Histone chaperones and modifiers cooperate to maintain heterochromatin integrity following DNA damage.

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SUMMARY

Heterochromatin is a critical chromatin compartment, whose integrity governs genome stability and cell fate transitions. How heterochromatin features, including higher-order chromatin folding and histone modifications associated with transcriptional silencing, are maintained following a genotoxic stress challenge is unknown. Here, we establish a system for targeting UV damage to pericentric heterochromatin in mammalian cells and for tracking the heterochromatin response to UV in real time. We uncover profound heterochromatin compaction changes during repair, orchestrated by the UV damage sensor DDB2. Importantly, the restoration of heterochromatin folding is uncoupled from the maintenance of heterochromatin-specific histone modifications. We also unveil a central role for the methyltransferase SETDB1 in the maintenance of heterochromatic histone marks after UV, SETDB1 coordinating H3 methylation with new histone deposition by histone H3 chaperones in damaged heterochromatin. Our data thus shed light on fundamental molecular mechanisms safeguarding higher-order chromatin integrity following DNA damage.

Keywords: chromatin compaction, DDB2, H3K9me3, heterochromatin, histone chaperones, histone modifications, nucleotide excision repair, SETDB1, UV damage

HIGHLIGHTS

• The damage sensor DDB2 governs heterochromatin unfolding and refolding after UV
• Restoration of heterochromatin folding and silencing histone marks are uncoupled
• The methylase SETDB1 promotes H3K9me3 maintenance in damaged heterochromatin
• SETDB1 coordinates H3K9me3 with new H3 deposition by histone chaperones
INTRODUCTION

Eukaryotic cell identity and function are governed by the epigenetic information stored in the form of chromatin inside the cell nucleus, where DNA wraps around histone proteins (Allis and Jenuwein, 2016). This information encompasses multiple layers of regulation, from histone modifications (Bannister and Kouzarides, 2011) and histone variants (Buschbeck and Hake, 2017), up to higher-order folding of the chromatin fiber into nuclear domains (Yu and Ren, 2017) which, in concert, control gene expression. Among higher-order chromatin domains, heterochromatin covers a significant fraction of metazoan genomes (Ho et al., 2014; Roadmap Epigenomics Consortium et al., 2015) and plays a central role in the maintenance of genome stability. Highly concentrated at pericentromeric and subtelomeric regions, heterochromatin is indeed crucial for chromosome segregation and integrity, and alterations of heterochromatin features are commonly associated with aging and cancer (Janssen et al., 2018). Furthermore, heterochromatin formation is instrumental for silencing repetitive elements and preventing their illegitimate recombination (Janssen et al., 2018; Padeken et al., 2015). Heterochromatin silencing is mediated by specific patterns of histone post-translational modifications (Allshire and Madhani, 2017; Janssen et al., 2018). For instance, pericentric heterochromatin domains (Saksouk et al., 2015) carry a distinct chromatin signature, including trimethylation on H3 lysine 9 (H3K9me3) and on H4 lysine 20 (H4K20me3) (Martens et al., 2005), which contribute to epigenetic silencing of major satellite repeats. H3K9me3 heterochromatin also plays a pivotal role in defining cell identity by silencing lineage-specific genes during development (Becker et al., 2016; Ninova et al., 2019).

Considering the profound influence of heterochromatin on genome stability and cell fate transitions, much effort has been devoted to understanding how heterochromatin domains are established during development and maintained through cell divisions (Allshire and Madhani,
2017). One of the most persistent challenges to heterochromatin maintenance is the response to DNA damage, which can arise at any time, anywhere in the genome (Hoeijmakers, 2009; Tubbs and Nussenzweig, 2017) and poses a major threat to epigenome stability (Dabin et al., 2016; Hauer and Gasser, 2017). Indeed, substantial rearrangements affect chromatin during the repair response, including histone exchange and chromatin mobility (Dabin et al., 2016; Dion and Gasser, 2013), changes in histone post-translational modifications (Dantuma and van Attikum, 2016), and alterations in chromatin compaction (Khurana et al., 2014; Kruhlak et al., 2006; Luijsterburg et al., 2012; Sellou et al., 2016; Smeenk and van Attikum, 2013; Smith et al., 2019; Strickfaden et al., 2016). These rearrangements are accompanied by transient changes in chromatin transcriptional activity (Capozzo et al., 2017; Geijer and Marteijn, 2018; Marnef et al., 2017). The destabilization of chromatin organization upon genotoxic stress is followed by a restoration of chromatin structure (Polo and Almouzni, 2015; Smerdon, 1991). However, our knowledge of this fundamental process is still largely incomplete and little is known about the maintenance of higher-order heterochromatin domains following DNA damage.

Moreover, due to high compaction and to the abundance of repeated sequences prone to ectopic recombination, heterochromatin represents a challenging environment for the DNA damage response. Heterochromatic regions indeed pose a barrier to DNA damage signaling (Lemaître et al., 2014) and repair, as described for nucleotide excision repair (Adar et al., 2016; Han et al., 2016; Zheng et al., 2014), DNA double-strand break (DSB) repair (Kallimasioti-Pazi et al., 2018; Lemaître and Soutoglou, 2014) and mismatch repair (Supek and Lehner, 2015) in mammalian cells. In line with this, higher mutation rates are found in heterochromatin in human cancer genomes (Schuster-Böckler and Lehner, 2012; Zheng et al., 2014).
In recent years, exciting progress has been made in understanding how DNA damage repair proceeds in heterochromatin, as mostly studied in response to DNA breaks (Amaral et al., 2017). In drosophila and mouse cells, DSBs elicit a decompaction of pericentric heterochromatin and relocate to the periphery of heterochromatin domains for the completion of recombinational repair, which is thought to prevent illegitimate recombination between pericentromeric repeats (Chiolo et al., 2011; Jakob et al., 2011; Janssen et al., 2016; Tsouroula et al., 2016). However, beyond the restoration of genome integrity, the mechanisms underlying the maintenance of heterochromatic features during the repair response remain uncharacterized (reviewed in Fortuny and Polo, 2018). In particular, how heterochromatin compaction and silencing histone marks are preserved following DNA damage is still unknown, and whether they are maintained in a concerted manner also remains elusive.

Here, we explore these mechanisms by inflicting UV damage to pericentric heterochromatin domains in mammalian cells. We reveal that the maintenance of heterochromatin-specific histone marks and of heterochromatin folding are uncoupled and we uncover a critical role for the UV damage sensor DDB2 in orchestrating heterochromatin compaction changes during the repair response. Our findings also unveil a tight cooperation between histone chaperones and histone modifying enzymes in the maintenance of heterochromatic histone marks following UV damage.

RESULTS

Heterochromatin integrity is maintained in response to UV damage

In order to study heterochromatin maintenance in response to DNA damage, we first established an appropriate cellular model where heterochromatin domains could be easily
distinguished and where DNA repair events and histone deposition into chromatin could be tracked. For this purpose, we selected NIH/3T3 mouse embryonic fibroblasts, characterized by a clustering of pericentric heterochromatin domains into chromocenters (Probst and Almouzni, 2011) (Figure 1A), and we focused on the cell response to UVC damage (Figure S1A). Noteworthy, mouse fibroblasts express the UV damage sensor DNA Damage-Binding Protein 2 (DDB2) at very low levels, which impairs both UVC damage repair (Tang et al., 2000) and repair-coupled histone dynamics (Adam et al., 2016; 2013). To overcome these defects, NIH/3T3 stable cell lines were engineered to ectopically express GFP-tagged human DDB2 (GFP-hDDB2) at close to physiological levels as compared to human cells (data not shown). These cells also stably express SNAP-tagged H3.3, which allows specific tracking of newly synthesized H3.3 histones (Adam et al., 2015) (Figure S1A, see Figure S1B-D for a complete characterization of the cell lines). The ectopic expression of DDB2 and H3.3 did not affect pericentric heterochromatin organization as judged by immunostaining for H3K9me3 and Heterochromatin protein 1 α (HP1α) (Figure 1A). We verified that GFP-hDDB2 expression rescued UVC damage repair and associated histone dynamics in mouse cells, by analyzing the recruitment of the nucleotide excision repair (NER) factor Xeroderma Pigmentosum complementation group B (XPB) and the deposition of newly synthesized H3.3 histones at sites of UVC damage (Figure S1E).

