

## **The histone chaperones FACT and ANP32E reshape the chromatin landscape during DNA damage repair through H2A.Z/H2A.X histone variant exchange**

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## SUMMARY

Safeguarding cell function and identity following a genotoxic stress challenge entails a tight coordination of DNA damage signaling and repair with chromatin maintenance. How this coordination is achieved and with what impact on chromatin integrity remains elusive. Here, by investigating the mechanisms governing the distribution of H2A.X in mammalian chromatin, we demonstrate that this histone variant, which plays a central role in damage signaling, is deposited *de novo* at damage sites in a repair synthesis-coupled manner. Our mechanistic studies further identify the histone chaperone FACT (Facilitates Chromatin Transcription) as responsible for the deposition of newly synthesized H2A.X and orchestrating, together with ANP32E (Acidic Nuclear Phosphoprotein 32 Family Member E), a H2A.Z/H2A.X exchange reaction, which reshapes the chromatin landscape at repair sites. We propose that this mechanism promotes chromatin accessibility to repair machineries and helps tailoring damage signaling to repair progression.

## HIGHLIGHTS

- H2A.X, but not H2A.Z, is deposited *de novo* at sites of DNA damage repair
- FACT promotes new H2A.X deposition coupled to repair synthesis
- FACT-mediated H2A.X deposition is dispensable for DNA damage repair
- FACT and ANP32E chaperones orchestrate H2A.Z/H2A.X exchange in UV-damaged chromatin

## INTRODUCTION

Cells are constantly exposed to genotoxic stress, and respond by activating dedicated DNA damage signaling and repair pathways that safeguard genome stability (Aguilera and García-Muse, 2013; Ciccia and Elledge, 2010; Hoeijmakers, 2009; Jackson and Bartek, 2009). Damage signaling consists in the initiation of cell cycle checkpoint kinase cascades upon DNA damage detection/processing to coordinate cell cycle progression with DNA repair (Lazzaro et al., 2009; Shaltiel et al., 2015). Adding another layer of complexity to these highly orchestrated cellular responses, damage signaling and repair machineries operate on chromatin substrates in eukaryotic cell nuclei, where DNA wraps around histone proteins (Luger et al., 2012). Importantly, chromatin landscapes, characterized by specific patterns of histone variants (Buschbeck and Hake, 2017; Talbert and Henikoff, 2017), post-translational modifications (Bannister and Kouzarides, 2011) and various degrees of chromatin folding, convey epigenetic information that instructs cell function and identity through the regulation of gene expression programs (Allis and Jenuwein, 2016). While the importance of maintaining epigenome integrity is widely recognized, it is still unclear whether and how this is achieved during the DNA damage response.

DNA damage signaling and repair indeed elicit profound chromatin rearrangements, challenging epigenome maintenance (Dabin et al., 2016), with a transient destabilization of chromatin organization, accompanied by DNA damage-induced changes in histone modifications (Dantuma and van Attikum, 2016), followed by restoration of chromatin structure (Polo and Almouzni, 2015; Smerdon, 1991). It is still elusive if chromatin restoration after damage is an entirely faithful process or if genotoxic stress responses contribute to significant reshaping of the epigenetic landscape, leaving a signature of DNA damage. Although not fully characterized yet, restoration of damaged chromatin involves the deposition of newly synthesized histone variants, as shown for H2A, H3.1 and H3.3 at sites of

UVA and UVC damage in human cells (Adam et al., 2013; Dinant et al., 2013; Luijsterburg et al., 2016; Polo et al., 2006). Mechanistically, new histone deposition employs dedicated histone chaperones (Hammond et al., 2017), including the H3 variant-specific chaperones HIRA (Histone Regulator A) and CAF-1 (Chromatin Assembly Factor-1), which are recruited to UV-damaged chromatin by distinct repair proteins (Adam et al., 2013; Polo et al., 2006). Less is known about the factors that govern H2A dynamics in damaged chromatin, apart from a role for the histone chaperone FACT in promoting H2A-H2B turnover at UVC damage sites (Dinant et al., 2013). Regarding the histone variant H2A.Z, its dynamic exchange at sites of DNA double-strand breaks in human cells involves the concerted action of the histone chaperone ANP32E and of the chromatin remodeling factors p400 and INO80 (Inositol-requiring 80) (Alatwi and Downs, 2015; Gursoy-Yuzugullu et al., 2015; Xu et al., 2012).

Here, we investigate how such repair-coupled chromatin rearrangements may leave an imprint on the epigenetic landscape and cross-talk with damage signaling by focusing on the H2A.X histone variant, which represents the ancestral form of H2A, conserved in all eukaryotes (Talbert and Henikoff, 2010). Making up only 10 to 25% of total H2A, H2A.X is nevertheless central to the DNA damage response, owing to a particular carboxyl-terminal Serine, in position 139 in mammals, that is targeted by DNA damage responsive kinases (Bonner et al., 2008; Rogakou et al., 1998). Importantly, H2A.X S139 phosphorylation spreads at distance from the damage (Rogakou et al., 1999), which is key for amplifying the DNA damage signal through the coordinated recruitment of DNA damage checkpoint mediators (Altmeyer and Lukas, 2013; Smeenk and van Attikum, 2013). Best described in response to DNA double-strand breaks, this signaling cascade also operates following other types of genomic insults including UV irradiation. Indeed, UV damage processing triggers checkpoint signaling (Marini et al., 2006) by activating the ATR (ataxia telangiectasia-mutated and Rad3-related) kinase, which phosphorylates H2A.X (Hanasoge and Ljungman, 2007).

A salient feature of H2A.X phosphorylation is that it occurs *in situ* in damaged chromatin (Rogakou et al., 1998). The original distribution of H2A.X in chromatin is thus a critical determinant of the damage response, as it will govern the distribution of the phosphorylated form, known as  $\gamma$ H2A.X. A commonly held view is that H2A.X is phosphorylated at DNA damage sites but incorporated ubiquitously in chromatin, independently of DNA damage. However, recent ChIP-seq studies in mammalian cells have challenged this view by revealing a non-random distribution of H2A.X, with enrichments at active transcription start sites, and sub-telomeric regions in activated human lymphocytes (Seo et al., 2014; 2012) and at extra-embryonic genes in mouse pluripotent stem cells (Wu et al., 2014). Yet, the mechanisms underpinning the non-random distribution of H2A.X in chromatin are unknown, as is their potential connection to the DNA damage response.

In this study, by investigating the mechanisms governing H2A.X distribution in chromatin during the repair of UVC damage in mammalian cells, we reveal that H2A.X histones are deposited *de novo* in damaged chromatin by the histone chaperone FACT in a repair synthesis-coupled manner. We also uncover a H2A.Z/H2A.X exchange reaction orchestrated by FACT and ANP32E chaperones, which reshapes the chromatin landscape by altering the histone variant pattern at UV damage sites.

## RESULTS

### ***De novo* deposition of H2A histone variants at repair sites**

In order to characterize H2A.X deposition pathways, we monitored *de novo* histone deposition using the SNAP-tag technology (Bodor et al., 2012) in human U2OS cell lines stably expressing SNAP-tagged H2A variants (Figure 1A and Figure S1 for a characterization of the cell lines). Specific labeling of newly synthesized histones combined with local UVC irradiation did not reveal any detectable accumulation of new H2A variants at UVC damage sites, contrary to what we observed with newly synthesized H3.3 (Adam et al., 2013) (Figure 1B). We reasoned that this discrepancy might be due to differences in mobility between inner and outer core histones, the latter being more readily exchanged (Kimura and Cook, 2001; Louters and Chalkley, 1985). Since outer core histone mobility is partly transcription-dependent (Jackson, 1990; Kimura and Cook, 2001), we tracked new histones in the presence of transcription inhibitors - Flavopiridol or DRB (Bensaude, 2011). Note that short-term transcription inhibition reduces but does not abolish histone neosynthesis. Thus, we revealed new H2A.X accumulation at sites of UVC damage in the vast majority of cells (Figure 1D-E). We recapitulated our observations in mouse embryonic fibroblasts (Figure S2A-D) and obtained similar results upon transcription inhibition with the RNA Polymerase II inhibitor  $\alpha$ -amanitin (Figure S3A). Importantly, new H2A.X accumulation at UVC damage sites was not an artifact of transcription inhibition as it was also detectable in the absence of transcription inhibitors upon exposure to higher UVC doses (Figure S3B). In contrast to new H2A.X, no significant enrichment was observed at damaged sites when staining for total H2A.X (Figure 1D), showing that this accumulation was specific for newly synthesized histones and most likely reflects histone exchange at damage sites. Noteworthy, *de novo* accumulation of H2A.X was also observed at sites of UVA laser micro-irradiation (Figure S3C), arguing that it is not unique to the UVC damage response.

