

Is PCNA unloading the central function of the Elg1/ATAD5 replication factor C-like complex?

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Maintaining genome stability is crucial for all cells. The budding yeast Elg1 protein, the major subunit of a replication factor C-like complex, is important for genome stability, since cells lacking Elg1 exhibit increased recombination and chromosomal rearrangements. This genome maintenance function of Elg1 seems to be conserved in higher eukaryotes, since removal of the human Elg1 homolog, encoded by the ATAD5 gene, also causes genome instability leading to tumorigenesis. The fundamental molecular function of the Elg1/ATAD5-replication factor C-like complex (RLC) was, until recently, elusive, although Elg1/ATAD5-RLC was known to interact with the replication sliding clamp PCNA. Two papers have now reported that following DNA replication, the Elg1/ATAD5-RLC is required to remove PCNA from chromatin in yeast and human cells. In this Review, we summarize the evidence that Elg1/ATAD5-RLC acts as a PCNA unloader and discuss the still enigmatic relationship between the function of Elg1/ATAD5-RLC in PCNA unloading and the role of Elg1/ATAD5 in maintaining genomic stability.

found to be shared by its human homolog, called ATAD5 (ATPase family AAA domain-containing protein 5). Budding yeast cells lacking Elg1 exhibit pleiotropic chromosome instability phenotypes including increased recombination rate, gross chromosomal rearrangements, elongated telomeres, cohesion defects, and sensitivity to the DNA-alkylating drug MMS.¹⁻⁹ The role of Elg1 in maintaining chromosome stability seems to be conserved in higher eukaryotes, since mutating the ATAD5 gene caused genomic instability and tumorigenesis in mice and human cells.^{10,11} Elg1 and ATAD5 were identified as the major subunits of a replication factor C-like complex (RLC) having structural similarity to replication factor C (RFC), which loads the polymerase clamp PCNA at replication forks. The Elg1/ATAD5-RLC physically interacts with PCNA, but its physiological role in PCNA transactions for years remained obscure, making the Elg1-RLC a particularly mysterious guardian of genomic stability. Two recent papers have illuminated the *in vivo* effect of Elg1/ATAD5 on PCNA. Anne Donaldson's group reported that the Elg1-RLC functions in PCNA unloading from chromatin during replication in yeast,¹² while Kyungjae Myung's groups showed that ATAD5 is needed for proper removal of PCNA and disassembly of replication factories in human cell lines.¹³ In this review, we focus on these recent advances and discuss how loss of Elg1 function might cause genome instability. We begin by briefly introducing PCNA and previous investigations of the effects of RLCs. We then summarize advances in the 2 recent papers

Keywords: Elg1, ATAD5, RFC, PCNA, PCNA unloading, genome stability

Abbreviations: RLC, replication factor C-like complex; RFC, replication factor C

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The Elg1/ATAD5 Replication Factor C-like Complex: A Mysterious Guardian of Genomic Stability

The *ELG1* gene (enhanced level of genomic instability 1) was first identified in *Saccharomyces cerevisiae* as required for accurate chromosome maintenance, a role

and outline possible PCNA unloading mechanisms. Finally, we discuss how failure of PCNA unloading might impact on genomic stability.

Role of PCNA in DNA Replication

A central coordinator of DNA replication, PCNA is a homotrimeric, ring-shaped molecule that encircles DNA to act as a polymerase clamp as well as a sliding platform for recruitment of other replication and repair proteins, including DNA helicase, nuclease, ligase, and histone chaperones.¹⁴ On the lagging strand, PCNA cooperates with DNA polymerase δ to synthesize DNA discontinuously in a series of Okazaki fragments, 100–200 nucleotides in length. Ligation of the Okazaki fragments into a continuous daughter strand then depends on PCNA-mediated recruitment of the flap endonuclease FEN-1 and DNA ligase I.¹⁵

On the lagging strand, PCNA must be loaded repeatedly onto the DNA to synthesize each Okazaki fragment. PCNA is loaded by RFC, a hetero-pentameric complex consisting of Rfc1–5 subunits.^{16–18} The five subunits show sequence similarity to each other and are members of the AAA+ ATPase family. On binding ATP, RFC interacts with PCNA and opens its trimeric ring. The open PCNA-RFC complex then recognizes and loads at the 3' end of a primer–template junction. On hydrolysis of ATP, RFC ejects PCNA, leaving it on the DNA in closed form. The loading process is discussed in detail in excellent recent reviews.^{19,20}

RFC-Like Complexes

All eukaryotic cells contain a series of 3 RFC-like complexes (RLCs), which share with RFC the Rfc2–5 subunits, but in place of Rfc1 have an alternative subunit: Rad24 (called Rad17 in human), Ctf18, or Elg1 (ATAD5 in human) (Fig. 1).²¹ The structural resemblance of these RLCs with RFC suggested they may mediate interactions of ring-shaped clamps with DNA—and, indeed, Rad24-RLC, the best-understood of the three RLCs, acts to load the trimeric PCNA-like “9-1-1” complex at damaged DNA for checkpoint activation.^{22–25} The 9-1-1 complex is conserved, with subunits

called RAD9-HUS1-RAD1 in human and Ddc1-Mec3-Rad17 in *S. cerevisiae*. Ctf18-RLC is unique in the RLC family in forming a heptamer containing 2 additional subunits, Dcc1 and Ctf8.^{26,27} Ctf18-RLC is important for sister chromatid cohesion and replication checkpoint activation^{26,28,29} and is also required for proper telomere length regulation and telomere intranuclear positioning.³⁰ Using in vitro assays in biochemically defined systems, Ctf18-RLC was shown to be capable of loading PCNA onto DNA and also of unloading it,³¹ but whether the Ctf18-RLC loads or unloads PCNA in vivo remains unclear.

