Deregulated origin licensing leads to chromosomal breaks by rereplication of a gapped DNA template

Kai J. Neelsen,1,3,4 Isabella M.Y. Zanini,1,3,5 Sofija Mijic,1 Raquel Herrador,1 Ralph Zellweger,1 Arnab Ray Chaudhuri,1,6 Kevin D. Creavin,2 J. Julian Blow,2 and Massimo Lopes1,7

1Institute of Molecular Cancer Research, University of Zurich, CH-8057 Zurich, Switzerland; 2Centre for Gene Regulation and Expression, University of Dundee, Dundee DD1 5EH, United Kingdom

Deregulated origin licensing and rereplication promote genome instability and tumorigenesis by largely elusive mechanisms. Investigating the consequences of Early mitotic inhibitor 1 (Emi1) depletion in human cells, previously associated with rereplication, we show by DNA fiber labeling that origin reactivation occurs rapidly, well before accumulation of cells with >4N DNA, and is associated with checkpoint-blind ssDNA gaps and replication fork reversal. Massive RPA chromatin loading, formation of small chromosomal fragments, and checkpoint activation occur only later, once cells complete bulk DNA replication. We propose that deregulated origin firing leads to undetected discontinuities on newly replicated DNA, which ultimately cause breakage of rereplicating forks.

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The activation of DNA replication origins is a tightly regulated mechanism, entailing two main steps: [1] “origin licensing,” restricted to late mitosis and early G1, when essential replication initiation proteins (ORC1, Cdc6, Cdt1, and Mcm2–7) are sequentially loaded on origin DNA sequences, forming the “prereplicative complex” (preRC), and [2] “origin firing,” occurring throughout S phase, when additional proteins are recruited to the preRC and start unwinding and DNA synthesis [Arias and Walter 2007]. As relicensing and thus rereplication are detrimental to genome stability, several cyclin-dependent kinase (CDK)-dependent and -independent mechanisms have evolved to coordinate these steps with cell cycle progression [Blow and Dutta 2005; Arias and Walter 2007]. Although several preRC components are targets of regulation, the major mechanism by which metazoans prevent origin licensing during S phase is inactivation of Cdt1 by ubiquitin-mediated degradation or binding to its inhibitor Geminin. Cdt1 proteolysis is tightly linked to the cell cycle, as ubiquitylation requires CDK-dependent phosphorylation [Li et al. 2003; Sugimoto et al. 2004; Nishitani et al. 2006]. Moreover, CUL4/DDB1-mediated ubiquitylation of Cdt1 occurs in S phase or in response to DNA damage [Arias and Walter 2006; Nishitani et al. 2006; Senga et al. 2006]. Geminin exerts its inhibitory function on Cdt1 in S, G2, and early M phase and is inactivated in late M phase by anaphase-promoting complex (APC/C)-dependent polyubiquitylation, leading to reactivation of origin licensing [McGarry and Kirschner 1998; Wohlschlegel et al. 2000; Tada et al. 2001; Li and Blow 2004]. Accordingly, Geminin depletion induces rereplication and activation of the DNA damage response (DDR) [Melixetian et al. 2004; Zhu et al. 2004].

By direct control of Geminin and indirect control of Cdt1 proteolysis via regulation of CycA–CDK activity, APC/C plays a pivotal role coordinating origin licensing with cell cycle progression [Hook et al. 2007]. APC/C activity is inhibited by Early mitotic inhibitor 1 [Emi1] [Wang and Kirschner 2013], which thereby stabilizes APC/C substrates like Geminin and Cyclin A [Di Fiore and Pines 2007]. Thus, inactivation of Emi1 leads to degradation of both inhibitors of Cdt1 activity, resulting in massive rereplication and DDR activation [Machida and Dutta 2007].

As many origin licensing genes are overexpressed in cancer cells and several oncogenes are known to affect origin licensing, it is suspected that deregulated licensing contributes to genome instability and tumorigenesis [Hook et al. 2007; Blow and Gillespie 2008]. However, our understanding of how rereplication challenges genome stability is very limited. Studies with Xenopus laevis egg extracts provided the first insight into the effects of rereplication. Addition of recombinant Cdt1 to G2-arrested egg extracts was shown to trigger DNA breaks, proposed to arise from head-to-tail collision of rereplicating forks [Davidson et al. 2006]. However, little information is available on the mechanisms leading to DNA damage and DDR activation in rereplicating human cells.