In order to assess whether new H2A.X deposition at UVC damage sites was specific for the damage-responsive histone H2A.X, we extended our analyses to other H2A variants, namely canonical H2A and another replacement variant conserved in all eukaryotes, H2A.Z, considering both H2A.Z.1 and H2A.Z.2 forms, which display different dynamics in response to UVA laser damage in human cells (Nishibuchi et al., 2014). For this, we established U2OS cell lines that stably express comparable levels of SNAP-tagged H2A variants (Figure S1). Similar to what observed for H2A.X, we detected *de novo* accumulation of the canonical replicative variant H2A, but not of the replacement variants H2A.Z.1 and H2A.Z.2, at UVC damage sites (Figure 1E). Similar results were obtained without transcription inhibition (Figure S3D), pointing to a specific histone deposition mechanism that is not general to all H2A variants.

Collectively, these data reveal a transcription-independent deposition of newly synthesized H2A and H2A.X but not H2A.Z histone variants at repair sites.

### ***De novo* deposition of H2A histone variants at replication foci**

In addition to its accumulation at sites of DNA damage, new H2A.X displayed a punctuate pattern in a subset of cells upon transcription inhibition, which corresponded to replication foci, as shown by co-staining with the replication-coupled histone chaperone CAF-1 (Figure 2A, S3A). We observed new H2A.X accumulation at replication foci throughout S phase (Figure 2B). We recapitulated these results in mouse embryonic fibroblasts, where transcription inhibitors could be omitted to visualize new H2A.X accumulation in replicating heterochromatin domains, which are poorly transcribed by nature (Figure S2E-F). Similar to H2A.X accumulation at repair sites, the enrichment observed at replication foci was specific for newly synthesized H2A.X (Figure 2C) and only detected for H2A and H2A.X, but not H2A.Z variants (Figure 2D).

Altogether, these findings demonstrate that newly synthesized H2A and H2A.X are deposited both at repair sites and at replication foci.

### **New H2A.X deposition is independent of S139 phosphorylation**

We next examined the importance of H2A.X S139 phosphorylation status for new H2A.X deposition. We first verified that the SNAP-tag in C-terminal did not prevent H2A.X phosphorylation (data not shown) and we compared the dynamics of H2A.X wild-type to S139 phospho-mimetic and phospho-deficient mutants. These experiments consistently showed that new H2A.X accumulation at UV sites (Figure 1E) and replication foci (Figure 2D) occurred irrespective of H2A.X S139 phosphorylation status. Since SNAP-tagged H2A.X proteins are expressed in the context of wild-type endogenous H2A.X, we used a complementary approach to address the importance of H2A.X phosphorylation: we treated cells with an ATR kinase inhibitor, which inhibits UV-induced H2A.X phosphorylation as reported (Hanasoge and Ljungman, 2007), but does not impair the *de novo* deposition of H2A.X at UVC-damage sites (Figure S4A). Thus, *de novo* deposition and phosphorylation of H2A.X in response to DNA damage are mechanistically uncoupled.

### **New H2A.X deposition is coupled to replicative and repair synthesis**

To gain insights into the mechanisms underlying new H2A.X deposition at repair sites and replication foci, we investigated a possible dependency on DNA synthesis, which is a common feature of replication and repair. We prevented repair synthesis by down-regulating the repair factor XPG (Xeroderma Pigmentosum, group G), an endonuclease involved in the excision of the UVC-damaged oligonucleotide (Figure 3A). XPG knock-down markedly reduced the accumulation of newly synthesized H2A.X at sites of UVC damage (Figure 3B) with no detectable effect on new H2A.X deposition at replication foci (data not shown). We



confirmed that new H2A.X deposition at repair sites was dependent on the DNA synthesis machinery by knocking-down of the DNA polymerase-associated factor PCNA (Proliferating Cell Nuclear Antigen, Figure 3C). In line with these observations at repair sites, we also uncovered a dependency on DNA synthesis for new H2A.X deposition occurring at replication foci. Indeed, when we inhibited replicative synthesis with aphidicolin, the focal patterns of new H2A.X deposition in S phase were strongly reduced (Figure 3D). These results indicate that the deposition of newly synthesized H2A.X at replication foci and UVC damage sites is dependent on replicative and repair synthesis.

### **New H2A.X deposition is not coupled to new H3 deposition**

We next investigated whether new H2A.X deposition was coupled to the deposition of newly synthesized H3 variants that occurs at replication and repair sites (Adam et al., 2013; Polo et al., 2006; Ray-Gallet et al., 2011). Knocking down the histone chaperone CAF-1, responsible for new H3.1 deposition at UV sites (Polo et al., 2006), did not significantly affect new H2A.X accumulation at sites of UVC damage repair (Figure S4B). Similar results were obtained upon down-regulation of the histone chaperone HIRA, which deposits new H3.3 at UV sites (Adam et al., 2013) (Figure S4B) and upon loss-of-function of both pathways simultaneously (data not shown). These data demonstrate that new H2A.X deposition occurs independently of new H3 deposition by CAF-1 and HIRA at UVC damage sites.

### **The histone chaperone FACT promotes new H2A.X deposition at UV damage sites**

To uncover the molecular determinants of new H2A.X accumulation at sites of UVC damage repair, we examined the effect of knocking down candidate H2A histone chaperones (Gurard-Levin et al., 2014; Hammond et al., 2017) and chromatin remodelers (Zhou et al., 2016). We focused on the histone chaperone FACT (Belotserkovskaya et al., 2003) and on the chromatin

remodeler INO80 (Shen et al., 2000). Indeed, the latter is known to maintain the levels of chromatin-bound H2A.X in human cells (Seo et al., 2014) and to promote H2A deposition in yeast (Papamichos-Chronakis et al., 2006), while FACT incorporates H2A and H2A.X but not H2A.Z into chromatin (Heo et al., 2008) and stimulates H2A turnover at UVC damage sites in human cells (Dinant et al., 2013). Moreover, FACT associates with replisome components in human cells (Alabert et al., 2014; Tan et al., 2006), is involved in replication-coupled nucleosome assembly in yeast (Yang et al., 2016), and is critical for replisome progression through chromatin *in vitro* (Kurat et al., 2017). Interestingly, we observed that FACT accumulation at sites of DNA replication and UV damage repair exhibited dependencies to replicative and repair synthesis similar to new H2A.X deposition (Figures 4A-C and S5A). Furthermore, FACT trapping to chromatin by the intercalating agent curaxin CBL0137 (Gasparian et al., 2011; Safina et al., 2017) (Figure S5B) impaired FACT accumulation and new H2A.X deposition at UVC damage sites (Figure S5C-D). In line with these findings, siRNA-mediated down-regulation of FACT subunits, but not of the remodeler INO80, led to a marked reduction of newly deposited H2A.X at UVC damage sites (Figure 4D). The decrease in new H2A.X deposition was also observed in the whole nucleus, most likely reflecting replication- and transcription-coupled deposition of H2A.X by FACT, and cannot be explained by reduced histone neo-synthesis as we verified that FACT knock-down does not inhibit nascent transcription (Figure S6A). *De novo* accumulation of H2A was similarly reduced in FACT-knocked-down cells (Figure S6B). These results establish that new H2A and H2A.X deposition at repair sites is promoted by the histone chaperone FACT.

### **New H2A.X deposition by FACT does not stimulate DNA damage repair**

Having identified FACT as a key factor for new H2A.X deposition, we set out to determine the functional relevance of new H2A.X deposition by FACT in damaged chromatin. We first

tested whether FACT down-regulation might impact UVC damage repair. We observed that FACT knock-down did not impair the recruitment of the repair factor XPA (Xeroderma Pigmentosum, group A) to UVC damage sites (Figures 4D and S6B). In addition, we found no significant effect of FACT loss-of-function on repair synthesis (Figure 5A), in line with previous data (Dinant et al., 2013) and on the timely removal of UV photoproducts (Figure 5B), ruling out any major effect of this histone chaperone on UVC damage repair. Similarly, preventing new H2A.X synthesis by siRNA (Figure 5C) did not detectably impair UVC damage repair synthesis (Figure 5D). Together, these experiments do not reveal any significant impact of FACT-mediated deposition of new H2A.X on UVC damage repair.