The Elg1/ATAD5-RLC was for years the most mysterious of the 3 RLCs. Although shown to physically interact with PCNA, in vitro biochemical assays suggested that Elg1-RLC could not load PCNA onto or unload it from DNA,^{6,31} so its fundamental molecular function remained elusive.

Investigation of PCNA Unloading by RFC and RLCs In Vitro

PCNA must be loaded for synthesis of each Okazaki fragment and unloaded on its completion, as well as at replication fork termination. It was proposed that RFC acts to unload PCNA as well as loading it; here, we summarize the in vitro studies that led to this suggestion. In 1996, the Hurwitz and O'Donnell groups reported that human RFC unloads PCNA from singly nicked circular plasmid, as tested using an in vitro system which uses gel filtration to measure free PCNA released from a PCNA-plasmid DNA complex.^{32,33} However, in a later investigation of PCNA ring opening by yeast RFC, the O'Donnell group made the surprising observation that a tetrameric Rfc2-Rfc5 subassembly (lacking Rfc1) is also capable of opening and unloading PCNA from nicked circular plasmid.³⁴ Since all RLCs contain Rfc2–5, this finding might suggest that all the RLCs as well as RFC could potentially open and unload PCNA, although such a lack of specificity seems unlikely in vivo. PCNA unloading by Elg1-RLC was not tested in this system.

Shibahara and Stillman also showed that human RFC can mediate unloading

of PCNA.³⁵ Using cell extract and a plasmid containing the SV40 origin, they found that in the presence of RFC, PCNA is removed from fully replicated SV40 plasmid DNA in an ATP-dependent manner.³⁵ The regulation of PCNA transactions in this SV40 replication system might, however, differ from those in normal genome replication.

The above investigations suggest that RFC can unload PCNA from DNA. However, using an assay similar to that used by the O'Donnell group, Peter Burgers' group could not detect significant unloading of PCNA by yeast RFC, even on testing several DNA substrates with various concentrations of proteins.³¹ The Rfc2–5 subcomplex was also unable to unload PCNA in their assay. Their failure to measure significant unloading suggests that, at least in the case of yeast, the efficiency of RFC as a PCNA loader far outweighs its efficiency as an unloader. While it is possible that RFC catalyzes cycles of PCNA loading and unloading, the current evidence does leave significant doubt about whether the canonical RFC complex is a major PCNA unloader in vivo.

In the same in vitro investigation, Bylund and Burgers found that neither Rad24-RLC nor Elg1-RLC were able to mediate significant unloading of PCNA.³¹ Remarkably, they did find that yeast Ctf18-RLC can unload PCNA from DNA in a reaction driven by ATP hydrolysis.³¹ However, Ctf18-RLC is probably not the main PCNA unloader in vivo, since *ctf18 Δ* cells or human cells subject to siRNA knockdown of *CTF18* expression show approximately normal levels of PCNA on chromatin in an unperturbed S phase.^{13,29} Its effectiveness in unloading PCNA from DNA in vitro³¹ does hint that Ctf18-RLC may play this role under specific in vivo circumstances yet to be elucidated.

These results are summarized in Figure 1. Taken together, previous studies certainly do not exclude that RFC might unload PCNA during DNA replication, but they have equally failed to demonstrate that RFC acts as the major PCNA unloader in the normal in vivo situation. The need for RFC to load PCNA has made it difficult to design experiments that might test for unloading of PCNA by RFC in vivo.

PCNA Unloading by *S. cerevisiae* Elg1-RLC

Several lines of evidence from yeast suggested a role for Elg1-RLC in PCNA regulation during DNA replication.^{6,29,36} Martin Kupiec's group demonstrated that yeast Elg1-RLC preferentially interact with SUMOylated PCNA, and in cells lacking Elg1, they found SUMOylated PCNA accumulated on chromatin in the presence of the DNA-alkylating drug MMS,³⁶ suggesting that Elg1-RLC might unload SUMOylated PCNA from

DNA. Using quantitative proteomics, Anne Donaldson's group showed that in yeast cells lacking Elg1, PCNA accumulates abnormally on chromatin in mid-S phase even without MMS treatment.^{29,37} These studies suggested PCNA unloading as a possible role for Elg1-RLC *in vivo*, but could not exclude the possibility that PCNA accumulation on chromatin might be an indirect effect of the *elg1Δ* mutation.

