

Disordered regions and folded modules in CAF-1 promote histone deposition in *S. pombe*

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Abstract

Genome and epigenome integrity in eukaryotes depends on the proper coupling of histone deposition with DNA synthesis. This process relies on the evolutionary conserved histone chaperone CAF-1, for which the links between structure and function are still a puzzle. While studies of the *S. cerevisiae* CAF-1 complex enabled to propose a model for the histone deposition mechanism, we still lack a framework to demonstrate its generality and in particular, how its interaction with the polymerase accessory factor PCNA is operating. Here, we reconstituted a complete SpCAF-1 from fission yeast. We characterized its dynamic structure using NMR, SAXS and molecular modeling together with *in vitro* and *in vivo* functional studies on rationally designed interaction mutants. Importantly, we identify the unfolded nature of the acidic domain which folds up when binding to histones. We also show how the long KER helix mediates DNA binding and stimulates SpCAF-1 association with PCNA. Our study highlights how the organization of CAF-1 comprising both disordered regions and folded modules enables the dynamics of multiple interactions to promote histone deposition coupled to DNA synthesis, and is essential for CAF-1 functions in DNA replication, heterochromatin maintenance, and genome stability.

eLife assessment

This **important** study advances our understanding of the machinery that couples DNA synthesis with the deposition of histone proteins onto newly synthesized DNA. A **solid** array of experiments combines NMR, protein biochemistry, and *in vivo* analyses of Chromatin Assembly Factor-1 of fission yeast. The work is of interest to researchers in the field of chromosome/chromatin biology as well as epigenetics.

Introduction

In eukaryotes, genomic DNA is packaged in a dynamic nucleoprotein complex, the chromatin, which protects DNA and regulates its accessibility. The fundamental repeat unit of chromatin, the nucleosome core particle, comprises 146 bp of DNA wrapped around a histone octamer including a tetramer of histone H3–H4 flanked by two dimers of H2A–H2B (Luger et al. 1997 [\[1\]](#)). Histone chaperones are critical players in ensuring histone traffic and deposition. Without energy consumption, they escort histones, facilitate their transfer and deposition on DNA, and provide links with DNA based-processes such as DNA replication, repair and gene transcription (Gurard-Levin et al. 2014 [\[2\]](#)). In line with these key properties, perturbations of histone chaperones are associated with defects in genome and epigenome maintenance and function as found in cancer, aging and viral infections (Yi and Kim 2020 [\[3\]](#), Sultana et al. 2021 [\[4\]](#), Ray-Gallet and Almouzni 2022 [\[5\]](#)). Discovered over thirty years ago (Smith and Stillman 1989 [\[6\]](#)), conserved in all eukaryotes (Loyola and Almouzni 2004 [\[7\]](#)), the histone chaperone Chromatin Assembly Factor 1 (CAF-1) is central and unique in promoting the deposition of replicative histones H3–H4 in a manner coupled to DNA synthesis, i.e. during DNA replication and repair, and is also involved in heterochromatin maintenance (see (Ridgway and Almouzni 2000 [\[8\]](#), Gurard-Levin et al. 2014 [\[2\]](#)) for review). The unique feature of CAF-1 is that it provides a link with DNA synthesis via its association with the trimeric DNA polymerase processivity factor, Proliferation Cell Nuclear Antigen (PCNA), through PCNA Interacting Protein motifs (PIP) (Martini et al. 1998 [\[9\]](#), Moggs et al. 2000 [\[10\]](#), Rolef Ben-Shahar et al. 2009 [\[11\]](#), Pietrobon et al. 2014 [\[12\]](#), Gopinathan Nair et al. 2022 [\[13\]](#)), but the precise function of this interaction remains to be examined.

CAF-1 comprises three subunits (Figure 1a [\[14\]](#)) (Smith and Stillman 1989 [\[6\]](#), Kaufman et al. 1997 [\[15\]](#), Dohke et al. 2008 [\[16\]](#)). While progress in uncovering its molecular/genetic properties derives from work in *Saccharomyces cerevisiae*, and biochemical work in human cells, there is still a lack of atomic information for the complex. In *cerevisiae*, CAF-1 is a hetero-trimer that binds to a single H3–H4 dimer (Mattioli et al. 2017 [\[17\]](#), Sauer et al. 2017 [\[18\]](#)). The current model states that histone binding induces a conformational rearrangement promoting its interaction with the DNA. Two complexes must co-associate to ensure the deposition of H3–H4 tetramers on DNA in the first step of nucleosome assembly (see (Sauer et al. 2018 [\[19\]](#)) for review). Two domains of the large subunit Cac1 contribute to DNA binding (Zhang et al. 2016 [\[20\]](#), Sauer et al. 2017 [\[18\]](#)), the conserved low complexity region called KER (for Lys, Glu and Arg rich) and the C-terminal Winged Helix Domain (WHD). These features are conserved in human CAF-1 (Gopinathan Nair et al. 2022 [\[13\]](#)). However, a comprehensive and dynamic view of the CAF-1 complex is still missing, and most critically understanding how the functional domains cooperate to ensure efficient histone deposition coupled to DNA synthesis remains to be determined. Moreover, the degree of conservation of these properties across species and possible connections with heterochromatin regulation elements such as histone H3K9 trimethylation and HP1 recruitment remain unknown.

To tackle these issues, we isolated the fission yeast complex (Pcf1-Pcf2-Pcf3) and investigated its binding mode with its three main partners, histones H3–H4, DNA and PCNA. Based on novel structural insights, we designed targeted mutations to specifically identify the key features promoting Pcf1 interactions with DNA, PCNA and histones H3–H4. We next monitored functional impacts of these mutations *in vitro* and *in vivo* by analysing the phenotypes of the corresponding mutants in fission yeast. We show that disordered or partly disordered segments of the large subunit of CAF-1 are key for interactions with H3–H4 and DNA, underscoring the dynamic nature of the binding interface between CAF-1, histones H3–H4, and DNA. Upon histone binding, the acidic domain (ED) of the large subunit folds and this conformational changes impact CAF-1-PCNA interaction *in vivo*. We propose that such conformational changes upon histone binding contribute to PCNA/CAF1 recycling during DNA replication. We further show that PCNA binding accelerates nucleosome assembly *in vitro* and is also essential for the proper targeting of the complex to the

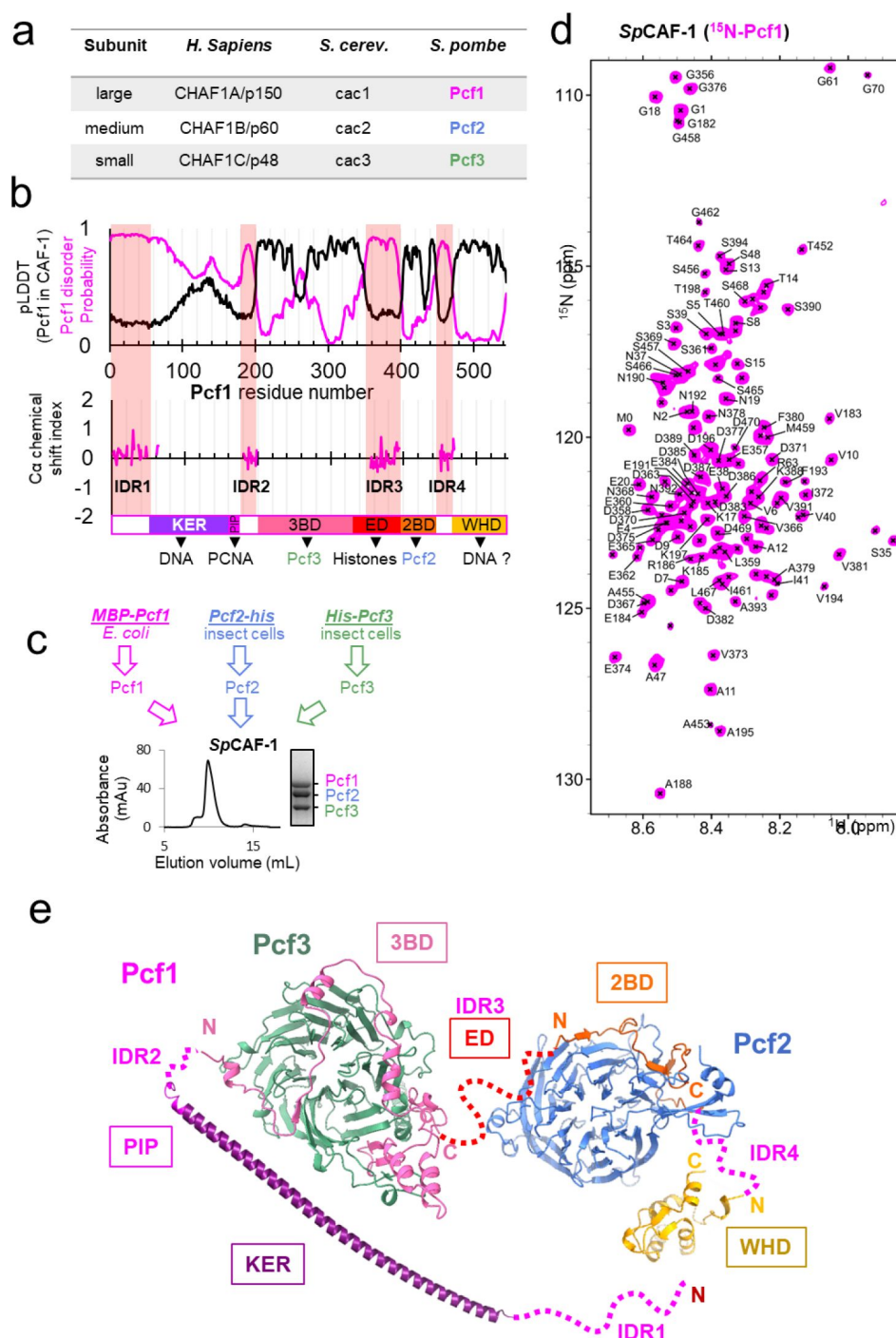


Figure 1

The large *SpCAF-1* subunit includes four intrinsically disordered regions (IDR)

a Names for the large, the medium and the small subunits of CAF-1 in *H. sapiens*, *S. cerevisiae* and *S. pombe*. **b** Upper panel: The magenta line shows the predicted disorder of Pcf1 (spot disorder software) and the black line the Ca Local Distance Difference Test (pLDDT) calculated for Pcf1 residues by the AlphaFold2 model of the full *SpCAF-1* complex. Lower panel: Ca chemical shift index calculated for the 101 assigned residues. This Ca chemical shift index is consistent with their disordered nature. The four IDR regions are highlighted with pink semi-transparent vertical bars. The predicted domains of Pcf1 are labeled. **c** General strategy for the production of *SpCAF-1*. The lower panel shows the purification SEC profile and the SDS-PAGE purity of the sample. **d** ^1H - ^{15}N SOFAST-HMQC spectrum of the FL *SpCAF-1* complex composed of uniformly labeled ^{15}N -Pcf1 and unlabeled Pcf2 and Pcf3 (*SpCAF-1*(^{15}N -Pcf1)). The assigned signals are labeled. **e** AF2 model of the *SpCAF-1* complex. The four IDR segments are shown with a dashed line. The relative orientation of the four modules is arbitrary.

chromatin *in vivo*. Finally, we found that the WHD C-terminal domain does not bind DNA but allows to uncouple the unique functions of CAF-1 in replication-dependent chromatin assembly, genome stability and heterochromatin maintenance. We suggest that this domain contributes to specify CAF-1 functions.

Results

Global organization of the full-length *SpCAF-1* complex

The large subunit of CAF-1, present in all major groups of eukaryotes, exhibits significant sequence divergence (16% sequence identity between *SpPcf1* and *ScCac1* and 21% sequence identity between *SpPcf1* and *HsCHAF1A/p150*, **Figure S1a**). Given this high sequence divergence, conserved biochemical properties between *ScCAF-1* and *SpCAF-1* should reveal important functional features. Of particular interest, *S. pombe* heterochromatin shares with human a regulation based on the recruitment of Swi6, the Heterochromatin Function 1 (HP1) orthologue, via histone H3K9 trimethylation. From sequence alignments, the 6 main conserved regions previously proposed to contribute to the nucleosome assembly activity of CAF-1 can be inferred in *SpPcf1* sequence, a KER domain, a single PIP motif, an acidic domain (ED domain), the domains predicted to bind Pcf2 (2BD) and Pcf3 (3BD) and a C-terminal WHD domain (**Figure S1b**). Although *SpPcf1* is shorter than *ScCac1* and *HsCHAF1A/p150* (544 residues instead of 606 and 956, respectively) its sequence includes a remarkable high abundance of predicted Intrinsically Disordered Regions (IDRs) (**Figure 1b**). These IDRs include the predicted histone-binding domain (Pcf1_ED), the PCNA (PIP motif) and the DNA binding domain (Pcf-1 KER).

We produced and purified the three subunits of *SpCAF-1* separately from bacteria and insect cells (**Figure 1c**, materials & methods). When isolated, both Pcf2 and Pcf3 are monomeric while Pcf1 forms large soluble oligomers (**Figure S1c**). Mixing the subunits by pairs, we observed stable complexes for Pcf1-Pcf2 and Pcf1-Pcf3 by size exclusion chromatography (SEC). Pcf2 and Pcf3 did not interact with each other (**Figure S1c-d**) suggesting that the large subunit Pcf1 mediates the complex assembly. We next reconstituted and isolated the recombinant full-length (FL) *SpCAF-1* complex by SEC (**Figure 1c**). An experimental molecular weight of 179 kDa was calculated using Small Angle X-ray Scattering (SAXS), consistent with a 1:1:1 stoichiometry (**Figure S1e**). These data are in agreement with a globular complex with a significant flexibility (**Figure S1f**).

To determine the extent of disorder in the large subunit of *SpCAF-1*, Pcf1 was produced with uniform ^{15}N (or ^{15}N - ^{13}C) labeling. The CAF-1(^{15}N -Pcf1) complex with unlabeled Pcf2 and Pcf3 was reconstituted and SEC-purified. Given the size of this complex (167 kDa), we expected that only amide signals from residues in long disordered regions could be observed by Nuclear Magnetic Resonance (NMR) spectroscopy. The ^{15}N - ^1H spectrum shows about 140 amide signals, revealing that up to a quarter of Pcf1 residues are intrinsically disordered in the full *SpCAF-1* complex (**Figure 1d**). These residues are located in four continuous segments of Pcf1 and define intrinsically disordered regions that we labeled IDR1 to IDR4 (**Figure 1b**). IDR1 corresponds to the ~50 N-terminal residues of the protein, IDR2 (181-198) is located between the PIP motif and the 3BD region, IDR3 (355-394) overlaps a large segment of the acidic ED domain and IDR4 (451-470) is located between the 2BD region and the C-terminal WHD domain. The boundaries of the four IDRs are in agreement with the segments of Pcf1 predicted to harbor disorder with a high probability (**Figure 1b**).