We combined cell/molecular biology and in vivo single-molecule approaches to investigate how deregulated origin licensing by Emi1 depletion affects replicating chromosomes. We show that cells experience mild DNA replication stress and ssDNA accumulation during the first replication round upon licensing deregulation, which may act as precursor for DNA breaks, when rereplicating forks approach ssDNA gaps on the template. Extending the analysis to other experimental systems of deregulated...
licensing [Gemmin depletion and Cdt1 addition in *X. laevis* extracts], we propose a new model for replication-induced chromosomal breakage, which may contribute to cancer-relevant genome rearrangements.

Results and Discussion

Emi1 depletion affects DNA synthesis prior to accumulation of >4N DNA, chromosomal breakage, and checkpoint activation

To gain mechanistic insight into how deregulated origin licensing affects the replication process, leading to DNA breaks and DDR activation, we depleted Emi1 in U2OS cells, a condition previously associated with rereplication and DNA damage [Machida and Dutta 2007]. Using flow cytometry, we monitored cell cycle progression [DNA content], DNA synthesis [EdU incorporation], and DDR activation [phosphorylation of H2AX [γH2AX]] (Supplemental Fig. S1) after Emi1 depletion. Sixteen hours to 24 h after siEmi1 transfection, we noticed accumulation of cells in S phase and a reduced incorporation rate in mid–late S phase (Fig. 1A). In synchronized cells, the impact of deregulated origin licensing on DNA synthesis was detected from the onset of the first S phase (Supplemental Fig. S2). At these time points, γH2AX was only detected in cells close to having completed a first round of bulk DNA replication (Fig. 1A,B). Later [32–40 h], γH2AX and a markedly reduced rate of DNA synthesis were detected in cells displaying >4N DNA, a commonly used readout for rereplication (Fig. 1A,C). Only at 32–40 h did cells accumulate detectable levels of double-strand breaks (DSB) and display activation of ATM and ATR pathways [phosphorylation of KAP1/RPA2-S4/S8 and CHK1/RPA2-S33, respectively], as expected for DSB-induced DDR (Fig. 1D,E). A relevant fraction of chromosomal fragments induced by Emi1 depletion is significantly smaller [20–100 kb] than camptothecin-induced DSB [0.5–2 Mb] (Fig. 1E; Supplemental Fig. S1B; Hanada et al. 2007), suggesting that rereplication-induced DSBs are clustered. DNA breakage at 32–40 h was also confirmed by colocalization of γH2AX and 53BP1, particularly evident in cells with “giant nuclei,” a sign of extensive rereplication (Supplemental Fig. S1C–E; Zhu et al. 2004). Altogether, these data indicate that mild replication stress during the first S phase after Emi1 depletion precedes cell cycle arrest, DNA breakage, and DDR activation, which are coupled to overt rereplication [DNA content >4N]. Similar observations were made in untransformed human epithelial cells [RPE-1] (Supplemental Fig. S3), showing that the stepwise impact on DNA replication and genome stability is a general consequence of Emi1 depletion.

Progressive RPA accumulation on chromatin precedes rereplication-associated DNA damage

To further characterize DNA replication stress early after Emi1 depletion, we monitored chromatin loading of the human ssDNA-binding protein [RPA] (Forment et al. 2012). Limited amounts of ssDNA are present during DNA replication, leading to RPA chromatin loading in S phase (Fig. 2A; Supplemental Fig. S4A; Forment et al. 2012). While this signal is rapidly lost as control cells complete S phase, Emi1 depletion leads to progressive accumulation of RPA on chromatin and unusually high RPA levels in mid–late S-phase cells (Fig. 2A). RPA foci colocalized with γH2AX foci at late time points [32–40 h] (Supplemental Fig. S4A), presumably marking processing of the detected DSBs (Fig. 1E; Supplemental Fig. S1C,D). However, in both U2OS and RPE-1 cells, some RPA accumulation was already observed at earlier time points.
Emi1 depletion does not detectably impair fork progression but induces refiring of clustered origins before accumulation of >4N DNA content

