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PARP1 promotes nucleotide excision repair through DDB2 stabilization and recruitment of ALC1

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ABSTRACT

The WD40-repeat protein DDB2 is essential for efficient recognition and subsequent removal of ultraviolet (UV)-induced DNA lesions by nucleotide excision repair (NER). However, how DDB2 promotes NER in chromatin is poorly understood. Here, we identify poly(ADP-ribose) polymerase 1 (PARP1) as a novel DDB2-associated factor. We demonstrate that DDB2 facilitated poly(ADP-ribosyl)ation of UV-damaged chromatin through the activity of PARP1 resulting in the recruitment of the chromatin remodeling enzyme ALC1. Depletion of ALC1 rendered cells sensitive to UV and impaired repair of UV-induced DNA lesions. Additionally, DDB2 itself was targeted by poly(ADP-ribosyl)ation, resulting in increased protein stability and a prolonged chromatin retention time. Our in vitro and in vivo data support a model in which poly(ADP-ribosyl)ation of DDB2 suppresses DDB2 ubiquitylation and outline a molecular mechanism for PARP1-mediated regulation of NER through DDB2 stabilization and recruitment of the chromatin remodeler ALC1.
INTRODUCTION

Nucleotide excision repair (NER) is a multistep process that mediates the removal of structurally and chemically diverse DNA lesions including ultraviolet (UV) light-induced cyclobutane pyrimidine dimers (CPD) and 6-4 pyrimidine pyrimidone photoproducts (6-4PP). The importance of NER in protecting organisms against solar UV-induced DNA damage is underscored by the hereditary disease xeroderma pigmentosum (XP), which is clinically characterized by hypersensitivity to sunlight and predisposition to skin cancer [Cleaver et al., 2009]. XP has been linked to defects in seven proteins (XP-A through XP-G) which, with the exception of XPC and XPE (hereafter named DDB2), function in the core NER reaction. The proteins encoded by the XPC and XPE genes are involved in the global genome NER sub-pathway (GG-NER), but are dispensable for transcription-coupled NER (TC-NER) [Cleaver et al., 2009].

Reconstitution of the NER reaction with purified proteins has defined the minimal set of proteins required for GG-NER in vitro [Aboussekhra et al., 1995]. The initial step of DNA damage recognition depends on the XPC-Rad23 complex and subsequently results in local DNA unwinding and damage verification by the basal transcription factor TFIIH, the single-stranded DNA-binding complex RPA and XPA. Dual incision of the damaged DNA strand is carried out by the 5’and 3’ structure-specific endonucleases XPF-ERCC1 and XPG, respectively, followed by gap filling and DNA ligation [Aboussekhra et al., 1995].

DNA damage recognition by XPC involves the detection of unpaired bases [Min and Pavletich, 2007;Clement et al., 2010], which renders lesion recognition of minor helix-distorting lesions such as CPD very inefficient [Sugasawa et al., 2001]. In addition to XPC, efficient repair of CPDs therefore requires the heterodimeric UV-DDB protein complex consisting of the DDB1 and DDB2 subunits [Fitch et al., 2003;Moser et al., 2005]. The crystal structure of UV-DDB bound to a 6-4PP-containing DNA duplex revealed the direct and exclusive binding of DDB2 to the photodimer [Scrima et al., 2008]. XP-E cells lacking functional DDB2 are deficient in repair of CPD but competent in repair of 6-4PP albeit at reduced rates [Hwang et al., 1999;Moser et al., 2005]. This partial requirement for UV-DDB in GG-NER is reflected in the relative mild sensitivity of XP-E cells to UV-induced cell death [Tang and Chu, 2002]. Although UV-DDB deficiency impairs repair of photolesions in vivo, it is dispensable for NER in vitro [Aboussekhra et al., 1995;Mu et al., 1995;Rapic, V et al., 1998] suggesting that UV-DDB is important for the repair of DNA lesions in a chromatin context.

The UV-DDB complex interacts with several factors known to modulate chromatin structure such as histone acetyltransferase p300, the STAGA complex [Datta et al., 2001;Rapic-Otrin et al., 2002;Martinez et al., 2001] and the Cullin-RING ubiquitin ligase (CRL4) complex CUL4A-RBX1 [Shiyano et al., 1999;Groisman et al., 2003]. The CRL4-DDB2 complex ubiquitylates DDB2 and XPC in response to UV irradiation, which facilitates efficient recognition of photolesions by XPC [Sugasawa et al., 2005]. Moreover, the CRL4 complex also ubiquitylates histones H2A, H3 and H4 [Kapetanaki et al., 2006] of which H3 and H4 ubiquitylation affects nucleosome stability [Wang et al., 2006].

Despite these studies, the molecular mechanisms through which UV-DDB facilitates recognition of DNA damage in chromatin remain poorly understood. Here we purified DDB2...
and associated factors from human cells and identified poly(ADP-ribose) polymerase 1 (PARP1) as a novel component of the UV-DDB complex. We provide evidence for a central role of DDB2-associated PARP1 in mediating poly(ADP-ribose) (PAR) synthesis and recruitment of the SWI/SNF chromatin remodeler ALC1 to UV-damaged DNA. Moreover, we show that poly(ADP-ribose)ylation of DDB2 itself regulates the stability as well as the chromatin retention time of DDB2. Interfering with either PARP1 or ALC1 function impairs CPD repair and renders cells highly sensitive to UV irradiation. Together, these findings outline a novel molecular mechanism for the DDB2-mediated and PARP1-executed regulation of NER.

RESULTS
PARP1 is a component of the UV-DDB complex
To identify novel factors involved in the DNA damage recognition step of GG-NER we isolated DDB2-associated protein complexes by chromatin immunoprecipitation (figure 1A-B) and analysed purified proteins by mass spectrometry (MS). MRC5 cells expressing FLAG-tagged DDB2 were either mock-treated or irradiated with UV-C light (20 J/m²) and incubated for five minutes prior to chromatin isolation and immunoprecipitation using FLAG antibody. MS analysis identified several proteins known to interact with DDB2, including DDB1, CUL4A, CUL4B, and components of the COP9 signalosome (supplementary table 1) showing that native DDB2 is indeed isolated from cells using this approach. Interestingly, multiple DDB2 peptides were identified by MS with protein weights exceeding 50 kDa in UV-irradiated cells but not in mock treated cells (figure 1A), suggesting the presence of UV-specific posttranslational modifications of DDB2. In addition to these known factors, we also identified poly (ADP-ribose) polymerase 1 (PARP1) as a novel DDB2-associated factor. We confirmed the interaction between endogenous DDB2 and endogenous PARP1 by reciprocal immunoprecipitation experiments (figure 1C) using chromatin prepared from UV-irradiated or mock-treated normal human fibroblasts (NHF). These results show that UV irradiation stimulates the interaction between the UV-DDB complex and PARP1 on chromatin. Moreover, an interaction between recombinant UV-DDB and recombinant PARP1 could indeed be detected in vitro supporting a direct interaction between these factors (supplementary figure 1A-B). To test which subunit of the UV-DDB complex interacts with PARP1, we purified GFP-DDB1 or GFP-DDB2 under denaturing conditions from cells. The results revealed that both DDB1 and DDB2 interact with PARP1 (figure 1D). Consistent with this notion, we found that GFP-DDB2<sup>307Y</sup>, which is unable to form a complex with DDB1 (Luijsterburg et al., 2012), also binds PARP1, indicating that DDB2-PARP1 interaction does not require DDB1. To corroborate these findings, we isolated GFP-DDB1 or GFP-DDB2 from polyacrylamide gels and found both extracted proteins to interact with PARP1 in vitro (supplementary figure 1C). Finally, far-western blotting also revealed that both DDB1 and DDB2 avidly bind to PARP1 (supplementary figure 1B). Together, our findings reveal a novel and direct interaction between the UV-DDB complex and PARP1, which prompted us to assess the involvement of PARP1 in modifying DDB2 and regulating NER.
Figure 1. (A) SDS-PAGE electrophoresis and Coomassie staining of FLAG-DDB2 immunoprecipitates obtained from FLAG-DDB2 expressing MRC5 cells mock treated or irradiated with 20 J/m² UV-C. Negative control (NC) indicates the elute obtained from agarose beads incubated with MRC5 FLAG-DDB2 chromatin. The arrows in the zoom-in window indicate the position of the gel where DDB2 and the respective unique peptides detected by MS [A unique peptide is defined as a peptide, irrespective of its length, that exists only in one protein of a proteome of interest]. (B) Western blot of DDB2 and PARP1 immunoprecipitates from NHF cells mock treated or irradiated with 20 J/m² UV-C followed by 5 minutes incubation and immunoblotted against DDB1, DDB2 or PARP1. (C) GFP-DDB2-PARP-1 binding assay. U2OS cells transfected with the indicated GFP constructs were lysed in denaturing buffer and subjected to immunoprecipitation with GFP-TRAP beads and then incubated with 100 ng purified recombinant PARP-1. The beads were then processed for immunoblotting.
Poly(ADP-ribose) chains are synthesized at UV-induced DNA lesions