We next built a model of the *SpCAF-1* complex using the AlphaFold2 multimer software (AF2) with one copy of each full-length protein (**Figure 1e**, **Figures S1g-m**). The model is consistent with our biochemical data showing that Pcf1 mediates the complex assembly. Also, in agreement with their disordered nature, low values around 0.2-0.3 of the local quality of the model as calculated by the Local Distance Difference Test (pLDDT) were obtained in the four IDR segments with a

remarkable match for the delimitations of the four IDR segments by pLDDT values and NMR data (**Figure 1b**). Accordingly, these segments are symbolized with a dashed line in **Figure 1e**. In contrast, significantly high pLDDT was obtained for the 3BD, 2BD and WHD domains of Pcf1 and for the two subunits Pcf2 and Pcf3. (**Figure 1b**, **Figures S1g**). These data allowed to identify four independent modules, not predicted to interact with each other. The first module corresponds to the KER domain of Pcf1, predicted to form a long helix ending by the PIP motif. The second module contains the Pcf2 subunit, composed of 7 WD repeats arranged in a circular fold, and a segment of Pcf1 corresponding to the 2BD domain forming three short beta strands and a short helix (**Figure S1h-j**). In the third module, the 3BD domain of Pcf1 composed of 7 helices and 3 beta strands establishes a large interface with Pcf3, composed of 7 circular WD repeats (**Figure S1k-m**). The fourth domain is the WHD domain. We next used these models to fit our SAXS data allowing flexibility between the four modules. The best model fitted the experimental data with a high accuracy and is in agreement with a relatively globular complex. Superimposing the generated models did not define a unique orientation between the four modules, suggesting that the complex has an inter-module flexibility (**Table S1**, **Figure S1n-o**).

Taken together, our findings indicate that the large subunit Pcf1 mediates the (1:1:1) complex assembly. Pcf1 includes four IDR, and can organize its key regions (KER, PIP, 3BD, ED, 2BD and WHD) allowing them to be exposed and bound by Pcf2 and Pcf3 simultaneously.

In the FL *SpCAF-1* complex, the acidic ED domain is disordered but folds upon histone binding

We next investigated the interaction of *SpCAF-1* with histones H3–H4. A stable complex was isolated by SEC at low (150 mM NaCl) and high (1 M NaCl) salt concentrations, confirming that the reconstituted *SpCAF-1* complex tightly binds histones (**Figure 2a**). SAXS measurements at low salt allowed to calculate an experimental molecular weight of 193 kDa for this complex, showing that *SpCAF-1* binds a dimer of histones H3–H4 (**Figure S1e**). In addition, these data are compatible with a more extended shape compared to *SpCAF-1* alone (**Figure S1f**).

Addition of histones to *SpCAF-1*(¹⁵N-Pcf1) led to a drastic decrease in intensity of the NMR signal specifically for residues in the IDR3 segment (**Figure 2b**, **Figure S2a**). To further characterize this domain, we designed a short construct of Pcf1 (325–396), called Pcf1_ED, corresponding to the IDR3 segment extended in its N-terminus with the conserved acidic segment (325–355) (**Figure 2c**, **Figure S1b**). We confirmed the fully disordered nature of Pcf1_ED by NMR (**Figure S2b-c**). Signals corresponding to residues 355–394 (IDR3) remarkably overlap in the spectra of Pcf1_ED and *SpCAF-1*(¹⁵N-Pcf1) (**Figure S2a**), showing that this segment was fully flexible in *SpCAF-1*, and did not interact with other regions of the complex. Upon binding of unlabeled histones H3–H4, we observed the vanishing of almost all NMR signals of ¹⁵N Pcf1_ED as in the full *SpCAF-1* complex (**Figure 2d**). In contrast, a large part of signals (338–396) did not vanish anymore upon addition of a histone complex preformed with two other histone chaperones known to compete with CAF-1 for histone binding (Sauer et al. 2017), Asf1–H3–H4–Mcm2(69–138) (**Figure 2d**). This region of the ED domain, is indeed in direct competition with Asf1 and Mcm2 whose histone binding modes are well established (**Figure 2e**) (Huang et al. 2015, Richet et al. 2015). Fully consistent with this NMR competition experiment, this segment of Pcf1_ED domain was predicted by AlphaFold2 to interact with histones H3–H4 through the same surface as the one bound by Mcm2 (**Figure 2f**, **Figure S2d**). Two highly conserved positions in Pcf1, L359 and F380, are thus proposed to mediate histone H3–H4 binding in the same region as Mcm2 (**Figure 2f**). We next used these AlphaFold2 models to fit the SAXS curve of the *SpCAF-1*–H3–H4 complex allowing reorientation of the different modules (**Figure S1p-q**). Remarkably, all generated models show a significant exclusion of the KER domain from the complex, suggesting that the KER domain of *SpCAF-1* becomes more accessible upon histone binding.

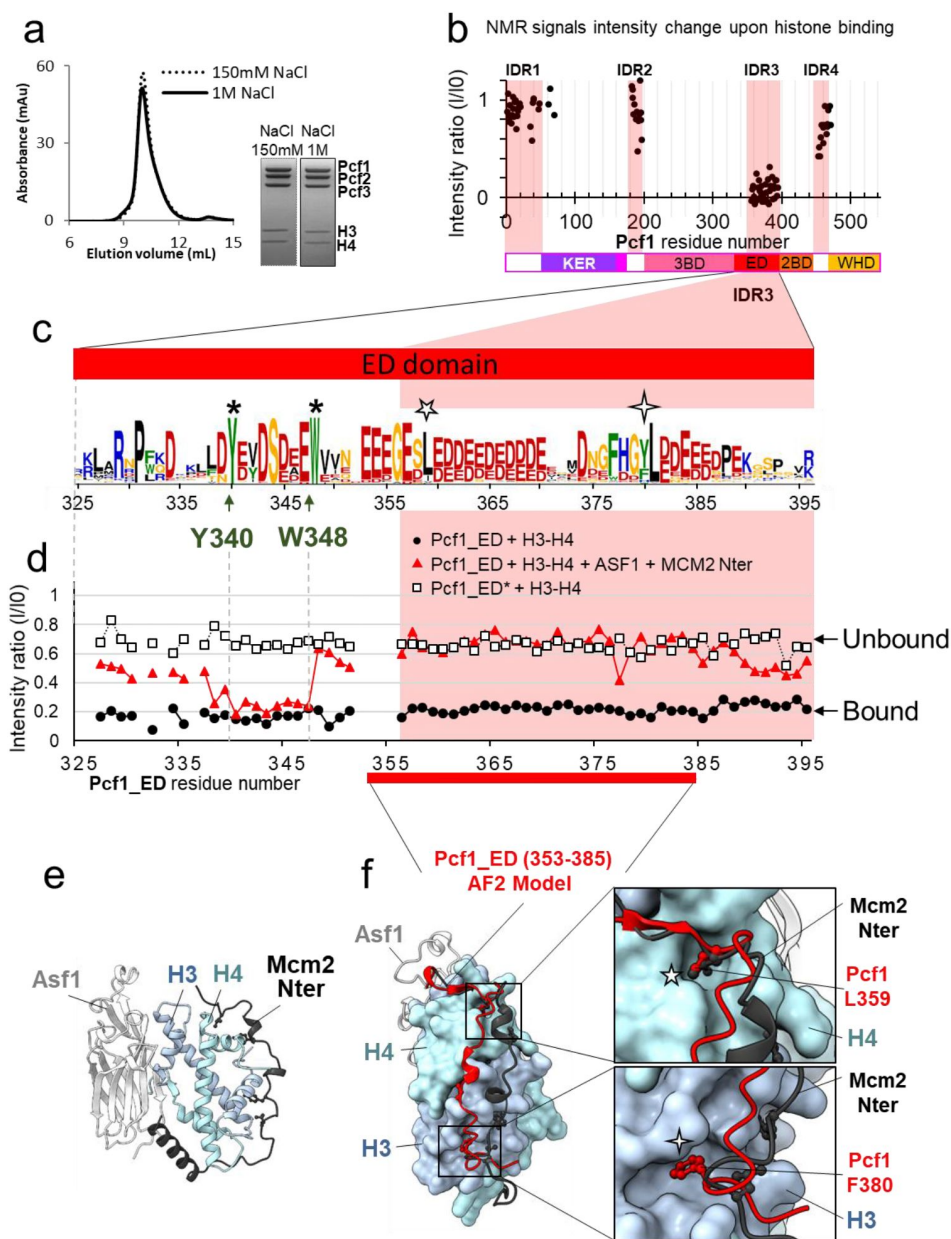


Figure 2

The acidic ED domain binds histones alone and in the full *SpCAF-1* complex

a SEC profile and the SDS-PAGE purity of *SpCAF-1*-H3-H4 histones at 150 mM NaCl and 1 M NaCl. **b** Mapping of the interaction between *SpCAF-1* (^{15}N -Pcf1) and *SpH3*-H4 histones, using the intensities ratio (I/I_0), where I and I_0 are the intensity of the signals ^1H - ^{15}N SOFAST-HMQC spectra before and after addition of histones, respectively. **c** Sequence Logo of the ED domain generated with a large data set of Pcf1 homologues. The position of the two conserved residues Y340 and W348, mutated in ED* are indicated with stars and conserved Pcf1 L359 and F380 residues with five and four branch stars respectively. **d** Mapping of the interaction between Pcf1_ED or Pcf1_ED* with *SpH3*-H4 histones using the intensities ratio (I/I_0) as in **b**. Histones were added alone or previously complexed with histones chaperones. **e** Cartoon representation of the complex between human histones H3-H4 (light blue and cyan), Asf1 (light grey) and Mcm2 (dark grey) (PDB: 5BNX). **f** AlphaFold2 model of Pcf1 (353-385) (as red cartoon), corresponding to the segment of the ED domain indicated in red, in complex with histones H3-H4 (light blue and cyan surface) superimposed with Mcm2 and Asf1 as in panel **e**. The two insets represent zoomed views of the sidechains of the conserved Pcf1 L359 and F380 residues (red sticks) binding into H4 and H3 pockets, respectively. The same four and five branch stars are used to label these positions in the logo panel **c**.

The NMR competition experiment also reveals that an additional region of Pcf1_ED domain (338-351) is involved in the interaction with H3-H4 but is not competing with the Asf1-Mcm2 module. (**Figure 2d** [↗](#)). In order to alter the interaction of the ED domain with histones without modifying its charge and without interfering with Asf1 or Mcm2 binding, we identified from sequence alignments in this segment (**Figure 2c** [↗](#)), two invariant hydrophobic residues, Y340 and W348, that were mutated into alanines (mutant called ED*, **Figure S1b**). As expected, the isolated Pcf1_ED* domain showed almost no histone binding as observed by the intensity of ^1H - ^{15}N NMR signals (**Figure 2d** [↗](#)). We next monitored the impact of the ED* mutations in the context of the full *SpCAF-1* complex. To do so, the mutations Y340A-W348A were introduced in the full length Pcf1, and the complex reconstituted with the uniformly ^{15}N labeled Pcf1-ED* and unlabeled Pcf2 and Pcf3 (**Figure S2e**). The ^1H - ^{15}N NMR spectrum of this mutant was similar that of the WT complex, but upon addition of unlabeled histones H3-H4 no major change was observed (**Figure S2a**), which strongly suggest an alteration of the histone binding of this mutant.

In summary, we identified critical amino-acids in the ED domain involved in H3-H4 binding. We also showed that addition of histones leads to a conformational change in the *SpCAF-1* complex with less disorder in the ED domain and an increased accessibility of the KER domain.

***SpCAF-1* binds dsDNA longer than 40bp**

We next analyzed the DNA binding properties of *SpCAF-1*. Electrophoretic mobility shift assays (EMSA) were performed with a DNA ladder as substrate in order to determine the minimal DNA size for *SpCAF-1* binding. The complex *SpCAF-1* showed significant binding for DNAs longer than 40bp (**Figure S3a**). EMSAs with a double-stranded 40bp DNA fragment confirmed the homogeneity of the bound complex. When increasing the *SpCAF-1* concentration, additional mobility shifts suggest, a cooperative DNA binding (**Figure 3a** [↗](#)). Micro-Scale Thermophoresis (MST) measurements were next performed using an alexa-488 labeled 40bp dsDNA (**Figure 3b** [↗](#), **Table 1** [↗](#)). The curves were fitted with a Hill model with a EC_{50} value of $0.7 \pm 0.1 \mu\text{M}$ and a cooperativity (Hill coefficient, h) of 2.7 ± 0.2 , in line with a cooperative DNA binding of *SpCAF-1*.

The KER domain is the main DNA binding region of *SpCAF-1*

The KER and WHD domains of the CAF-1 large subunit were shown to be involved in DNA binding in *ScCAF-1* and *HsCAF-1* (Zhang et al. 2016 [↗](#), Sauer et al. 2017 [↗](#), Ayoub et al. 2022 [↗](#), Gopinathan Nair et al. 2022 [↗](#)). We were thus interested to explore the conservation of these features in Pcf1. We first isolated the KER domain (Pcf1_KER, **Figure S1b**), and an extended fragment we called Pcf1_KER-PIP (**Figure S1b**), which includes the PIP motif (Q₁₇₂-L-K-L₁₇₅-N-N-F₁₇₈-F₁₇₉). These domains are predicted by AlphaFold2 to form a long helix with partial disorder at both ends and possible extension over the first half of the PIP motif (**Figure 1d** [↗](#), **Figure 3c** [↗](#)). Using a combination of circular dichroism (CD) (**Figure S3b-c**), SEC-SAXS (**Figure S1e-f**, **Figure S3d**) and NMR (**Figure S3e-f**), we confirmed that the isolated KER domain of Pcf1 domain forms a straight monomeric helix, partially continuing over the PIP motif. This long helix exhibits a strong bias in amino acid composition and remarkably, almost all basic residues are positioned on the same side of the helix (**Figure 3d** [↗](#)) providing a suitable interface for DNA binding (Gopinathan Nair et al. 2022 [↗](#)). We performed EMSA using a DNA ladder as substrate and we found that Pcf1_KER domain binds DNA that are longer than 40bp, as observed with the full *SpCAF-1* complex (**Figure S3e**). EMSAs with double strand 40bp DNA fragment showed the presence of multiple bands for Pcf1_KER bound DNA, indicating a possible cooperative DNA binding of this fragment (**Figure 3d** [↗](#)). Affinity measurements by MST led to a EC_{50} of $1.1 \pm 0.2 \mu\text{M}$ for Pcf1_KER with a cooperativity around 3, consistent with EMSA experiments (**Figure 3d** [↗](#), **Table 1** [↗](#)). The DNA binding properties of Pcf1_KER-PIP are comparable to that of Pcf1_KER (**Figure 3e** [↗](#), **Table 1** [↗](#), **Figure S3f**). The EC_{50} obtained for the isolated Pcf1_KER are also close that of the full *SpCAF-1* complex ($0.7 \pm 0.1 \mu\text{M}$) suggesting that the KER domain constitutes the principal DNA binding domain of *SpCAF-1*.