Using the mammalian cellular model described above, we first assessed the importance of heterochromatin integrity for the cellular response to UV damage. We impaired heterochromatin integrity by knocking-down the histone methyltransferases SUV39H1 and 2 (Suppressor Of Variegation 3-9 Homolog 1/2), which are the main drivers of H3K9me3 in pericentric heterochromatin (Peters et al., 2001) (Figure 1A), and tested the ability of SUV39H1/2-depleted cells to survive UVC damage. We observed that SUV39H1/2 knockdown sensitized cells to global UVC irradiation (Figure 1A). Loss of heterochromatin
integrity thus correlates with reduced cell viability following UV damage.

To determine whether heterochromatin integrity was preserved following a genotoxic stress challenge, we developed an innovative approach for targeting UVC damage to pericentric heterochromatin domains in live cells and for tracking the response to heterochromatin damage in real time. We employed the live-cell DNA stain Hoechst 33258 to visualize chromocenters in mouse cells and then inflicted UVC damage specifically to chromocenters of interest by using a UVC laser coupled to a confocal microscope (Figure 1B). Using this approach, we observed a pronounced and rapid decompaction of damaged heterochromatin within minutes after UVC laser damage, reaching a maximum (up to 6-fold) 30 min to 1 h after irradiation (Figure 1B and Movie S1). Heterochromatin decompaction was restricted to UVC-damaged chromocenters (Figure S2A), and was further confirmed by DNA-fluorescence in situ hybridization (FISH) analysis of mouse major satellite sequences (Figure S2B). Importantly, damaged heterochromatin decompaction was followed by a slower recompaction phase taking several hours, which restored heterochromatin compaction close to its original state (Figure 1B and Movie S2). Furthermore, immunostaining for H3K9me3 in cells fixed 1 h after UVC laser damage revealed that damaged heterochromatin decompaction was not associated with a reduction of this heterochromatin-specific histone mark, which instead appeared slightly increased on damaged chromocenters (Figure 1C). Similar results were obtained when staining for H4K20me3 (Figure S2C). The observed increase of silencing marks was restricted to damaged chromocenters, with no detectable increase of H3K9me3 in damaged euchromatin regions (Figure S2D). In addition, a modest but reproducible increase in H3K9me3 levels was detected by western blot on total extracts from cells exposed to global UVC irradiation (Figure S2E), which confirms the above findings and excludes the possibility of increased H3K9me3 detection due to increased antibody accessibility in decompacted heterochromatin. Together, these experiments demonstrate that
heterochromatin-specific histone marks are maintained, and even slightly increased, in UVC-damaged heterochromatin. In line with these findings, damaged heterochromatin decompaction was not accompanied by a burst of aberrant transcription. Indeed, the staining of nascent transcripts with Ethynyl-Uridine (EU) and the quantification of nascent transcript levels in heterochromatin domains before and after UV irradiation revealed that transcription was even further reduced in UV-damaged heterochromatin (Figure 1D, left graph). This UV-induced transcriptional arrest in heterochromatin was confirmed by RT-qPCR of pericentric major satellite transcripts (Figure 1D, right graph). From these observations, we conclude that UVC damage challenges heterochromatin integrity and that maintenance mechanisms operate to restore heterochromatin compaction and to reinforce heterochromatin-specific histone marks and heterochromatin silencing following UV damage.

**The UV damage sensor DDB2 regulates heterochromatin compaction**

To characterize the mechanisms underlying heterochromatin maintenance following UVC damage, we first sought to identify the molecular trigger for damaged heterochromatin decompaction. For this, we examined the potential contribution of the UV damage sensor DDB2, whose binding to chromatin was shown to promote histone redistribution and chromatin relaxation in human cells (Adam et al., 2016; Luijsterburg et al., 2012). Noteworthy, we observed decompaction of UV-damaged heterochromatin domains only in the engineered cell line expressing hDDB2 and not in the parental mouse cell line (DDB2-deficient) (Figure 2A), supporting the idea that DDB2 is required for heterochromatin decompaction following UVC damage.

To directly test whether DDB2 could drive heterochromatin decompaction, we tethered GFP-hDDB2 to mouse pericentric heterochromatin in the absence of DNA damage by co-expressing catalytically dead Cas9 (dCas9) fused to a GFP nanobody and a guide RNA
targeting major satellite repeats (Anton and Bultmann, 2017; Tsouroula et al., 2016) (Figure 2B and S3A-B). DDB2 tethering led to substantial changes in the shape and size of pericentric heterochromatin domains, which were enlarged and less spherical compared to control cells, indicative of a decompaction of pericentric heterochromatin domains. This effect was specific to DDB2 tethering as it was not observed upon targeting of another early NER factor, Xeroderma Pigmentosum complementation group C (XPC), to chromocenters (Figure 2C and S3C). Noteworthy, when we induced the release of tethered DDB2 from major satellite repeats with the anti-Cas9 bacteriophage protein AcrIIA4, thus mimicking the release of DDB2 from damaged chromatin that occurs during repair progression (Rapić-Otrin et al., 2002), the typical size and shape of chromocenters were restored, showing that DDB2 release allows pericentric heterochromatin recompaction (Figure 2D and S3D). Collectively, these findings establish that the UV damage sensor DDB2 is both necessary and sufficient for driving changes in heterochromatin compaction following UVC damage.

**UV damage repair operates within heterochromatin domains**

DDB2-mediated decompaction of damaged heterochromatin could facilitate access of downstream repair factors to the core of heterochromatin domains. We thus examined the recruitment to UVC-damaged heterochromatin of repair proteins acting downstream of DDB2 in the NER pathway, namely the intermediate repair factor XPB, which contributes to opening the damaged DNA double-helix, and the late repair factor Proliferating Cell Nuclear Antigen (PCNA), involved in repair synthesis after damage excision (Figure 3A). Similar to what observed for GFP-hDDB2, we detected the accumulation of endogenous XPB and PCNA in damaged heterochromatin upon cell exposure to local UVC irradiation (Figure 3B). Importantly, we noticed that PCNA accumulated within heterochromatin domains during DNA damage repair, as observed both in our mouse cell line model and in human MCF7 cells.
that endogenously express DDB2 (Fig 3C and S4A). The recruitment of PCNA to the core of heterochromatin domains following UV damage contrasts with PCNA peripheral localization during heterochromatin replication (Quivy et al., 2004). This indicates that, unlike replicative synthesis, UV damage repair synthesis takes place inside heterochromatin domains. Altogether, these results establish that pericentromeric heterochromatin is fully permissive for NER factor recruitment up to late repair steps.

**Repair-coupled deposition of new H3 histones in heterochromatin domains**

UV damage repair elicits the deposition of newly synthesized histones in human cells (Adam et al., 2013; Dinant et al., 2013; Piquet et al., 2018; Polo et al., 2006), including the H3 histone variants H3.1 and H3.3. To investigate whether such repair-coupled histone deposition was taking place in damaged heterochromatin, we examined the recruitment of H3 variant-specific chaperones, starting with the H3.1 histone chaperone Chromatin Assembly Factor-1 (CAF-1), which is known to interact with PCNA during repair (Moggs et al., 2000) and to deposit new H3.1 histones at UVC damage sites (Polo et al., 2006). Similar to PCNA (Figure 3B-C), we observed that CAF-1 accumulated in damaged heterochromatin upon local UVC irradiation both in mouse NIH/3T3 GFP-hDDB2 cells and in human MCF7 cells (Figure 4A and S4B). Regarding H3.3 histone chaperones, both HIRA (Histone Regulator A) and DAXX (Death Domain Associated Protein) can drive H3.3 deposition (reviewed in Sitbon et al., 2017). HIRA deposits H3.3 within transcribed euchromatin in mammalian cells (Goldberg et al., 2010; Ray-Gallet et al., 2011) and in UVC-damaged chromatin in human cells (Adam et al., 2013), while DAXX promotes H3.3 enrichment at repeated sequences including pericentric heterochromatin (Drané et al., 2010; Goldberg et al., 2010; Wong et al., 2010). We thus examined whether one or both of these chaperones were recruited to UVC-damaged heterochromatin. We observed that while HIRA accumulated in a comparable
manner in damaged euchromatin and heterochromatin domains (Figure 4B), DAXX was specifically recruited to damaged heterochromatin (Figure 4C). Thus, although originally considered to operate in distinct chromatin domains (Elsaesser and Allis, 2010), HIRA and DAXX chaperones co-exist within damaged heterochromatin, which we confirmed in human MCF7 cells (Figure S4C). In light of these findings, we explored the possibility of a co-recruitment of these two chaperones to damaged heterochromatin. However, siRNA-mediated depletion of HIRA did not impair DAXX recruitment and reciprocally, showing that both H3.3 histone chaperones are independently recruited to UVC-damaged heterochromatin (Figure S5A).

