasmic egg extract (NPE).

**Nascent Strand Analysis**

Nascent strand analysis was performed as previously described (Räschle et al., 2008). Briefly, purified DNA was digested with the indicated restriction enzymes followed by addition of 0.5 volumes of Gel Loading Dye II (Denaturing PAGE) (Life Technologies, Grand Island, NY, USA). DNA fragments were subsequently separated on 5% or 7% denaturing polyacrylamide gels, transferred to filter paper, dried, and visualized using a phosphorimager. The sequencing ladders were generated with the indicated primers using Cycle Sequencing kit (USB Corporation, Cleveland, OH, USA). Radioactive signal was quantified using ImageJ (NIH, USA).

**Antibodies**

The following antibodies were described previously: FANC D2 (Räschle et al., 2008), MCM7 (Walter and Newport, 2000), Rev7 (Räschle et al., 2008) and CDT1 (Arias and Walter, 2005), p-Chk1 (S345) and ubiquitin antibodies were purchased from Cell Signaling (#2341) (Danvers, MA, USA) and Santa Cruz (SC-8017), respectively. FANCI antibody was raised against full-length FANCI expressed in insect cells and was shown to recognize and immunoprecipitate the FANCI protein in Xenopus egg extracts (data not shown).

**SUPPLEMENTAL INFORMATION**

Supplemental Information includes Extended Experimental Procedures and seven figures and can be found with this article online at http://dx.doi.org/10.1016/j.cell.2014.09.024.

**AUTHOR CONTRIBUTIONS**

J.M.D. generated and characterized the lac repressor replication fork barrier; H.Y. purified M.Hpall-biotin; J.P.D. and J.C.W. designed and analyzed the experiments; J.P.D. performed the experiments; J.P.D. and J.C.W. prepared the manuscript.

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**REFERENCES**