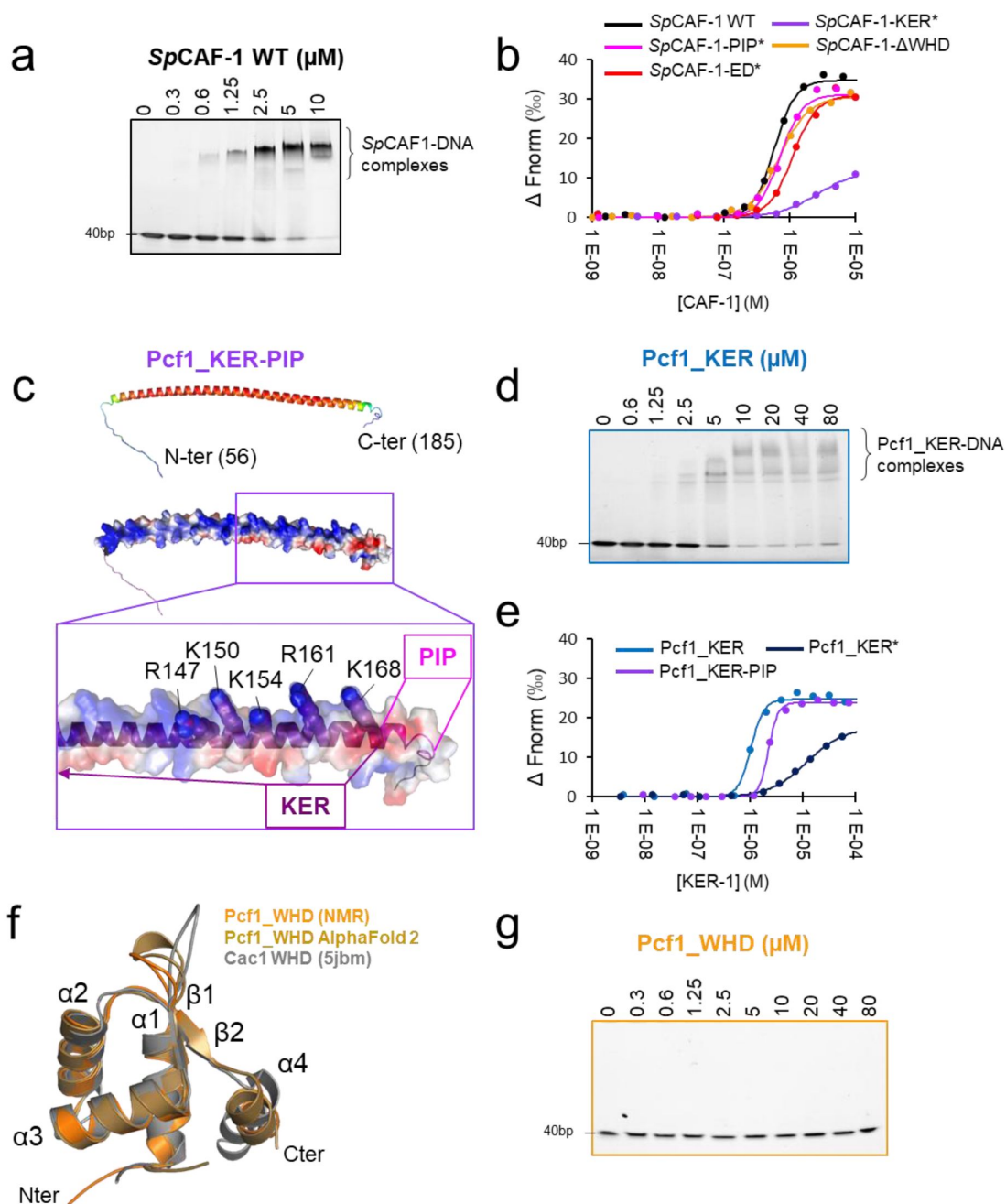


Figure 3

Pcf1_KER is the main DNA binding domain of *SpCAF-1*:

a EMSA with *SpCAF-1* and 40 dsDNA (1 μ M) revealed with SYBR SAFE staining. **b** Microscale thermophoresis (MST) fitted curves of *SpCAF-1* WT and mutants with 40bp dsDNA. **c** Upper panel: Modelled structure of the Pcf1_KER-PIP domain (56-185) rainbow coloured according to the pLDDT of each residue. Red corresponds to pLDDT values of 1 and dark blue of 0. Middle panel same model represented with its electrostatic surface. Lower panel: zoom of the C-terminus of the KER domain and the PIP motif. The five mutated residues are labeled and highlighted with spheres. **d** EMSA of Pcf1_KER binding with a 40bp dsDNA (1 μ M) revealed with SYBR SAFE staining. **e** MST fitted curves of Pcf1_KER constructs and mutants with 40bp dsDNA. **f** Overlay of the calculated model of the WHD domain obtained with the CS-rosetta software (light orange) using NMR assignments of the domain (Figure S3n), with AlphaFold2 (gold) and the structure of Cac1 WHD from budding yeast (PDB 5jbm, in grey)(Liu et al. 2016)(Grey). **g** EMSA revealed with SYBR SAFE staining of Pcf1_WHD domain with a 40 dsDNA (1 μ M).

Construct	EC50 (μM)	Hill coeff., h
Pcfl KER	1.1 ± 0.2	3.3 ± 0.5
Pcfl KER*	12.2 ± 0.7	1.5 ± 0.3
Pcfl KER-PIP	1.9 ± 0.3	5.2 ± 0.9
Pcfl WHD	Not detected	Not detected
<i>SpCAF-1</i> WT	0.7 ± 0.1	2.7 ± 0.2
<i>SpCAF-1</i> -ΔWHD	0.7 ± 0.1	2.3 ± 0.3
<i>SpCAF-1</i> -KER*	2.8 ± 0.4	1.3 ± 0.3
<i>SpCAF-1</i> -ED*	1.0 ± 0.1	2.3 ± 0.1
<i>SpCAF-1</i> -PIP*	0.7 ± 0.1	2.7 ± 0.3

Table 1

Experimental affinities of different *SpCAF-1* constructs with a 40bp dsDNA measured by Microscale thermophoresis (MST).

Based on these results, we designed a mutant called Pcf1_KER* with a charge inversion for five positive residues at the C-terminus of the potential DNA binding face of the KER helix (R147E-K150E-K154E-R161E-K168E) (**Figure 3c**, **Figure S1b**). The CD analysis of Pcf1_KER* shows this mutant is mainly helical (**Figure S3g**). MST quantification confirmed that the mutation of Pcf1_KER* impaired DNA binding by a factor of 10 even though residual DNA binding remained (**Figure 3e**, **Figure S3h-j**, **Table 1**). The KER* mutation was then introduced in the full complex SpCAF-1-KER* (**Figure S1b**, **Figure S2e**) and we confirmed by MST and EMSA its lower affinity for dsDNA (**Figure 3b**, **Figure S3k**, **Table 1**). Importantly, the NMR signals of all IDR for this mutant with or without histones were close to that of the WT (**Figure S3l-m**) indicating that the KER* mutation did not impair histone binding.

The C-terminal of Pcf1 folds as a WHD domain but does not bind DNA

We next isolated the Pcf1_WHD domain (**Figure S1b**) and confirmed by NMR and AlphaFold2 that its global fold is similar to ScWHD (Liu et al. 2016, Zhang et al. 2016) (**Figure 3f**, **Figure S3n**). Unexpectedly, Pcf1_WHD does not interact with DNA of any size (**Figure 3g**, **Figure S3n**). The residues involved in DNA binding in ScWHD, K564 and K568, correspond to S514 and G518 in SpWHD, respectively, leading to a different electrostatic surface, probably not favorable for DNA binding (**Figure S3p-q**). To further investigate the role of the WHD domain of SpCAF-1, the WHD domain was deleted in the reconstituted SpCAF-1-ΔWHD complex (**Figure S1b**, **Figure S2e**) and analyzed by EMSA and NMR. We observed the similar DNA binding property and IDR properties for SpCAF-1-ΔWHD and the WT complex (**Table 1**, **Figure 3b**, **Figure S3k-m**).

Together our results show that the KER domain constitutes the main DNA binding region of SpCAF-1 and that the WHD domain does not contribute to this binding.

Crosstalk between DNA and PCNA binding

The PIP motif of Pcf1 was found crucial for SpCAF-1 interaction with PCNA *in vivo* (Pietrobon et al. 2014). Given its proximity with the KER domain, we further investigated potential cross-talks between PCNA and DNA binding. We first measured by isothermal microcalorimetry (ITC) an affinity of 7.1 ± 1.3 μM between SpPCNA and a short PIP motif segment (**Figure S4a**, **Table 2**). This affinity is in the same range (2-fold less affine) as a peptide isolated from the replicative polymerase delta from *S. pombe* Cdc27 (**Figure S4a**, **Table 2**). In agreement with its consensus sequence, the binding mode of Pcf1_PIP motif to SpPCNA is predicted by AlphaFold to be canonical (**Figure S4b**). Consistently, no binding was observed for the Pcf1_PIP* peptide with 4 Alanine mutations, previously designed to disrupt the PIP motif (**Figure S4a**, **Table 2**) (Pietrobon et al. 2014). We next measured the affinity of the longer fragment Pcf1_KER-PIP for SpPCNA and observed an affinity gain of a factor 10 (0.7 ± 1.3 μM) (**Figure S4c**, **Table 2**), revealing interactions between the KER domain and PCNA. ITC also fits a stoichiometry of ~2 Pcf1_KER-PIP per PCNA trimer, suggesting that, in each PCNA trimer, one monomer remains unbound and potentially accessible for binding to other partners. Pcf1_KER-PIP* did not interact with PCNA confirming the importance of the PIP motif for this association (**Figure S4c**, **Table 2**). The KER* mutation impaired the interaction of Pcf1_KER*-PIP with PCNA of a factor 10 reaching the affinity of the short isolated Pcf1 PIP peptide (**Figure S4c**, **Table 2**). Collectively these results show that both the PIP motif and the C-terminal part of the KER domain are involved in PCNA binding.

To reveal possible crosstalk between CAF-1 binding to PCNA and DNA, we analysed, in the presence or absence of dsDNA, the binding of the full SpCAF-1 complexes (WT SpCAF-1, SpCAF-1-PIP*, SpCAF-1-ED*, SpCAF-1-KER* and SpCAF-1-ΔWHD) with recombinant SpPCNA, using EMSA (**Figure 4a**, **Figure S4d**). For all combinations tested, we quantified binding by monitoring the disappearance of free PCNA (**Figure 4b**) and free DNA (**Figure 4c**). In this assay, only 20% of free PCNA intensity was lost by addition of DNA (**Figure 4b**), probably because the PCNA trimer can slide along the linear DNA and dissociates during the migration. In the absence of DNA, we

Ligand	Kd (μM)	ΔG (kCal.M ⁻¹)	N*	ΔH (kCal.M ⁻¹)	-TΔS (kCal.M ⁻¹)
Pcf1_PIP	7.1 ± 1.3	-6.9 ± 0.1	0.97 ± 0.08	-2.9 ± 0.2	-0.39 ± 0.3
Pcf1_PIP*	undetectable	ND	ND	ND	ND
Pcf1_KER-PIP	0.7 ± 0.2	-8.2 ± 0.2	0.64 ± 0.04	+2.9 ± 0.6	-11.2 ± 0.8
Pcf1_KER*-PIP	7.1 ± 1.5	-6.9 ± 1.2	0.7 ± 0.2	+1.0 ± 0.5	-7.9 ± 0.7
Pcf1_KER-PIP*	undetectable	ND	ND	ND	ND
Cdc27_PIP	3.5 ± 0.3	-7.3 ± 0.1	0.9 ± 0.1	-4.8 ± 0.02	-2.4 ± 0.1

Table 2

Interactions parameter with SpPCNA measured by isothermal microcalorimetry (ITC): *The stoichiometry (N) is calculate as a molar ratio of monomeric PCNA.

observe a small but significant decrease of free PCNA upon addition of WT *SpCAF-1*, in agreement with the relatively low binding affinity between Pcf1_KER helix and PCNA (**Table 2** [↗](#)). In contrast, in the presence of dsDNA, addition of an excess of WT *SpCAF-1* leads to the complete disappearing of the free PCNA band and to a large shift of the band corresponding to *SpCAF-1*-DNA, corresponding to a larger complex engaging CAF-1, PCNA and DNA (**Figure 4a** [↗](#)). *SpCAF-1*-ED* and *SpCAF-1*-ΔWHD show similar binding properties for PCNA and DNA compared to WT CAF-1. In contrast, *SpCAF-1*-PIP* binds DNA like the WT, but is strongly impaired for PCNA binding alone and in the presence of DNA, while *SpCAF-1*-KER* is impaired for binding both DNA and *SpPCNA*. In agreement, the large shifted band corresponding to a *SpCAF-1*-PCNA-DNA complex is not observed for these two mutants (**Figure S4d**).

Altogether, our data show the stabilization of the CAF-1-PCNA interaction by DNA that requires both the KER domain and the PIP motif but not the ED and WHD domain. Conversely, the capacity of CAF-1 to bind PCNA does not impair its interaction with DNA.