We next monitored the effect of Emi1 depletion on fork progression by DNA fiber spreading [Jackson and Pombo 1998]. Ongoing forks were identified by a red-green pattern [Supplemental Fig. S5A]. Fork progression appeared unaffected by Emi1 depletion when we used a 10-min labeling time [Fig. 3A]. However, in Emi1-depleted cells, track length increased more markedly than in control cells with longer labeling times [Fig. 3A; Supplemental Fig. S5C,D]. This suggests that Emi1 depletion does not affect progression of individual forks but that deregulated activation of clustered replication origins leads to more frequent fork fusion and thus longer tracts. We then adapted the labeling protocol to detect DNA rereplication events, modifying a published protocol [Dorn et al. 2009]. A 120-min CldU pulse followed by a 30-min IdU pulse allowed us to follow fork progression and reactivation of replication origins in previously replicated tracts [Supplemental Fig. S5A]. “Rereplication fork breakage on a gapped template”

Figure 3. Rereplication is detectable by a DNA fiber-spreading assay before completion of bulk DNA synthesis. (A) Length of newly replicated tracts [IdU; green] in mock-depleted U2OS cells [siLuc] and after Emi1 depletion [siEmi1 #1], using 10-min or 20-min labeling pulses. [B,C] Representative DNA tracts labeled with CldU for 2 h and IdU for 30 min to identify termination and rereplication events. [B] A replication “termination” event. [C] Two “rereplication” events in close proximity. [D] Quantification of rereplication/termination events as shown in B and C after mock [siLuc] or Emi1 depletion. The percentage indicated represents the fraction of rereplication events in the total population of “red-green-red” tracts analyzed. [Whiskers] 10–90 percentile; [***] P < 0.0001; [*] P < 0.005; [ns] not significant, Mann-Whitney test; n = 100 in A. Bar, 10 μm. See Supplemental Figure S5, A and B, for Emi1 levels and labeling protocols to study fork progression [A] and rereplication events [B,C].
Deregulation of origin licensing induces ssDNA gaps on replicated duplexes, which can be template for rereplication.

To gain additional insight into the molecular consequences of deregulated origin licensing, we investigated in vivo replication fork structure by electron microscopy (EM) [Neelsen et al. 2014]. Already 20 h after siRNA transfection (Fig. 3D). At 40 h, when rereplication has led to >4N DNA content [Fig. 1A,C], rereplication events were more frequent than fork fusions and were occasionally clustered on the same DNA fiber [Fig. 3C,D]. These data demonstrate that origin reactivation can be detected by DNA fiber spreading before it is detectable by flow cytometry and that refiring of clustered origins occurs already during a first round of replication with deregulated origin licensing.

Figure 4. Emi1 depletion leads to ssDNA gaps on the replicated duplex, which persist as a template for rereplicating forks. (A,C) Electron micrographs of representative replication forks from U2OS cells 40 h after transfection with siEmi1. Black arrows indicate ssDNA gaps. The insets show magnified ssDNA gaps and schemes of fork structure, indicating parental (P) and replicated (R) duplexes. Gaps are on a replicated duplex in A and on the parental duplex in C. Black and gray lines describe parental and newly synthesized DNA strands in the replicated duplexes, respectively. Bars: 100 nm (250 base pairs [bp]; inset, 50 nm). (B) Frequency of replication forks with ssDNA gaps in mock-depleted cells (siLuc) and after Emi1 depletion (siEmi1 #1). #RI is the number of analyzed replication intermediates. (D) Sperm nuclei replication assays in Xenopus interphase extracts. For S-phase experiments, extracts were optionally supplemented with 10 ng/μL Cdt1 at the time of sperm and [α-32P]dATP addition and incubated for 60 min. For G2 experiments, Cdt1 was optionally added with [α-32P]dATP 90 min after sperm addition and incubated for a further 60 min. After incubation, DNA was isolated, separated by neutral agarose gel electrophoresis, and autoradiographed. The dashed line indicates sperm DNA fragmentation. The asterisk indicates branched replicating DNA molecules retained in the well. (E) Frequency of replication forks with ssDNA gaps recovered after sperm nuclei incubation in S-phase or G2-phase extracts [see D], with optional addition of Cdt1. #RI is the number of analyzed replication intermediates. [P] Model for the formation of chromosomal breaks upon deregulation of origin licensing by Emi1 depletion. Excessive firing of clustered origins leads to replication stress during the first S phase and accumulation of ssDNA gaps. Uncontrolled reactivation of replication origins in this context triggers chromosomal breakage by replication of a discontinuous template.
These data strongly suggest that gaps accumulating during the first round of replication after Emi1 depletion persist and present a damaged template for new replication rounds.