We first assessed whether poly(ADP-ribose) (PAR) chains are synthesized in chromatin containing UV-induced DNA lesions. To this end, we locally irradiated G0/G1 synchronized telomerase-immortalized human fibroblasts with UV-C light (254 nm) through a polycarbonate mask (Mone et al., 2001). Staining with specific antibodies revealed the presence of PAR chains at sites of DNA damage marked by the recruitment of the p89 subunit of TFIIH or replication factor PCNA known to be involved in NER (figure 2A-B). Moreover, PAR staining in Ki67-negative cells confirmed that PAR synthesis occurred at DNA lesions in non-proliferating cells underlining the replication-independent nature of these events (figure 2C). Finally, chemical inhibition of PARP1 impaired the formation of PAR chains at damaged sites (figure 2D) indicating that the activity of PARP1 is responsible for PAR synthesis at sites of local UV damage.

PARP and DNA synthesis inhibition modulate UV-dependent PARylation

We noted during our experiments that the synthesis of PAR chains at DNA lesions was only detectable in a subset of cells (figure 2A-B). To gain insight into this phenomenon, we locally UV irradiated human fibroblasts with different doses (30 or 100 J/m²) and subsequently monitored the formation of PAR chains 30 minutes after irradiation (figure 2E). The percentage of cells with PAR chains at sites of local damage significantly increased between 30 J/m² and 100 J/m², but did not exceed 50% of the cells (figure 2E-F). It is known that the transient and highly dynamic nature of PAR chains is due to the rapid reversal of this modification by the activity of the poly (ADP-ribose) glycohydrolase (PARG) (Slade et al., 2011). To increase the steady-state level of UV-induced PAR chains, we lowered the levels of PARG by siRNAs. Indeed, depletion of PARG resulted in a significantly elevated percentage of cells that displayed PAR chains at DNA lesions, which was roughly 60% at 30 J/m² and 75% at 100 J/m² (figure 2E-F). These findings show that UV-induced PAR chains are formed in the majority of cells, but are rapidly reversed by the activity of PARG. We then hypothesized that single-stranded DNA gaps transiently generated during NER, might elicit the synthesis of PAR chains. In support of this we found that inhibition of DNA synthesis and ligation by hydroxyurea (HU) and cytosine-β-arabinofuranoside (AraC), known to result in the formation of persistent single-stranded DNA gaps (Overmeer et al., 2011), resulted in robust PAR synthesis in all UV-irradiated cells (figure 2E-F).

DDB2 mediates PARylation during the pre-incision stage of NER

To evaluate whether UV-induced PAR synthesis was exclusively dependent on the presence of a single-stranded DNA repair intermediate, we examined PARylation in XP-A cells that are unable to perform incision and hence do not accumulate ssDNA (Friedberg, 2001) (figure 3C). Indeed, PAR synthesis could not be detected in XP-A cells even following treatment with HU and AraC (figure 3A-B), consistent with a role of dual incision in triggering these events. Strikingly, however, the formation of PAR chains was still detected in XP-A cells upon the depletion of PARG (figure 3A-B), suggesting that PAR chain formation is not solely dependent on the formation of ssDNA. Given our finding that UV-DDB interacts with PARP1, we addressed whether DDB2 contributes to PAR synthesis at sites of DNA damage.
Figure 2. [A-B-C-D] NHF cells were locally UV irradiated (100 J/m²), fixed after the indicated time and stained with an antibody-recognizing PAR, TFIIH, PCNA or Ki67. PAR colocalizes with the damage markers TFIIH and PCNA [A-B] including non-cycling cells [Ki67 negative staining] [C]. Treatment with a specific PARPi (10μM) resulted in a complete loss of PAR signal [D]. Arrows indicate local damage sites. Scale bar represent 20µm. [E] NHF cells were transfected with indicated siRNA or treated with HU/AraC. 48 hours after transfection the cells were locally UV exposed (30 or 100 J/m²), fixed after the indicated time and stained with an antibody recognizing PAR or TFIIH. Scale bar represent 20µm. [F] The percentage of colocalization of PAR with TFIIH in NHF cells is plotted for the different siRNA transfections and HU/AraC treatment. The results are from three independent experiments in which about 100 cells per condition were analysed; Error bars indicate standard deviation. The data shown are from a single representative experiment out of three repeats.
Figure 3. (A) XP-A cells expressing sh-Control or shDDB2 were transfected with indicated siRNA or treated with HU/AraC. 48 hours after transfection the cells were locally exposed to 30 J/m², fixed after the indicated time and stained with an antibody recognizing PAR or TFIIH. Scale bar represent 20µm. (B) The percentage of colocalization of PAR with TFIIH in XP-A cells expressing sh-Control or shDDB2 is plotted for the different siRNA transfections and HU/AraC treatment. The results are from three independent experiments in which about 100 cells per condition were analysed; Error bars indicate standard deviation. (C) Scheme of the early stage of Nucleotide Excision Repair (D) XP-E cells were transfected with indicated siRNA or treated with HU/AraC. The cells were locally UV exposed with 100 J/m², fixed after the indicated time and stained with an antibody-recognizing PAR or TFIIH. Scale bar represent 20µm. (E) The percentage of colocalization of PAR with TFIIH in XP-E cells is plotted for the different siRNA transfections and HU/AraC treatment. Error bars indicate standard deviation. The data shown are from a single representative experiment out of three repeats.
At later time-points after UV irradiation (30 minutes after 100 J/m²), when stretches of ssDNA have been generated, we detected a substantial difference in PAR synthesis between normal human cells and DDB2-deficient XP-E cells as only 10% of the XP-E cells displayed clear PARylation at damaged sites when compared to normal cells (figures 3D-E). However, similar to repair-proficient cells we found inhibition of PAR turnover by PARG depletion to increase the percentage of PAR-positive cells to about 40%, while inhibition of DNA synthesis resulted in PAR synthesis at all locally damaged sites (figure 3D-E). These findings are consistent with the notion that XP-E cells are impaired in dual incision due to deficient repair of CPD and underscore the role of ssDNA formation in UV-induced PAR synthesis. During the time period of 30 minutes after UV-irradiation, a substantial part of repair represents removal of 6-4PP being repaired much more rapidly than CPD. XPE cells display efficient repair of 6-4PP under the conditions described in figure 3 (Moser et al., 2005; Nishi et al., 2009) and all these repair events (in the presence of HU/AraC) provoke PAR synthesis although the absolute number of events is lower than in NHF.