***In vitro* histone deposition properties of *SpCAF-1* mutants**

We next examined the ability of the full *SpCAF-1* complex reconstituted with the four Pcf1 mutants (*SpCAF-1*-PIP*, *SpCAF-1*-ED*, *SpCAF-1*-KER*, *SpCAF-1*-ΔWHD) to promote nucleosome assembly mediated by CAF-1 in a complex environment closer to physiological conditions. We used *Xenopus* high speed egg extract (HSE) that are powerful systems competent for chromatin assembly and effective to exploit depletion/complementation assays (Ray-Gallet and Almouzni 2004 [↗](#)). We depleted HSE for the endogenous *Xenopus* CAF-1 largest subunit (xp150) and assessed the capacity of *SpCAF-1*-PIP*, *SpCAF-1*-ED*, *SpCAF-1*-KER* and *SpCAF-1*-ΔWHD to complement these xp150-depleted extracts (Ray-Gallet and Almouzni 2004 [↗](#), Sauer et al. 2017 [↗](#)) (**Figure S5**). We monitored nucleosome assembly coupled to DNA synthesis using as a template a circular UV-damaged plasmid enabling to analyse by supercoiling assay and nucleotide incorporation simultaneously both repair synthesis and nucleosome formation (**Figure 5** [↗](#)) (Moggs et al. 2000 [↗](#)). We verified that p150-depleted HSE lacked the capacity to promote nucleosome assembly on labeled DNA when compared to mock depleted HSE, and that the recombinant WT *SpCAF-1* complex efficiently rescued the loss of xp150 as attested by the detection of supercoiled form I. In contrast, when we complemented the depleted extract with *SpCAF-1* mutant complexes *SpCAF-1*-ED*, *SpCAF-1*-KER*, *SpCAF-1*-ΔWHD we did not detect the supercoiled form I. This indicates that these mutants cannot promote nucleosome assembly (**Figure 5** [↗](#)). When we used the *SpCAF-1*-PIP* mutant, we did not detect supercoiling on labeled DNA at 45 minutes, yet at 2 hours supercoiling ultimately reached levels achieved using the WT *SpCAF-1* (**Figure 5** [↗](#), bottom, synthesized DNA). Interestingly both for 45 and 2 hours of assembly *SpCAF-1*-PIP* mutant yielded more supercoiling than any of the *SpCAF-1*-ED*, *SpCAF-1*-KER*, *SpCAF-1*-ΔWHD mutants. Thus, while mutation in the PIP motif of Pcf1 impaired chromatin assembly at a short time, when more time is given, it allows ultimately to catch up with the wild type. In contrast, none of the *SpCAF-1*-ED*, *SpCAF-1*-KER*, *SpCAF-1*-ΔWHD mutants could catch up, leading to a *SpCAF-1* complex deficient for nucleosome assembly even after longer incubation time. Therefore, these data validate the important role of the amino-acids Y340 and W348 within the ED domain in Pcf1 and the importance to preserve the integrity of the KER and WHD domain to ensure a proper *SpCAF-1* mediated nucleosome assembly on synthesized DNA.

Together, these results indicate that the PIP domain provides Pcf1 with the ability to accelerate nucleosome assembly, yet the integrity of the ED, KER and WD domain proved absolutely mandatory for an efficient *SpCAF-1* mediated nucleosome assembly.

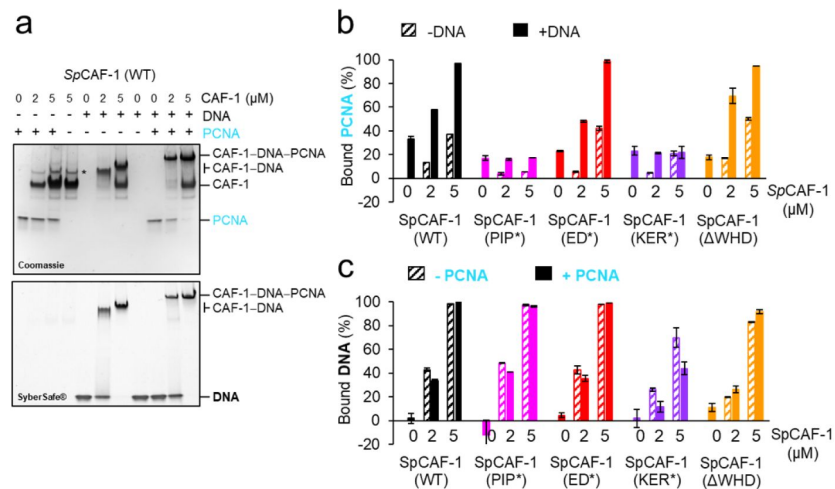


Figure 4

The *SpCAF-1*-KER* mutant is affected for PCNA binding.

a EMSA showing interactions of purified *SpCAF-1* (at the indicated concentrations), with or without recombinant *SpPCNA* (3μM) in the presence and absence of 40bp dsDNA (1μM), revealed with Coomassie blue (upper panel) and with SYBR SAFE staining (lower panel). **b** Quantification of bound *SpPCNA* in the EMSA shown in panel **a** and in **Figure S4d** for *SpCAF-1* and mutants. Values are indicated in % compared to the free PCNA reference (PCNA alone in line 1 in panel **a**) after addition of *SpCAF-1* (WT or mutant) at the indicated concentration and in the presence (filled bars) or absence (dashed bars) of 40bp dsDNA (1μM). **c** Quantification of bound DNA for EMSA shown in panel **a** and in **Figure S4d** for *SpCAF-1* and mutants. Bound DNA in % is compared to the free DNA reference (line 5 in panel **a**) after addition of *SpCAF-1* (WT or mutant) at the indicated concentration and in the presence (filled bars) or absence (dashed bars) of *SpPCNA* (3μM). All experiments were done in duplicates. Mean values are indicated and error bars shows their standard deviation.

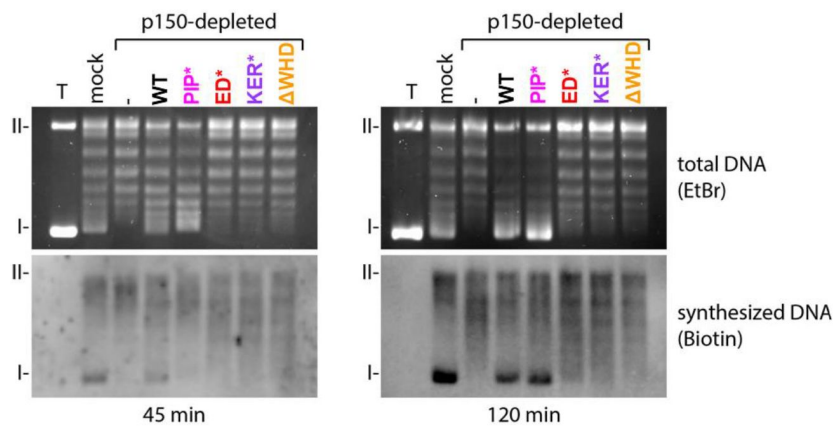


Figure 5

Rescue using Y340 and W348 in the ED domain, the intact KER DNA binding domain and the C-terminal WHD of *Pcf1* in *SpCAF-1* mediated nucleosome assembly.

Supercoiling analysis after 45 (left) and 120 (right) minutes incubation to monitor chromatin assembly in control mock and xenopus p150-depleted HSE. Total DNA visualized by EtBr staining (top) and synthesized DNA visualized by biotin detection (bottom) are shown. The xenopus p150-depleted HSE is either mock complemented (-) or complemented using *SpCAF-1* complex composed of wild type *Pcf1*(WT) or mutants *Pcf1*-PIP*, *Pcf1*-ED*, *Pcf1*-KER*, or *Pcf1*-ΔWHD- as indicated. T: pBS plasmid incubated without extract run in parallel serves as a migration control to locate supercoiled DNA. The position of relaxed (II) and supercoiled (I) DNA are indicated.

Association of *SpCAF-1* with histones impacts PCNA interaction *in vivo*

We next investigated the consequences of the four Pcf1 mutations previously characterized *in vitro*, on *SpCAF-1* function *in vivo* by introducing the respective mutations at the endogenous *pcf1* gene. Both *WT* and mutants were FLAG tagged in their N-terminal part. Immuno-blot of total cell extract with anti-flag antibody showed that all mutated forms of Pcf1 were expressed to the same level than *WT* Pcf1 (**Figure S6a**).

We first tested PCNA–Pcf1 interaction by co-immunoprecipitation of FLAG–Pcf1 and found that Pcf1-ΔWHD showed a similar PCNA interaction than *WT* Pcf1 (**Figure 6a-b** and **Figure S6b**). No interactions were detected with Pcf1-PIP* and Pcf1-KER*, in line with the requirement of the KER and PIP domains for PCNA binding (**Figure 4**, **Table 2**). Surprisingly, we found that Pcf1-ED* binds eight times more to PCNA than the *WT* Pcf1 (**Figure 6a-b**) although the corresponding *SpCAF-1*-ED* bound PCNA with or without DNA *in vitro*, similarly to *WT* (**Figure 4**, **Figure S4d**). This suggest an interplay *in vivo* between the binding of CAF-1 to PCNA and its capacity to bind histones.

We next analyzed the ability of CAF-1 mutated forms to form foci during S-phase (Pietrobon et al. 2014). Since previously reported GFP-tagged forms of Pcf1 are not fully functional, we made use of cells expressing Pcf2-GFP, a functional tagged form (Hardy et al. 2019) (**Figure 6c**). As expected, Pcf2-GFP formed discrete foci during the bulk of S-phase (in bi-nucleated cells with septum) but not during G2 phase (mono-nucleated cells) in a Pcf1-dependent manner (**Figure 6c-d** and **S6c**). The *pcf1-ΔWHD* mutation behaved like the *WT* in this assay whereas, S-phase Pcf2 foci were undetectable when Pcf1–PCNA interaction is impaired (in *pcf1-KER** and *pcf1-PIP** (**Figure 6c** and **6d**). Interestingly, Pcf2-GFP foci were more frequent in all cell cycle phases in *pcf1-ED** mutated cells compared to *WT*. Simultaneous acquisition of GFP fluorescence in living *WT* and mutated *pcf1* cells revealed that Pcf2-GFP foci were more abundant and brighter in *pcf1-ED** cells compared to *WT* (**Figure 6c**), suggesting a higher concentration of CAF-1 within replication factories. In conclusion, CAF-1 foci in S-phase correlate with its association with PCNA *in vivo*, possibly modulated by the histone binding.

The WHD domain specifies CAF-1 function in distinct cellular processes

In *S. pombe*, CAF-1 is involved in the replication-coupled maintenance of heterochromatin (Dohke et al. 2008). We employed a strain in which *ura4*⁺ is inserted at the peri-centromeric heterochromatin of the chromosome I (**Figure 7a**, top panel). The expression of *ura4* is repressed by the surrounding heterochromatin resulting in a poor growth on uracil-depleted media and resistance to 5-fluoro-orotic acid (5FOA) (**Figure 7a**, bottom panel). As previously reported, the deletion of *pcf1* resulted in a better cell growth on uracil-depleted media compared to *WT* cells, showing that the heterochromatin is not properly maintained, leading to the derepression of *ura4*⁺. All mutants, excepted *pcf1-ΔWHD*, exhibited defects in *ura4* silencing, similar to the one observed in the null mutant. This shows that the inability to interact with histone, PCNA and DNA results in a complete lack of CAF-1 function in maintaining heterochromatin. Interestingly, the WHD domain, while required for chromatin assembly *in vitro* (**Figure 5**), is dispensable for the maintenance of heterochromatin. We thus investigated further the role of this domain.

We analyzed the accumulation of Rad52-GFP foci as a readout of global accumulation of DNA damage (**Figure 7b**). The deletion of *pcf1* led to a modest but significant increase in the frequency of cells showing Rad52-GFP foci. A similar effect was observed in *pcf1-ED** mutated cells, while the presence of a CAF-1 complex unable to interact with PCNA resulted in a greater

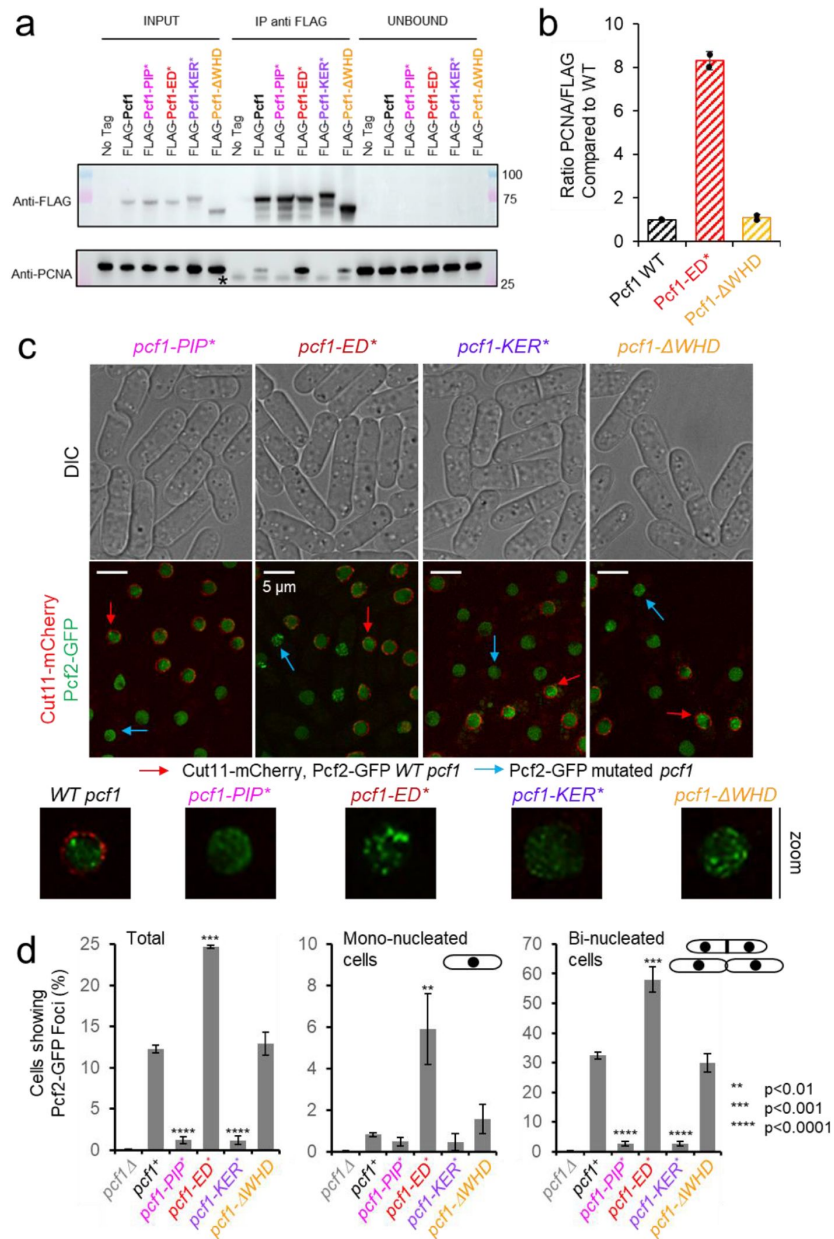


Figure 6

Association of CAF-1 with histone modulates PCNA interaction *in vivo* and foci formation.

a Anti-FLAG Pulldown to address PCNA–CAF-1 interaction *in vivo* in indicated strains. **b** Quantification of bound PCNA from (a). **c** Simultaneous acquisition of Pcf2-GFP in WT (red arrow) and *pcf1* mutated strains (blue arrow). Strains were grown separately and equally mixed before to process them for cell imaging. WT *pcf1* cells expressed the fusion Cut11-mCherry, a component of the nuclear pore complex, leading to a red labelling of the nuclear periphery. Under same illumination and acquisition conditions, Pcf2-GFP foci are detected in WT and *pcf1*-ΔWHD cells, but not in *pcf1*-PIP* and *pcf1*-KER* cells. In contrast, Pcf2-GFP foci are brighter and more abundant in *pcf1*-ED* cells. **d** Quantification of cells showing Pcf2-GFP foci, according to cell morphology in indicated strains. Mono-nucleated cells mark G2-phase cells and bi-nucleated cells with septum mark S-phase cells. Values are means of at least 3 independent experiments ± standard error of the mean (sem). At least 1000 nuclei were analysed per strain. P values are indicated with stars and were calculated using the student test.