DAXX recruitment being heterochromatin-specific, we further investigated the underlying mechanisms. We noticed a co-enrichment on damaged heterochromatin of the DAXX-binding partner and the heterochromatin-associated protein ATRX (Alpha Thalassemia/Mental Retardation Syndrome X-Linked) (Figure S5B). ATRX knock-down revealed that ATRX was driving DAXX recruitment to damaged heterochromatin (Figure S5C). Noteworthy, we observed DAXX accumulation in damaged heterochromatin both in and outside S-phase (Figure S5D), ruling out the possibility that DAXX recruitment could be coupled to heterochromatin replication, owing to enhanced chromatin accessibility during this process. Analogous to what observed for HIRA recruitment to UV sites in human cells (Adam et al., 2013), DAXX accumulation in UVC-damaged heterochromatin domains was dependent on the UV damage sensor DDB2 (Figure S5E).

In line with the recruitment of H3.3-specific histone chaperones, we observed an accumulation of newly synthesized H3.3-SNAP histones within damaged heterochromatin domains, comparable to neighboring euchromatin regions (Figure 4D). Loss-of-function experiments revealed that only HIRA depletion markedly inhibited new H3.3 deposition in UV-damaged chromocenters (Figure 4E). Although we cannot exclude a minor contribution
of DAXX, HIRA thus appears to be the main driver for new H3.3 deposition in damaged heterochromatin. Collectively, these findings demonstrate that UVC damage drives the recruitment of H3 histone chaperones and new H3 deposition in heterochromatin domains.

**SETDB1 coordinates new H3 histone deposition and H3K9me3 maintenance in damaged heterochromatin**

Newly synthesized H3 histones do not carry the same post-translational modifications as nucleosomal H3 and are largely devoid of trimethylation marks (Alabert and Groth, 2012; Loyola et al., 2006). Thus, we wondered whether and how the newly synthesized H3 histones deposited in damaged heterochromatin would acquire heterochromatin-specific modifications, including H3K9me3, which we observed was maintained after UVC damage (Figure 1C). Interestingly, when we examined the recruitment of H3K9 trimethyltransferases, we found that SETDB1 (SET Domain Bifurcated 1) was specifically recruited to damaged pericentric heterochromatin (Figure 5A) while SUV39H1 displayed only a slight enrichment on UVC-damaged compared to undamaged chromocenters (Figure S6A). The H3K27 trimethyltransferase EZH2 (Enhancer Of Zeste Homolog 2) in contrast did not show any significant accumulation in damaged chromocenters (Figure S6B).

Given the specific recruitment of SETDB1 to damaged heterochromatin, we further investigated its potential role in methylating newly deposited H3 histones. SETDB1 was dispensable for new H3.3 deposition (Figure S6C). However, abrogation of new H3 histone deposition, achieved by simultaneous depletion of H3.3 and of the H3.1-chaperone CAF-1, impaired SETDB1 recruitment to UVC-damaged heterochromatin (Figure 5B). Single depletion of H3.3 or CAF-1 had no or a very modest effect (Figure S6D). Furthermore, erasing parental H3K9me3 by knocking down SUV39H1/2 prevented SETDB1 recruitment to damaged heterochromatin (Figure 5C). These results indicate that SETDB1 recruitment to
damaged heterochromatin domains is driven by the deposition of newly synthesized H3 histones and also by SUV39H1/2 enzymes, which maintain parental H3K9me3. An attractive possibility is thus that SETDB1 may trimethylate newly deposited H3 histones in UV-damaged heterochromatin by copying the K9me3 mark from neighboring SUV39H1/2-modified parental histones.

To further characterize the possible role of SETDB1 in H3K9me3 maintenance following UV damage, we first analyzed H3K9me3 total levels post UV by western blot (Figure 5D). These experiments revealed that, contrary to control cells where H3K9me3 levels increase moderately following UV irradiation (Figure S2C and 5D), SETDB1-knocked down cells displayed a significant reduction in H3K9me3 levels post UV (Figure 5D). Similar results were obtained by focusing on pericentric heterochromatin, as observed by chromatin immunoprecipitation (ChIP) of H3K9me3 on major satellite repeats (Figure 5E).

Together, these findings demonstrate that de novo deposition of H3 histones stimulates the recruitment to UV-damaged heterochromatin of the histone modifying enzyme SETDB1, which promotes H3K9me3 maintenance.

DISCUSSION

By assessing the consequences of UVC damage on mammalian pericentric heterochromatin domains, we provide important novel insights into the mechanisms for heterochromatin maintenance following DNA damage. We describe damage-mediated alterations in heterochromatin compaction with the retention of silencing histone marks, which may facilitate repair in compact regions of the genome while preserving heterochromatin identity. We also unveil a repair-coupled deposition of newly synthesized histones in damaged heterochromatin, and propose that histone chaperones and chromatin modifiers cooperate to maintain heterochromatin integrity following DNA damage (Figure 6).
Regulation of heterochromatin compaction following UV damage

Chromatin reorganization coupled to the early stages of the DNA damage response is considered to be critical for efficient DNA repair (Dabin et al., 2016; Dion and Gasser, 2013). This is particularly relevant in compact heterochromatin domains. Indeed, decompaction of pericentromeric heterochromatin has been reported in response to radiation- and nuclease-induced breaks both in flies (Chiolo et al., 2011) and in mouse embryonic fibroblasts (Jakob et al., 2011; Tsouroula et al., 2016), as well as following heat stress in plants (Pecinka et al., 2010). Here, we provide evidence for pericentric heterochromatin decompaction following UVC damage in mouse fibroblasts, together with the maintenance of silencing histone marks in decondensed heterochromatin. We propose that by retaining their histone marks, these chromatin domains also maintain their identity, which could be crucial for the re-establishment of the original chromatin state once DNA repair is complete. Our observation that UV damaged chromatin decompacts while retaining heterochromatic histone marks is in line with previous studies in response to DSBs (Natale et al., 2017; Tsouroula et al., 2016), and highlights an uncoupling between chromatin structural and molecular determinants during DNA damage repair. Reciprocally, heterochromatic histone marks can be erased without any significant effect on heterochromatin decompaction, as observed upon SUV39H1/2 knockdown.

In addition to heterochromatin histone marks, Heterochromatin Protein 1 alpha (HP1 α), which is recruited to UVC lesions (Luijsterburg et al., 2009), could contribute to preserve heterochromatin identity following DNA damage. Owing to its liquid-liquid demixing properties (Larson et al., 2017; Strom et al., 2017), HP1α could stimulate heterochromatin recompaction at later time points post-UVC.
Mechanistically, the regulation of damaged heterochromatin compaction differs in response to distinct types of DNA damage. In response to UV lesions, we have identified the UV damage sensor DDB2 as a master regulator of heterochromatin compaction. It is not yet clear if DDB2 promotes decompaction by itself or by interacting with other factors. It has been shown that the DDB2-binding partners DDB1 and Cul4A/B are not involved in controlling chromatin decompaction (Luijsterburg et al., 2012). Recent structural data indicate that the DDB2 complex can expose UV lesions occluded in nucleosomal DNA by promoting DNA shifting (Matsumoto et al., 2019). However, such local activity at the nucleosome level is unlikely to sustain larger scale chromatin decompaction. Given that DDB2 does not harbor known chromatin remodeling activity or motifs, we hypothesize that it induces chromatin decompaction indirectly, by promoting the recruitment of chromatin remodelers or, alternatively, the release of factors involved in chromatin compaction like linker histones. Further studies will be needed to fully dissect the molecular bases of DDB2-mediated chromatin decompaction. Importantly, the function of DDB2 in regulating chromatin compaction likely extends to other types of DNA lesions than those processed by the NER pathway considering that DDB2 also detects oxidative damage and contributes to base excision repair (Jang et al., 2019; Reardon et al., 1997).

It will also be of major interest to assess the impact of damage-mediated chromatin decompaction on the three-dimensional organization of chromatin in the nuclear space (Rowley and Corces, 2018). Indeed, it is not known whether decompaction entails only local chromatin movement with the loss or enlargement of chromatin loops within topologically associated domains, or more profound and global alterations of chromatin topology.