We then took advantage of the finding that DDB2 is very rapidly recruited to UV-induced DNA lesions (Luijsterburg et al., 2007) compared to the assembly rates of the pre-incision factors and especially those of the post-incision factors (Luijsterburg et al., 2010). We therefore examined PAR synthesis very shortly after UV exposure when DDB2 readily accumulates, but PCNA recruitment cannot yet be detected (Luijsterburg et al., 2010). At five minutes after UV irradiation (30 or 100 J/m²), we could not detect PAR synthesis in normal human, XP-A or XP-E fibroblasts, not even when DNA synthesis was inhibited by HU/AraC treatment (figure 4). Strikingly, whereas the stabilization of PAR chains by PARG depletion (supplementary figure 3E) resulted in clearly detectable PARylation at UV-damaged regions in wild-type and XP-A fibroblasts (figure 4A-C-D-E), PAR synthesis at these early time-points was completely abolished in XP-E cells even at 100 J/m² (figure 4B-D). Taken together, our results suggest that two temporally distinct waves of PARylation take place at sites of UV-induced DNA damage. The early [pre-incision] wave of PAR synthesis is fully dependent on functional DDB2 whereas the late [post-incision] wave of PAR formation requires the generation of single-stranded DNA gaps resulting from dual incision. In concordance, we found that PAR synthesis was completely abolished in DDB2 depleted XP-A cells even when PARG was depleted [figures 3A-B and 4C-E, supplementary figure 3C].

PARylation regulates DDB2 release from UV-induced DNA lesions

To gain insight into the role of PARylation in NER complex formation, we investigated the assembly kinetics of GFP-tagged DDB2 at UV-C laser-induced DNA lesions. The kinetics of GFP-DDB2 accumulation were not affected by PARPi or depletion of PARG (supplementary figure 2A-B) indicating that the recruitment of DDB2 is not regulated by PAR chains. We subsequently applied fluorescence loss in photobleaching (FLIP) to measure the dissociation rate of DDB2 from UV-damaged DNA. Although the dissociation of DDB2 (t½ = 19 seconds) was not affected by PARPi, we measured a prolonged chromatin retention time (t½ = 27 seconds) upon knock-down of PARG (figure 5A) suggesting that PAR synthesis positively affects the retention of DDB2 on UV-damaged chromatin. Likewise, the immobilization of GFP-DDB2 following global UV irradiation, as measured by fluorescence
Figure 4. (A-B) NHF(A) and XP-E(B) cells were transfected with indicated siRNA or treated with HU/AraC. 48 hours after transfection the cells were locally UV exposed with 30 or 100 J/m², fixed after the indicated time and stained with an antibody recognizing PAR or TFIIH. Scale bar represent 20µm. (C) XP-A cells expressing shControl or shDDB2 were transfected with indicated siRNA or treated with HU/AraC. 48 hours after transfection the cells were locally UV exposed to 30 J/m², fixed after the indicated time and stained with an antibody-recognizing PAR or TFIIH. Scale bar represent 20µm. (D) The percentage of colocalization of PAR with TFIIH in NHF and XPE cells is plotted for the different siRNA transfections and HU/AraC treatment. The results are from three independent experiments in which about 100 cells per condition were analysed; Error bars indicate standard deviation. (E) The percentage of colocalization of PAR with TFIIH in XP-A cells expressing sh-Control or shDDB2 is plotted for the different siRNA transfections and HU/AraC treatment. The results are from three independent experiments in which about 100 cells per condition were analysed; Error bars indicate standard deviation. The data shown are from a single representative experiment out of three repeats.
recovery after photo-bleaching (FRAP), was significantly reduced following treatment with PARP inhibitors (figure 5B), suggesting that PAR synthesis positively affects the retention of DDB2 on UV-damaged chromatin. Consistent with these findings, western blot analysis revealed that the UV-induced degradation of DDB2 was significantly retarded by PARG depletion (figure 5C) when cells were exposed to a UV dose \(100\text{J/m}^2\) comparable with the UV-laser treatment (figure 5A). Inhibition of PARP activity resulted in accelerated degradation of DDB2 following UV irradiation most clearly seen at 30 J/m². This result indicates that PARylation of DDB2 affects its UV-induced degradation presumably by affecting ubiquitylation. To address how PARylation modulates the chromatin binding and stability of DDB2, we biochemically examined whether UV-DDB is modified by PARP1. In vitro PARylation experiments using purified components revealed that both DDB2 and DDB1 are directly modified by PARP1 (figure 6A, and supplementary figure 2C). Conversely, human DDB2 lacking its first 40 N-terminal amino acids including 7 lysines, (Fischer et al., 2011) failed to undergo PARylation (figure 6A). This finding was further substantiated by the lack of in vitro PARylation of the zebrafish orthologue of DDB2 (drDDB2) lacking the first 93 N-terminal residues (Scrima et al., 2008) showing that the N-terminus of DDB2 is targeted for PARylation. Both Cul4A or Rbx1 were not modified by PARP1 (figure 6A) suggesting that DDB1 and DDB2 are specific targets of PARP1. To assess PARylation of DDB2 in vivo, we expressed double-tagged DDB2 in human cells followed by its isolation using two consecutive purifications under denaturing conditions in order to disrupt protein-protein interactions while preserving post-translational modifications (Figure 6B). Using this purification approach we detected robust PARylation of DDB2 in response to UV irradiation, while PARylation was virtually absent in mock-treated cells showing that DDB2 is modified in a DNA damage-specific manner. Strikingly, inhibition of PARP activity, which resulted in suppressed DDB2 PARylation, was accompanied by increased level and altered spectrum of ubiquitylation of DDB2 (figure 6B). However, it is obvious that lysine residues on the N-terminus of DDB2 are the major target of ubiquitylation (Fischer et al., 2011) and these might have more impact on DDB2 degradation than modification of lysine residues toward the C-terminus when PARP activity is inhibited. These findings identify DDB2 as a novel target for PARP1-mediated PARylation and suggest that poly(ADP-ribose)ylation of DDB2 directly suppresses DDB2 auto-ubiquitylation providing a molecular explanation for the PAR-dependent stabilization of DDB2 in response to UV irradiation.

**DDB2-dependent and -independent recruitment of the chromatin remodeling enzyme ALC1 to UV-induced photolesions**

Recent studies uncovered that PAR chains mediate the recruitment of PAR-binding proteins to single-stranded and double-stranded DNA breaks [Timinszky et al., 2009; Gottschalk et al., 2009; Ahel et al., 2009]. In particular, the macrodomain-containing chromatin remodeling enzyme ALC1 promotes PAR-dependent nucleosome remodeling in vitro and is recruited to sites of DNA breaks, which prompted us to test whether ALC1 is involved in the repair of UV-induced DNA lesions. Staining with a specific antibody revealed that endogenous ALC1 was readily recruited to UV-induced DNA lesions shortly after UV exposure (figure 7A). Live cell imaging of GFP-tagged ALC1-expressing cells confirmed the rapid, but transient
Figure 5. (A) NHF cells stably expressing GFP-DDB2 were transfected with the indicated siRNA or treated with PARPi (10μM). 48 hours after transfection cells were UV irradiated using a UV-C (266 nm) laser. To determine the dissociation kinetics of DDB2 from UV-damaged DNA, the undamaged nucleus was continuously bleached and the fluorescence decrease in the local damage was monitored. Relative fluorescence was normalized at 100% (before bleach at maximum level of accumulation). The half-time ($t_{1/2}$) of a FLIP curve corresponds to the residence time of a protein molecule in the locally damaged area. Error bars indicate SEM. (B) VH10-tert cells stably expressing GFP-DDB2 were incubated in CO2-independent microscopy medium supplemented with 1‰ DMSO (mock treatment) or 10 µM PARP inhibitor dissolved in DMSO three hours prior to FRAP analysis. Cells were mock-treated or globally UV-C irradiated (10 J.m^-2) and transferred to the microscope chamber in microscopy medium. Cells were incubated on the microscope chamber at 37° for 10 minutes to allow repair proteins to accumulate at UV-induced DNA lesions after which the mobility of GFP-tagged NER factors was analyzed by strip-FRAP. The data were normalized to pre-bleach intensity (set to 1) and bleach depth (set to 0). Three independent experiments were performed for each condition. (C) Western blot of normal fibroblasts transfected with the indicated siRNA or treated with PARPi (10μM). Whole cell extracts of non-irradiated and UV-irradiated cells (30–100 J/m²) after the indicated time were probed with antibodies against DDB2, PAR or H2B.
recruitment of ALC1 to sites of UV-C laser-induced DNA damage (figure 7B). GFP-ALC1 was rapidly recruited to UV-induced DNA lesions in wild-type (supplementary figure 3B) and XP-A cells (figure 7B) shortly after exposure to the UV-C laser and could be completely suppressed by addition of the PARPi (supplementary figure 3A). Strikingly, knockdown of DDB2 significantly reduced the recruitment of GFP-ALC1 in XPA-deficient cells (figure 7B) as well as in repair-proficient cells at early time-points after UV irradiation (supplementary figure 3B), suggesting an important role for DDB2 in the recruitment of chromatin remodeler ALC1 through PARP1-mediated PAR synthesis. Consistent with our findings that two mechanistically distinct PARylation waves exist in response to UV irradiation, we found that single-stranded DNA gaps also triggered robust GFP-ALC1 recruitment at later time-points after UV irradiation in normal human as well as in XP-E cells (supplementary figure 4) whereas recruitment of ALC1 was absent in dual incision-defective XP-A cells not even in the presence of HU/AraC (supplementary figure 4). In contrast the stabilization of PAR chains by PARG depletion (supplementary figure 4) resulted in clearly detectable GFP-ALC1 recruitment at UV-damaged regions in XP-A fibroblasts. In summary, our results