To further test this hypothesis, we analyzed by EM a different experimental system associated with rereplication and DNA breakage, i.e., addition of Cdt1 to replication sperm nuclei in X. laevis egg extracts [Davidson et al. 2006]. In line with published results, addition of Cdt1 (Ferenbach et al. 2005) induced [α-32P]dATP incorporation in G2 extracts, particularly visible on branched DNA molecules retained in the well (Fig. 4D, Supplemental Fig. S6E). Moreover, rereplication was associated with DNA breakage when Cdt1 was added to S-phase and G2 extracts (Fig. 4D). The analysis of replication intermediates confirmed an accumulation of ssDNA gaps upon Cdt1 addition, particularly marked in G2 extracts (Fig. 4E) where multiple rounds of rereplication have been reported [Davidson et al. 2006]. Similar to Emi1 depletion, ssDNA gaps were also observed ahead of the replication forks, showing that rereplication was impaired by template discontinuities [Fig. 4E].

A new model for chromosomal breakage associated with deregulated origin licensing and rereplication

Our data strongly suggest that deregulated origin firing rapidly induces ssDNA gaps during DNA replication and that these persist in the template, where they cause stalling and eventually breakage of rereplicating forks [Fig. 4F]. Rereplicating forks could break by simply impacting ssDNA gaps on the template (“runoff”). Alternatively, they could stall upstream of the gap and later be resolved into DSBs by slow “runoff,” nuclease processing, or head-to-tail collision with forks generated in following rounds of rereplication. In support of transient stalling and remodeling, small replication bubbles accumulated upon Emi1 depletion, indicative of early fork stalling from reactivated origins. Moreover, the frequency of reversed forks remained high 40 h after Emi1 depletion, when rereplicating forks are overrepresented in our EM samples (Fig. 3D, Supplemental Fig. S6C). Fork reversal was associated with fork slowing during replication of a nicked template, thereby protecting forks from breakage [Ray Chaudhuri et al. 2012]. Thus, rereplicating forks could transiently arrest and reverse at ssDNA gaps before eventually undergoing breakage.

A prediction of our model is that the genotoxicity of rereplication correlates with the extent of origin firing deregulation in the previous replication round, as this creates the template discontinuities for rereplication. We tested this hypothesis by comparing the described effects for Emi1 depletion with Geminin depletion, a genetic condition associated with mild overreplication [Melixetian et al. 2004; Zhu et al. 2004], in which CDK-dependent Cdt1 inactivation partially restrains deregulated origin firing [Hook et al. 2007; Neelsen et al. 2013]. Thus, rereplicating forks could transiently arrest and reverse at ssDNA gaps before eventually undergoing breakage.

Materials and methods

Cell culture and transfections

U2OSs and hTERT RPE-1 retinal pigmented epithelial cells were grown in DMEM + 10% FCS. Cells were transfected with the indicated siRNAs using RNAiMAX (Invitrogen) according to the manufacturer’s instructions: siLuc (10 nM; 5'-GGUACGCGGAAUACUGAGdTdT-3'), siEmi1 #1 (10 nM; 5'-GAUUGUGACCUUCUUHUAAAdTdT-3'), siEmi1 #2 (10 nM; 5'-GAGAUUUCGGCGACAGUCU UAAdTdT-3'), and siGeminin (20 nM; 5'-UGCCCAUUCUGGAAUCAAAdTdT-3').

Methods

Flow cytometry was essentially performed as described previously for γH2AX/EdU/DAPI in Neelsen et al. (2013) and for γH2AX/RPA/DAPI in Forment et al. (2012). DNA fiber spreadings were performed according to Ray Chaudhuri et al. (2012) with the modifications outlined in the text. Pulse-field gel electrophoresis, immunofluorescence, and sample preparation for EM have been described in Neelsen et al. (2013, 2014). Protocols for Cdt1 purification, replication assays in X. laevis egg extracts, and isolation of genomic DNA for electron microscopic analysis can be found in the Supplemental Material. Detailed protocols for all other methods and a list of antibodies are included in the Supplemental Material.

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