Functionally, whether heterochromatin decompaction facilitates the access of repair factors to damaged DNA is not entirely clear. Here, we have shown that the NER machinery can access UV lesions in pericentric heterochromatin and that repair can be completed within these
domains. This contrasts with the relocalization of repair foci to the periphery of heterochromatin domains for late steps of DSB recombination (Jakob et al., 2011; Tsouroula et al., 2016), as also observed for replication foci (Quivy et al., 2004), thus showing that not all pathways that involve DNA synthesis are excluded from the core heterochromatin domain. Our results are consistent with several studies showing that, even if volume exclusion and moderate diffusive hindrance occur in heterochromatin domains (Bancaud et al., 2009), heterochromatin is accessible to large proteins (Verschure et al., 2003), including non-homologous end joining (NHEJ), single-strand annealing (SSA) and early homologous recombination (HR) factors (Tsouroula et al., 2016). Considering that NER, unlike HR of DSBs, does not pose a risk for ectopic recombination between heterochromatic repeats, there would be no need for a relocalization of the NER machinery to the heterochromatin periphery and thus no spatial segregation of UV damage repair events. Instead, there is a temporal regulation of NER in heterochromatin, with slower kinetics of UV damage repair (Han et al., 2016), likely due to the necessary decompaction to promote access to lesions buried in heterochromatin.

**Histone deposition in UV-damaged heterochromatin: role of histone chaperones**

By assessing the recruitment of H3 variant-specific histone chaperones to UVC damaged heterochromatin, we have identified the histone chaperone HIRA as the main driver of new H3.3 deposition at UVC-damaged heterochromatin. While we cannot formally exclude that the DAXX-ATRX complex has a minor contribution to this process, we can envision alternative roles for this complex, such as stimulating parental H3.3 recovery during the repair response. Known as a promiscuous histone chaperone, the DAXX-ATRX complex could also regulate the dynamics of other histone variants, including CENP-A (Lacoste et al., 2014) and macroH2A.1 (Kim et al., 2019; Ratnakumar et al., 2012). Another potential role would be the control of heterochromatin recompaction at late time points post-UV, in light of recent data.
revealing that DAXX-ATRX-mediated deposition of H3.3 is key for chromocenter clustering during myogenic differentiation (Park et al., 2018). Finally, it has been proposed that this histone chaperone complex could regulate repair synthesis as observed during HR (Juhasz et al., 2018), but our preliminary data do not support such a function during NER (data not shown).

**Maintenance of silencing histone marks in UV-damaged heterochromatin**

We have established that the histone methyltransferase SETDB1 promotes H3K9me3 maintenance in UV-damaged heterochromatin. In line with these findings, SETDB1 has been involved in DNA damage-induced H3K9me3 leading to sex chromosome inactivation in meiosis (Hirota et al., 2018). We have found that the histone methyltransferase SETDB1 is specifically recruited to UVC-damaged heterochromatin. Although the underlying mechanisms are still unclear, they may involve SETDB1 tandem Tudor domains, reported to bind specifically to dually modified histone H3 containing both K14 acetylation (H3K14ac) and K9 methylation (H3K9me1/2/3) (Jurkowska et al., 2017). Given that newly synthesized H3 histones are enriched in K14ac (Alabert et al., 2015), and are not optimal substrates for trimethylation by SUV39H1/2 (Rea et al., 2000), we hypothesize that SETDB1 could bind, via its Tudor domains, the new H3 histones deposited in damaged heterochromatin, and then trimethylate these histones, thus mirroring SUV39H1/2-dependent trimethylation on parental H3. Future studies will determine whether SETDB1 indeed promotes trimethylation of newly deposited H3 histones in UV-damaged heterochromatin. While H3K9me3 is maintained, and even slightly increased, in UV-damaged heterochromatin, we did not find evidence for ectopic H3K9me3 formation in damaged euchromatic regions. This contrasts with the increase in H3K9me3 reported after DSB induction, which contributes to transcriptional repression (Ayrapetov et al., 2014; Marnef et al., 2019).

Collectively, our work sheds new light on the processes safeguarding pericentric
heterochromatin integrity following DNA damage. It would be of interest to determine if similar or distinct mechanisms operate in other heterochromatin domains characterized by different patterns of epigenetic marks, such as telomeric chromatin and facultative heterochromatin. Beyond the DNA damage response, our findings may also provide a molecular framework for understanding heterochromatin maintenance during other disruptive events in both normal and pathological conditions, like DNA replication, cell differentiation, aging and disease.

ACKNOWLEDGMENTS

We thank members of our laboratory for stimulating discussions and P-A. Defossez and C. Rougeulle for critical reading of the manuscript. We thank J. Bondy-Denomy, S. Bultmann and R. Nishi for sharing plasmids. We acknowledge the ImagoSeine facility (Institut Jacques Monod, France BioImaging) for confocal microscopy and the imaging platform of the Epigenetics and Cell Fate Center for epifluorescence microscopy. This work was supported by the European Research Council (ERC starting grant ERC-2013-StG-336427 “EpIn” and consolidator grant ERC-2018-CoG-818625 “REMIND”), the French National Research Agency (ANR-12-JSV6-0002-01 and ANR-18-CE12-0017-01), 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-INBS-04). S.P. is an EMBO Young Investigator. A.F was recipient of a PhD fellowship from University of Paris.

AUTHOR CONTRIBUTIONS

A.F., A.C., P.C. and S.E.P. designed and performed experiments, analyzed the data and wrote the manuscript. O.C. provided technical assistance and established mouse stable cell lines.
O.L. and O.R. implemented the UVC laser technology and helped with image analyses. S.E.P. supervised the project.

**COMPETING INTERESTS STATEMENT**

The authors declare no competing financial interests.
FIGURE LEGENDS

Figure 1. Heterochromatin integrity is maintained in response to UV damage

(A) Schematic representation of pericentric heterochromatin domains in mouse cells and delocalization of heterochromatin marks (H3K9me3, HP1α) upon knock-down of SUV39H1/2 methyltransferases in NIH/3T3 GFP-DDB2 cells. Clonogenic survival of the same cell line treated with the indicated siRNAs (siLUC, negative control; siXPG, positive control) and exposed to global UVC irradiation.

(B) Technical approach for targeting UVC damage to pericentric heterochromatin domains (HC) in live murine cells. Heterochromatin compaction changes upon UVC laser micro-irradiation are analyzed by live imaging in NIH/3T3 GFP-DDB2 cells stained with Hoechst. White arrowheads point to UVC-damaged heterochromatin domains.

(C) H3K9me3 levels in damaged heterochromatin (white arrowheads) analyzed by immunofluorescence in NIH/3T3 GFP-DDB2 cells 1h after UVC laser micro-irradiation. Scatter plots represent DAPI and H3K9me3 levels measured on reconstructed 3D images in damaged heterochromatin domains compared to undamaged heterochromatin in the same nucleus.

(D) Heterochromatin transcription analyzed 1h30 after global UVC damage in NIH/3T3 GFP-DDB2 cells by EU staining (fluorescence images and left graph) and by RT-qPCR for major satellite transcripts (right graph).

Error bars, s.d. (s.e.m. for (B) panel only) from n cells or from at least three independent experiments. a.u., arbitrary units. All microscopy images are confocal sections. Scale bars, 10 µm. Zoomed in views of heterochromatin domains (x2.6). See also, Figures S1-S2, Movies S1-S2.
Figure 2. The UV damage sensor DDB2 regulates heterochromatin compaction

(A) Decompaction of damaged pericentric heterochromatin domains (white arrowheads) one hour after UVC laser micro-irradiation analyzed by live imaging in the indicated cell lines. CPD staining in fixed cells highlights the damaged chromocenter. The scatter plots represent the area of the damaged chromocenters normalized to the same chromocenters before UVC laser.

(B) Procedure for targeting GFP-tagged DDB2 to major satellites sequences in pericentric heterochromatin.

(C, D) Confocal sections showing the aspect of pericentric heterochromatin domains upon tethering of the indicated GFP-tagged proteins in NIH/3T3 (C) or NIH/3T3 GFP-DDB2 cells (D). Heterochromatin tethering is relieved by expressing an anti-Cas9 peptide (D). The scatter plots show changes in volume and sphericity of heterochromatin domains quantified on reconstructed 3D images. Error bars, s.d. from n cells scored in at least three independent experiments. Scale bars, 10 µm. Zoomed in views of heterochromatin domains (x2.6). See also, Figure S3.

Figure 3. The NER pathway operates within heterochromatin domains

(A) Scheme of the Global Genome Nucleotide Excision Repair factors studied.

(B) Recruitment to UVC damage (CPD) of early (DDB2), intermediate (XPB) and late (PCNA) repair factors, analyzed by immunofluorescence 30 min after local UVC irradiation through micropore filters in NIH/3T3 GFP-DDB2 cells. Cells with damaged heterochromatin domains (white arrowheads) were selected for the analysis. PCNA accumulation to damaged heterochromatin was analyzed outside S-phase. XPB and PCNA were not stained in green because the cells express GFP-DDB2, but are presented in green for simplicity. Scatter plots represent log2 fold enrichments of repair proteins in damaged heterochromatin (HC) and
damaged euchromatin (EC) compared to the whole nucleus.