### **H2A.Z/H2A.X exchange in UVC-damaged chromatin**

The *de novo* deposition of H2A and H2A.X but not H2A.Z that we observed at sites of UV damage repair (Figure 1E) parallels the reported specificity of FACT towards H2A variants (Heo et al., 2008). We thus decided to examine whether this selective histone variant deposition by FACT may result in an altered histone variant pattern in UVC-damaged chromatin. Supporting this idea, when we stained for total histones, we observed that while H2A.X total levels were not detectably altered at sites of UVC irradiation (as also seen in Figure 1D), H2A.Z total levels were reduced by about 10% in damaged chromatin (Figure 6A-B). Importantly, the local depletion of H2A.Z was detectable with different antibodies targeting H2A.Z (amino- and carboxy-termini) and upon fluorescent labeling of total H2A.Z-SNAP (Figure 6A-B), ruling out the possibility of impaired detection due to post-translational modification of H2A.Z. Mechanistically, such selective depletion affecting the H2A.Z histone variant and not H2A.X at damage sites is unlikely to result from a local decompaction of chromatin. Furthermore, H2A.Z loss from UVC-damaged chromatin is not a mere consequence of transcription inhibition at UV sites because transcription inhibition with

flavopiridol does not significantly reduce the levels of chromatin-bound H2A.Z (Fig. S7A), suggesting that H2A.Z is actively removed from UVC-damaged chromatin. When exploring the molecular determinants of H2A.Z loss, we found that the local depletion of H2A.Z occurred independently of FACT and INO80 but was dependent on the histone chaperone ANP32E, previously characterized for its ability to remove H2A.Z from chromatin (Mao et al., 2014; Obri et al., 2014). Notably, ANP32E-mediated removal of H2A.Z was dispensable for FACT recruitment and new H2A.X deposition at UVC damage sites (Figure S7B-C). Thus, ANP32E and FACT independently orchestrate a histone variant exchange reaction in UVC-damaged chromatin resulting in the maintenance of H2A.X and the loss of H2A.Z.

## DISCUSSION

By analyzing the dynamics of H2A histone variants in damaged chromatin, we have uncovered a H2A.Z/H2A.X exchange reaction controlled by two histone chaperones, ANP32E and FACT (Figure 6C). Chromatin restoration after UVC damage repair thus entails reshaping of the chromatin landscape through a change in histone variant pattern. We also reveal that H2A.X is deposited *de novo* at sites of DNA damage repair, a process which should be integrated in the DNA damage response, in addition to the well-characterized phosphorylation of H2A.X. We propose that repair-coupled deposition of new H2A.X constitutes an important additional regulation level for fine-tuning the DNA damage response. By ensuring that the phosphorylation target for DNA damage checkpoint kinases is present in sufficient amount and at the right place, new H2A.X deposition may allow an efficient and timely response to genotoxic insults in chromatin regions that are susceptible to DNA damage.

### **New H2A.X deposition at sites of DNA synthesis during replication and repair**

In our study, we uncover a selective deposition of newly synthesized H2A variants at sites of UVC damage repair, with H2A and H2A.X being deposited *de novo* in damaged chromatin while H2A.Z is not. This parallels our previous findings on H3 variants showing that H3.1 and H3.3 are deposited *de novo* in UVC-damaged chromatin in contrast to CENP-A (Adam et al., 2013; Polo et al., 2006). Nevertheless, we demonstrate that new H2A.X deposition is not coupled with new H3.1/H3.3 deposition at repair sites. These results are in line with pioneering observations that newly synthesized H2A and H3 do not deposit within the same nucleosomes in replicating cells (Jackson, 1987), and suggest that new H2A.X-H2B associate with parental H3-H4 in repaired chromatin. We have also observed a selective deposition of

H2A.X and H2A but not H2A.Z variants on replicating DNA, which is consistent with a recent study in mouse pericentric heterochromatin domains showing a replication-dependent deposition of new H2A and not H2A.Z (Boyarchuk et al., 2014), and with recent proteomic analyses of histone variants associated with newly replicating DNA in human cells (Alabert et al., 2015). The coupling between new H2A.X deposition and DNA replication is intriguing considering that H2A.X is not a prototypical replicative histone variant. Indeed, H2A.X is synthesized at basal levels throughout the cell cycle in mammalian cells, in a replication-independent manner (Wu and Bonner, 1981). Nevertheless, the lack of introns and the existence of a non-polyadenylated form of H2A.X mRNA - both characterizing replicative histone transcripts - place H2A.X in a unique position between replicative and replacement histone variants (Mannironi et al., 1989). Functionally, the deposition of newly synthesized H2A.X at replication foci may ensure that H2A.X is not diluted out during replication, which would happen if only the bona fide replicative variant H2A was deposited. Such replication-coupled deposition of H2A.X could also explain the increase in soluble H2A.X detected in human cells treated with replication inhibitors (Liu et al., 2008). Furthermore, the *de novo* deposition of H2A.X that we have uncovered at repair sites provides a likely molecular basis for the H2A.X hotspots found in chromatin regions that are susceptible to endogenous damage (Seo et al., 2014). We can speculate that the observed enrichment of H2A.X may be a consequence of repeated genomic insults. H2A.X could thus mark chromatin regions that are more susceptible to damage to facilitate subsequent damage responses. Noteworthy, the H2A.X protein was also shown to be rapidly stabilized following DNA damage, contributing to a local enrichment at repair sites (Atsumi et al., 2015). However, this stabilization required H2A.X phosphorylation on Ser139, while we found that new H2A.X deposition at repair sites occurred independently of H2A.X phosphorylation. H2A.X protein stabilization and *de novo*

deposition thus appear to be two distinct and complementary mechanisms contributing to the distribution of H2A.X in chromatin, which reflects DNA damage experience.

### **Role of the histone chaperone FACT in new H2A.X deposition**

We have identified FACT as the responsible histone chaperone for new H2A and H2A.X deposition at repair sites. Interestingly, FACT also promotes the deposition of another H2A variant, macroH2A1.2, at sites of replication stress in mammalian cells (Kim et al., 2017). However, FACT does not incorporate H2A.Z in chromatin (Heo et al., 2008), which is consistent with the lack of new H2A.Z deposition at sites of DNA synthesis. We demonstrate that FACT-mediated deposition of new H2A.X at damage sites is coupled to repair synthesis and dependent on the DNA polymerase sliding clamp PCNA. It will be interesting to examine whether FACT directly associates with PCNA. This is an attractive hypothesis considering that the FACT subunit SSRP1 (Structure-Specific Recognition Protein 1) harbors a non-canonical PIP (PCNA interacting protein) box (Mailand et al., 2013). Alternatively, FACT interaction with PCNA could be mediated by PCNA-binding factors like XRCC1 (X-ray repair cross-complementing protein 1)(Fan et al., 2004), which contributes to UVC damage repair (Moser et al., 2007), and was recently reported to associate with the FACT complex subunit SSRP1 (Gao et al., 2017). In addition, other factors associated with the DNA synthesis machinery and shown to cooperate with FACT, like MCM (MiniChromosome Maintenance) proteins (Tan et al., 2006), could also contribute to FACT recruitment to sites of UV damage repair.

### **Impact of new H2A.X deposition on DNA damage signaling and repair**

Although H2A.X is deposited *de novo* at sites of UVC damage repair, we have observed that this histone variant is not required for repair of UVC lesions. In this respect, our findings are

in line with previous studies in mouse embryonic stem cells knocked-out for H2A.X, which display increased genomic instability but are not particularly sensitive to UV irradiation (Bassing et al., 2002). Similar to H2A.X, the histone chaperone FACT is dispensable for repair synthesis at UV damage sites. This contrasts with FACT requirement for replicative DNA synthesis (Abe et al., 2011; Kurat et al., 2017; Okuhara et al., 1999), suggesting that FACT-mediated chromatin disassembly/re-assembly activity is more critical for replication fork progression through chromatin than for DNA synthesis *per se*.

Besides an effect on DNA repair, FACT-mediated deposition of new H2A.X could impact DNA damage signaling. We can hypothesize that FACT may either potentiate damage signaling by bringing in more phosphorylation targets for DNA damage responsive kinases or contribute to turn off damage signaling at the end of repair by replacing  $\gamma$ H2A.X by unphosphorylated H2A.X and H2A. In this respect, the histone chaperones ASF1 and CAF-1 are required for checkpoint termination in response to DNA double-strand breaks (Chen et al., 2008; Diao et al., 2017; Kim and Haber, 2009), suggesting that the re-establishment of nucleosomal arrays contributes to turning off DNA damage signaling after repair of DNA damage. It is tempting to speculate that FACT-mediated histone deposition at repair sites may similarly help coordinate repair synthesis with checkpoint termination. Furthermore, we cannot exclude that the repair-coupled deposition of H2A.X may be functionally important for signaling during a subsequent damage response by maintaining critical levels of H2A.X in chromatin regions that are susceptible to DNA damage.