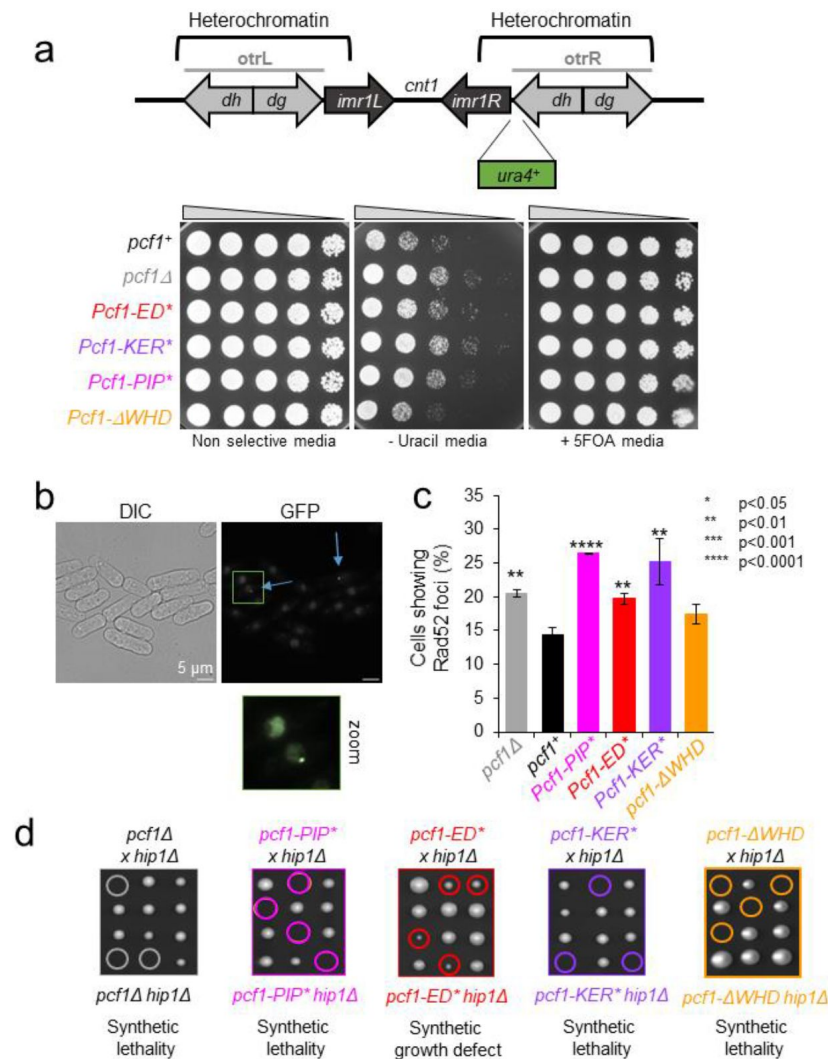


Figure 7

The WHD domain of SpCAF-1 specifies CAF-1 function.

a Top: Schematic representation of the silencing assay used. Otr: outer repeats, imr; inner repeats: *cnt1*; central core of the centromere 1. Bottom: Serial fivefold dilution on indicated strains on indicated media. **b** Example of Rad52-GFP foci in *WT* cells. Blue arrows indicate Rad52 foci-positive cells. **c** Quantification of Rad52-GFP foci in indicated strains. Values are means of at least 3 independent experiments ± sem. P values are indicated as stars and were calculated with the student test. At least 1000 nuclei were analyzed per strain. **d** Co-lethality assay. Tetrad dissections of cells deleted for *hip1* (*hip1Δ*) crossed with cells deleted for *pcf1* (*pcf1Δ*) (grey) or harbouring *pcf1-PIP⁺* (magenta), *pcf1-ED⁺* (red), *pcf1-KER⁺* (purple) or *pcf1-ΔWHD⁺* (orange). Spores with double mutations are surrounded and were deducted from the analysis of viable spores from each tetrad (see material and method). At least 18 tetrads were analysed per cross.

increase (in *pcf1-PIP** and *pcf1-KER** mutants). In contrast, no significant increase was observed in *pcf1-ΔWHD* cells. Thus, both CAF-1 interaction with histone and PCNA prevent the accumulation of DNA damage, but histone deposition is not absolutely required.

The deletion of *pcf1* is synthetic lethal with the deletion of *hip1*, the gene encoding one subunit of the fission yeast HIRA complex (Hardy et al. 2019 [\[1\]](#)), indicating that in the absence of replication-coupled histone deposition by CAF-1, cell viability relies on H3–H4 deposition by HIRA as suggested in human (Ray-Gallet et al. 2011 [\[2\]](#)). We found that *pcf1-KER**, *pcf1-ΔWHD* or *pcf1-PIP** are co-lethal with *hip1* deletion (Figure 7d [\[3\]](#)). Spores harboring *pcf1-ED** were viable when combined with *hip1* deletion, but exhibited a severe growth defect (Figure 7d [\[3\]](#) and Figure S7), suggesting that CAF1-ED* complexes can still perform some histone deposition *in vivo*. These genetic interactions indicate that binding of CAF-1 to PCNA, DNA and histones are critical determinants for its function *in vivo*, as well as the WHD C-terminal domain.

Discussion

In the present work, we provide a comprehensive and dynamic view of key structural features of the histone chaperone CAF-1 from *S. pombe*. Despite the low sequence conservation between orthologues of the large subunit of CAF-1 (Figure S1a), Pcf1 from *SpCAF-1* mediates the heterotrimer complex that binds dimeric histones H3–H4, as does ScCAF-1 (Liu et al. 2012 [\[4\]](#), Mattioli et al. 2017 [\[5\]](#), Sauer et al. 2017 [\[6\]](#)). Using AlphaFold2, we propose a structural model of the *SpCAF-1*, fully supported by our experimental data (Figure 1 [\[7\]](#), Figure S1). This structure defines the 2BD and 3BD regions in Pcf1 as involved in the binding of Pcf2 and Pcf3, respectively. This matches remarkably the corresponding segments identified by HDX in ScCAF-1 (Mattioli et al. 2017 [\[5\]](#)). In line with previous observations in HsCAF-1 (Kaufman et al. 1995 [\[8\]](#)) and ScCAF-1 (Liu et al. 2016 [\[9\]](#), Mattioli et al. 2017 [\[5\]](#), Mattioli et al. 2017 [\[5\]](#)), the ED domain of *SpCAF-1* is crucial for histone binding (Figure 2 [\[10\]](#)). Mainly disordered in the free chaperone, we show that this domain folds upon histone binding, promoting a conformational change with increased accessibility of the KER domain (Figure S1). *SpCAF-1* binds dsDNA longer than 40bp in the micromolar affinity range (Figure 3 [\[11\]](#), Table 1 [\[12\]](#), Figure S3) through the KER domain forming a long monomeric helix with a positively charged face. Interestingly, the helix length roughly corresponds to the size of 40bp dsDNA, suggesting that it could lie on DNA and act as a DNA ruler to sense free DNA for histone deposition (Sauer et al. 2017 [\[6\]](#), Gopinathan Nair et al. 2022 [\[13\]](#), Rosas et al. 2023 [\[14\]](#)). Together, our findings highlight the conservation of CAF-1 properties in histone deposition mechanism *in vitro*, and thus unifies the current model (Sauer et al. 2018 [\[15\]](#)).

This work revealed strong interdependency between histone deposition by CAF-1 and its association with PCNA. The PIP* mutation did not compromise DNA binding of *SpCAF-1 in vitro* (Figure S4). Conversely, upon interaction with DNA, *SpCAF-1* interacted tighter with PCNA (Figure 4 [\[16\]](#)), consistently with a recent study in budding yeast (Rouillon et al. 2023 [\[17\]](#)). We show that *SpCAF-1-PIP** is still able to assemble histones *in vitro*, although slower than WT-*SpCAF-1*. In contrast, *in vivo*, *pcf1-PIP** phenocopy the deletion of *pcf1*. From these results, we conclude that the binding of *SpCAF-1* to PCNA though the PIP motif is required for *SpCAF-1* functions *in vivo*, by allowing its recruitment and efficient histone deposition at DNA synthesis sites.

*SpCAF-1-ED** showed a stronger interaction with PCNA than WT-*SpCAF-1 in vivo*, and formed more abundant and intense foci during S-phase. (Figure 6 [\[18\]](#)). This default may not result from a direct competition between PCNA and histones for CAF-1 association since *SpCAF-1-ED** and WT-*SpCAF-1* show similar interaction with DNA and PCNA *in vitro* (Figure 4 [\[16\]](#), Figure S4). In human cells lacking new histones, PCNA accumulates on newly synthesized DNA, and PCNA unloading has recently linked to histone deposition in budding yeast (Mejlvang et al. 2014 [\[19\]](#), Janke et al. 2018 [\[20\]](#), Thakar et al. 2020 [\[21\]](#)). We propose that the accumulation of CAF-1 at replication foci in the ED* mutant may reflect PCNA recycling defects. This cannot be attributed to the inability of *SpCAF-1-*

ED* to deposit histones otherwise we should have observed the same accumulation for *SpCAF-1-ΔWHD* also defective for histone deposition. The ED* mutation could rather interfere with other interactions, or with Post-Translational Modifications (PTMs) contributing to recycle PCNA.

Deletion of the WHD domain allowed separating *SpCAF-1* functions in chromatin assembly, heterochromatin maintenance and the prevention of DNA damage. Unlike *ScCAF-1* (Liu et al. 2016 [↗](#), Zhang et al. 2016 [↗](#), Mattioli et al. 2017 [↗](#), Mattioli et al. 2017 [↗](#)) and *HsCAF-1*²⁸, *Pcf1_WHD* did not bind DNA nor the ED domain (which remains fully disordered) in the free chaperone. Nevertheless, on the NMR spectra of the free and histone bound *SpCAF-1*(¹⁵N-*Pcf1*), the resonances of the isolated WHD domain are not present (**Figure 1d** [↗](#), **S3n**), in agreement with a restricted movement of this domain that could likely interact with other folded parts of the complex. *In vitro*, we found no impact of the WHD deletion on CAF-1 interaction with DNA, histones or PCNA, but the *SpCAF-1-ΔWHD* was deficient for histone deposition. Thus, the synthetic lethality of this mutant with *hip1* most likely reflects a replication-coupled assembly defect. Unexpectedly, this defect does not cause a problem of heterochromatin maintenance or damage accumulation, showing that the multiple functions of CAF-1 in replication-dependent nucleosome assembly, genome stability and heterochromatin maintenance can be uncoupled thanks to the WHD domain contributing to specify CAF-1 functions. Further investigations will be necessary to understand the role of this domain.

We reveal that disorder is a fundamental feature of *Pcf1* supporting its molecular functions. First, the ED domain is disordered in the FL complex and folds upon histone binding. Second, four IDRs demarcate specific domains within *Pcf1*. We believe that these unfolded regions provide unique ‘plasticity’ properties to *Pcf1* allowing these domains to bind concomitantly their multiple specific partners (*Pcf1*, *Pcf3*, PCNA, DNA and histones). We also reveal that although these domains individually bind their specific partners, there is an important crosstalk between them as exemplified by the fact that DNA stabilizes the CAF-1–PCNA interaction. Such plasticity and cross-talks provided by structurally disordered domains might be key for the multivalent CAF-1 functions.

Materials and Methods

Plasmid preparation for recombinant protein production

The cDNA sequence of WT *Pcf1* (codon optimized for *E.coli* expression) was synthesized and inserted into the pCM153 plasmid to obtain the recombinant MBP–6His–TEV cleavage site–*Pcf1* protein (named MBP-*Pcf1* below). The cDNA sequence of WT *Pcf2* and WT *Pcf3* (codon optimized for insect cells expression) were synthesized and introduced into a pKL plasmid for protein expression in insect cells (MultiBac approach (Bieniossek et al. 2012 [↗](#))) with either a C-terminal (for *Pcf2*) or a N-terminal (for *Pcf3*) 6His tag with a TEV cleavage site between the protein and the His tag. *Pcf1_ED* (325–396) and *Pcf1_WHD* (471–544) were sub cloned in frame into pET28A-B18R plasmid for expression with a N-terminal 6His–SUMO tag. *Pcf1_KER* (56–170) and *Pcf1-KER-PIP* (56–185) were inserted in frame into pCM153 plasmid (Miele et al. 2022 [↗](#)) for expression with a N-terminal 6His–MBP–TEV tag. The cDNA sequence of *S. pombe* histones H3–H4 (codon optimized for *E.coli* expression) were introduced in the 6His–dAsf1 from the pET28 plasmid (generous gift from R.N. Dutnall) in place of histones *DmH3-H4* (Anderson et al. 2010 [↗](#)). With this vector, histones H3–H4 are coexpressed with the chaperone ASF1, leading to soluble untagged free-histones, and ASF1-bound histones. The cDNA of *SpPCNA* (codon optimized for *E.coli* expression) was synthesized and inserted into the pET28A-B18R plasmid for expression with an N-terminal 6His–SUMO. *Pcf1* mutants were generated by PCR. All plasmids for recombinant protein expression were constructed by GenScript.