(C) Accumulation of PCNA within heterochromatin domains upon local UVC irradiation and confined to the periphery of replicating heterochromatin in mid-late S-phase.

Error bars, s.d. from n cells scored in at least two independent experiments. All microscopy images are confocal sections. Scale bars, 10 µm. Insets show zoomed in views of heterochromatin domains (x2.3). See also, Figure S4.

**Figure 4. Histone H3 deposition in UVC-damaged heterochromatin**

(A-C) Recruitment of the H3.1 histone chaperone CAF-1 (p150 subunit) (A), and of the H3.3 histone chaperones HIRA (B) and DAXX (C) to UVC-damaged regions, analyzed by immunofluorescence in NIH/3T3 GFP-DDB2 cells 1h30 after local UVC irradiation through micropore filters. Zoomed in views (x2.6) show damaged regions (delineated by green dotted lines) containing heterochromatin domains (delineated by white dotted lines).

(D, E) Accumulation of newly synthesized H3.3 histones in UVC-damaged heterochromatin regions (white arrowheads) analyzed in NIH/3T3 GFP-DDB2 H3.3-SNAP cells 45 min after local UVC irradiation through micropore filters. H3.3 chaperones are knocked down by siRNA (siLUC, control) (E). siRNA efficiencies are controlled by western blot (Tubulin, loading control).

Scatter plots represent log2 fold enrichments of histone chaperones or new H3.3-SNAP histones in damaged heterochromatin (HC) and damaged euchromatin (EC) compared to the whole nucleus (A-D) or normalized to the corresponding siLUC experiment (E). Error bars, s.d. from n cells scored in at least three independent experiments. All microscopy images are confocal sections. Scale bars, 10 µm. See also, Figures S4-S5.
Figure 5: SETDB1 is recruited to UVC-damaged heterochromatin and promotes H3K9me3 maintenance

(A) Recruitment of SETDB1 to damaged heterochromatin (white arrowheads) analyzed by immunofluorescence 1h30 after local UVC irradiation through micropore filters in NIH/3T3 GFP-DDB2 cells.

(B, C) SETDB1 recruitment to damaged heterochromatin in cells treated with the indicated siRNAs (siLUC, control).

(D) Changes in total H3K9me3 levels 1h30 post global UVC irradiation, detected by western blot on total extracts from NIH/3T3 GFP-DDB2 cells treated with the indicated siRNAs (siLUC, control). Tubulin, loading control; γH2A.X, damage marker.

(E) H3K9me3 abundance on major satellites detected by ChIP before and 1h30 after global UVC irradiation in NIH/3T3 GFP-DDB2 cells treated with the indicated siRNAs (siLUC, control). Mock, no antibody.

Scatter plots represent log2 fold enrichments of SETDB1 in damaged heterochromatin (HC) and damaged euchromatin (EC) compared to the whole nucleus (A) or normalized to the corresponding siLUC experiment (B, C). Bar graphs represent H3K9me3 abundance in damaged/undamaged conditions (D, E right) or H3K9me3/total H3 levels normalized to the input (E left). Error bars, s.d. from n cells scored in at least three independent experiments.

All microscopy images are confocal sections. Scale bars, 10 µm. See also, Figure S6.

Figure 6: Model for heterochromatin maintenance following UVC damage

Recognition of UVC damage by the sensor protein DDB2 (1) triggers decompaction of damaged pericentric heterochromatin (2), thus facilitating access of downstream repair factors and histone chaperones (3) to the core of the domain. Histone chaperones promote the incorporation of newly synthesized H3 histones (in red), which subsequently acquire heterochromatin-specific modifications through the action of histone methyltransferases (4).
DDB2 release during repair progression allows heterochromatin recompaction (5).
METHODS

Cell culture

U2OS (ATCC HTB-96, human osteosarcoma, female), MCF7 (ATCC HTB-22, human breast adenocarcinoma, female), and NIH/3T3 cells (ATCC CRL-1658, mouse embryonic fibroblast, male) were grown at 37°C and 5% CO₂ in Dulbecco’s modified Eagle’s medium (DMEM, Invitrogen) supplemented with 10% fetal bovine serum (EUROBIO) and antibiotics (100 U/ml penicillin and 100 µg/ml streptomycin, Invitrogen) and the appropriate selection antibiotics (Euromedex, Supplementary Table 1). For seeding NIH/3T3 cells on coverslips, coverslips were first coated with 20 µg/ml Collagen Type I (MERCK Millipore) and 2 µg/ml Fibronectin (Sigma-Aldrich) to increase cell adhesion.

| Stable cell lines          | Selection antibiotics         |
|----------------------------|--------------------------------|
| NIH/3T3 GFP-DDB2           | 200 µg/ml Hygromycin          |
| NIH/3T3 GFP-DDB2 H3.3-SNAP | 200 µg/ml Hygromycin + 250 µg/ml G418 |
| NIH/3T3 H3.3-SNAP          | 100 µg/ml G418                |
| U2OS H3.3-SNAP             | 100 µg/ml G418                |

Supplementary Table 1: Stable cell lines.
Ectopically expressed proteins are of human origin and are more than 80% similar to mouse proteins. Antibiotics: Hygromycin, G418 (Euromedex)

siRNA and plasmid transfections

siRNA purchased from Eurofins MWG Operon (Supplementary Table 2) were transfected into cells using Lipofectamine RNAiMAX (Invitrogen) following manufacturer’s instructions. The final concentration of siRNA in the culture medium was 50-80 nM. Cells were harvested 48-72 hr after transfection.

| Designation | Target species | Target sequence | Working conditions |
|-------------|----------------|-----------------|--------------------|
| siATRX      | Mouse          | 5’-GTACAGAAATCTCGCTCAA3’ | 50 nM – 72 hours   |
| siCAF-1 p150 | Mouse         | 5’-AAGGAGAAGGGAGGAGGAGCAG3’ | 30 nM – 48 hours   |
**siRNA Sequences**

| siRNA Name | Species | Sense Oligo Sequence | Concentration Range |
|------------|---------|----------------------|---------------------|
| siDAXX    | Mouse   | 5’TGACCTTACAAACACTGAA^3’ | 50 nM – 72 hours   |
| siH3.3    | Mouse   | 1:1 combination of siH3.3A: 5’CTACAAAAGCCGCTCGAA^3’ and siH3.3B: 5’GCCAAGAGAGTCACCATCA^3’ | 50 nM – 48 to 72 hours |
| siHIRA    | Mouse   | 5’GGAAGGTTTGTGATCTGGAA^3’ | 50 nM – 72 hours   |
| siLUC     | Firefly | 5’CGTACGCGGAATCTTCCA^3’ | 50 nM – 48 to 72 hours |
| siSETDB1  | Mouse   | 5’GCCGAGGCTTTGCTCTTA^3’ | 50 nM – 72 hours   |
| siSETDB1  | Mouse   | 5’CCCGAGGCTTTGCTCTTA^3’ | 50 nM – 72 hours   |
| siSU93H1/2| Mouse   | 5’ACCTCTTTGACCTGGACTA^3’ | 50 nM – 72 hours   |
| siXPG     | Mouse   | 5’TGATGATAACGATGAGAAA^3’ | 50 nM – 72 hours   |

**Supplementary Table 2: siRNA sequences.**

Cells were transfected with plasmid DNA (Supplementary Table 3) using Lipofectamine 2000 (Invitrogen) according to manufacturer’s instructions. For stable cell line establishment (Supplementary Table 1), plasmid DNA was transfected into cells at 1 µg/ml final, 48 hr before antibiotic selection of clones. For transient transfections, each plasmid was at 0.5 µg/ml final and cells were fixed 48 hr post transfection. For DDB2 tethering to major satellites, plasmids encoding GFP-tagged proteins, GBP-dCas9-mRFP and major satellite gRNA were co-transfected into NIH/3T3 cells 48 hr before cell analysis. For DDB2 detachment from major satellites, NIH/3T3 GFP-DDB2 cells were transfected with GBP-dCas9-mRFP and major satellite gRNA plasmids and 24 hr later with anti-Cas9 plasmid. Cells were fixed 24 hr after the second transfection.