### **H2A.Z/H2A.X exchange at sites of DNA damage repair**

Not only have we shown that H2A.X is deposited *de novo* at sites of UV damage repair but we have also observed that this is paralleled by the removal of H2A.Z, resulting in a local change in histone variant pattern. It still remains to be determined if the histone variant



exchange takes place at the level of the nucleosome, H2A.X replacing H2A.Z, or at the level of the damaged chromatin domain. H2A.Z is removed from UVC-damaged chromatin by the histone chaperone ANP32E, showing interesting similarities with what has been reported in response to UVA laser damage (Gursoy-Yuzugullu et al., 2015). However, in this case ANP32E activity results in H2A.Z returning to basal levels in damaged chromatin after a transient accumulation while we observe a local loss of H2A.Z at UVC damage sites. We did not find any significant role for the remodeler INO80 in H2A.Z removal from UVC-damaged chromatin, in contrast to what observed at sites of UVA laser damage in human cells (Alatwi and Downs, 2015), suggesting that different types of DNA lesions may engage different chromatin remodeling machineries.

Given that H2A.Z promotes chromatin folding *in vitro* (Fan et al., 2002; 2004), the local depletion of H2A.Z in damaged chromatin could contribute to the chromatin relaxation observed at UVC damage sites (Adam et al., 2016; Luijsterburg et al., 2012). In line with this hypothesis, H2A.Z removal by ANP32E is required for histone H4 acetylation at sites of DNA double-strand breaks (Gursoy-Yuzugullu et al., 2015).

### **Dynamics of H2A variants at repair sites and transcription regulation**

While we have shown that the *de novo* deposition of H2A and H2A.X at repair sites was not transcription-mediated, it could impact transcription regulation in damaged chromatin. In particular, given the reported function of FACT in regulating transcription recovery after UVC damage in human cells (Dinant et al., 2013), FACT-mediated deposition of new H2A variants in UVC-damaged chromatin could facilitate the coordination of repair synthesis with transcription restart. Tipping the balance towards H2A and H2A.X as opposed to H2A.Z could also help keeping transcription in check and avoiding transcription interference with repair because H2A.Z was proposed to poise genes for transcription activation (Zhang et al.,

2005) and to promote unscheduled/cryptic transcription when mislocalized in yeast cells (Jeronimo et al., 2015). It is thus tempting to speculate that H2A.Z/H2A.X histone variant exchange at UV sites may contribute to mitigate transcription-repair conflicts by maintaining transcription inhibition during repair synthesis. In this respect, it is worth mentioning that FACT was shown to limit the formation of RNA:DNA hybrids in yeast and human cells (Herrera-Moyano et al., 2014). It would thus be of major interest to investigate whether this function of FACT relies on the *de novo* deposition of H2A histone variants on newly synthesized DNA.

New H2A.X deposition in damaged chromatin could also have a broader impact on transcription regulation. Indeed, several recent studies have uncovered a critical role for H2A.X in regulating gene expression patterns during cell differentiation. In particular, H2A.X is involved in silencing of extra-embryonic genes in mouse embryonic stem cells (Wu et al., 2014), and also represses genes governing the epithelial-mesenchymal transition in human cells (Weyemi et al., 2016). Therefore, alterations in H2A.X distribution as a consequence of DNA damage experience may impact cell fate determination via the rewiring of transcriptional programs, thus contributing to the reported effect of DNA damage repair on cell differentiation and reprogramming (Rocha et al., 2013).

## **EXPERIMENTAL PROCEDURES**

### **Cell culture and drug treatments**

All U2OS and NIH/3T3 cell lines were grown at 37°C and 5% CO<sub>2</sub> in Dulbecco's modified Eagle's medium (DMEM, Invitrogen) supplemented with 10% fetal calf serum (EUROBIO), 100 U/mL penicillin and 100 µg/mL streptomycin (Invitrogen) and the appropriate selection antibiotics (Table S1).

Transcription inhibition was performed either by adding 5,6-Dichlorobenzimidazole 1-beta-D-ribofuranoside (DRB, 100 µM final concentration, Sigma-Aldrich), alpha-Amanitin (AMA, 20 µg/mL final concentration, Sigma-Aldrich) or Flavopiridol hydrochloride hydrate (FLV, 10 µM final concentration, Sigma-Aldrich) to the culture medium at 37°C before the analysis.

Inhibition of replicative synthesis was performed by adding Aphidicolin (Aphi, 1 µg/mL final concentration, Sigma-Aldrich) to the culture medium at 37°C 5 h before harvesting the cells.

For ATR inhibition, cells were incubated with 2 µM ATR inhibitor AZ20 (Selleckchem) 1h before subsequent cell treatment. For FACT trapping, cells were incubated with 2 µM CBL0137 (Bertin Pharma) 15 min before subsequent cell treatment.

### **SNAP-tag labeling of histone proteins**

For specific labeling of newly-synthesized histones, pre-existing SNAP-tagged histones were first quenched by incubating cells with 10 µM non-fluorescent SNAP reagent (SNAP-Cell Block, New England Biolabs) for 30 min (quench) followed by a 30 min-wash in fresh medium and a 2 h-chase. The SNAP-tagged histones neo-synthesized during the chase were then pulse-labeled by incubation with 2 µM red-fluorescent SNAP reagent (SNAP-Cell TMR-Star, New England Biolabs) for 15 min followed by 1h to 1h30-wash in fresh medium. Cells were pre-extracted with Triton detergent before fixation (see Immunofluorescence section for

details). If cells were subject to local UVC irradiation, irradiation was performed immediately before the pulse. When transcription inhibitors were used, they were added to the medium at the quench step and kept throughout the experiment. For total labeling of SNAP-tagged histones, the quench step was omitted and cells were pulsed with SNAP-Cell TMR-Star immediately before harvesting.

### **siRNA and plasmid transfections**

siRNAs purchased from Eurofins MWG Operon (Table S2) were transfected into cells using Lipofectamine RNAiMAX (Invitrogen) following manufacturer's instructions. The final concentration of siRNA in the culture medium was 50 nM. Cells were analyzed and/or harvested 48 to 72 h post-transfection. For stable cell line establishment, cells were transfected with plasmid DNA (1 µg/ml final) using Lipofectamine 2000 (Invitrogen) according to manufacturer's instructions 48 h before antibiotic selection of clones. All constructs were verified by direct sequencing and/or restriction digests. Cloning details and primer sequences (Sigma-Aldrich) are available upon request. Plasmids are described in Table S3.

### **Local UVC irradiation**

Cells grown on glass coverslips (Menzel Gläser) were irradiated with UVC (254 nm) using a low-pressure mercury lamp. Conditions were set using a VLX-3W dosimeter (Vilbert-Lourmat). For local irradiation (Katsumi et al., 2001; Moné et al., 2001), cells were covered with a polycarbonate filter (5 µm pore size, Millipore) and irradiated with 150 J/m<sup>2</sup> UVC unless indicated otherwise.

### **UVA laser micro-irradiation**

Cells grown on glass coverslips (Menzel Gläser) were presensitized with 10  $\mu$ M 5-bromo-2'-deoxyuridine (BrdU, Sigma-Aldrich) for 24 hr at 37°C. Damage was introduced with a 405 nm laser diode (3 mW) focused through a Plan-Apochromat 63X/1.4 oil objective to yield a spot size of 0.5-1  $\mu$ m using a LSM710 NLO confocal microscope (Zeiss) and the following laser settings: 40% power, 50 iterations, scan speed 12.6  $\mu$ sec/pixel.

### **Cell extracts and western blot**

Total extracts were obtained by scraping cells in Laemmli buffer (50 mM Tris HCl pH 6.8, 1.6% SDS (Sodium Dodecyl Sulfate), 8% glycerol, 4%  $\beta$ -mercaptoethanol, 0.0025% bromophenol blue) followed by 5 min denaturation at 95°C. Cytosolic and nuclear extracts were obtained as previously described (Martini et al., 1998). The chromatin fraction was prepared by addition of benzonase (Novagen) to the pellet after nuclear extraction.