Figure 6. (A) The N-terminus of DDB2 is targeted for PARylation. In vitro PARylation experiments using purified components reveal that both DDB2 and DDB1 are directly modified by PARP1. Human DDB2 lacking its first 40 N-terminal amino acids including 7 lysines (sUVDDB), failed to undergo PARylation. The zebrafish orthologue of DDB2 (drDDB) lacking the first 93 N-terminal residues is also not PARylated in vitro. (B) 6His StreptI-tag DDB2 isolation using tandem purifications under denaturing conditions. NHF cells stably expressing 6His StreptI-tag DDB2 were irradiated with UV-C light (100 J/m²) in presence or in absence of PARPi (10µM) or mock-irradiated and incubated for 30 minutes. The final Strept-Tactin column purifications were separated on SDS-PAGE gels, and proteins were visualized with antibodies against DDB2, PAR or Ubiquitin.
Figure 7. (A) NHF cells were locally UV irradiated (100 J/m²), fixed after the indicated time and stained with an antibody recognizing ALC1 or TFIIH. ALC1 colocalizes with the damage marker TFIIH. Scale bars represent 20µm. (B) XP-A cells stably expressing GFP-ALC1 were infected with the indicated short hairpin RNA. The cells were UV damaged using UV-C (266 nm) laser irradiation. GFP fluorescence intensities at the site of UV damage were measured by real time imaging until they reached a maximum. Assembly kinetic curves were derived from at least six cells for each protein. Error bars indicate SEM. Scale bar represent 20µm. (C) Clonal survival of UV-irradiated NHF cells expressing shControl or shALC1 and XPA cells. The percentage of surviving cells is plotted against the applied UV-C dose (J/m²). The results are from three independent experiments; Error bars indicate standard deviation. Scale bar represent 20µm. (D) NHF cells expressing shControl or shALC1 RNAi or treated with PARPi (10μM) were irradiated with 10 J/m² UV-C, fixed immediately, at 8 or 24 hours after UV treatment and stained with anti-CPD antibody. (asterisk indicates p<0.05, ANOVA) (E) NHF cells expressing shControl or shALC1 or treated with PARPi (10μM) were irradiated with 10 J/m² UV-C, fixed immediately, at 1 or 2 hours after UV treatment and stained with an anti-6-4PP antibody. The total fluorescence intensity of the nucleus was quantified and divided by the surface area, resulting in a specific fluorescence intensity expressed in arbitrary units. Values are the result of three independent experiments (100 cells per time point).
reveal that the chromatin remodeling enzyme ALC1 is recruited to PAR chains synthesized during repair by NER through distinct molecular mechanisms.

**PARP1 inhibition and ALC1 depletion impair CPD repair and render cells sensitive to UV irradiation**

Having established that ALC1 is recruited to sites of local damage, we subsequently addressed the biological impact of this finding. To this end, we generated a cell line stably expressing a shRNA targeting endogenous ALC1. Knock-down of ALC1 rendered cells sensitive to UV irradiation compared to control cells [figure 7C], indicating that ALC1 protects cells against UV-induced cytotoxicity. Likewise, chemical inhibition of PARP also rendered cells UV sensitive [supplementary figure 3D], underscoring an important role for PAR synthesis in NER. Finally, we directly measured the repair of UV-induced DNA lesions following a UV dose of 10 J/m² by immunostaining using antibodies against 6-4 PPs or CPDs. While the repair of 6-4 PP was not significantly affected, we measured a significant reduction in CPD repair upon ALC1 depletion or chemical inhibition of PARP1 [figure 7D-E]. Corroborating these findings, an ELISA-based assay confirmed that knock-down of ALC1 conferred a significant reduction in CPD repair [supplementary figure 5]. These findings reveal an unanticipated role of PAR synthesis and ALC1 in efficient repair of CPDs by human NER.

**DISCUSSION**

Despite detailed insights into the NER reaction and the core proteins involved (Gillet and Scharer, 2006; Sugasawa, 2010), the regulatory pathways that govern NER activity in living cells are still poorly understood. Among others, these pathways involve the post-translational modifications of NER proteins and the activity of chromatin remodelling enzymes to optimize repair of DNA damage embedded in chromatin. DDB2 is the first NER factor to be recruited to UV-induced DNA lesions (Luijsterburg et al., 2007) and it regulates NER by direct DNA lesion recognition (Scrima et al., 2008) and modulation of chromatin structure (Palomera-Sanchez and Zurita, 2011). We identified PARP1 as a novel DDB2-associated factor in UV–irradiated cells. The fact that the interaction between these factors occurred in non-diving UV-irradiated human fibroblasts excludes the possibility that involvement of PARP1 in NER is merely related to the stalling of replication forks (Bryant et al., 2009). Although we found PARP1 as a novel DDB2–associated factor, in vitro assays with purified proteins revealed that PARP binds to both DDB1 and DDB2 but with higher presence for DDB2.

We found robust synthesis of PAR chains in nuclear regions containing UV-induced DNA lesions that was completely suppressed by chemical PARP inhibition. These findings directly link PARP1 to the repair of photolesions and fit with previous observations that UV irradiation triggers both stimulation of poly(ADP-ribose) synthesis (Cleaver et al., 1983) and association of PARP1 with UV-photolesions in chromatin (Vodenicharov et al., 2005). Although these findings clearly implicate PARP1 activity in response to UV irradiation, we and others (Bryant et al., 2009; Schultz et al., 2003) failed to detect the recruitment of endogenous PARP1 to UV-induced DNA lesions or UV-induced stalled replication forks,
which is possibly due to transient nature of its interaction or the abundance of PARP1 in
the nucleus (Krishnakumar and Kraus, 2010).

Our study identified two distinct molecular mechanisms that orchestrate the synthesis
of PAR chains at UV-induced DNA damage. Firstly, we show that persistent single-stranded
DNA gaps generated by inhibition of repair synthesis, elicit DDB2-independent PARylation
at NER sites. However, a possible role of PARylation in regulation of post-incision repair
has not yet dissected although recruitment of post-incision factor XRCC1, disassembly
kinetics of NER complexes [Moser et al., 2007] or sealing of UV-induced single-stranded
DNA gaps was not impaired by PARP inhibition [Cleaver et al., 1983]. Secondly and more
importantly, we show that DDB2 regulates fast and transient PARylation at sites of UV-
induced DNA damage during the pre-incision stage of NER. One target of PARylation is
DDB2 itself as shown by PARylated DDB2 purified from UV-irradiated cells. It is likely that
this modification underlies among others the UV-specific occurrence of DDB2 peptides
of larger than 50 KDa molecular weight in immunoprecipitates of chromatin bound
DDB2. The initial DDB2-mediated wave of PAR synthesis does not require incision and is
regulated by the activity of PARG. Indeed, there is increasing evidence that DNA breaks
are not an absolute requirement for PARP1 activation [Krishnakumar and Kraus, 2010].
Several alternative mechanisms to activate PARP1 in the absence of DNA breaks have
been proposed, including interaction with other proteins [Cohen-Armon et al., 2007],
or post-translational modifications such as phosphorylation, acetylation [Hassa et al.,
2003; Rajamohan et al., 2009], SUMOylation and ubiquitylation [Martin et al., 2009;Messner
et al., 2009]. In this light, it is feasible that the DDB2-associated E3 ubiquitin ligase activity
[Shiyanov et al., 1999;Groisman et al., 2003] might activate PARP1. At the same time, it
is possible that PARP1 activation is modulated by DDB2-mediated acetylation through
its interaction with histone acetyltransferases p300 and the STAGA complex [Datta et al.,
2001;Rapic-Otrin et al., 2002;Martinez et al., 2001].