**Plasmid details**

| Plasmid                          | Construct details                                      | Reference/Provider                                      |
|----------------------------------|--------------------------------------------------------|---------------------------------------------------------|
| anti-Cas9 (pJH376-AcrIIA4)       | Bacteriophage AcrIIA4 sequence inserted into pcDNA3.1(+) | Gift from Joseph Bondy-Denomy. Addgene plasmid #86842 (Rauch et al., 2017) |
| Plasmid | Description | Source |
|---------|-------------|--------|
| GBP-dCas9-mRFP | GFP-binding nanobody (GBP) and mRFP coding sequences cloned into pCAG-dCas9 | Gift from Sebastian Butlmann (Anton and Bultmann, 2017) |
| GFP | pEGFP-C1 | Clontech #6084-1 |
| GFP-DDB2 | Human *DDB2* coding sequence (Montpellier Genomic Collections) subcloned into GFP-XPC plasmid replacing XPC | (Adam et al., 2016) |
| GFP-XPC | cDNA encoding GFP-human XPC cloned into pIREShyg vector (Clontech) | Gift from Ryotaro Nishi (Nishi et al., 2009) |
| H3.3-SNAP | Human *H3F3B* coding sequence cloned into pSNAPm (New England Biolabs) | (Dunleavy et al., 2011) |
| HIRA-YFP | Human HIRA sequence from (Hall et al., 2001) edited by Genscript and subcloned into pEYFP-N1 (Clontech) | (Adam et al., 2013) |
| MajSat gRNA | Major satellite guide RNA sequence cloned into pEX-A-U6-gRNA | Gift from Sebastian Butlmann (Anton et al., 2014) |

**Supplementary Table 3: Plasmids.**

**UVC irradiation**

Cells grown on glass coverslips (12 mm diameter, thickness No.1.5, Thorlabs) were irradiated with UVC (254 nm) using a low-pressure mercury lamp. Conditions were set using a VLX-3W dosimeter (Vilbert-Lourmat). For global UVC irradiation, cells in Phosphate Buffer Saline (PBS) were exposed to UVC doses ranging from 4 to 12 J/m² for survival assays and to 10 J/m² in other experiments. For local UVC 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 or 300 J/m² UVC. Irradiated cells were allowed to recover in culture medium for the indicated times before fixation.

For UVC laser micro-irradiation (Dinant et al., 2007), cells were grown on quartz coverslips (25 mm diameter, thickness No.1, SPI supplies) and nuclei were stained by adding Hoechst 33258 (10 µg/mL final, Sigma-Aldrich) to the culture medium 30 min before UVC irradiation.
irradiation. Quartz coverslips were transferred to a Chamlide magnetic chamber on a custom stage insert (Live Cell Instrument) and cells were irradiated for 50 ms using a 2 mW pulsed diode-pumped solid-state laser emitting at 266 nm (RappOptoElectronics, Hamburg GmbH) directly connected to a Zeiss LSM 700 confocal microscope adapted for UVC transmission with all-quartz optics. The laser was attenuated using a neutral density filter OD1 and focused through a 40x/0.6 Ultrafluor glycerol objective with quartz lenses.

Cell extracts and western blot
Total extracts were obtained by scraping cells on plates or resuspending cell pellets in Laemmli buffer (50 mM Tris-HCl pH 6.8, 1.6% Sodium Dodecyl Sulfate (SDS), 8% glycerol, 4% β-mercaptoethanol, 0.0025% bromophenol blue) followed by 5 min denaturation at 95°C. Alternatively, cell pellets were resuspended in lysis buffer (1 M Tris-HCl pH 6.8, 50 mM NaCl, 0.5% NP-40, 1%, Sodium Deoxycholate, 1% SDS, 5 mM MgCl2) before addition of Laemmli buffer with 0.25 U/µL benzonase (final concentration, Merck Millipore) for 10 min followed by 5 min denaturation at 95°C.

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). Proteins were transferred onto nitrocellulose membranes (Amersham Protran) for 30 min at 15V with a Trans-Blot SD semidy transfer cell (Bio-Rad) or in transfer buffer (25 mM Tris, 200 mM glycine, 20% ethanol) for 2 h at 52V with a liquid transfer system (Bio-Rad). Total proteins were revealed by Pierce® Reversible Stain (Thermo Scientific). Proteins of interest were probed using the appropriate primary and Horse Radish Peroxidase (HRP)-conjugated secondary antibodies (Supplementary Table 4), detected using SuperSignal West Pico or Femto chemiluminescence substrates (Pierce) on hyperfilms MP (Amersham). When fluorescence detection was used instead of chemi-luminescence, secondary antibodies were conjugated to IRDye 680RD or
800CW (Supplementary Table 4), membranes were scanned with an Odyssey Fc-imager (LI-COR Biosciences) and analyzed with Image Studio Lite software using total protein stain for normalization.

**Flow cytometry**

For cell cycle analysis, 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 and 0.5 mg/ml RNase A (USB/Affymetrix). DNA content was analyzed by flow cytometry using a BD FACScalibur flow cytometer (BD Biosciences) and FlowJo software (TreeStar).

**SNAP-tag labeling of newly synthesized histones**

For specific labeling of newly synthesized histones (Adam et al., 2015; Bodor et al., 2012), cells were grown on glass coverslips and pre-existing SNAP-tagged histones were first quenched by incubating cells with 10 µM of the non-fluorescent substrate SNAP-cell Block (New England Biolabs) for 30 min followed by a 30 min-wash in fresh medium and a 2 hr-chase. The new SNAP-tagged histones synthesized during the chase were fluorescently labeled with 2 µM of the red-fluorescent reagent SNAP-cell TMR star or SiR-647 (New England Biolabs) during a 15 min-pulse step followed by 30 min wash in fresh medium. Cells were subsequently permeabilized with Triton X-100, fixed and processed for immunostaining. Cells were irradiated with a UVC lamp before the pulse step.

**EdU-labeling of replicating cells and repair sites**

To visualize replication foci, 10 µM Ethynyl-deoxyUridine (EdU) was incorporated into cells on glass coverslips during 15 min at 37°C and revealed using the Click-It EdU Alexa Fluor 647 Imaging kit (Invitrogen) according to manufacturer’s instructions. To localize the sites of
UV damage repair, cells were incubated with 10 μM EdU for 1h30 after local UVC irradiation and EdU was revealed using the Click-It EdU Alexa Fluor 488 Imaging kit (Invitrogen).

**Nascent RNA labeling**

Cells on glass coverslips were incubated in medium supplemented with 0.5 mM Ethynyl-Uridine (EU) for 45 min at 37°C, and EU incorporation was revealed with Click-iT RNA Alexa Fluor 594 Imaging kit (Invitrogen) according to manufacturer’s instructions. Coverslips were mounted in Vectashield medium with DAPI (Vector laboratories). EU fluorescence intensity in heterochromatin was measured using ImageJ software. Heterochromatin segmentation was based on DAPI staining.

**Immunofluorescence**

Cells grown on coverslips were either fixed directly with 2% paraformaldehyde (Electron Microscopy Sciences) for 20 min and permeabilized for 5 min with 0.5% Triton X-100 in PBS or cells were pre-extracted before fixation 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 MgCl₂) for 5 min at room temperature to remove soluble proteins. For PCNA staining, cells were fixed with 100% ice-cold methanol for 15 min. For the detection of UVC photoproducts (CPD), DNA was denatured with 2N HCl for 10 min at 37°C (Cosmo Bio antibody, Supplementary Table 4) or with 0.5 M NaOH for 5 min at room temperature (Kamiya antibody, Supplementary Table 4). Since this denaturation quenches GFP fluorescence, when CPD detection was combined with the visualization of GFP-DDB2, immunofluorescence was performed in two steps starting with GFP immunodetection using a rat anti-GFP antibody (Supplementary Table 4) followed by fixation, denaturation and CPD immunodetection.
Samples were blocked for 10 min in 5% BSA (Bovine Serum Albumin, Sigma-Aldrich) in PBT (PBS 0.5% Tween-20), followed by 45 min incubation with primary antibodies and 30 min incubation with secondary antibodies coupled to AlexaFluor dyes (Supplementary Table 4) diluted in blocking buffer. Coverslips were mounted in Vectashield medium with DAPI (Vector laboratories).