The new paper from the Donaldson group provides more direct evidence that yeast Elg1-RLC functions in PCNA unloading during DNA replication

(Fig. 1).¹² This study used an improved degron system to test the immediate effects of Elg1 depletion, revealing that PCNA accumulates on chromatin during the first S phase without Elg1. This accumulated PCNA is unloaded rapidly *in vivo* if Elg1 expression is switched back on, either during or after completion of DNA replication. Overexpression of Rfc1, Ctf18, or Rad24 could not substitute for Elg1 in removing accumulated PCNA. These results implicated Elg1 in PCNA unloading *in vivo*. The investigators then proceeded to demonstrate PCNA

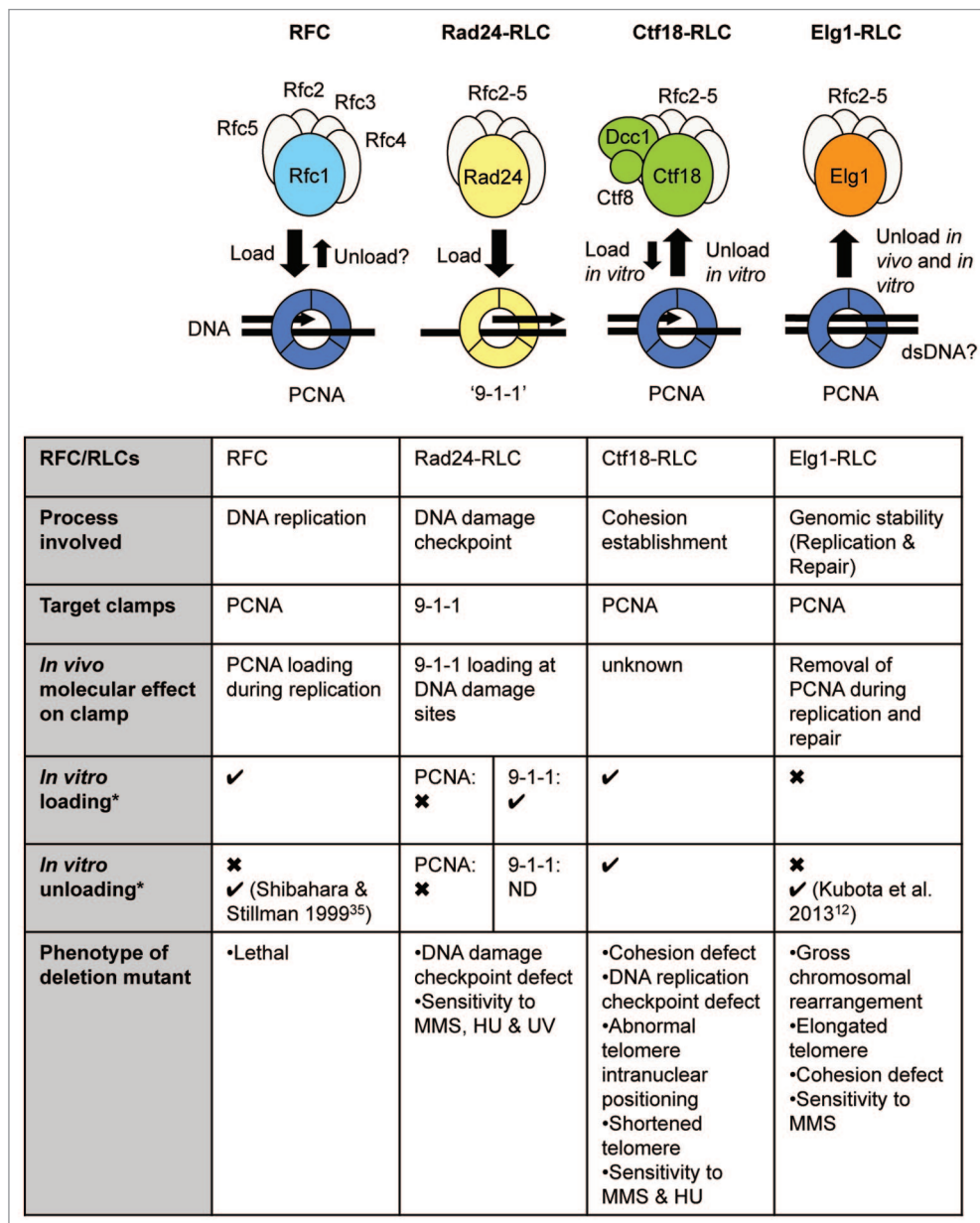


Figure 1. Summary of known functions of replication factor C and replication factor C-like complexes. *Except where indicated, *in vitro* loading and unloading results as derived from Bylund and Burgers 2005,³¹ Majka and Burgers 2003.²⁴ ND, not determined.

unloading by Elg1-RLC using a novel in vitro assay, in which chromatin prepared from an *elg1Δ* mutant was treated with affinity-purified Elg1-RLC, resulting in removal of PCNA. While formal identification of Elg1-RLC as a PCNA unloader will require reconstitution of the reaction with biochemically pure ingredients, this study provides the first coupled in vivo and in vitro evidence for PCNA removal by a clamp unloader.

The complex nature of the chromatin substrate used by Kubota et al.¹² leaves important questions about the biochemistry of PCNA unloading by Elg1-RLC. The fact that Elg1-RLC could not unload PCNA in the earlier biochemical assay using defined components³¹ suggests that there are additional requirements still to be identified that are needed for Elg1-RLC to unload PCNA from DNA. The earlier assay used purified PCNA loaded at a synthetic 3' primer–template junction,³¹ while use of native chromatin from *elg1Δ* as substrate may provide PCNA in a form that is more suitable for unloading. In particular, in the Donaldson group's system, the PCNA may be modified, histones can be deposited onto DNA, and any co-factors supporting Elg1-dependent PCNA unloading are likely to be present on chromatin. Alternatively, DNA structure might be critical for PCNA unloading by Elg1-RLC. For example, Elg1-RLC might be unable to unload PCNA from a nicked DNA substrate and may remove PCNA only from fully double-stranded DNA, as present after Okazaki fragment processing. It will be of interest to investigate the specific components or requirements needed for PCNA unloading by Elg1-RLC.