Molecular mechanisms for DDB2-mediated and PARP1-executed regulation of NER
Our data provide mechanistic insights into how DDB2 promotes NER in chromatin through
two novel mechanisms. On one hand, DDB2 is directly targeted by poly(ADP-ribosyl)
ation and ubiquitylation [Fischer et al., 2011] in response to UV irradiation. As PARylation
[Messner et al., 2010] and ubiquitylation are targeted to lysine and both modifications
appear to occur in the same N-terminal region of DDB2, competition between PARylation
and ubiquitylation of target lysine residues might constitute an important mechanism of
DDB2 mediated regulation of NER. The in vivo data supports a competition model in which
the poly(ADP-ribosyl)ation of DDB2 results in increased protein stability and a prolonged
chromatin retention time on the UV lesion. At the same time poly(ADP-ribosyl)ation of DDB2
suppresses its UV-induced ubiquitylation and consequently leads to reduced degradation
of DDB2. Together our data disclose a mechanism by which two opposing modifications
regulate the steady-state levels and retention time of DDB2 at sites of UV-photolesions.

On the other hand, DDB2-dependent PARylation events also stimulate the pre-incision
step of NER. We show that DDB2-dependent PARylation through PARP1 at UV-induced DNA
lesions targets chromatin remodeler ALC1 to these sites. ALC1 belongs to the Swi2/Snf2
ATPase superfamily and through its macrodomain, interacts transiently with chromatin that is modified by PARP1. We propose that these protein modifications serve to locally modulate chromatin structure through PARP1-stimulated nucleosome sliding to promote NER [Gottschalk et al., 2009; Ahel et al., 2009]. Consistent with this scenario, we show that loss of ALC1 or chemical inhibition of PARP1 resulted in defects in the repair of CPDs concomitantly with increased sensitivity to UV exposure, underscoring the importance of the DDB2-PARP1-ALC1 pathway in promoting NER. We noted that the defects in CPD repair due to loss of ALC1 or PARP1 activity are less pronounced than repair defects caused by loss of DDB2 [Pines et al., 2009], suggesting that the essential role of DDB2 in CPD repair is not solely due to its recruitment of PARP1-mediated activities, but also involves other functions of DDB2 such as its ubiquitin ligase activity. Interestingly, XPC contains a putative PAR binding sequence [Gagne et al., 2008], suggesting that UV-DDB dependent PAR may promote the accessibility of UV lesions through remodeling of the chromatin structure as well as providing an enhancer signal for the recruitment of preincision NER proteins.

**A model of DDB2 and PARP1 dependent regulation of NER**

The high affinity of DDB2 for DNA and its preference for UV-damaged DNA makes UV-DDB the most important DNA damage recognition factor for 6-4PP and CPD [Wittschieben et al., 2005]. UV-DDB is the first NER factor to be recruited to UV damage [Luijsterburg et al., 2007; Nishi et al., 2009] as part of the Cullin-RING ubiquitin ligase [CRL4] complex CUL4A-RBX1 [Figure 7]. As shown for 6-4PP [Scrima et al., 2008], the CUL4A-RBX1 complex binds to photolesions by the WD40 domain of DDB2 and in concert with PARP1 tightly regulates steady-state levels and retention time of DDB2 by opposing modifications [PARylation and ubiquitylation] of the same N-terminal region of DDB2. The enhanced extension time of PARylated DDB2 on UV-damage might be particularly important for CPD photolesions that induce much less disruption of base pairing interactions than 6-4PP [Kim and Choi, 1995] and fully depend on functional DDB2 for their repair. Purified [nonPARylated] DDB2 recognizes CPD and 6-4PP with a 5 and 80 fold higher affinity respectively than nondamaged DNA [Wittschieben et al., 2005]; this affinity of DDB2 for CPD might too low for productive repair i.e. recruitment of XPC. We speculate that extended binding of PARylated DDB2 to CPD will provoke the induction of chromatin modifications at the site of DNA damage to allow productive interaction and ubiquitylation of XPC and UV-DDB required for NER [Sugasawa et al., 2005]. Underscoring the importance of PAR synthesis in the assembly of the pre-incision NER complex, we found that PARP inhibition leads to reduced recruitment of the pre-incision factor XPC as shown in our recent work [Luijsterburg et al., 2012] whereas depletion of PARG stimulates binding of XPC [Luijsterburg et al., 2012] and TFIIH [data no shown]. Whether UV-DDB activates PARP1 through its E3 ubiquitin ligase activity or whether PARP1 activation occurs in parallel with or precedes ubiquitylation is not clear. PARP1 mediated increase of retention time of DDB2 at UV damage and DDB2 protection by suppressing its ubiquitylation-dependent degradation argue for PARylation as the initiating event. Moreover, PARP1 might create accessibility for recruitment of NER factors by its ability to disrupt chromatin structure by PARylation of histones and destabilizing nucleosomes [Krishnakumar and Kraus, 2010]. Additionally
PARylation of chromatin effectuates recruitment of NER promoting factors such as the Swi2/Snf2 chromatin remodeler ALC1 to UV damaged DNA to locally modulate chromatin structure through nucleosome sliding (Figure 8) thereby stimulating the recruitment of XPC to assemble a functional repair complex. Our study also identified DDB2-independent PARylation and recruitment of ALC1 at NER sites that is triggered by transient single-stranded DNA gaps generated by the dual incision step of NER (figure 8); this process of PARylation is amplified by inhibition of DNA repair synthesis. The role of PARylation and ALC1 in regulation of post-incision step of NER remains to be resolved.

Figure 8: Model of DDB2 and PARP 1 dependent regulation of NER. UV-DDB is the first NER factor to be recruited to UV damage as part of the Cullin-RING ubiquitin ligase (CRL4) complex CUL4A-RBX1. This complex binds to UV damage and both DDB1 and DDB2 might be involved in binding of PARP1. In concert with PARP1, the CUL4A-RBX1 complex tightly regulates steady-state levels and retention time of DDB2 by opposing modifications (PARylation and ubiquitylation) of the same N-terminal region of DDB2. Additionally, PARP1-dependent PARylation of chromatin also effectuates recruitment of the Swi2/Snf2 chromatin remodeler ALC1 to UV damaged DNA in order to locally modulate chromatin structure through nucleosome sliding thereby stimulating the recruitment of XPC. The second distinct waves of PARylation and ALC1 recruitment requires the generation of single-stranded DNA gaps resulting from dual incision.
**EXPERIMENTAL PROCEDURES**

**Cell culture and UV-C irradiation**

The following cell lines were used for this study: VH10 hTert (normal human fibroblast, NHF), XP25RO hTert (XP-A), GM01389 hTert (XP-E), U2OS and MRC5 fibroblast. Cells were grown in DMEM supplemented with 10% fetal calf serum, penicillin and streptomycin. Two days prior to experiments medium was changed to DMEM supplemented with 0.2% serum fetal calf serum, penicillin and streptomycin. UV-radiation of cells was carried out using a 254 nm UV-C source. Local irradiation was performed using 5 μm filters as described previously [Volker et al., 2001]. UV lamp–induced damage was inflicted using a 254 nm UV source [TUV PL-S 9W; Philips]. For induction of global UV damage, cells were rinsed with PBS and irradiated with 8 or 16 J/m2. For induction of local UV damage, cells were UV irradiated through a polycarbonate mask (Millipore) with pores of 8 μm and subsequently irradiated with 30, or 100 J/m2.