| Type | Antibody target | Species | Supplier | Dilution/amount | Application |
|------|-----------------|---------|----------|-----------------|-------------|
|      | ATRX            | Mouse   | Santa Cruz Biotechnology (sc-15408) | 1:400 | IF |
|      |                 |         |          | 1:500 | WB |
|      | CAF-1 p60       | Mouse   | Active Motif (39996) | 1:500 | IF |
|      | CAF-1 p150      | Goat    | Santa Cruz Biotechnology (sc-10206) | 1:50 | IF |
|      |                 |         |          | 1:250 | WB |
| Primary | CPD             | Mouse   | Kamiya Biomedical Company (MC-062, clone KTM53) | 1:1000 | IF |
|      |                 |         | Cosmo Bio (CAC-NM-DND-001, clone TDM2) | 1:1000 | IF |
|      | DAXX            | Rabbit  | Santa Cruz Biotechnology (sc-7152) | 1:250 | IF |
|      |                 |         | Sigma-Aldrich (HPA008736) | 1:100 | IF |
|      |                 |         | Ozyme (4533) | 1:100 | WB |
|      | DDB2            | Mouse   | Abcam (ab51017) | 1:200 | WB |
|      | EZH2            | Mouse   | BD-Biosciences (612666) | 1:100 | IF |
|      | GFP             | Rat     | Santa Cruz Biotechnology (sc-101536) | 1:50 | IF |
|      |                 | Mouse   | Roche Applied Science (11814460001) | 1:1000 | WB |
| Antibody          | Species | Source                                      | Dilution | Format |
|-------------------|---------|---------------------------------------------|----------|--------|
| γH2A.X            | Mouse   | MERCK Millipore (05-636, clone JBW301)      | 1:1000   | IF     |
|                   |         |                                             | 1:1000   | WB     |
| H3.3              | Rabbit  | MERCK Millipore (09-838)                    | 1:1000   | WB     |
| H3K4me3           | Rabbit  | MERCK Millipore (07-473)                    | 1:5000   | IF     |
| H3K9me3           | Rabbit  | Active Motif (39765)                        | 1:500    | IF     |
|                   |         | Abcam (ab8898)                              | 3 µg     | ChIP   |
|                   |         |                                             | 1:1000   | WB     |
| H4K20me3          | Rabbit  | Abcam (ab9053)                              | 1:500    | IF     |
| H3                | Rabbit  | Abcam (ab1791)                              | 2 µg     | ChIP   |
|                   |         |                                             | 1:5000   | WB     |
| HIRA              | Mouse   | Active Motif (39557)                        | 1:100    | IF     |
|                   |         |                                             | 1:200    | WB     |
| HP1α              | Mouse   | Millipore (MAB3584)                         | 1:500    | IF     |
| PCNA              | Rabbit  | Santa Cruz Biotechnology (sc-7907)          | 1:50     | IF     |
|                   | Mouse   | Dako (M0879)                                | 1:1000   | IF     |
| SETDB1            | Mouse   | Thermo scientific (MA515722)                | 1:200    | IF     |
|                   | Rabbit  | Santa Cruz Biotechnology (sc-66884)         | 1:200    | IF     |
|                   | Mouse   | Abcam (ab107225)                            | 1:1000   | WB     |
| SNAP              | Rabbit  | Pierce Antibodies (CAB4255)                 | 1:500    | IF     |
|                   |         |                                             | 1:1000   | WB     |
| SUV39H1           | Rabbit  | Cell signaling technology (8729)            | 1:25     | IF     |
|                   |         |                                             | 1:1000   | WB     |
| Tubulin           | Mouse   | Sigma-Aldrich (T9026)                       | 1:10000  | WB     |
| XPA               | Mouse   | BD Biosciences (556453)                     | 1:500    | IF     |
| Antibody | Species | Company | Concentration | Detection |
|----------|---------|---------|---------------|-----------|
| XPB | Rabbit | Santa Cruz Biotechnology (sc-293) | 1:400 | IF |
| Goat HRP | Donkey | Santa Cruz Biotechnology (sc-2020) | 1:10000 | WB |
| Mouse HRP | Goat | Jackson ImmunoResearch Laboratories (115-035-068) | 1:10000 | WB |
| Rabbit HRP | Donkey | Jackson ImmunoResearch Laboratories (711-035-152) | 1:10000 | WB |
| Anti-Rabbit IRDye 680RD Conjugated | Goat | LI-COR Biosciences (926-68071) | 1:15000 | WB |
| Anti-Rabbit IRDye 800CW Conjugated | Goat | LI-COR Biosciences (926-32211) | 1:15000 | WB |
| Anti-Mouse IRDye 680RD Conjugated | Goat | LI-COR Biosciences (926-68070) | 1:15000 | WB |
| Anti-Mouse IRDye 800CW Conjugated | Goat | LI-COR Biosciences (926-32210) | 1:15000 | WB |
| Goat Alexa Fluor 594 | Donkey | Invitrogen (A11058) | 1:1000 | IF |
| Mouse Alexa Fluor 488 | Goat | Invitrogen (A11029) | 1:1000 | IF |
| Mouse Alexa Fluor 568 | Goat | Invitrogen (A11031) | 1:1000 | IF |
| Mouse Alexa Fluor 594 | Goat | Invitrogen (A11032) | 1:1000 | IF |
| Mouse Alexa Fluor 647 | Goat | Invitrogen (A21236) | 1:1000 | IF |
| Rabbit Alexa Fluor 568 | Goat | Invitrogen (A11036) | 1:1000 | IF |
| Rabbit Alexa Fluor 594 | Goat | Invitrogen (A11037) | 1:1000 | IF |
| Rabbit Alexa Fluor 647 | Goat Invitrogen (A21245) | 1:1000 | IF |
|------------------------|--------------------------|--------|----|
| Rat Alexa Fluor 488    | Goat Invitrogen (A11006) | 1:1000 | IF |

**Supplementary Table 4: Antibodies**

IF: Immunofluorescence; WB: Western-Blot, ChIP: Chromatin immunoprecipitation

**DNA-Fluorescence In Situ Hybridization (DNA-FISH) of mouse major satellites**

Cells on quartz coverslips were fixed in 2% paraformaldehyde (Electron Microscopy Sciences). 5’TYE563-labeled locked nucleic acid probes (Exiqon) against mouse major satellite sequences (Supplementary Table 5) were precipitated with mouse Cot-1 DNA (Invitrogen) and Salmon Sperm DNA (Thermo Fisher Scientific), resuspended in formamide and denatured for 7 min at 75°C. The probes were then diluted in an equal volume of 2X Hybridization Buffer (4X SSC, 20% Dextran Sulfate, 2 mg/ml BSA). Coverslips were dehydrated in 80%, 90% and 100% ethanol and equilibrated in 2X SCC at 80°C. Coverslips were then denaturated for 10 min at 80°C in 70% formamide/2X SSC (pH=7.2), dehydrated in 70%, 80%, 90% and 100% ethanol and incubated overnight with the major satellite probes at 37°C. After three 5-min washes in 50% formamide/2X SSC at 45°C, and three 4-min washes in 2X SSC at 45°C, coverslips were mounted in Vectashield medium with DAPI (Vector laboratories). Since sample denaturation quenches GFP fluorescence, immunofluorescence against GFP (to detect GFP-DDB2) was performed prior to DNA-FISH.

**Image acquisition and analysis**

Fluorescence imaging was performed with a Leica DMI6000 epifluorescence microscope using a Plan-Apochromat 40x/1.3 or 63x/1.4 oil objective. Images were captured using a CCD camera (Photometrics) and Metamorph software. Images were assembled with Adobe Photoshop. For confocal imaging, samples were observed on a Zeiss LSM710 confocal microscope using a Plan-Apochromat 63x/1.4 oil objective. Live cell imaging coupled to
UVC laser micro-irradiation was performed using a 40x/0.6 Ultrafluar Glycerol objective on a Zeiss LSM700 confocal microscope. Images were captured using Zen software, and analysed with ImageJ (U. S. National Institutes of Health, Bethesda, Maryland, USA, http://imagej.nih.gov/ij/). Nuclei and heterochromatin domains were segmented on 2D confocal images based on DAPI or Hoechst staining, and UVC-damaged regions based on GFP-DDB2 fluorescence using custom-made ImageJ macros. The volume, sphericity and GFP intensity of heterochromatin domains as well as H3K9me3 intensity in damaged heterochromatin were analyzed on 3D images reconstructed from z-stacks using Imaris (Bitplane, http://www.bitplane.com/imaris).

Colony forming assays
Cells were replated 48h after siRNA transfection and exposed to global UVC irradiation (4, 8 and 12 J/m²) the following day. Colonies were stained 12 days later with 0.5% crystal violet/20% ethanol and counted. Results were normalized to plating efficiencies.