PCNA Removal by Human Elg1/ATAD5-RLC

Remarkably, an independent, simultaneous study provided compelling evidence that the function of Elg1/ATAD5 in PCNA unloading is conserved in human cells. This investigation from Kyungjae Myung's group demonstrated that human ATAD5 is critical for correct regulation of the lifespan of DNA replication factories.¹³ Specifically, knocking down ATAD5 in human cell lines caused PCNA

to accumulate on chromatin and remain there even after replication at that site had finished—analogueous to the retention of PCNA on chromatin seen by Kubota et al.¹² in a yeast *elg1Δ* mutant and consistent with the idea that Elg1/ATAD5-RLC unloads PCNA from DNA. Prolonged PCNA retention on chromatin caused by the knockdown of ATAD5 resulted in an extended lifespan of DNA replication factories, with all replication proteins enriched in the long-lived factories. This effect correlated with an accumulation on chromatin of other replisome proteins, including MRN complex and the chromatin-remodeling protein, CAF1. This accumulation was mediated by the excess PCNA retention, since simultaneous ATAD5 and PCNA knockdown mitigated the retention of other replisome components. As well as half-life, the intensity and number of DNA replication factories were increased upon knockdown of ATAD5. Interestingly, in the absence of ATAD5, many DNA replication factories were inactive for DNA synthesis, remaining even during G₂ phase of cell cycle. Conversely, ectopic ATAD5 expression caused a reduction of PCNA on chromatin, consistent with the idea that ATAD5 drives PCNA unloading. Although PCNA unloading by human ATAD5-RLC has yet to be directly demonstrated in vitro, the 2 recent studies by Donaldson's and Myung's groups together provide good evidence that the Elg1/ATAD5-RLC acts to unload PCNA during DNA replication, this function being conserved from yeast to human.

Does the Elg1/ATAD5-RLC Preferentially Unload SUMOylated PCNA?

Studies from Kupiec's group suggest the interesting idea that yeast Elg1-RLC is an unloader specific to SUMOylated PCNA. This idea is based on their observations that Elg1-RLC preferentially interacts with SUMOylated PCNA, and that SUMOylated PCNA accumulates on chromatin in *elg1Δ* mutant, especially in the presence of MMS.³⁶ Kubota et al. also observed an increase of SUMOylated PCNA on chromatin in *elg1Δ* mutant during unperturbed S phase.¹² However, an unSUMOylatable PCNA Pol30-K164R/

K127R still accumulates on chromatin when Elg1 is deleted,¹² implying that PCNA SUMOylation is not essential for PCNA to be unloaded by Elg1-RLC. Although SUMOylation is not absolutely required for PCNA unloading by Elg1-RLC, the observations by Kubota et al.¹² are still consistent with the idea that Elg1-RLC has a preference for unloading SUMOylated PCNA. DNA association leads to PCNA SUMOylation,³⁸ suggesting that the increase of SUMOylated PCNA in an *elg1Δ* mutant is probably a direct consequence of the prolonged retention of PCNA on chromatin when its normal unloader is absent.

SUMOylation of PCNA has been observed in higher eukaryotes, such as chicken DT40 cells, *X. laevis* egg extracts, and recently mammalian cells.^{39–43} In human cell lines, the modification is present at very low levels.⁴³ Human ATAD5-RLC could potentially prefer to interact with SUMOylated PCNA, since ATAD5 has a SUMO interaction motif in its N-terminal domain.^{44,45} However, accumulation of SUMOylated PCNA on chromatin was not observed in ATAD5-knockdown cells, even though both unmodified and ubiquitinated PCNA increased on chromatin (the latter due to defects in recruiting USP1-UAF1 deubiquitinating enzymes to PCNA).^{13,44} Overall, it seems likely that yeast Elg1-RLC has a preference, but not an absolute requirement, to unload SUMOylated PCNA, but any such preference appears not to be conserved in human ATAD5-RLC.

Structural Domain Requirements for PCNA Unloading

What are the structural features of Elg1/ATAD5 that might permit it to act as a PCNA unloader? As illustrated in **Figure 2**, Elg1 has a central domain showing similarity to Rfc1 and Ctf18 and containing a conserved ATPase domain, flanked by unique N-terminal and C-terminal regions. Below we discuss our understanding of the Elg1/ATAD5 structure in relation to its proposed role in PCNA unloading.

ATPase domain

All subunits of RFC/RLC subunits (excluding Dcc1 and Ctf8) are members

of the AAA+ ATPase family and contain ATP-binding Walker A and B motifs (Fig. 2). Walker A motifs contain an invariant lysine that is critical for ATP binding.⁴⁶ In human Rfc1, and in yeast Rad24 and Ctf18, this invariant lysine is critical for function,^{31,47-50} but surprisingly, the lysine in yeast Rfc1 is not essential for PCNA loading *in vivo* or *in vitro*.^{51,52}

Human Elg1/ATAD5 also contains this invariant lysine within a consensus Walker A motif (Fig. 2). Myung's group has tested if the ATPase domain in human Elg1/ATAD5 is important for its function by mutating the invariant lysine (K1138). An ATAD5 K1138E mutant protein could not prevent PCNA retention on chromatin or the defective replication factories of

ATAD5 knockdown cells, suggesting the invariant lysine is important for removal of PCNA from chromatin.¹³ However, since the K1138E mutation also affects the stability of ATAD5 protein and its interaction with small Rfc subunits,¹³ failure of PCNA removal may not be solely due to loss of ATP binding.