Recombinant protein production

Pcf1 was overexpressed in *E. coli*. After fresh transformation of *E. coli* BL21 (DE3) Star cells (Thermo Fisher Scientific), cells were grown in an auto-induction rich medium Terrific Broth (12 g/L tryptone, 24 g/L yeast extract) containing 50 µg/mL of Kanamycin for 30 hours at 20°C, under agitation. SpHistones H3-H4, SpPCNA and the all domains of Pcf1 were overexpressed in *E. coli*. The plasmid for expressing the desired protein was freshly transformed in *E. coli* strain BL21 DE3 STAR (Thermo Fisher Scientific). Cells were grown at 37 °C in LB medium containing 50 µg/mL of Kanamycin until OD reached 0.7 and recombinant protein expression was induced for 16 hours at 20 °C under agitation by adding 1 mM isopropyl β-D-1-thiogalactopyranoside IPTG, or cells were grown 30 hours at 20 °C in a ZY auto-inducible medium. For ¹⁵N or ¹³C uniformly labeled proteins, the expression was made in minimal media with 0.5g/L of ¹⁵NH₄Cl and/or 2g/L of ¹³C-glucose. Pcf2 and Pcf3 were produced in insect cells. Sf9 Insect cells were infected with an MOI of 5*10⁻³ virus/cell and incubated for 5 days at 27 °C at 130 rpm. After centrifugation, cell pellets stored at -70°C until further use.

Protein purifications

Purification of Pcf1

Cells were pelleted by centrifugation and resuspended in the lysis buffer LB1 for 30 minutes (50 mM Tris-HCl pH 8, 500 mM NaCl, 5% glycerol, 0.1% Triton X-100, 2 mM DTT, 5 mM MgCl₂, 0.5 mM PMSF, 1X cOmplete™ EDTA-free Protease Inhibitor Cocktail, 1.2 mg/mL lysozyme and 70 U/mL of benzonase). Cells were lysed by sonication at 4°C, the lysate was clarified by centrifugation at 5°C at 18 500 rpm for 30min and loaded onto gravity flow amylose resin (NEB) previously equilibrated with buffer WB1_1 (50 mM Tris-HCl pH 8, 500 mM NaCl, 2 mM DTT). After loading the cell lysate onto the resin, the resin was washed with 5 column volumes of buffer WB1_1 to ensure complete passage of the cell lysate through the resin. Then, the resin was further washed with 10 column volumes of buffer WB1_2 (50 mM Tris-HCl pH 8, 1000 mM NaCl, 2 mM DTT) to remove non-specific binding, before re-equilibration with 10 column volumes of buffer WB1_1. MBP-Pcf1 was eluted with 10 column volumes of buffer EB1 (50 mM Tris-HCl pH 8, 500 mM NaCl, 0.5 mM TCEP, 10 mM maltose and 1X cOmplete™ EDTA-free Protease Inhibitor Cocktail). After addition of 1 mM MgCl, the eluate containing MBP-Pcf1 was incubated 16 hours at 5°C with TEV protease (added with a ratio 1/20 in mass). The elate was then concentrated to 300µL (with Amicon® Ultra-15 30kDa filter concentrators), 2000 U of benzonase were added and incubated for 2 hours. The concentrated eluate was injected into a column Superose™ 6 increase 10/300 GL (Cytiva) previously equilibrated with the final buffer FB1 (50 mM Tris-HCl pH 8, 500 mM NaCl, 1 mM DTT). The Pcf1-containing fractions were pooled. 1X cOmplete™ EDTA-free Protease Inhibitor Cocktail, 0.5 mM TCEP and 30% glycerol was added and samples were snap-frozen and stored at -70°C.

Purification of Pcf2

Cells pellets were resuspended into lysis buffer LB2 (50 mM Tris-HCl pH 8, 500 mM NaCl, 5% glycerol, 0.1% Triton X-100, 10 mM imidazole, 0.5 mM PMSF, cOmplete™ EDTA-free Protease Inhibitor Cocktail and 70 U/mL of benzonase) and sonicated at 4°C. Lysates were clarified by centrifugation at 5 °C at 18 500 rpm for 30 min and loaded to gravity flow Ni-NTA agarose resin (QIAGEN) previously equilibrated with wash buffer WB2_1 (50 mM Tris-HCl pH 8, 500 mM NaCl, 10 mM imidazole). Resin was then washed with 10 column volumes of wash buffer WB2_1 followed by 10 column volumes of wash buffer WB2_2 (50 mM Tris-HCl pH 8, 1M NaCl, 10 mM imidazole). Pcf2-6His was eluted with 5 column volumes of the elution buffer EB2 (50 mM Tris-HCl pH 8, 500 mM NaCl, 250 mM imidazole, 0.5 mM TCEP, and 1X cOmplete™ EDTA-free Protease Inhibitor Cocktail). The eluate was then concentrated to 300 µL (with Amicon® Ultra-15 30 kDa filter concentrators). 1 mM DTT, 1 mM MgCl and ~2000 U of benzonase were added to the concentrated eluate and the sample was incubated 2 hours at 4 °C and injected into a Superdex 200 increase 10/300 (Cytiva) previously equilibrated with the final buffer FB2 (50 mM Tris-HCl pH 8, 500 mM

NaCl and 1 mM DTT). Pcf2-6His containing fractions were pooled and directly used for CAF-1 reconstitution or stored at -70 °C with cOmplete™ EDTA-free Protease Inhibitor Cocktail, 0.5 mM TCEP and 30% glycerol.

Purification of Pcf3

Cells pellets were resuspended into lysis buffer LB3 (50 mM Tris-HCl pH 8, 200 mM NaCl, 5% glycerol, 0.1% Triton X-100, 10 mM imidazole, 0.5 mM PMSF, cOmplete™ EDTA-free Protease Inhibitor Cocktail and 70 U/mL of benzonase) and sonicated at 4°C. Lysate was clarified by centrifugation at 5 °C at 18500 rpm for 30 min and loaded to gravity flow Ni-NTA agarose resin (QIAGEN) previously equilibrated with wash buffer WB3_1 (50 mM Tris-HCl pH 8, 200 mM NaCl, 10 mM imidazole). Resin was then washed with 5 column volumes of wash buffer WB3_1 and 10 column volumes of wash buffer WB3_2 (50 mM Tris-HCl pH 8, 200 mM NaCl, 30 mM imidazole). his-Pcf3 was then eluted with EB3 (50 mM Tris-HCl pH 8, 200 mM NaCl, 250 mM imidazole, 1 mM DTT, 1X cOmplete™ EDTA-free Protease Inhibitor Cocktail). After addition of 1 mM MgCl₂, 6His-TEV protease (with a ratio 1/10 in mass), ~2000 U of benzonase the eluate was dialysed o/n at 5 °C in the final buffer FB3 (50 mM Tris-HCl pH 8, 200 mM NaCl and 1 mM DTT). Because of their similar size, Pcf3 and 6His-TEV protease cannot be completely separated by size-exclusion chromatography. Therefore, to remove the 6His-TEV protease and uncleaved His-Pcf3, 30 mM of imidazole was added to the dialysate, which was then loaded to gravity flow Ni-NTA agarose resin (QIAGEN) previously equilibrated with wash buffer WB3_2. The Flow through was concentrated to 300μL (with Amicon® Ultra-15 30 kDa filter concentrators) and injected into a Superdex 200 increase 10/300 (Cytiva) previously equilibrated with the final buffer FB3. Pcf3-containing fractions were pooled and stored at - 70 °C after adding 1X cOmplete™ EDTA-free Protease Inhibitor Cocktail, 0.5 mM TCEP and 30% glycerol.

Reconstitution of CAF-1 complexes

CAF-1 complexes were formed by mixing the isolated proteins Pcf1 (WT or mutant), Pcf2-6His and Pcf3 previously purified as described above. Isolated Pcf2-6His and Pcf3 were added in small excess compared to Pcf1. Tris 50 mM pH 8 was added to the Pcf1/Pcf2-his/Pcf3 mix to reach a final NaCl concentration of 150mM. After addition of 1 mM MgCl₂, 1X cOmplete™ EDTA-free Protease Inhibitor Cocktail, 6His-TEV protease (with a ratio 1/10 in mass) and ~2000 U of benzonase, the mixture was incubated over night at 4 °C and applied on a HiTrap® heparin FF column (Cytiva) previously equilibrated with EB4_1 (50 mM Tris-HCl pH 8, 100 mM NaCl). A gradient was applied with the high salt buffer EB4_2 (50 mM Tris-HCl pH 8, 1M NaCl). Fractions containing the full SpCAF-1 were pooled, concentrated to 300μL (with Amicon® Ultra-15 30 kDa filter concentrators) and injected into a Superdex 200 increase 10/300 (Cytiva) previously equilibrated with the final buffer FB4_1 (50 mM Tris-HCl pH 8, 150 mM NaCl and 1 mM DTT). The SpCAF-1-containing fractions were pooled and directly used for MST or EMSA analysis or stored at -70°C with 1X cOmplete™ EDTA-free Protease Inhibitor Cocktail, 0,5 mM TCEP and 30% glycerol.

The SpCAF-1(¹⁵N-¹³C-Pcf1) and SpCAF-1(¹⁵N-Pcf1) was reconstituted by co-lysing the pellets of ¹⁵N-¹³C-MBP-Pcf1 or ¹⁵N-MBP-Pcf1 (WT or mutants), Pcf2-6His and his-Pcf3. Cell pellets from Pcf2-6His and Pcf3-his were added in excess compared to labeled MBP-Pcf1, based on the yield previously obtained for the isolated proteins. The pellets were resuspended and mixed in the lysis buffer LB4 (50 mM Tris-HCl pH 8, 150 mM NaCl, 5% glycerol, 0.1% Triton X-100, 10 mM imidazole, 0.5 mM PMSF, cOmplete™ EDTA-free Protease Inhibitor Cocktail and 70 U/mL of benzonase), sonicated and centrifuged as described before. The clarified lysate was applied to gravity flow Ni-NTA agarose resin (QIAGEN) previously equilibrated with wash buffer WB4_1 (50 mM Tris-HCl pH 8, 150 mM NaCl, 10 mM imidazole). Beads were washed with 5 column volume of WB4_1 buffer, followed by 10 column volumes of WB4_2 (50 mM Tris-HCl pH 8, 1 M NaCl, 10 mM imidazole). Elution was performed with EB4 (50 mM Tris-HCl pH 8, 150 mM NaCl, 250 mM imidazole, 1X cOmplete™ EDTA-free Protease Inhibitor Cocktail) and applied to an anion exchange column HiTrap® Q FF (Cytiva) previously equilibrated with buffer EB4_1. A gradient was applied with the high salt buffer EB4_2.

The tagged CAF-1-containing fractions were pooled, and dialyzed overnight against buffer 4 DB4 (Tris 50 mM pH 8, 150 mM NaCl, 1 mM DTT) after addition of 1 mM DTT, 1 mM MgCl₂, 1X cOmplete™ EDTA-free Protease Inhibitor Cocktail, 6His-TEV protease (with a ratio 1/10 in mass) and ≈2000 U of benzonase. The mixture was applied on a HiTrap® heparin FF column (Cytiva) using the same buffers (EB4_1 and EB4_2). The SpCAF-1(¹⁵N-¹³C/¹⁵N-Pcf1)-containing fractions were concentrated to 300 μL (with Amicon® Ultra-15 30 kDa filter concentrators) and injected into a Superdex 200 increase 10/300 (Cytiva) previously equilibrated with buffer FB4_2 (10 mM Tris-HCl, 50 mM HEPES pH 7, 300 mM NaCl and 0.5 mM TCEP). The SpCAF-1(¹⁵N-¹³C-Pcf1)-containing fractions were pooled and immediately used for NMR measurements.

Purification of histones SpH3–SpH4

Cells expressing SpH3, SpH4 with 6His-dAsf1 were pelleted by centrifugation and resuspended in the lysis buffer LB5 (50 mM Tris-HCl pH 8, 500 mM NaCl, 5% glycerol, 1% Triton X-100, 1 mM DTT, 10 mM MgCl₂, 0.5 mM PMSF, 1X cOmplete™ EDTA-free Protease Inhibitor Cocktail) and flash frozen in liquid nitrogen. After thawing, lysozyme and benzonase were added at a final concentration of 0.25 mg/mL and 70 U/mL respectively. After incubation 20 minutes at 4°C, cells were lysed by sonication. Soluble 6His-Asf1 was removed on a NiNTA column (Qiagen) equilibrated in the LB5 buffer. The flow through (containing soluble free histones) was filtered with 0.22 μ filters and loaded on a cation exchange Resource S column (GE Healthcare) equilibrated with the dilution buffer EB5_1 (50 mM Tris-HCl pH8). Histones H3–H4 were eluted with a NaCl gradient in a buffer EB5_2 (50 mM Tris-HCl pH8, 2 M NaCl). The H3–H4-containing fractions were pooled, the salt concentration adjusted to 2 M NaCl, and concentrated in a 3 kDa concentrator (Millipore), flash freezed in liquid nitrogen and stored at -70°C.