For induction of global UV damage, cells were rinsed with PBS and irradiated with 8 or 16 J/m2. For induction of local UV damage, cells were UV irradiated through a polycarbonate mask (Millipore) with pores of 8 μm and subsequently irradiated with 30, or 100 J/m2. The AZ12640831-009 PARPi was used at a final concentration of 10 μM and was a gift from AstraZeneca. Cells were pretreated 30 minutes before irradiation.

**Generation of cell lines**

ALC1 and DDB2 cDNA were cloned into vector pENTR4-GFP-C1 [E.Campeau; addgene: w392-1] and were subsequently recombined into pLenti6.3 V5-DEST [Invitrogen] using gateway recombination. VH10 hTert or XP25RO hTert fibroblasts were transduced with pLenti6.3 GFP-ALC1 or pLenti6.3 GFP-DDB2 lentiviral particles and cultured with 5 μg/ml blasticidin [Invivogen] to select for integrands.

For DDB2 isolation a 6 His and strepII-tag were fused to the N-terminus of DDB2. A synthetic oligo coding for 6His StrepII-tag was inserted into pENTR4 [invitrogen] and DDB2 cDNA was subsequently cloned in. Lentiviral particles were generated after recombination of this vector to pLenti6.3 V5-DEST and used for transducing VH10 hTert cells.

NHF and XP-A fibroblasts stably expressing short hairpin RNA (shRNA) were generated by lentiviral transduction of control, ALC1 or DDB2 targeting constructs followed by 1 μg/ml puromycin selection. The following shRNA vectors were used: TRCN0000013471 (ALC1); TRCN00000083995 (DDB2) and SHC002 [non-targeting control] from the RNAi Consortium [Sigma-Aldrich].

**RNA interference**

Short interfering RNA [siRNA] duplexes used were as follows: smartpool siRNA targeting the PARG transcript and smartpool non-targeting siRNA [Dharmacon]. Cells were transfected using Hiperfect [Qiagen] according to the manufacturer’s protocol. For PARG knockdown two sequential transfections were performed. Immunostaining and western-blot experiments were performed 48 hours after the final transfection.

**Complex isolation**

Isolation of DDB2 complex was performed according to published procedures with some modifications [Groisman et al., 2003]. Briefly, cells were irradiated with UV-C at 20 J/m², and incubated for 5 minutes. To prepare nuclear extracts, cells were suspended in hypotonic buffer [10 mM Tris-HCl pH 7.3, 10 mM KCl, 1.5 mM MgCl2, 10
mM β-mercaptoethanol, and 0.2 mM PMSF) and disrupted by Dounce homogenization. Nuclei were collected by centrifugation at 2000g for 15 min at 4°C and resuspended in extraction buffer (15 mM Tris-HCl pH 7.3, 1 mM EDTA, 0.21 M NaCl, 1 mM MgCl2, 10% glycerol, 10 mM β-mercaptoethanol, and 0.2 mM PMSF). After incubating on ice for 30 minutes, the samples were centrifuged at 20,000g for 30 min at 4°C, and the supernatant was used as the nuclear extract fraction. The nuclear pellet fraction was washed and resuspended in the micrococcal nuclease buffer (20 mM Tris-HCl pH 7.5, 100 mM KCl, 2 mM MgCl2, 1 mM CaCl2, 0.3 M sucrose, 0.1% Triton X-100, and complete protease inhibitor cocktail [Roche]). Micrococcal nuclease was added at 3 U/ml and the samples incubated for 10 min at room temperature, whereupon the reaction was terminated by adding 5mM EGTA and 5mM EDTA. The samples were centrifuged at 2000g for 5 min at 4°C, and the supernatant was used as the solubilized chromatin fraction. The UV-DDB complex was immunoprecipitated from solubilized chromatin prepared from MRC5 cells expressing FLAG-DDB2 by incubating with M2 anti-FLAG agarose overnigh with rotation. After an extensive wash with wash buffer (0.1 M KCl, 20 mM Tris-HCl pH 8.0, 5 mM MgCl2, 10% glycerol, 1 mM PMSF, 0.1% Tween 20, 10 mM β-mercaptoethanol), the bound proteins were eluted from M2 agarose by incubation for 30 min with FLAG peptide (0.2 mg/ml in PBS). The elution procedure was repeated three times.

**In-gel tryptic digestion**

Immunoprecipitates were separated on SDS-PAGE gels, and proteins were visualized with Coomassie [SimplyBlue, Invitrogen]. Gel lanes were sliced into 25–30 bands, cut into small pieces and washed with 25 mM NH4HCO3 followed by two rounds of dehydration with 100% acetonitrile for 10 min. For reduction and alkylation, gel particles were first incubated with 10 mM dithiothreitol for 30 minutes at 56°C. Following dehydration with acetonitrile, gel plugs were subsequently incubated in 55 mM iodoacetamide for 20 minutes at room temperature. After two rounds of washing with 25 mM NH4HCO3 and dehydration with 100% acetonitrile, the gel particles were completely dried in a centrifugal vacuum concentrator [Eppendorf, Hamburg, Germany]. Dried gel particles were re-swollen for 15 min. on ice by addition of 15 µl of a trypsin solution [12.5 ng/µl in 25 mM NH4HCO3, Sequencing grade modified trypsin, Promega, Madison, WI]. Following this, 20 µl of 25 mM NH4HCO3 was added and samples were kept on ice for an additional 30 min. Tryptic digestion was subsequently performed overnight at 37 °C. Following tryptic digestion, the overlaying digestion-solution was collected. Two additional rounds of extraction with 20 µl 0.1% TFA were used to extract peptides from the gel plugs and all extracts were pooled.

**Nano LC ESI MS/MS**

Nanoflow LC was performed on an Ultimate LC system [Dionex, Sunnyvale, CA]. A volume of 10 µL of sample was injected onto a C18 PepMapTM 0.3 mm×5 mm trapping column [Dionex] and washed with 100% A [2% acetonitrile in 0.1% formic acid in MQ water, v/v] at 20 µL/min for 15 min. Following valve switching, peptides were separated on a C18 PepMap 75 µm×150 mm column [Dionex] at a constant flow of 200 nL/min. The peptide elution gradient was from 10 to 60% B [95% acetonitrile in 0.1% formic acid in MQ water v/v] over
50 min. The nanoflow LC system was coupled to an HCTultra IonTrap (Bruker Daltonics, Bremen, Germany) using a nano-electrospray ionisation source. The spray voltage was set at 1.2 kV and the temperature of the heated capillary was set to 165 °C. Eluting peptides were analyzed using the data dependent MS/MS mode over a 300–1500 m/z range. The five most abundant ions in an MS spectrum were selected for MS/MS analysis by collision-induced dissociation using helium as the collision gas.

**Mass spectrometry data analysis**

Peak lists were generated using DataAnalysis 4.0 software [Bruker Daltonics] and exported as Mascot Generic (MGF) files. These files were searched against the human IPI database using the Mascot (version 2.2.1) search algorithm [Matrix Science, London, UK] An MS tolerance of 0.6 Da (with # 13C = 1) and a MS/MS tolerance of 0.5 Da was used. Trypsin was designated as the enzyme and up to one missed cleavage site was allowed. Carbamidomethylcysteine was selected as a fixed modification and oxidation of methionine as a variable modification.