Yeast Elg1 lacks some of the consensus residues of the Walker A motif (Fig. 2). The invariant lysine is present in yeast Elg1 but does not appear to be required for its *in vivo* function, in contrast with human ATAD5. Davidson and Brown mutated the invariant lysine (Elg1 K343) and found the cells exhibited wild-type levels of sensitivity to MMS (unlike the *elg1Δ* mutant which shows greatly elevated MMS sensitivity).⁵³ Their study

suggests that, similar to yeast Rfc1, ATP binding to yeast Elg1 is not essential for its *in vivo* function. One possibility is that ATP binding to the smaller subunits Rfc2–5 is sufficient for PCNA loading by RFC and unloading by Elg1-RLC in budding yeast. Overall, current results suggest the requirement for ATP binding to Elg1/ATAD5-RLC may differ among species. It will be of interest to test directly whether ATP binding by yeast Elg1 is required for PCNA unloading.

N-terminal domain

The unique N-terminal domain of Elg1 may make a crucial contribution to PCNA unloading, as it interacts with PCNA and SUMOylated PCNA and is known to be important for *in vivo* function of Elg1.^{36,53} Budding yeast Elg1 has

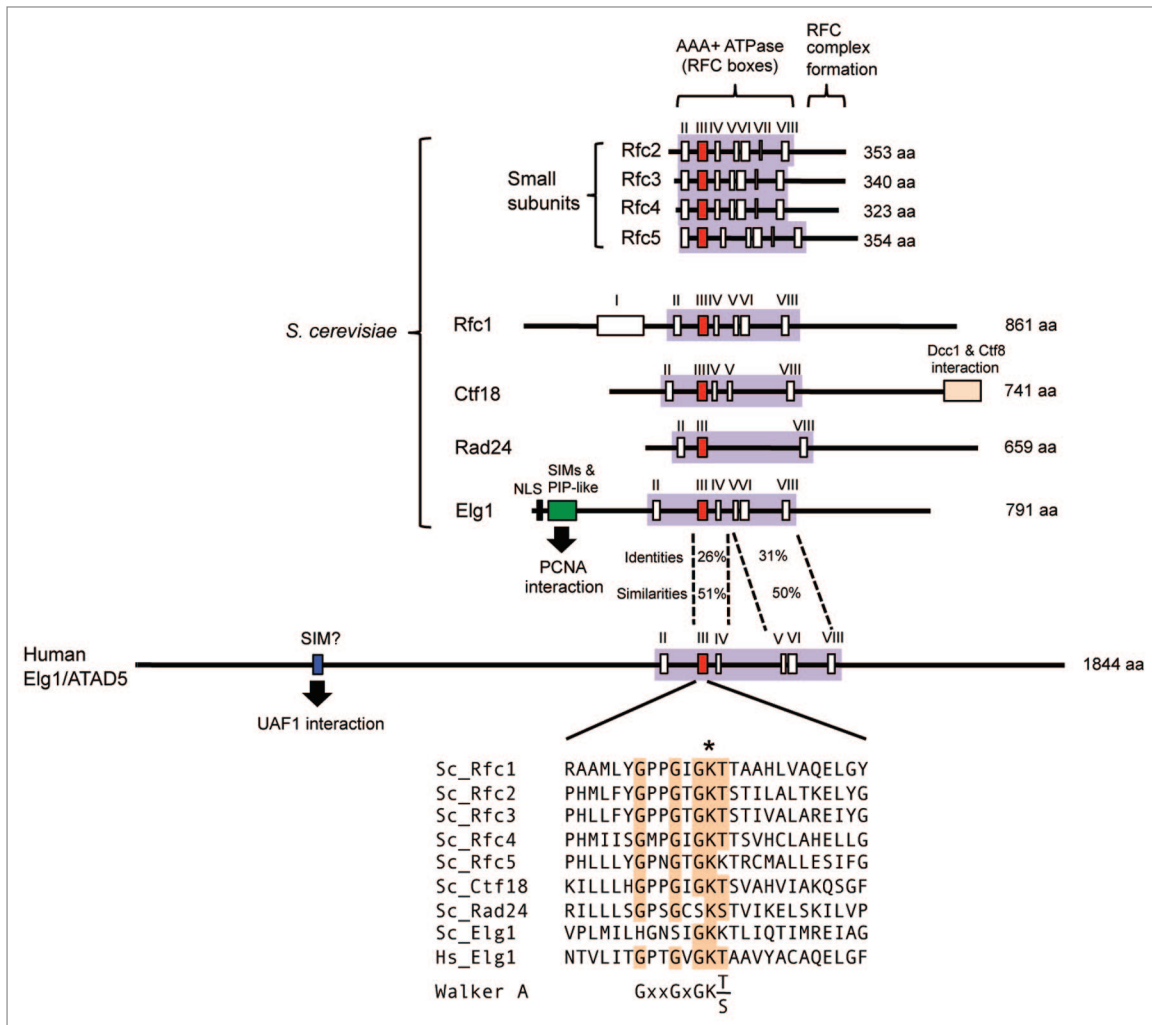


Figure 2. Structure of RFC/RLCs. Within the AAA+ ATPase domain (colored lilac), box III (colored red) contains Walker A sequence, box V contains Walker B sequence, box VII (in Rfc2–5 subunits) contains Arginine finger. Detailed sequence comparison of Walker A motifs shown below. *Indicates conserved lysine. NLS, nuclear localization signal; SIM, SUMO interaction motif; PIP-like, PCNA interacting peptide-like.