For western blot analysis, extracts were run on 4%–20% Mini-PROTEAN TGX gels (Bio-Rad) in running buffer (200 mM glycine, 25 mM Tris, 0.1% SDS) and transferred onto nitrocellulose membranes (Amersham) with a Trans-Blot SD semidry transfer cell (Bio-Rad). Total proteins were revealed by reversible protein stain (Pierce). Proteins of interest were probed using the appropriate primary and HRP (Horse Radish Peroxidase)-conjugated secondary antibodies (Jackson ImmunoResearch), detected using Super-Signal West Pico or Femto chemi-luminescence substrates (Pierce). Alternatively, when fluorescence detection was used instead of chemi-luminescence, total proteins were revealed with REVERT total protein stain, secondary antibodies were conjugated to IRDye 680RD or 800CW and imaging was performed with Odyssey Fc-imager (LI-COR Biosciences) (see Table S4 for the list of antibodies).

### **Immunofluorescence, image acquisition and analysis**

Cells grown on sterile round glass coverslips 12 mm diameter, thickness No.1.5 (Menzel Gläser) were either fixed directly with 2% paraformaldehyde (PFA, Electron Microscopy Sciences) and permeabilized with 0.5% Triton X-100 in PBS or first pre-extracted with 0.5% Triton X-100 in CSK buffer (Cytoskeletal buffer: 10 mM PIPES pH 7.0, 100 mM NaCl, 300 mM sucrose, 3 mM MgCl<sub>2</sub>) and then fixed with 2% PFA to remove soluble proteins (not bound to chromatin). For CPD staining, DNA was denatured with 0.5 M NaOH for 5 min. For PCNA staining, cells were treated with Methanol (Prolabo) for 15 min at -20°C. Samples were blocked in 5% BSA (Bovine Serum Albumin, Sigma-Aldrich) in PBS supplemented with 0.1% Tween 20 (Euromedex) before incubation with primary antibodies and secondary antibodies conjugated to Alexa-Fluor 488, 594, 647 or 680 (Invitrogen) (see Table S4 for a description of the antibodies).

Coverslips were mounted in Vectashield medium with DAPI (Vector Laboratories) and observed with a Leica DMI6000 epifluorescence microscope using a 40x or 63x oil objective. For confocal images, we used a Zeiss LSM710 confocal microscope with a Plan-Apochromat 63x/1.4 oil objective. Image J (U. S. National Institutes of Health, Bethesda, Maryland, USA, <http://imagej.nih.gov/ij/>) and ICY (Institut Pasteur, Paris, France, [icy.bioimageanalysis.org/](http://icy.bioimageanalysis.org/)) softwares were used for image analysis.

### **Visualization of Replicative and Repair Synthesis**

To visualize replicative synthesis, EdU (Ethylnyl-deoxyUridine) was incorporated into cells during 15 min (10 μM final concentration) and revealed using the Click-iT EdU Alexa Fluor 594 (or 647) Imaging kit (Invitrogen) according to manufacturer's instructions. To visualize repair synthesis, EdU (10 μM final concentration) was incorporated into cells during 4 h immediately after local UVC irradiation and revealed using the same kit, before CPD labeling by immunofluorescence.

### **Nascent RNA labeling**

Cells were incubated in DMEM supplemented with 0.5 mM EU (EthylnylUridine) for 45 min, rinsed in cold medium and in PBS before fixation in 2% paraformaldehyde. EU incorporation was revealed with Click-iT RNA Imaging kits (Invitrogen) using Alexa Fluor 594 dye according to manufacturer's instructions.

### **Flow cytometry**

Cells were fixed in ice-cold 70% ethanol before DNA staining with 50 µg/mL propidium iodide (Sigma-Aldrich) in PBS containing 0.05% Tween 20 and 0.5 mg/mL RNaseA (USB/Affymetrix). DNA content was analyzed by flow cytometry using a FACS Calibur Flow Cytometer (BD Biosciences) and FlowJo Software (TreeStar).

### **Statistical analyses**

Percentages of positively stained cells were obtained by scoring at least 100 cells in each experiment unless indicated otherwise. In experiments using transcription inhibitors, only cells showing above background TMR levels were scored. Statistical tests were performed using GraphPad Prism. Loss or enrichment of H2A.Z at UV sites relative to the whole nucleus was compared to a theoretical mean of 1 by one-sample t-tests. P-values for mean comparisons between two groups were calculated with a Student's t-test with Welch's correction when necessary. Multiple comparisons were performed by one-way ANOVA with Bonferroni post-test. ns: non-significant, \*:  $p < 0.05$ , \*\*:  $p < 0.01$ , \*\*\*:  $p < 0.001$ .

## **AUTHOR CONTRIBUTIONS**

S.P., F.L.P., S-K.B. and S.E.P designed and performed experiments and analyzed the data. O.C. provided technical assistance and established mouse cell lines. S.A. generated the H2A.X-SNAP plasmids and the H2A-SNAP and H2A.Z.1-SNAP cell lines. S.E.P. supervised the project and wrote the manuscript with inputs from all authors.

## **ACKNOWLEDGMENTS**

We thank members of our laboratory for stimulating discussions and critical reading of the manuscript. We thank E. Dunleavy and S. Jackson for sharing reagents. We acknowledge the ImagoSeine core facility (Institut Jacques Monod, France BioImaging) for assistance with confocal microscopy. This work was supported by the European Research Council (ERC starting grant ERC-2013-StG-336427 “EpIn”), the French National Research Agency (ANR-12-JSV6-0002-01), the “Who am I?” laboratory of excellence (ANR-11-LABX-0071) funded by the French Government through its “Investments for the Future” program (ANR-11-IDEX-0005-01), EDF Radiobiology program RB 2014-01, the Fondation ARC and France-BioImaging (ANR-10-INSB-04). S.P. is an EMBO Young Investigator. S.A. was recipient of PhD fellowships from University Pierre et Marie Curie and La Ligue contre le Cancer.

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## FIGURE LEGENDS

### Figure 1: *De novo* accumulation of H2A variants at UVC damage sites.

(A) Scheme of the assay for monitoring the accumulation of newly synthesized histones at UVC damage sites in cultured human cells stably expressing SNAP-tagged histones. Pre-existing SNAP-tagged histones are quenched with a non-fluorescent substrate (block) so that only the histones neo-synthesized during the chase period are labeled with the red fluorescent substrate tetramethylrhodamine (TMR)-star during the pulse step. Local UVC damage is induced before the pulse step and damage sites are detected by immunostaining for the repair factor XPA.

(B) New histone accumulation at sites of UVC damage marked by XPA analyzed in U2OS cell lines stably expressing the indicated SNAP-tagged histone variants.

(C) New H2A.X accumulation at sites of UVC damage marked by XPA in U2OS cells stably expressing SNAP-tagged H2A.X and treated with the indicated RNAPII inhibitors (left panel). The efficiency of RNAPII inhibition is measured in the same cell line by incorporating 5-Ethynyl Uridine (EU) into nascent transcripts (right panel).

(D) New H2A.X accumulation at sites of UVC damage marked by XPA analyzed in the presence of flavopiridol (+FLV) in U2OS cells stably expressing SNAP-tagged H2A.X. Total H2A.X is detected by immunostaining for SNAP or H2A.X.

(E) *De novo* accumulation of H2A variants at sites of UVC damage marked by XPA analyzed in the presence of flavopiridol (+FLV) in U2OS cell stably expressing the indicated SNAP-tagged H2A variants. Percentages of cells accumulating new H2A variants at UV sites are shown on the graph. Error bars represent SD from two independent experiments. Scale bars, 10  $\mu$ m.

### Figure 2: *De novo* accumulation of H2A variants at replication foci.



(A) New H2A.X accumulation at sites of replication foci marked by CAF-1 p150 analyzed in U2OS cells stably expressing SNAP-tagged H2A.X and treated with the indicated RNAPII inhibitors. *Right*, confocal images.

(B) New H2A.X accumulation at replication foci throughout S-phase analyzed in U2OS cells stably expressing SNAP-tagged H2A.X.

(C) New H2A.X accumulation at replication foci marked by CAF-1 p150 analyzed in U2OS cells stably expressing SNAP-tagged H2A.X. Total H2A.X is detected by immunostaining for SNAP or H2A.X.

(D) *De novo* accumulation of H2A variants at replication foci marked by CAF-1 p150 analyzed in U2OS cell stably expressing the indicated SNAP-tagged H2A variants. Percentages of cells accumulating new H2A variants at replication foci are shown on the graph. Error bars represent SD from two independent experiments. +FLV: flavopiridol. Scale bars, 10  $\mu$ m.