Reverse Transcription- quantitative Polymerase Chain Reaction (RT-qPCR)
Total RNA was extracted from cells with TRIzol™ reagent following manufacturer’s instructions (Invitrogen) and precipitated in isopropanol. RNA samples were subject to DNA digestion with Turbo DNA-free (Invitrogen) before reverse transcription with Superscript III RT using random primers (200 ng/reaction, Invitrogen). Quantitative PCR reactions were carried out with the indicated primer pairs (Eurofins MWG Operon, 500 nM final concentration, Supplementary Table 5) and Power SYBR® Green PCR Master Mix (Applied Biosystems) and read in MicroAmp® Fast Optical 96-well plates (Applied Biosystems) using an ABI 7500 Fast detection system (Applied Biosystems). Results were normalized to the amount of the GAPDH housekeeping gene product.
### Supplemental Table 5: qRT-PCR primers and FISH probes

| Designation         | Sequence                   | Application     |
|---------------------|----------------------------|-----------------|
| Major satellite_F  | 5′ GACGACTTGAATAATGACGAAATC 3′ | RT-qPCR         |
| Major satellite_R  | 5′ CATATTCCAGGTCCTTCAGTGTC 3′ | RT-qPCR         |
| GAPDH_F             | 5′ TGCACCACCAACTGCTTAGC 3′  | RT-qPCR         |
| GAPDH_R             | 5′ GGCATGGACTGTGGTCATGAG 3′  | RT-qPCR         |
| Major satellite_F  | 5′ ACGTGAATATGGCGAGGAA 3′   | ChIP            |
| Major satellite_R  | 5′ CAAGTCGTCAAGTGGATGTT 3′  | ChIP            |
| Major satellite_F  | 5′ TCTTGCCATATCCACGTCC 3′   | DNA-FISH        |
| Major satellite_R  | 5′ GCGAGGAAAAACTGAAAAAGG 3′ | DNA-FISH        |

**Chromatin immunoprecipitation (ChIP)**

ChIP was performed as described in (Tytca et al., 2006) with few modifications. Cells were crosslinked for 15 min with 1% formaldehyde (Sigma-F8775). The fixation reaction was stopped by adding glycine (0.125 M final concentration) for 5 min. Cells were collected and resuspended in cell lysis buffer (5 mM Pipes, 85 mM KCl, 0.5% NP-40). Lysates were homogenised with a tight Dounce homogeniser (DWK Life Sciences) and nuclei were collected by centrifugation. Samples were resuspended in nucleus lysis buffer (50 mM Tris pH 8.1, 10 mM EDTA, 1% SDS) and sonicated using the Bioruptor plus water bath system (Diagenode, 16 cycles at high power, 30 sec ON/30 sec OFF) to an average fragment size of 0.5-1 kb as assessed by agarose gel electrophoresis. Chromatin was diluted in dilution buffer (0.01% SDS, 1.1% Triton X-100, 1.2 mM EDTA, 16.7 mM Tris pH 8.1, 167 mM NaCl) and 25 µg of chromatin was incubated overnight at 4°C with primary antibodies (Supplementary table 4) or without antibody as negative control. Immune complexes were recovered with 20 µL blocked protein A/G magnetic beads (Thermo Scientific) during 4 h at 4°C. Beads were washed once in dialysis buffer (2 mM EDTA, 50 mM Tris pH 8.1, 0.2% Sarkosyl) and four times in wash buffer (100 mM Tris pH 8.1, 500 mM LiCl, 1% NP-40, 1% Sodium...
Deoxycholate). Samples were resuspended in TE buffer (10 mM Tris pH 8.1, 1 mM EDTA pH 8.1), treated with RNase A and DNA was eluted from the beads by adding SDS (1% final concentration) overnight with constant mixing at 60°C. After proteinase K treatment, DNA was purified by phenol:chloroform:isoamyl alcohol extraction (Invitrogen) and resuspended in TE buffer.

Quantitative PCR reactions were performed as described in the RT-qPCR section. All experiments included a standard curve and all samples were analyzed in triplicates. Results were normalized to the input.

**Statistical analyses**

Percentages of positively stained cells were obtained by scoring at least 150 cells in each experiment. Statistical tests were performed using GraphPad Prism. H3K9me3 relative abundance +/-UV was compared to a theoretical mean of 1 by one-sample t-test. 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. Comparisons of clonogenic survival were based on non-linear regression with a polynomial quadratic model. ns: non-significant, *: p < 0.05, **: p < 0.01, ***: p < 0.001.
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Figure 1

A

Pericentric heterochromatin (HC) Mouse acrocentric chromosome Mouse cell

H3K9me3 HP1α DAPI

siLUC

siSUV39H1/2

Mean intensity in HC / siLUC

H3K9me3 HP1α

siLUC siSUV39

n=21 n=21

% survival

100 10

0 4 8 12

UVC dose (J/m^2)

*** p<0.0001

B

HC staining Targeted damage to HC

Hoechst UVC laser Live cell imaging

UVC laser

30 sec 1 min 20 min

Hoechst

Decompa

n=12-45

n=5-11

n=2

Recompa

Area of damaged HC dot (normalized to before laser)

Time post UVC (h)

C

UVC laser

Hoechst DAPI + GFP-DDB2 H3K9me3

Before UVC 1h after UVC

DAPI

H3K9me3

H3K9me3 / DAPI

n=20, p<0.0001

n=20, p=0.6811

n=20, p=0.0186

Mean intensity (a.u.)

Undamaged HC Damaged HC Undamaged HC Damaged HC Undamaged HC Damaged HC

RT-qPCR

EU levels in HC (relative to -UV)

Major satellite / GAPDH transcripts (relative to -UV mean)

-UV 1h30

n=45 n=55

p<0.0001

1.5

1.0

0.5

0.0
Figure 2

A

NIH/3T3
GFP-DDB2

Hoechst
UVC laser

Before UVC

NIH/3T3
GFP-DDB2

1h after UVC

DAPI + CPD

B

Pericentric heterochromatin

Major satellite DNA target

5' ... ATGCAAATCGCAAGAAGACCTGAAAATCAG... 3'

3' ... TACCTTATACGCTTTTGTACGTCTTTACTCT... 5'

5' GGCAAGAGGCTGAAATCA

guide RNA

mRFP

dCas9

αGFP

C

HC-tethered GFP proteins

GFP
GFP-DDB2
GFP-XPC

DAPI

D

Volume of HC domains (μm³)

Sphericity of HC domains

GFP
GFP-DDB2
GFP-XPC

n=40
n=38
n=40

p<0.0001

ns

anti-Cas9

GFP-DDB2

+ 

- 

p<0.0001

anti-Cas9

mRFP

DAPI

n=47
n=44

p<0.0001
Figure 3

A. Global genome NER

- Lesion detection
- DNA double helix opening
- Excision
- Repair synthesis & ligation

B. Local UVC

- DDB2
- XPB
- PCNA

C. REPAIR & REPLICATION

- Local UVC irradiation
  - PCNA
  - GFP-DDB2
  - DAPI
- Mid-late S-phase
  - PCNA
  - DAPI
  - PCNA + DAPI

Intensity in damaged chromatin/ nucleus (log2)

|        | HC | EC | p-value |
|--------|----|----|---------|
| DDB2   | 4.0| 3.5| <0.0001 |
| XPB    | 5.0| 4.0| 0.0583  |
| PCNA   | 5.5| 5.5| 0.2062  |

n=40, p<0.0001; n=40, p=0.0583; n=20, p=0.2062
Figure 4

A. CAF-1 complex

B. HIRA complex

C. DAXX complex

D. New H3.3-SNAP

E. siLUC, siHIRA, siDAXX

DAXX
HIRA
Tubulin
Figure 5

A

SETDB1

H3K9me3

SETDB1

GFP-DDB2

DAPI

SETDB1 in damaged chromatin / nucleus (log2)

HC   EC

n=30, p<0.0001

B

siLuc   siH3.3 + siCAF-1

SETDB1

GFP-DDB2

DAPI

SETDB1 in damaged HC / siLuc

p<0.0001

n=30

n=30

CAF-1 p150

H3.3

Tubulin

C

siLuc   siSUVC39H1/2

SETDB1

GFP-DDB2

DAPI

SETDB1 in damaged HC / siLuc

p<0.0001

n=28

n=39

SUVC39H1

H3K9me3

Tubulin

D

siLuc   siSETDB1

UVC  -  -  -  +  +  +

H3K9me3

SETDB1

γH2A.X

Tubulin

H3

p=0.0229

H3K9me3 fold change (4UVC / UVC)

E

H3K9me3 ChIP on major satellites

H3K9me3 relative abundance / H3

Mock

H3K9me3

UVC  -  -  +  +  +  +

siLuc   siSETDB1

H3K9me3 fold change (4UVC / UVC)