3 SUMO interaction motifs (SIMs) and a PCNA interacting peptide-like (PIP-like) motif in its N-terminal region, conferring a preferential interaction with SUMOylated PCNA (Fig. 2).³⁶ Cells expressing Elg1 lacking the N-terminal 215 amino acids exhibit greater MMS sensitivity than wild type, but less than an *elg1Δ* mutant.⁵³ These results suggest that the N-terminal region in Elg1 is important for its function in PCNA regulation, and probably in PCNA unloading. Consistent with this idea, yeast cells expressing Elg1 mutated in both SIMs

and PIP-like motif also show substantial MMS sensitivity.³⁶

One possibility is that 2 binding modes occur during Elg1-PCNA interaction. First, Elg1-RLC might search for chromatin-bound PCNA/SUMO-PCNA using its N-terminal region, which is probably extended from core “RFC” domain and could act as flexible “scanner”. After docking with its target (i.e., PCNA to be unloaded), Elg1-RLC may then bind PCNA strongly through all 5 subunits (in a manner similar to RFC), and then open the PCNA ring and release it from DNA.

Human ATAD5 has a much longer extended N-terminal region than yeast Elg1, and is therefore larger than yeast Elg1 (1844 a.a. and 791 a.a., respectively) (Fig. 2). Human ATAD5 has a SUMO interaction motif in its N-terminal region, shown to interact with a SUMO-like domain in the deubiquitination factor UAF1.^{44,45} Human ATAD5 regulates PCNA deubiquitination by recruiting the USP1-UAF1 complex to ubiquitinated PCNA.⁴⁴ The N-terminal region in human ATAD5 therefore likely confers additional functions in regulation of ubiquitination compared with yeast Elg1. Any PCNA-binding region in the ATAD5 N-terminus has yet to be investigated, and human ATAD5 may also interact with PCNA through its N-terminal region, like yeast Elg1.

C-terminal domain

There is less information about the function of C-terminal domain of yeast Elg1. Loss of the C-terminal 60 amino acids does not affect sensitivity to MMS, except in the context of an N-terminal truncation, where it does somewhat exacerbate MMS sensitivity.⁵³ The C-terminal domain of yeast Elg1 may also therefore contribute to its function, either in PCNA unloading or other cellular roles as discussed below. Interestingly, a C-terminal truncation mutation in ATAD5 found in endometrial tumor caused instability of protein,¹⁰ suggesting the ATAD5 C-terminal domain is important to suppress tumorigenesis.

Is Elg1-RLC the Main PCNA Unloader?

The observations described above suggest strongly that Elg1/ATAD5-RLC acts to unload PCNA, but do not address whether Elg1/ATAD5-RLC is the primary PCNA unloader during normal DNA replication. For example, Elg1/ATAD5-RLC might unload PCNA only at specific genomic sites. Three distinct models (Fig. 3) are consistent with the published data: (A) Elg1-RLC is the main PCNA unloader. If this is the case then other mechanisms must exist that can eventually unload PCNA, since most PCNA is eventually removed from chromatin after replication even in the absence

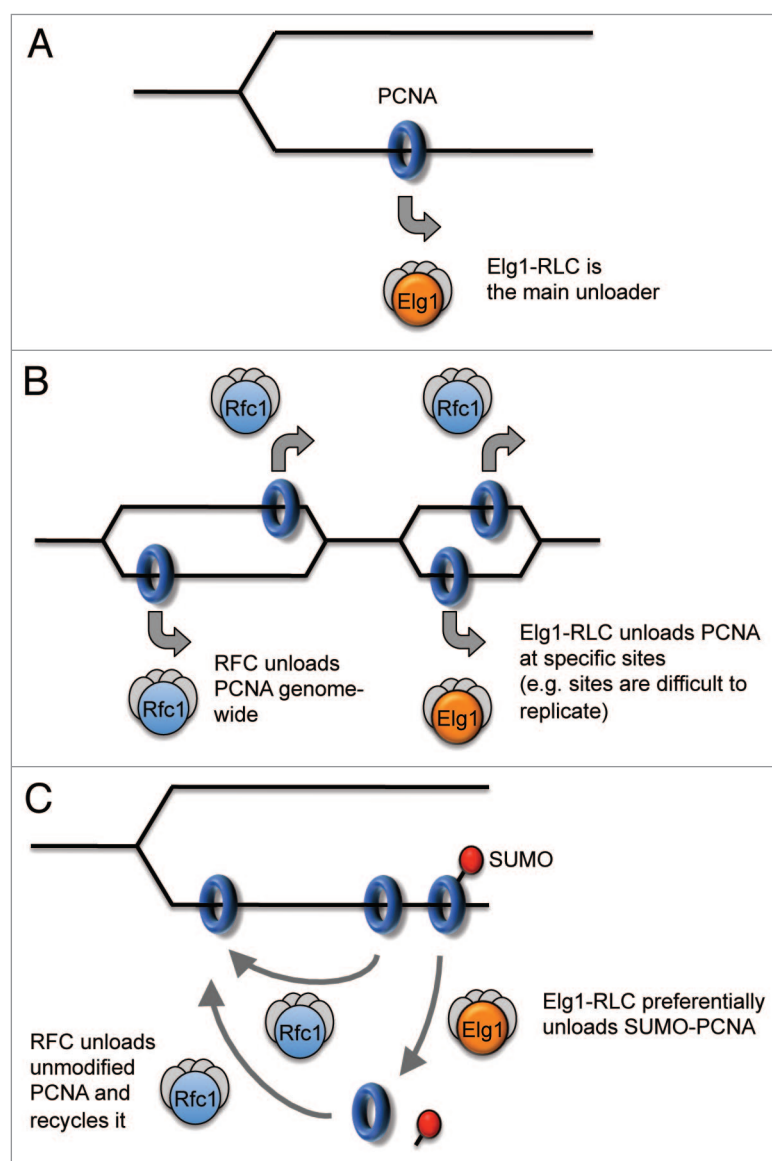


Figure 3. Three models of PCNA unloading. (A) Elg1-RLC is the main unloader genome-wide. (B) Elg1-RLC acts at specific sites. (C) Elg1-RLC primarily acts to unload SUMOylated PCNA. See text for further discussion