**Figure 3: *De novo* accumulation of H2A.X at UV sites and replication foci is coupled to DNA synthesis.**

(A) Schematic representation of the role of the endonuclease XPG in late steps of UVC damage repair (top). Repair synthesis in U2OS cells treated with two different siRNAs targeting XPG (or siLUC: control) is measured by incorporation of 5-Ethynyl-2'-deoxyUridine (EdU) at UVC damage sites (Cyclobutane Pyrimidine Dimers, CPD). The percentage of cells showing repair synthesis is indicated (at least 200 cells were scored in 2 independent experiments).

(B) New H2A.X accumulation at sites of UVC damage marked by XPA analyzed in the presence of flavopiridol (+FLV) in U2OS cells stably expressing SNAP-tagged H2A.X and transiently transfected with two different siRNAs targeting XPG (siLUC: control).

Percentages of cells accumulating new H2A.X at UV sites are shown on the graph. The efficiency of XPG knock-down is verified by western-blot and by the lack of CAF-1 p60 recruitment to UV damage sites.

**(C)** New H2A.X accumulation at sites of UVC damage marked by XPB analyzed as in (B) in cells treated with two different siRNAs targeting PCNA (siLUC: control). Percentages of cells accumulating new H2A.X at UV sites are shown on the graph. The efficiency of PCNA knock-down is verified by western-blot

**(D)** Scheme of the experiment for testing the effect of replicative synthesis inhibition by Aphidicolin (Aphi) on new H2A.X accumulation at replication foci in U2OS cells stably expressing SNAP-tagged H2A.X and treated with flavopiridol (+FLV). S-phase cells are labeled with EdU before Aphidicolin addition. The efficiency of replication inhibition by Aphidicolin is shown by the lack of CAF-1 p150 recruitment to replication foci. Percentages of cells accumulating new H2A.X at replication foci are shown on the graph. Error bars on the graphs represent SD from at least two independent experiments. Scale bars, 10  $\mu$ m.

**Figure 4: The histone chaperone FACT promotes new H2A.X deposition at UV damage sites**

**(A)** Recruitment of FACT subunits, SSRP1 and SPT16, to repair sites 2 h after local UVC irradiation at 500 J/m<sup>2</sup> in U2OS cells. The scheme represents FACT subunits bound to H2A-H2B.

**(B, C)** Recruitment of the histone chaperone FACT (SSRP1 subunit) to repair sites marked by XPB 2 h after local UVC irradiation at 500 J/m<sup>2</sup> in U2OS cells treated with the indicated siRNAs (siLUC: control). Knock-down efficiency is controlled on the western-blot panels. Percentages of cells recruiting FACT to repair sites are shown on the graphs. Error bars represent SD from at least two independent experiments.

**(D)** New H2A.X accumulation 1h after local UVC irradiation (500 J/m<sup>2</sup>) in U2OS H2A.X-SNAP cells treated with the indicated siRNAs (siLUC: control). Knock-down efficiencies are verified by western-blot. The intensity of new H2A.X signal (TMR fluorescence) at UV sites and in entire nuclei are shown on the graphs (bars: mean; error bars: SD from at least 100 cells). Similar results were obtained in three independent experiments and the results of a representative experiment are shown. Scale bar, 10  $\mu$ m.

**Figure 5. FACT-mediated deposition of new H2A.X does not promote UVC damage repair**

**(A)** Repair synthesis at UV damage sites (CPD) measured by EdU incorporation in U2OS cells treated with the indicated siRNAs (siLUC: control). XPG siRNA is used as a positive control. Percentages of cells incorporating EdU at UV damage sites are shown on the graph. Knock-down efficiencies are verified by western-blot.

**(B)** Kinetics of UV damage removal (CPD immunostaining at the indicated time points post global UVC irradiation) in U2OS cells treated with the indicated siRNAs (siLUC: control). XPG siRNA is used as a positive control.

**(C)** H2A.X neosynthesis analyzed in U2OS cells stably expressing H2A.X-SNAP and treated with the indicated siRNAs (siLUC: control).

**(D)** Repair synthesis at UV damage sites (CPD) measured by EdU incorporation in U2OS cells stably expressing H2A.X-SNAP and transiently transfected with the indicated siRNAs (siLUC: control). XPG siRNA is used as a positive control. Percentages of cells incorporating EdU at UV damage sites are shown on the graph.

Error bars on the graphs represent SD from two independent experiments. Scale bars, 10  $\mu$ m.

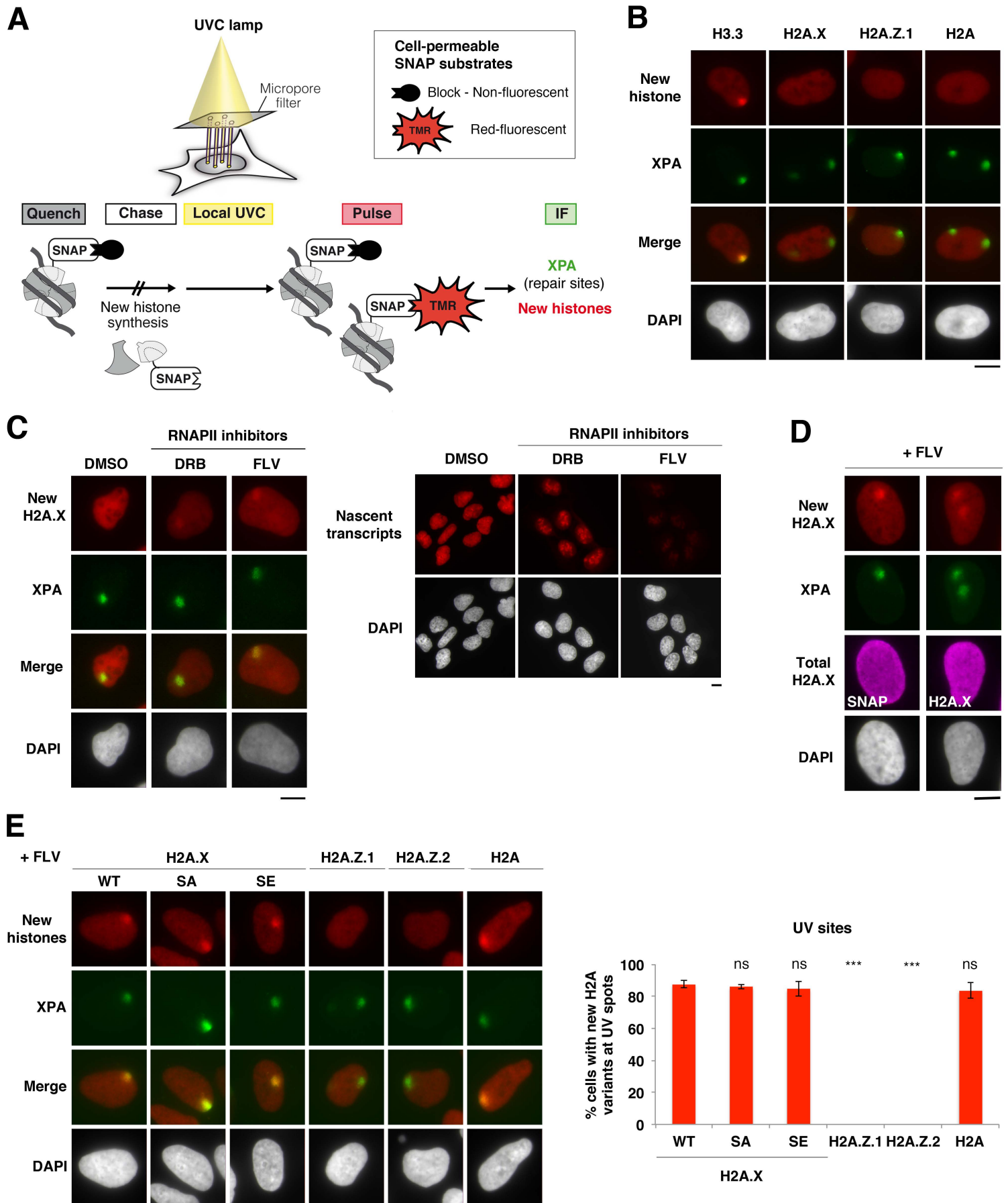
**Figure 6: The histone chaperone ANP32E removes H2A.Z from UV damage sites**

**(A)** Distribution of total H2A.X and H2A.Z variants in U2OS cells exposed to local UVC irradiation (500 J/m<sup>2</sup>). Total levels of SNAP-tagged histones are detected by a pulse with SNAP-Cell TMR-star (TMR) before cell fixation and total H2A.Z is also revealed by immunostaining with an H2A.Z-specific antibody recognizing H2A.Z amino-terminus (Nt). The arrowheads point to sites of UV irradiation. Fluorescence intensity profiles along the dotted lines are shown on the graphs.