**Immunofluorescent labelling (IF) and Western blotting (WB)**

The cells were fixed with methanol/acetone (50%/50%) for 10 minutes at 4°C. After an extensive wash with PBS the cells were incubated for 60 minutes at room temperature with buffer contains 0.5% BSA and 0.05% Tween-20 in PBS. Antibody incubations were performed at room temperature and cells were counterstained with DAPI. Images were captured with a Zeiss Axioplan2 microscope equipped with a Zeiss Axiocam MRm camera using either a Plan-NEOFLUAR 40×/1.30 or 63×/1.25 objective. Fluorescence intensity of randomly captured images was quantified using Zeiss Axiovision software. For total extract the cells were lysed directly in Laemli-SDS-sample buffer. Western blot analysis was performed as described previously [Fousteri et al., 2006] and protein bands were visualised via chemiluminescence [ECL-Plus, Amersham Biosciences] using Horseradish Peroxidase (HP)-conjugated secondary antibodies or via Odyssey Infrared Imaging System [LI-COR] using secondary antibodies labeled with IR fluorophores [LI-COR]. The following antibodies were used: mouse α-DDB2 at 1:500 (IF)- 1:1000 (WB) [MyBioSource]; mouse α-Parp1 at 1:1000 [WB] (Abnova); mouse α-Poly (ADP-Ribose) at 1:100 [IF and WB] (Abcam); mouse α-ALC1 at 1:500 (IF)- 1:1000 (WB) (Abcam); mouse α-GFP at 1:5000 [WB] (Roche); goat α-DDB1 at 1:1000 [WB] (Abcam); goat α-DDB2 at 1:1000 [WB] (Santa-Cruz); rabbit α-Poly (ADP-Ribose) at 1:100 [IF and WB] (BD pharmingen); rabbit α-H2B at 1:5000 [WB] (Santa-Cruz); rabbit α-PARG at 1:1000 [C-term] [WB] (Origene); mouse α-6-4PP and α-CPD at 1:1000 (IF) [CosmoBio] Alexafluor 488 and 555 conjugated antibodies were purchased from Invitrogen.

**Live cell confocal laser-scanning microscopy**

Confocal laser-scanning microscopy images were obtained using a confocal microscope [LSM 510 META] with a 63× oil Plan Apochromat 1.4 NA oil immersion lens [Carl Zeiss, Inc.] equipped with a cell culture microscopy stage. GFP fluorescence imaging was recorded after excitation with a 488-nm argon laser and a 515–540-nm band-pass
filter. Fluorescence loss in photobleaching (FLIP) was performed as described previously [Houtsmuller and Vermeulen, 2001; Zotter et al., 2006]. Kinetics of GFP-tagged ALC1, and DDB2 accumulation were performed using a UV-C (266 nm) laser irradiation as described previously [Dinant et al., 2007]. Briefly, VH10-tert cells stably expressing GFP-DDB2 and GFP-ALC1 were incubated in CO2-independent microscopy medium [137 mM NaCl, 5.4 mM KCl, 1.8 mM CaCl2, 0.8 mM MgSO4, 20 mM D-glucose, 20 mM HEPES and 10% FCS] and 2 mW pulsed (7.8 kHz) diode pumped solid state laser emitting at 266 nm (Rapp OptoElectronic, Hamburg GmbH) was used for local UV-C irradiation. To determine the dissociation kinetics of DDB2 from UV-damaged DNA, the undamaged nucleus was continuously bleached and the fluorescence decrease in the local damage was monitored. Relative fluorescence was normalized at 100% (before bleach at maximum level of accumulation). The half-time ($t_{1/2}$) of a FLIP curve corresponds to the residence time of a protein molecule in the locally damaged area. Images obtained with the confocal microscope were analyzed using AIM software (Zeiss). Fluorescence levels were determined for the specified region where damage was induced in addition to the complete nucleus. From these datapoints the relative amount of protein in the damaged area was determined in time.

**Fluorescence recovery after photobleaching (FRAP)**

VH10-tert cells stably expressing GFP-DDB2 were incubated in CO2-independent microscopy medium [137 mM NaCl, 5.4 mM KCl, 1.8 mM CaCl2, 0.8 mM MgSO4, 20 mM D-glucose, 20 mM HEPES and 10% FCS] supplemented with 1‰ DMSO (mock treatment) or 10 µM PARP inhibitor dissolved in DMSO three hours prior to FRAP analysis. Cells were subsequently rinsed with PBS, mock-treated or globally UV-C irradiated (10 J.m$^{-2}$) and transferred to the microscope chamber in microscopy medium (supplemented with DMSO or PARP inhibitor). Cells were incubated on the microscope chamber at 37° for 10 minutes to allow repair proteins to accumulate at UV-induced DNA lesions after which the mobility of GFP-tagged NER factors was analyzed by strip-FRAP. Briefly, FRAP analysis was performed by bleaching (5 iterations) a narrow strip [512 x 40 pixels at zoom 8] spanning the nucleus with maximal 488 nm laser intensity (AOTF 100%). The re-equilibration of bleached and non-bleached molecules was monitored in a region of 512 x 50 pixels (zoom 8) with low laser intensity (0.5% for GFP-DDB2) for at least 700 images with a 38 ms time-interval between images. The data were normalized to pre-bleach intensity [set to 1] and bleach depth [set to 0]. Three independent experiments were performed for each condition.

**UV survival**

Cellular survival of VH10 hTert shControl, VH10 hTert shALC1, and XP-A cells was determined using a colony assay. Cells were plated in 10-cm and after 16 hours cells were exposed to UV-C [254 nm; TUV lamp; Philips] and left to grow for 14 days, fixed, and stained with methylene blue. Colonies were counted to assess the colony-forming ability.

**In vitro polyADP-riboosylation assay**

The assay was performed according to published procedures [Deng et al., 2005] using recombinant proteins purified as described in Fischer et al. [Fischer et al., 2011].
**DDB2 purification**

VH10 hTert cells stably expressing 6His StreptII-tag DDB2 were irradiated with UV-C light (100 J/m²) or mock-irradiated and incubated for 30 minutes. Cells were collected and lysed in lysis buffer [8 M urea, 2 M NaCl, 25 mM Tris pH 8, 1 mM MgCl₂, 0.2 % Triton]. The lysates were diluted ≥ 7 times, after which 25 benzonase units were added per ml. After incubating at room temperature for 30 minutes samples were centrifuged at 16,000g for 10 minutes. TALON beads (Clontech) were added to the supernatants and incubated for 4 hr at room temperature. After an extensive wash with wash buffer [8 M urea, 25 mM Tris pH 8, 1 mM MgCl₂, 20 mM imidazole], the 6His StreptII-tag DDB2 was eluted by overnight incubation with the elution buffer [8 M urea, 25 mM Tris pH 8, 500 mM imidazole, 1 % SDS]. The elutes were concentrated by Vivaspin centrifugal concentrators (Sartorius) and diluted in Strep-Tactin buffer [100 mM Tris pH 8, 1 mM EDTA, 150 mM NaCl]. A second purification step was performed using Strep-Tactin spin columns according to the manufacturer’s protocol (IBA). Elutes were separated on SDS-PAGE gels, and proteins were visualized by Western-blot.

**GFP-DDB2-PARP-1 binding assay.**

U2OS cells transfected with GFP constructs for 24 hours were lysed in denaturing buffer [20 mM Tris, pH 7.5; 50 mM NaCl; 0.5% NP-40; 1% Sodium Deoxycholate; 1% SDS; 1 mM EDTA, Benzonase final conc. 0.25U/μl] containing protease inhibitor cocktails (Roche) and subjected to immunoprecipitation with GFP-TRAP beads (Chromotek) for 1 hours at RT. The beads were then washed extensively in a buffer [20 mM Tris, pH 7.5; 50 mM NaCl; 0.5% NP-40; 0.5% Sodium Deoxycholate; 0.5% SDS; 1 mM EDTA] that disrupts protein-protein interactions, followed by two washes in EBC buffer [50 mM Tris, pH 7.5; 150 mM NaCl; 0.5% NP-40; 1 mM EDTA], and incubated with 100 ng purified, recombinant PARP-1 [Sigma] for 2 hours at RT. The beads were then washed thoroughly in EBC buffer and processed for immunoblotting.

**In vitro co-immunoprecipitation**

UV-DDB, ΔUV-DDB (DDB2 lacking its first 40 N-terminal amino acids), GFP and PARP-1 recombinant protein were used to test direct interaction in vitro. The reaction volume was adjusted to 400 μL in EBC buffer [50 mM Tris, pH 7.5; 150 mM NaCl; 0.5% NP-40; 1 mM EDTA], and 0.5 μg of anti-PARP-1 antibody was added. The mixture was incubated and rotated at 4°C for 3 hours. The antigen-antibody complex was captured by incubation with 15 μl of protein A-agarose beads [GE Healthcare] for 2 hr in cold room. The beads were washed extensively in EBC buffer, and resuspended in 20 μL of Laemili sample buffer and processed for immunoblotting.