of Elg1/ATAD5. Without Elg1/ATAD5, PCNA-unloading function could conceivably be taken over by RFC and/or Ctf18-RLC, but with less efficient removal by these complexes resulting in the observed PCNA accumulation on chromatin. It is also possible that PCNA might eventually dissociate from DNA spontaneously without any active unloader.³⁴ (B) RFC could be the major PCNA unloader, with Elg1/ATAD5-RLC as a minor unloader acting at particular sites, for example, sites of cohesion binding or specific sequences that are difficult to replicate. In this case, the accumulation of PCNA on chromatin observed in the absence of Elg1/ATAD5 presumably represents PCNA retained at these particular locations. (C) A further possibility is that RFC is the main unloader of unmodified PCNA, with Elg1-RLC acting primarily to unload SUMOylated PCNA (even though it can also remove unmodified PCNA). In this case, RFC might unload unmodified PCNA from a newly ligated Okazaki fragment and immediately re-load it on a new primer-template junction, allowing rapid recycling of unmodified PCNA. Elg1-RLC might, in contrast, preferentially unload PCNA that has become SUMOylated but not re-load it (due to lack of loading activity). In this case, the PCNA could only be re-loaded after de-SUMOylation and subsequent recognition by RFC, providing a mechanism to prevent loading of SUMOylated PCNA. In fact Models (A) and (C) are not mutually exclusive, since PCNA may usually be SUMOylated by the time it is unloaded, and therefore unloaded by Elg1-RLC in most cases. Current results are consistent with all three possibilities, although the marked PCNA accumulation on chromatin when Elg1/ATAD5 are lacking might be construed as less compatible with its retention only at specific sites (Model B). It will be of interest to test these possibilities directly.

Is PCNA Unloading the Main Cellular Function of the Elg1/ATAD5-RLC?

Budding yeast lacking Elg1 shows pleiotropic phenotypes, including increased chromosomal rearrangements,

cohesion defects, elongated telomeres, and sensitivity to MMS (Fig. 1).¹⁻⁹ Similarly, low expression of ATAD5 in human or mouse cells caused increased chromosomal rearrangements, increased rate of sister chromatid exchange, and mild sensitivity to DNA damaging agents.^{10,11} In addition, homozygous null mutation of *Atad5* in mice caused embryonic lethality (Myung et al., unpublished results). Do all these phenotypes result from compromised PCNA unloading, or do they instead indicate that Elg1/ATAD5 has additional molecular roles beyond PCNA unloading? At this point the answer remains unclear, but the former is a possibility, since, as explained below, these *elg1Δ* phenotypes could conceivably all result from problems with PCNA transactions. ATAD5 has also been shown to regulate the removal of ubiquitin from PCNA: we also consider possible links between PCNA unloading by ATAD5-RLC, PCNA deubiquitination, and the genome instability observed in mammalian cells in the absence of ATAD5.

Increased chromosomal rearrangements

Could compromised PCNA unloading be an indirect cause of chromosome rearrangements? At least 2 pathways can be envisaged. First, failure of timely PCNA unloading may delay the recycling of PCNA and PCNA-binding proteins, in turn delaying lagging-strand synthesis and resulting in unprocessed lagging-strand replication intermediates that could stimulate inappropriate recombination events, leading ultimately to chromosome rearrangements. Consistent with this idea, cells lacking FEN-1, the flap endonuclease that processes Okazaki fragments, show increased recombination rates and gross chromosomal rearrangements.^{54,55}

A second possible pathway could involve the accumulation on chromatin of PCNA-binding proteins that promote unwanted recombination. One candidate is the AAA+ ATPase Mgs1 that interacts with PCNA. Mgs1 has both PCNA-binding and UBZ (ubiquitin-binding) domains and preferentially interacts with polyubiquitinated PCNA,⁵⁶ and in an *elg1Δ* mutant, both ubiquitinated PCNA and Mgs1 are abnormally increased on chromatin.^{12,29} Polyubiquitinated PCNA

is thought to direct error-free repair through template switching,^{57,58} and while the molecular function of Mgs1 is mysterious, either deletion or overexpression of the *MGS1* gene affects recombination and mutation rates.⁵⁹⁻⁶¹ It therefore seems possible that the PCNA retention on chromatin in *elg1Δ* leads to an imbalance of Mgs1 recruitment, eventually leading to chromosome rearrangements. In general, it will be of interest to investigate which of the many pathways coordinated by PCNA are affected by loss of Elg1-RLC activity.