**(B)** Distribution of total H2A.Z in U2OS H2A.Z.1-SNAP cells treated with the indicated siRNAs (siLUC, control) and exposed to local UVC irradiation (500 J/m<sup>2</sup>). Total levels of SNAP-tagged H2A.Z are detected by a pulse with SNAP-Cell TMR-star (TMR) before cell fixation and total H2A.Z is revealed by immunostaining with an H2A.Z-specific antibody recognizing H2A.Z carboxy-terminus (Ct). Intensities in the red channel (TMR) were adjusted on a cell-by-cell basis because of variable levels of SNAP-tagged H2A.Z in the cell population. The arrowheads point to sites of UV irradiation. H2A.Z levels at UV damage sites relative to the whole nucleus are shown on the graphs (bars: mean; error bars: SD from at least 40 cells). The significance of H2A.Z loss or enrichment at UV sites is indicated (compared to a theoretical mean of 1, dotted line). Similar results were obtained in three independent experiments. Knock-down efficiencies are verified by western-blot. Scale bars, 10  $\mu$ m.

**(C)** Model for chromatin reshaping during UVC damage repair through the concerted activities of ANP32E and FACT histone chaperones promoting H2A.Z/H2A.X histone variant exchange.

**Figure 1**



**Figure 2**

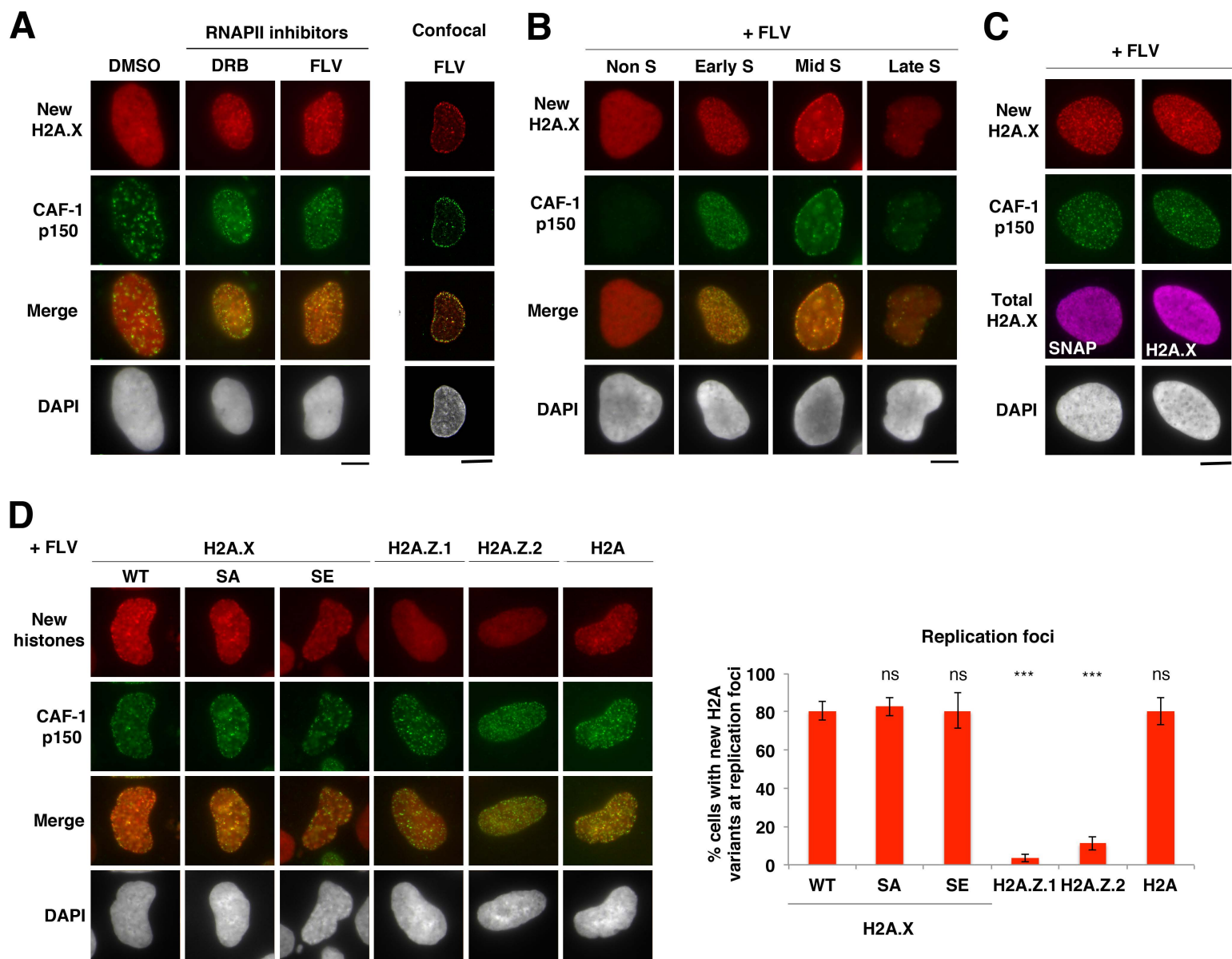
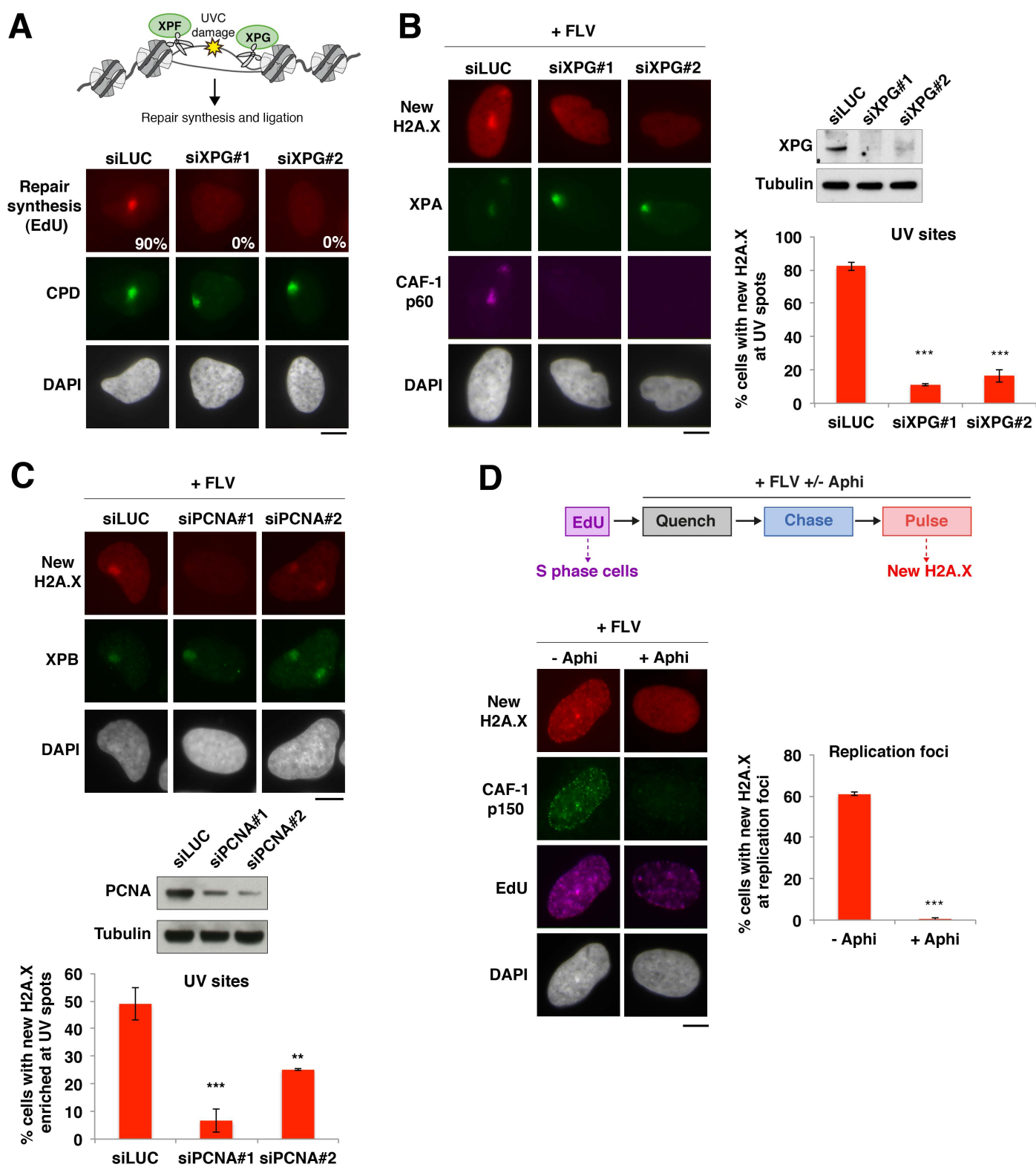
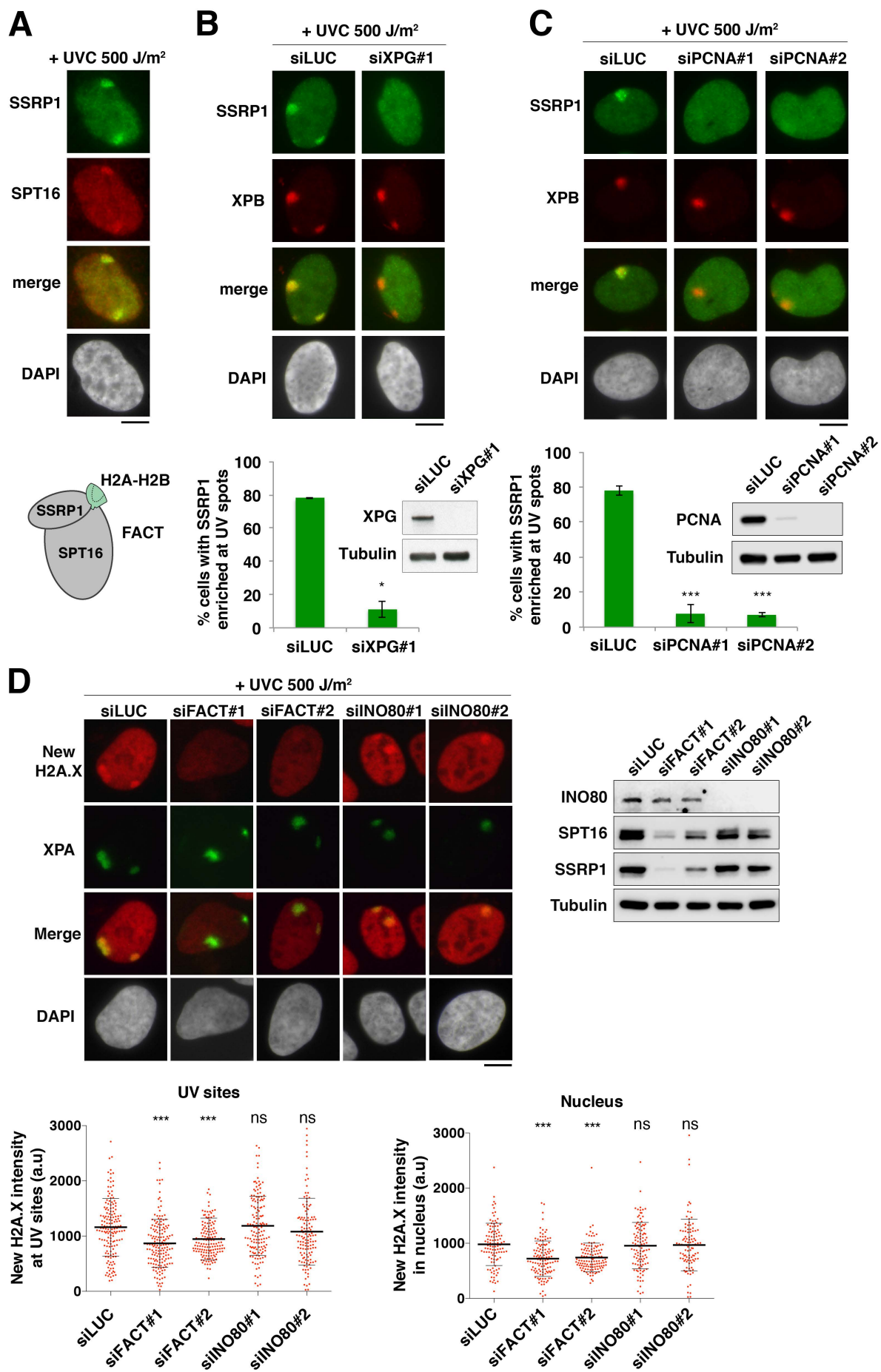