**Farwestern analysis**

100 ng of UV-DDB and 1000 ng of GFP recombinant proteins were separated by SDS-PAGE and transferred to a PVDF membrane. The proteins on the membrane were denaturated for 10 min with a 6 M guanidine hydrochloride [GuHCl] solution in HBB buffer [10 mM HEPES, pH 7.5, 60 mM KCl, 1 mM EDTA and 1 mM DTT]. Proteins were then renaturated in the same HBB buffer with progressively decreasing GuHCl concentration. The membrane was rinsed extensively in HBB and blocked for 1 hour in blocking solution. Following the
membrane was incubated with PARP-1 recombinant protein (10 µg/ml) in HBB for 16 h. Unbound proteins were removed with extensive washes for 30 min in the same buffer. The PARP-1 binding was visualised by Western-blot.

**CPD/6-4 PP ELISA**

Cells were plated in 96 well plates, irradiated with 10 J/m² UV and incubated for various periods to allow cells to repair DNA photolesions. The cells were fixed with methanol/acetone (50%/50%) for 10 minutes. After an extensive wash with PBS the cells were incubated for 3 minutes at room temperature with 10 mM NaOH. The cells were rinsed extensively in PBS and incubated for 60 minutes at room temperature with buffer contains 0.5% BSA and 0.05% Tween-20 in PBS. The plates were sequentially incubated with TDM-2 or 64M-2 antibodies specific for CDP or 6-4PP, respectively, and secondary antibody conjugated with horseradish peroxidase (HRP). After washings, substrate solution Turbo TMB-ELISA (Pierce) was added to the plates and incubated for 15–30 min. Absorbance at 490 nm was measured using a microplate reader after the addition of 2 M H₂SO₄.

**Online supplemental material**

Table 1 shows the proteins identified in Flag immunoprecipitates material from FLAG-DDB2 expressing MRC5 cells mock treated or irradiated with UV-C. Fig. S1 shows direct interaction in vitro between DDB2 and PARP-1. Fig. S2 shows that the kinetics of GFP-DDB2 accumulation were not affected by PARPi or depletion of PARG. Fig. S3 shows transient recruitment of GFP-ALC1 to sites of UV-C laser-induced DNA damage. Fig. S4 demonstrates that single-stranded DNA gaps also triggered robust GFP-ALC1 recruitment at later time-points after UV irradiation in normal human as well as in XP-E cells, whereas recruitment of ALC1 was absent in dual incision-defective XP-A cells. Fig. S5 shows a significant reduction in CPD repair upon ALC1 depletion or chemical inhibition of PARP1.

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PARP DEPENDENT REGULATION OF NER

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Table 1. Proteins identified in Flag immunoprecipitates material from FLAG-DDB2 expressing MRC5 cells mock treated or irradiated with 20 J/m² UV-C. A unique peptide is defined as a peptide, irrespective of its length, that exists only in one protein of a proteome of interest.

| Protein | Protein Score | Unique peptides | Protein | Protein Score | Unique peptides |
|---------|---------------|----------------|---------|---------------|----------------|
| DDB2    | 274           | 9              | DDB2    | 305           | 13             |
| DDB1    | 462           | 24             | DDB1    | 263           | 16             |
| PARP1   | 172           | 6              | CUL4B   | 221           | 9              |
| CUL4B   | 65            | 4              | PARP1   | 126           | 6              |
| GPS1    | 65            | 1              | COPS4   | 103           | 3              |
| CUL4A   | 44            | 1              | CUL4A   | 83            | 1              |
| GPS1    | 65            | 1              | COPS7A  | 64            | 1              |
Figure 1S. [A] In vitro co-immunoprecipitation. UVDDDB, ΔUVDDDB (ΔDDB2 lacking its first 40 N-terminal amino acids) and PARP-1 recombinant proteins were used to test direct interaction in vitro. GFP recombinant protein was used as negative control. [B] Far-western assay. UVDDDB recombinant proteins were separated by SDS-PAGE, transferred to a membrane and incubated with PARP-1 recombinant protein [10 µg/ml]. The PARP-1 binding was visualised by Western-blot. GFP recombinant protein was used as negative control. [C] GFP-DDB2-PARP-1 binding assay. U2OS cells transfected with the indicated GFP constructs were lysed in denaturing buffer and subjected to immunoprecipitation with GFP-TRAP beads. The elute was separated by SDS-PAGE and GFP-DDB1 or GFP-DDB2 were isolated from polyacrylamide gels [red square]. The proteins were subjected to immunoprecipitation with GFP-TRAP beads and incubated with 100 ng purified recombinant PARP-1. The beads were processed for immunoblotting.
Figure 2S. (A) Real-time recruitment of GFP-DDB2 in NHF at the site of DNA damage using UV-C (266 nm) laser irradiation. Scale bar represent 7.5µm. (B) NHF cells stably expressing the GFP-DDB2 were transfected with the indicated siRNA or treated with PARPi (10μM). 48 hours after transfection the cells were UV damaged using UV-C (266 nm) laser irradiation. GFP fluorescence intensities at the site of UV damage were measured by real time imaging until they reached a maximum. Assembly kinetic curves were derived from at least six cells for each protein. Relative fluorescence was normalized at 0 (before damage) and 100% (maximum level of accumulation). Error bars indicate SEM. (C) In vitro PARylation experiments using purified components. Antibody against the N-terminal of DDB2 revealed that DDB2 is directly modified by PARP1. DDB2 lacking its first 40 N-terminal amino acids (ΔUV-DDB) was not detectable.
Figure 3S. (A) Real-time recruitment of GFP-ALC1 in NHF in presence or absence of PARPi (10μM) at the site of DNA damage using UV-C (266 nm) laser irradiation. Scale bar represent 7.5μm. [B] NHF cells stably expressing GFP-ALC1 were infected with the indicated short hairpin RNA. The cells were UV damaged using UV-C (266 nm) laser irradiation. GFP fluorescence intensities at the site of UV damage were measured by real time imaging until they reached a maximum. Assembly kinetic curves were derived from at least six cells for each protein. Error bars indicate SEM. Scale bar represent 20μm. [C] Anti-DDB2 and histone H2B western blots of total lysates from NHF and XP-A cells expressing shRNAs targeting DDB2 or a non-targeting shControl [mock]. [D] Clonal survival of UV-irradiated NHF cells in presence or absence of PARPi (1μM). The percentage of surviving cells is plotted against the applied UV-C dose (J/m2). The results are from three independent experiments; Error bars indicate standard deviation. [E] Anti-PARG and histone H2B western blots of total lysates from XP-A, XP-E and NHF transfected with siRNA targeting PARG or a non-targeting siControl [mock].
Figure 4S. (A) NHF, XP-E and XP-A cells stably expressing GFP-ALC1 transfected with indicated siRNA or treated with HU/AraC were locally UV exposed to 30 J/m², fixed after the indicated time and stained with an antibody recognizing TFIIH. Scale bars represent 20µm. (B) The percentage of colocalization of GFP-ALC1 with TFIIH in NHF, XP-A and XP-E cells after 0.5 hours UV local damage is plotted for the different siRNA transfections and HU/AraC treatment. The results are from three independent experiments in which about 100 cells per condition were analysed; Error bars indicate standard deviation. (C) The percentage of colocalization of GFP-ALC1 with TFIIH in NHF, XP-A and XP-E cells after 5 minutes UV local damage is plotted for the different siRNA transfections and HU/AraC treatment. The results are from three independent experiments in which about 100 cells per condition were analysed; Error bars indicate standard deviation. The data shown are from a single representative experiment out of three repeats.
Figure S5. (A-B) NHF cells expressing shControl or shALC1 or treated with PARPi (10μM) were irradiated with 10 J/m² UV-C; CPDs and 6-4 PPs were detected immediately and 24 or 2 hours respectively after UV treatment by ELISA assay. Error bars indicate standard deviation. (C-D) XPE cells siRNA transfected or treated with PARPi (10μM) were irradiated with 10 J/m² UV-C; CPDs and 6-4 PPs were detected immediately and 24 or 2 hours respectively after UV treatment by ELISA assay. Error bars indicate standard deviation. The results are from three independent experiments.