Purification of Pcf1_ED and Pcf1_ED*

Cells expressing Pcf1_ED or Pcf1_ED* with a N-terminal 6His-SUMO tag were collected by centrifugation, resuspended in lysis buffer LB6 (50 mM Tris-HCl pH8, 500 mM NaCl, 5% glycerol, 1% Triton X-100, 1 mM PMSF, 1 μM aprotinin, 0.25 mM DTT) and flash frozen in liquid nitrogen. After thawing, lysosome was added at a final concentration of 1 mg/mL and cells were incubated 30 min at 4 °C and lysed by sonication. 6His-SUMO-Pcf1_ED was first purified on Histrap columns (Cytiva). Fractions containing the protein were pulled. SUMO protease was added at a final concentration 1/10 and the mixture was dialyzed overnight at 4 °C against the buffer DB6 (50 mM Tris-HCl pH 8, 150 mM NaCl, 10 mM imidazole) and applied on a NiNTA column (Qiagen) equilibrated in the DB6 buffer. The flow-through fraction containing Pcf1_ED or Pcf1_ED* was then purified by size exclusion chromatography using a Superdex 75 increase 10/300 column (Cytiva) previously equilibrated with the final buffer in FB6 (10 mM Tris-HCl, 50 mM HEPES pH 7, 300 mM NaCl). Fraction containing Pcf1_ED or Pcf1_ED* were concentrated using Amicon centrifuge filter units of 3 kDa cutoff (Millipore) flash freezed in liquid nitrogen and stored at -20 °C or -70 °C.

Purification of Pcf1_KER and Pcf1_KER-PIP

Cells expressing Pcf1_KER(56-170), or Pcf1-KER-PIP(56-185) (WT or mutant) with a N-terminal 6His-MBP-TEV tag were collected by centrifugation, resuspended in lysis buffer LB7 (50 mM Tris-HCl pH 8, 500 mM NaCl, 5% glycerol, 1% Triton X-100, 1 mM PMSF, 1 μM aprotinin, 0.25 mM DTT, 1X cOmplete™ EDTA-free Protease Inhibitor Cocktail) and flash frozen in liquid nitrogen. After thawing, 5 mM MgCl₂, 1 mg/mL lysozyme and 70 U/mL of benzonase were added and cells were lysed by sonication. Proteins were first purified on Histrap columns (Cytiva) including a wash step with WB7 (50 mM Tris-HCl pH 8, 1000 mM NaCl). 1 mM DTT and TEV protease (1/10 ratio) was added to the fractions containing the 6His-MBP-TEV_Pcf1_KER fragment and the mixture was incubated 2 hours at room temperature and injected on a resource S column (Cytiva) previously equilibrated with EB7_1 (50 mM Tris-HCl pH 8). A gradient was applied with the high salt buffer

EB7_2 (50 mM Tris-HCl pH 8, 2M NaCl). Fractions containing Pcf1_KER fragment were pooled and diluted to reach a concentration of 150 mM NaCl and concentrated (with Amicon® Ultra-10 kDa filter concentrators).

Purification of Pcf1_WHD

Cells expressing Pcf1_WHD with a N-terminal 6His-SUMO tag were resuspended in lysis buffer LB8 (50 mM Tris-HCl pH8, 500 mM NaCl, 5% glycerol, 1% Triton X-100, 1 mM PMSF, 1 μ M aprotinin, 0.25 mM DTT, 1X cOmplete™ EDTA-free Protease Inhibitor Cocktail) and flash frozen in liquid nitrogen. After thawing, 5 mM MgCl₂, 1 mg/mL lysozyme and 70 U/mL of benzonase was added and cells were further lysed by sonication. The lysate was loaded onto gravity flow amylose resin (NEB) previously equilibrated with buffer WB8_1 (50 mM Tris-HCl pH 8). Resin was then washed with 10 column volume of buffer WB8_1, 10 column volumes of buffer WB8_2 (50 mM Tris-HCl pH 8, 1000 mM NaCl), 10 column volumes of buffer WB8_1. 6His-SUMO-Pcf1_WHD was eluted with 10 column volume of buffer EB8 (50 mM Tris-HCl pH 8, 500 mM NaCl, 250 mM Imidazole). SUMO protease was added at a final concentration 1/10 and the mixture was incubated overnight at 4°C. The mixture was concentrated (with Amicon® Ultra-3 kDa filter concentrators) and applied on a Superdex 75 increase 10/300 size exclusion column (Cytiva) previously equilibrated with the final buffer FB8 (10 mM Tris-HCl, 50 mM HEPES pH7, 150 mM, NaCl). Finally, proteins were concentrated in a 3 kDa concentrator (Millipore).

Purification of SpPCNA

Cells expressing SpPCNA with a N-terminal 6His-SUMO tag were resuspended in lysis buffer LB9 (50 mM Tris-HCl pH8, 500 mM NaCl, 5% glycerol, 1% Triton X-100, 1 mM PMSF, 1 μ M aprotinin, 0.25 mM DTT, 1X cOmplete™ EDTA-free Protease Inhibitor Cocktail) and flash frozen in liquid nitrogen. After thawing, 1 mM MgCl₂, 1 mg/mL lysozyme and 70 U/mL of benzonase was added and cells were further lysed by sonication. The lysate was loaded onto gravity flow amylose resin (NEB) previously equilibrated with buffer WB9_1 (50 mM Tris-HCl pH 8). Resin was then washed with 10 column volume of buffer WB9_1, 10 column volumes of buffer WB8_2 (50 mM Tris-HCl pH 8, 2000 mM NaCl), 10 column volumes of buffer WB8_1. 6His-SUMO-SpPCNA was eluted with 3 column volumes of buffer EB9 (50 mM Tris-HCl pH 8, 250 mM Imidazole). 1mM DTT and SUMO protease was added at a final concentration 1/10 and the mixture was dialyzed overnight at 4°C against the buffer DB9 (50 mM Tris-HCl pH 8, 150 mM NaCl, 10 mM imidazole). The mixture was applied on a Histrap column (Cytiva), the flow through (containing SpPCNA) was concentrated (with Amicon® Ultra-15 3 kDa filter concentrators) and applied on a hiLoad 16/600 superdex 200 size exclusion column previously equilibrated with a FB9 (50 mM Tris-HCl pH8, 150 mM NaCl). In case the digestion of the tag was incomplete, the two last steps digestion with SUMO protease and gel filtration were repeated.

For all protein samples, depending on specific requirements of different techniques used, aliquots of concentrated protein were either maintained at 4°C or flash frozen in liquid nitrogen after addition or not of 30% glycerol and stored at -70°C for further use.

DNAs used to monitor protein-DNA interactions

The different DNAs were purchased from eurofins genomics. The sequences were derived from the 601 positioning sequence: ATCAATATCCACCTGCAGATACTACCAAAAGTGTATTTGG. For MST, the DNA were labeled with ALEXA488 at their 5' extremity. The ssDNA was annealed with the reverse-complementary sequence by heating at 90 °C and cooling slowly at room temperature.

Size-exclusion chromatography (SEC)

SpCAF-1 subunits interaction was performed by mixing 2.2 nmoles of each isolated protein together in a final volume of 1.26 mL and left o/n at 5°C. The complexes were then concentrated to 300µL (with Amicon® Ultra-15 30 kDa filter concentrators) and injected into a Superdex 200 increase 10/300 (Cytiva) for separation by size-exclusion chromatography previously equilibrated with the FB4_3 (50 mM Tris-HCl pH 7.5, 500 mM NaCl, 1 mM DTT). The different fractions were analyzed on mPAGE® 8% Bis-Tris Precast Gels (Sigma) with MOPS SDS running buffer. Interaction between CAF-1 and H3-H4 was carried out by incubating for 3 hours, 3 nmoles of *SpCAF-1* with 3 nmoles of *SpH3-H4* in a final buffer FB4_4 (50 mM Tris-HCl pH 7.5, 150 mM NaCl, 4 mM DTT, 1X cOmplete™ EDTA-free Protease Inhibitor Cocktail) or FB4_5 (50 mM Tris-HCl pH 7.5, 1 M NaCl, 4 mM DTT, 1X cOmplete™ EDTA-free Protease Inhibitor Cocktail). Samples were then concentrated to 300µL (with Amicon® Ultra-15 30 kDa filter concentrators) and injected into a Superdex 200 increase 10/300 (Cytiva) for separation by size-exclusion chromatography with their corresponding buffers. The different fractions were analyzed on mPAGE® 4-20% Bis-Tris Precast Gels (Sigma) with MES SDS running buffer.

Electrophoretic Mobility Shift Assay (EMSA)

The proteins and DNA were mixed to be in a final EMSA buffer EMB (25 mM Tris-HCl pH 8, 1 mM EDTA pH 8.0, 150 mM NaCl) and incubated at 4 °C for 30 min and heated at 37 °C for 5 min prior to analysis on precast “any KD” Mini-PROTEAN TGX (Bio-rad, Cat #4569033) polyacrylamide gels using 1x TBE as running buffer. The gels were stained with 1x of SYBR Safe (Thermo Fisher Scientific, Waltham, MA) then visualized with BIORAD EZ Imager. A second identical the gel was Coomassie Blue before being visualized with the BIORAD EZ Imager. Band intensities were quantified by ImageJ.

MicroScale Thermophoresis (MST)

DNAs labeled with ALEXA488 at their 5' extremity. The final dilution buffer was FB_MST (10 mM Tris-HCl, 40 mM HEPES pH 7, 150 mM NaCl). The labeled DNA was adjusted to 20 nM. Freshly prepared proteins or complexes were diluted in the same buffer with 16 serial dilution 1:2. Each protein dilution was mixed with one volume of labeled DNA and filled into Monolith NT standard treated capillaries (NanoTemper Technologies GmbH). Thermophoresis was measured using a Monolith NT.115 instrument (NanoTemper Technologies GmbH) at an ambient temperature of 20 °C with 3-s/20-s/1-s laser off/on/off times, respectively. Instrument parameters were adjusted with 80% LED power and 40% MST power. Data of two measurements were analyzed (MO.Affinity Analysis software, NanoTemper Technologies) using the signal from thermophoresis at 5s.

Circular Dichroism (CD)

Circular Dichroism (CD) measurements were carried out at 20 °C on a JASCO J-810 spectro-polarimeter. Temperature was controlled by a Peltier. Spectra from 190 to 250 nm were obtained using a 2 mm optical path length quartz cell (Hellma #100-2-40) containing Pcf1_KER or Pcf1_KER* (5µM) in 10 mM of phosphate buffer (pH 7.4).

Nuclear Magnetic Resonance (NMR)

NMR experiments were carried out on Bruker DRX-600 MHz, 700 MHz or 950MHz spectrometers equipped with cryo-probes. All NMR data were processed using Topspin (Bruker) and analyzed using Sparky (T.D. Goddard and D.G. Kneller, UCSF). Samples were prepared in 3 mm NMR tubes, in solution containing 5% D2O, 0.1% NaN3, 0.1 mM DSS with different buffer appropriate for different complex formations or reactions. Heteronuclear Multiple Quantum Correlation (sofast-HMQC) or best-HSQC spectra were all recorded at 283°K. The protein concentrations were between 9 µM and 500 µM. For backbone resonances assignments, 3D data were collected at 283°K using standard Heteronuclear Single Quantum Correlation (HSQC) spectra ¹H-¹⁵N HSQC, TOCSY-HSQC,

HNCA, HBHA(CO)NH, CBCA(CO)NH, HN(CA)CO, HNCO, HN(CO)CA, CBCANH and HN(CA)CO experiments. Proton chemical shifts (in ppm) were referenced relative to internal DSS and ^{15}N and ^{13}C references were set indirectly relative to DSS using frequency ratios (Wishart et al. 1995 [\[1\]](#)). Chemical shift index were calculated according to the sequence-specific random coil chemical shifts (Wishart et al. 1995 [\[1\]](#), Tamiola et al. 2010 [\[2\]](#)).

Structural models of the SpCAF-1 WHD domain were computed from NMR data with CS-ROSETTA (Shen et al. 2008 [\[3\]](#)) version 1.01. First, the MFR program from NMRpipe (Delaglio et al. 1995 [\[4\]](#)) was used to search a structural database for best matched fragments based on the protein backbone ^{15}N , ^{13}C , ^{13}CA , ^{13}CB and ^1HN chemical shifts. Then the ROSETTA 3.8 software was used to generate 27753 models by fragment assembly and full-atom relaxation. These models were rescored by comparing the experimental chemical shifts with the chemical shifts predicted by SPARTA (Shen and Bax 2007 [\[5\]](#)) for each model. The best model after rescoring was chosen as a representative NMR model of the WHD domain.

Small Angle Xray Scattering (SAXS)

SAXS data were collected at the SWING beamline on a EigerX 4 M detector using the standard beamline setup in SEC mode (Thureau et al. 2021 [\[6\]](#)). Samples were injected into a Superdex 5/150 GL (Cytivia) column coupled to a high-performance liquid chromatography system, in front of the SAXS data collection capillary. The initial data processing steps including masking and azimuthal averaging were performed using the program FOXTROT (Girardot et al. 2015 [\[7\]](#)) and completed using US-SOMO (Brookes et al. 2016 [\[8\]](#)). The final buffer subtracted and averaged SAXS profiles were analyzed using ATSAS v.3. software package (Manalastas-Cantos et al. 2021 [\[9\]](#)). To model the structures and improve the AlphaFold2 models, the program Dadimodo (Rudenko et al. 2019 [\[10\]](#)) (<https://dadimodo.synchrotron-soleil.fr>) that refines multidomain protein structures against experimental SAXS data was used (see **Table S1** in supplementary data for more information).

Structural Modeling

Sequences of *S. pombe* Pcf1 (Q1MTN9), Pcf2 (O13985), Pcf3 (Q9Y825), H3 (P09988), H4 (P09322) and PCNA (Q03392) were retrieved from UniProt database (UniProt 2021 [\[11\]](#)). These sequences were used as input of mmseqs2 homology search program (Steinegger and Soding 2017 [\[12\]](#)) used with three iterations to generate a multiple sequence alignment (MSA) against the uniref30_2103 database (Mirdita et al. 2022 [\[13\]](#)). The resulting alignments were filtered using hhfilter (Steinegger et al. 2019 [\[14\]](#)) using parameters ('id'=100, 'qid'=25, 'cov'=50) and the taxonomy assigned to every sequence keeping only one sequence per species. To increase the number of sequences in the alignment of *S. pombe* Pcf1 we independently generated MSA using mmseqs2 starting from the *S. cerevisiae* or the human homolog of Pcf1 (Q12495 and Q13111, respectively) and the resulting alignments were combined with the one of SpPcf1. Full-length sequences in the alignments were then retrieved and the sequences were realigned using MAFFT (Katoh and Standley 2013 [\[15\]](#)) with the default FFT-NS-2 protocol. To build the so-called mixed co-alignments, sequences in the alignment of individual partners were paired according to their assigned species and left unpaired in case no common species were found (Mirdita et al. 2022 [\[13\]](#)). A first global model with full-length Pcf1, Pcf2 and Pcf3 was generated to map the regions of Pcf1 binding to Pcf2 and Pcf3 and to obtain the pLDDT scores shown in **Figure 1b** [\[16\]](#) for Pcf1, **Figure S1g** for Pcf2 and Pcf3.