Cohesion defect

Cohesion establishment is coupled with DNA replication, and cohesion defects are reported in *S. cerevisiae* lacking replication proteins, including Ctf4, Ctf18, Mrc1, Tof1, Csm1, and FEN-1.^{28,62,63} Properly coordinated leading- and lagging-strand progression may be important for cohesion establishment, so that problems with lagging-strand synthesis in *elg1Δ* cells due to compromised PCNA unloading or recycling might result in cohesion defects. Robert Skibbens's group has proposed that cohesion establishment occurs in concert with lagging-strand synthesis⁶⁴—although Frank Uhlmann's group found no role of the lagging-strand processing enzymes FEN-1 and Dna2 in cohesion establishment.⁶⁵ A variant of this idea proposes that inefficient PCNA unloading causes progression problems when replication forks encounter cohesin rings, possibly affecting cohesion establishment.

Yet another possibility is that the accumulation of SUMOylated PCNA on chromatin interferes with function of the cohesion establishment factor Eco1. Eco1 normally promotes cohesion by acetylating the Smc3 cohesin subunit during S phase.⁶⁶⁻⁶⁹ Eco1 is recruited by PCNA, and PCNA SUMOylation appears to counteract Eco1 activity.⁷⁰ It is therefore possible that the excessive PCNA SUMOylation observed in an *elg1Δ* mutant (probably as a result of extended DNA association) interferes with the function of Eco1 in cohesion establishment. Notably, a *pol30-104* mutant (expressing PCNA A251V) that is over-SUMOylated also shows a cohesion defect.⁷⁰

Elongated telomeres

Again, the elongated telomere phenotype of *elg1Δ* cells might be connected to

lagging-strand synthesis problems caused by compromised PCNA unloading or recycling. Other mutations (*poll1*, *rfc1*, *pol32Δ*, and *rad27Δ*), affecting lagging-strand synthesis also cause elongated telomeres.⁷¹⁻⁷³ One idea is that delayed lagging-strand synthesis leaves single-stranded DNA exposed and unable to recruit the telomerase inhibitor complex Rap1-Rif1/Rif2 (which binds dsDNA), but still bound by the telomerase-recruiting complex Cdc13-Stn1-Ten1 (which binds to single-strand DNA), promoting telomere extension. Alternatively, PCNA accumulation itself may cause telomere extension by occupying the telomeric DNA and interfering with Rap1-Rif1/Rif2 recruitment, leading to inappropriate extension by telomerase.

MMS sensitivity

We suspect that the MMS sensitivity of *elg1Δ* cells may be connected to PCNA retention on chromatin, since we find that overexpression of PCNA also causes over-loading of PCNA on chromatin that is coupled with increased MMS sensitivity (unpublished data). The mechanism could result from an imbalance of PCNA modifications important for DNA repair.

Does loss of other activities of Elg1 cause genomic instability?

While speculative, the mechanisms described above suggest plausible pathways, consistent with established observations, through which compromised PCNA unloading might cause the *elg1Δ* mutant phenotypes. But of course molecular roles of Elg1 unrelated to PCNA unloading could account for some defects of an *elg1Δ* mutant. In particular, Elg1 interacts with SUMOylated and SIM-containing proteins through poly-SUMO chains, independent of SUMOylated PCNA.⁷⁴ Disturbing such interactions could affect recombination rate, cohesion establishment, and telomere extension, because SUMOylation is involved in all these processes.⁷⁵⁻⁷⁹ Clearly, further investigations will be needed to shed light on the issue of which *elg1Δ* phenotypes result from compromised PCNA unloading and which from other Elg1 molecular functions.

Human Elg1/ATAD5 promotes PCNA deubiquitination by recruiting the

USP1-UAF1 complex to ubiquitinated PCNA.⁴⁴ Can failure to deubiquitinate PCNA explain the increased chromosomal rearrangement in ATAD5-knockdown cells? Cells knocked down for USP1 expression show reduced homologous recombination, and a similar effect is observed upon knockdown of ATAD5.⁴⁴ Thus, the effect of ATAD5 in facilitating PCNA deubiquitination activity may be important during DNA double-strand break-induced homologous recombination. However, studies of mouse mutants suggest that ATAD5 plays a functional role extending well beyond its effect on the deubiquitination system. A null mutation of *atad5* in mice causes embryonic lethality (Myung lab, unpublished result), but *usp1*-null mice are in contrast born (although with an anemic phenotype). No tumorigenesis was observed in the *usp1*-null mice—in stark contrast to the high incidence of tumorigenesis in *atad5* heterozygous mice.^{10,80} These observations suggest that ATAD5 PCNA unloading activity is important for embryonic development and tumor suppression, in addition to its role in PCNA deubiquitination.

Concluding Remarks and Future Directions

Unloading of PCNA must take place repeatedly on completion of every Okazaki fragment, and also at replication fork termination. Two recent papers have reported that Elg1/ATAD5-RLC functions in removing PCNA from chromatin in yeast and human. These findings represent a substantial advance in understanding the operation of replication forks as well as Elg1/ATAD5 function. Compromised PCNA unloading could impact on multiple chromosome maintenance mechanisms to cause the genome instability of *elg1Δ* cells, through PCNA accumulation on chromatin, delayed recycling of PCNA, or altered recruitment of PCNA-interacting factors, including cohesion factors and DNA repair proteins.^{14,29,56} It will be of interest to investigate which PCNA-coordinated pathways are affected by loss of the Elg1-RLC, and the phenotypic consequences, especially in relation to the other activities of Elg1.⁷⁴

Disclosure of Potential Conflicts of Interest

No potential conflicts of interest were disclosed.

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