Figure 3

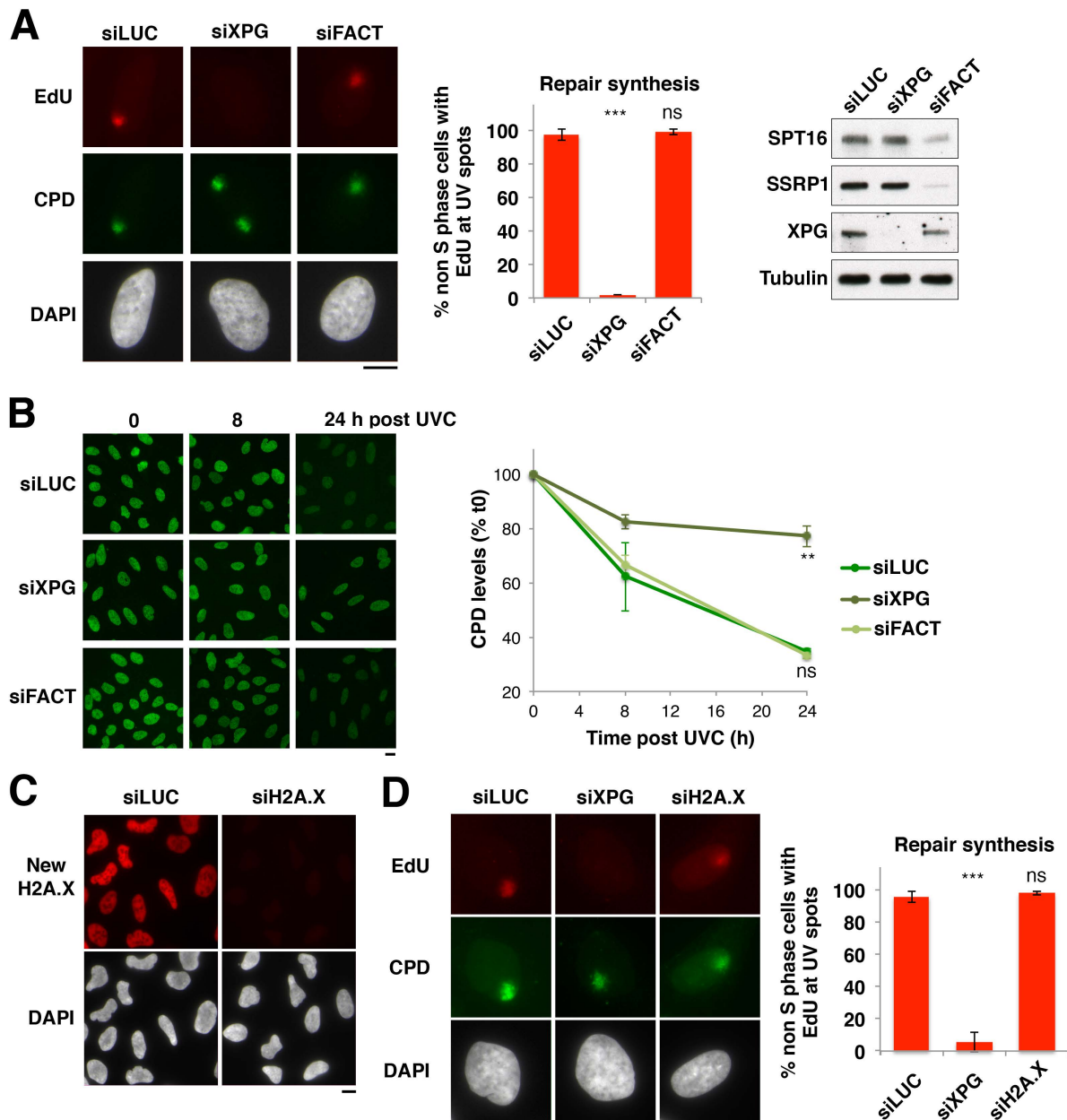


**Figure 4**





## Figure 5



**Figure 6**