Next, three models of the complex corresponding to independent modules of the complex were generated using different delimitations: model_1 (presented in **Figure S1h-j**) with Pcf1(403-450)-Pcf2(1-453) (MSA with 2180 species, 501 positions), model_2 (presented in **Figure S1k-m**) with Pcf1(200-335)-Pcf3(1-408) (MSA with 2148 species, 544 positions), model_3 Pcf1(352-383)-H3(60-136)-H4(25-103) (presented in **Figure 2f** [\[17\]](#) and **S2d**) (MSA with 3530 species, 188 positions). Concatenated mixed MSAs were generated using the delimitations defined above and used as input to run 5 independent runs of the AlphaFold2 algorithm with 6 iterations each (Jumper et al. 2021 [\[18\]](#)) generating 5 structural models using a local version of the ColabFold interface (Mirdita et al. 2021 [\[19\]](#)).

al. 2022 [\[1\]](#)) trained on the multimer dataset (Evans et al. 2022 [\[2\]](#)) on a local HPC equipped with NVIDIA Ampere A100 80Go GPU cards. The best models of each of the 5 runs converged toward similar conformations. They reached high confidence and quality scores with pLDDTs in the range [83.7, 84.3], [88.8, 89.8] and [86.5, 88.4] and the model confidence score (weighted combination of pTM- and ipTM-scores with a 20:80 ratio) (Evans et al. 2022 [\[2\]](#)) in the range [0.9, 0.93], [0.88, 0.89], [0.85, 0.87], for model_1, model_2 and model_3, respectively. The models with highest confidence score for each of the three models were relaxed using rosetta relax protocols to remove steric clashes (Leman et al. 2020 [\[3\]](#)) with constraints (std dev. of 2 Å for the interatomic distances) and were used for structural analysis. MSA web logos were generated with the weblogo server (<https://weblogo.berkeley.edu/logo.cgi> [\[4\]](#)).

Nucleosome assembly assay

Mock- and p150CAF-1-depleted *Xenopus* high speed egg extract (HSE) were prepared as previously (Ray-Gallet and Almouzni 2004 [\[5\]](#)). Nucleosome assembly was performed on pBS plasmid damaged by UV (500J/m²) to promote DNA synthesis as previously described (Ray-Gallet and Almouzni 2004 [\[5\]](#)) except that the reaction mixed contained 3.2 μM of biotin-14-dCTP (Invitrogen, Ref 19518-0189) instead of [α^{32} P]-dCTP. The p150CAF-1-depleted extracts were complemented with 50 ng of isolated/reconstituted spCAF-1 complex composed of WT or mutated Pcf1. After DNA purification, samples were processed for gel electrophoresis (1% agarose) to resolve topoisomers as previously described (Ray-Gallet and Almouzni 2004 [\[5\]](#)). After staining with Ethidium bromide to visualize total DNA and gel transfer on a Nylon N⁺ membrane (GE Healthcare Ref RPN203B) (Qiagen) for 45 min at 40 mbar in 10x SSC, the membrane was rinsed in PBS, air dried and DNA was crosslinked to the membrane using Stratalinker (Biorad). DNA synthesis was visualized by detecting biotin with the Phototope-Star detection kit (New England Biolabs Ref N7020S) and images acquired on a Chemidoc system (Biorad).

Standard yeast genetics

Yeast strains were freshly thawed from frozen stocks and grown at 30 °C using standard yeast genetics practices. All *pcf1* mutants were obtained using a two-step transformation approach in which the region of interest of the coding sequence was first replaced by the *ura4⁺* marker and then by synthetic DNA containing the appropriated mutations. Therefore, all mutations are marker-free and were followed in genetic analysis by PCR and sequencing. Yeast strains used in this study are listed in Supplementary **Table S2**. For tetrad analysis, the genotypes of viable spores were deduced by checking for resistance to Kanamycin (deletion) and sequencing, which made it possible to deduce the genotype of dead spores for each tetrad. At least 18 tetrads were analyzed for each cross.

Peri-centromeric silencing assay

5-FOA (EUROMEDEX, 1555) resistant colonies were grown on uracil-containing liquid media overnight and 10 μL of 5 fold serial dilutions (from 1.10⁷ cells/mL to 1.10⁵ cells/mL) were spotted on indicated media.

Co-immunoprecipitation

5.10⁸ cells from exponentially growing cultures were harvested with 10% NaN₃ and 1 mM PMSF, final concentration, and then washed twice in water and once in Lysis buffer (buffer (50 mM HEPES High salt, 50 mM KoAc pH7.5, 5 mM EGTA, 1% triton X100, 0.01mg/mL AEBSF, EDTA-free protease inhibitor cocktail). Cell pellets were resuspended in 800 μL of lysis buffer and were broken with a Precellys homogenizer (twice 4 cycles at 10 000 rpm, 20 sec-2 min pause). After lysate clarification (30 minutes at 13 000 rpm, 4°C), 2.5 mg of proteins were incubated with pre-washed Dynabeads protein G (Invitrogen, 10003D) coupled to anti-FLAG antibody (Sigma F7425) and incubated overnight at 4 °C on a wheel. Beads were washed three times for 5 minutes at 4 °C with 800 μL of lysis buffer, and then resuspended in 1X Laemmli buffer, and boiled at 95 °C for 10

minutes. INPUT and UNBOUND (both 10 % of initial protein extract) and BOUND (IP) fraction were resolved by electrophoresis on acrylamide gels (4-12% Invitrogen) and the transferred onto nitrocellulose membrane that were saturated for 1 hour, RT in TBS-0.075% tween-5% milk. Proteins of interest were detected with anti-FLAG antibody (Sigma F1805, 1:1000) and anti-PCNA antibody (Santa Cruz sc-8349, 1:500).

Live cell imaging

All image acquisition was performed on the PICT-IBiSA Orsay Imaging facility of Institut Curie. Cells were grown in filtered supplemented EMM-glutamate.. Exponentially growing cultures were centrifuged and resuspended in 50 μ L of fresh medium. 2 μ L from this concentrated solution was dropped onto a Thermo Scientific slide (ER-201B-CE24) covered with a thin layer of 1.4 % agarose in filtered EMMg. 13 z-stack pictures (each z step of 300 nm) were captured using a Spinning Disk Nikon inverted microscope equipped with the Perfect Focus System, Yokogawa CSUX1 confocal unit, Photometrics Evolve512 EM-CCD camera, 100X/1.45-NA PlanApo oil immersion objective and a laser bench (Errol) with 491 (GFP) and 561 (MmCherry) nm diode lasers, 100 mX (Cobolt). Pictures were collected with METAMORPH software and analyzed with ImageJ. For Pcf2-GFP and Rad52-GFP foci, a threshold (using the find maxima tool, > 400 for Pcf2-GFP foci and > 100 for Rad52-GFP) was setup at the same level for each genetic background analyzed within the same experiment. Images from **figure 6** [C](#) were deconvolved using the Huygens remote manager software.

Data availability

We deposited structural models generated by AlphaFold2 at the modelarchive repository site (<https://www.modelarchive.org/doi/10.5452/ma-1bb5w>, <https://www.modelarchive.org/doi/10.5452/ma-bxxkp>, <https://www.modelarchive.org/doi/10.5452/ma-htx0n>)

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Reviewer #1 (Public Review):

Summary:

This paper makes important contributions to the structural analysis of the DNA replication-linked nucleosome assembly machine termed Chromatin Assembly Factor-1 (CAF-1). The authors focus on the interplay of domains that bind DNA, histones, and replication clamp protein PCNA.

Strengths:

The authors analyze soluble complexes containing full-length versions of all three fission yeast CAF-1 subunits, an important accomplishment given that many previous structural and biophysical studies have focused on truncated complexes. New data here supports previous experiments indicating that the KER domain is a long alpha helix that binds DNA. Via NMR, the authors discover structural changes at the histone binding site, defined here with high resolution. Most strikingly, the experiments here show that for the *S. pombe* CAF-1 complex, the WHD domain at the C-terminus of the large subunit lacks DNA binding activity observed in the human and budding yeast homologs, indicating a surprising divergence in the evolution of this complex. Together, these are important contributions to the understanding of how the CAF-1 complex works.

Weaknesses:

1. There are some aspects of the experimentation that are incompletely described:

In the SEC data (Fig. S1C) it appears that Pcf1 in the absence of other proteins forms three major peaks. Two are labeled as "1a" (eluting at ~8 mL) and "1b" (~10-11 mL). It appears that Pcf1 alone or in complex with either or both of the other two subunits forms two different high molecular weight complexes (e.g. 4a/4b, 5a/5b, 6a/6b). There is also a third peak in the analysis of Pcf1 alone, which isn't named here, eluting at ~14 mL, overlapping the peaks labeled 2a, 4c, and 5c.

The text describing these different macromolecular complexes seems incomplete (p. 3, lines 32-33): "When isolated, both Pcf2 and Pcf3 are monomeric while Pcf1 forms large soluble oligomers". Which of the three Pcf1-alone peaks are oligomers, and how do we know? What is the third peak? The gel analysis across these chromatograms should be shown.

More importantly, was a particular SEC peak of the three-subunit CAF-1 complex (i.e. 4a or 4b) characterized in the further experimentation, or were the data obtained from the input material prior to the separation of the different peaks? If the latter, how might this have affected the results? Do the forms inter-convert spontaneously?

2. Given the strong structural predication about the roles of residues L359 and F380 (Fig. 2f), these should be mutated to determine effects on histone binding.

3. Could it be that the apparent lack of histone deposition by the delta-WHD mutant complex occurs because this mutant complex is unstable when added to the *Xenopus* extract?

Reviewer #2 (Public Review):

Summary:

The authors describe the structure-functional relationship of domains in *S. pombe* CAF-1, which promotes DNA replication-coupled deposition of histone H3-H4 dimer. The authors nicely showed that the ED domain with an intrinsically disordered structure binds to histone H3-H4, that the KER domain binds to DNA, and that, in addition to a PIP box, the KER domain also contributes to the PCNA binding. The ED and KER domains as well as the WHD domain are essential for nucleosome assembly in vitro. The ED, KER domains, and the PIP box are important for the maintenance of heterochromatin.

Strengths:

The combination of structural analysis using NMR and AlphaFold2 modeling with biophysical and biochemical analysis provided strong evidence on the role of the different domain structures of the large subunit of SpCAF-1, spPCF-1 in the binding to histone H3-H4, DNA as well as PCNA. The conclusion was further supported by genetic analysis of the various *pcf1* mutants. The large amounts of data provided in the paper support the authors' conclusion very well.

Reviewer #3 (Public Review):

Summary:

The study conducted by Ouasti et al. is an elegant investigation of fission yeast CAF-1, employing a diverse array of technologies to dissect its functions and their interdependence. These functions play a critical role in specifying interactions vital for DNA replication, heterochromatin maintenance, and DNA damage repair, and their dynamics involve multiple interactions. The authors have extensively utilized various in vitro and in vivo tools to validate their model and emphasize the dynamic nature of this complex.

Strengths:

Their work is supported by robust experimental data from multiple techniques, including NMR and SAXS, which validate their molecular model. They conducted in vitro interactions using EMSA and isothermal microcalorimetry, in vitro histone deposition using *Xenopus* high-speed egg extract, and systematically generated and tested various genetic mutants for functionality in in vivo assays. They successfully delineated domain-specific functions using in vitro assays and could validate their roles to large extent using genetic mutants. One significant revelation from this study is the unfolded nature of the acidic domain, observed to fold when binding to histones. Additionally, the authors also elucidated the role of the long KER helix in mediating DNA binding and enhancing the association of CAF-1 with PCNA. The paper effectively addresses its primary objective and is strong.

Weaknesses:

A few relatively minor unresolved aspects persist, which, if clarified or experimentally addressed by the authors, could further bolster the study.

1. The precise function of the WHD domain remains elusive. Its deletion does not result in DNA damage accumulation or defects in heterochromatin maintenance. This raises questions about the biological significance of this domain and whether it is dispensable. While in vitro assays revealed defects in chromatin assembly using this mutant (Figure 5), confirming these phenotypes through in vivo assays would provide additional assurance that the lack of function is not simply due to the in vitro system lacking PTMs or other regulatory factors.

2. The observation of increased Pcf2-gfp foci in pcf1-ED* cells, particularly in mono-nucleated (G2-phase) and bi-nucleated cells with septum marks (S-phase), might suggest the presence of replication stress. This could imply incomplete replication in specific regions, leading to the persistence of Caf1-ED*-PCNA factories throughout the cell cycle. To further confirm this, detecting accumulated single-stranded DNA (ssDNA) regions outside of S-phase using RPA as an ssDNA marker could be informative.

3. Moreover, considering the authors' strong assertion of histone binding defects in ED* through in vitro assays (Figure 2d and S2a), these claims could be further substantiated, especially considering that some degree of histone deposition might still persist in vivo in the ED* mutant (Figure 7d, viable though growth defective double ED*+hip1D mutants). For example, the approach, akin to the one employed in Fig. 6a (FLAG-IPs of various Pcf1-FLAG-tagged mutants), could also enable a comparison of the association of different mutants with histones and PCNA, providing a more thorough validation of their findings.

4. It would be valuable for the authors to speculate on the necessity of having disordered regions in CAF1. Specifically, exploring the overall distribution of these domains within disordered/unfolded structures could provide insightful perspectives. Additionally, it's intriguing to note that the significant disparities observed among mutants (ED*, PIP*, and KER*) in in vitro assays seem to become more generic in vivo, except for the indispensability of the WHD-domain. Could these disordered regions potentially play a crucial role in the phase separation of replication factories? Considering these questions could offer valuable insights into the underlying mechanisms at play.

Author Response

We thank the three reviewers and the reviewing editor for their positive evaluation of our manuscript. We particularly appreciate that they unanimously consider our work as “important contributions to the understanding of how the CAF-1 complex works”, “The large amounts of data provided in the paper support the authors' conclusion very well” and “The paper effectively addresses its primary objective and is strong”.

We also thank them for a careful reading and useful comments to improve the manuscript. We will build on this input to provide an improved version of the manuscript that will hope to submit soon to eLife along with our point by point answer.