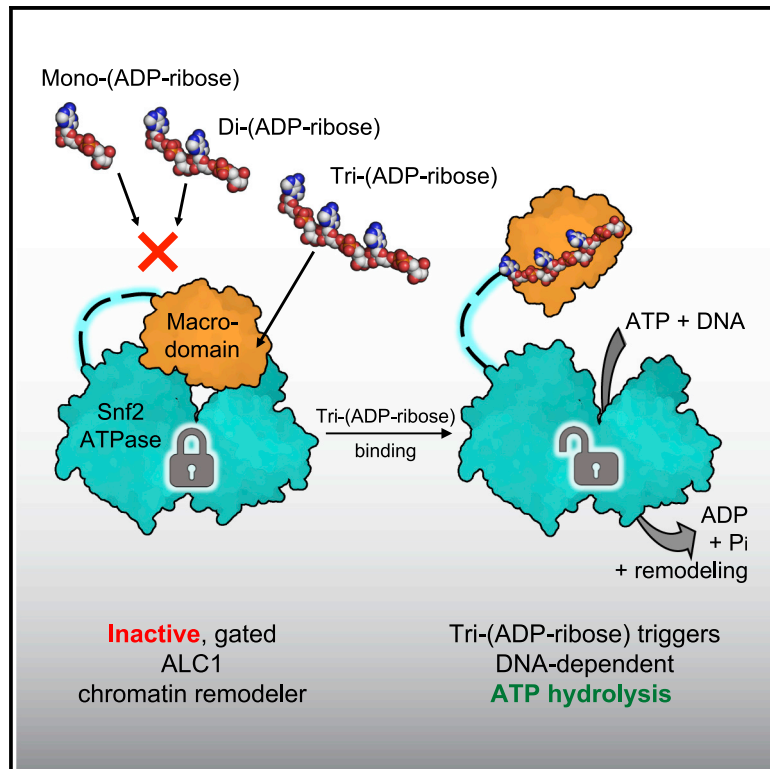


Molecular Cell

A Poly-ADP-Ribose Trigger Releases the Auto-Inhibition of a Chromatin Remodeling Oncogene

Graphical Abstract



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In Brief

The activity of the human oncogene and chromatin remodeler ALC1/CHD1L is strictly regulated by PARP1 activation. Singh et al. reveal how oligomers of ADP-ribose trigger the activation of ALC1 from an auto-inhibited state and identify cancer mutations that disrupt the NAD⁺-metabolite-regulated allosteric mechanism.

Highlights

- The ALC1 macrodomain mediates auto-inhibition of the remodeler's ATPase activity
- PARP1 hyper-activation suppresses the inhibitory protein-protein interaction
- Tri-ADP-ribose promotes an ungated ALC1 conformation and triggers ATPase activity
- Somatic cancer mutations disrupt ALC1's auto-inhibitory mechanism in living cells



A Poly-ADP-Ribose Trigger Releases the Auto-Inhibition of a Chromatin Remodeling Oncogene

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SUMMARY

DNA damage triggers chromatin remodeling by mechanisms that are poorly understood. The oncogene and chromatin remodeler ALC1/CHD1L massively decompacts chromatin *in vivo* yet is inactive prior to DNA-damage-mediated PARP1 induction. We show that the interaction of the ALC1 macrodomain with the ATPase module mediates auto-inhibition. PARP1 activation suppresses this inhibitory interaction. Crucially, release from auto-inhibition requires a poly-ADP-ribose (PAR) binding macrodomain. We identify tri-ADP-ribose as a potent PAR-mimic and synthetic allosteric effector that abrogates ATPase-macrodomain interactions, promotes an ungated conformation, and activates the remodeler's ATPase. ALC1 fragments lacking the regulatory macrodomain relax chromatin *in vivo* without requiring PARP1 activation. Further, the ATPase restricts the macrodomain's interaction with PARP1 under non-DNA damage conditions. Somatic cancer mutants disrupt ALC1's auto-inhibition and activate chromatin remodeling. Our data show that the NAD⁺-metabolite and nucleic acid PAR triggers ALC1 to drive chromatin relaxation. Modular allostery in this oncogene tightly controls its robust, DNA-damage-dependent activation.

INTRODUCTION

Chromatin structure safeguards the integrity of our genome. Distinct families of chromatin remodeling enzymes establish and maintain chromatin structure, for example by facilitating the binding of transcription factors to functional DNA elements or assisting the repair machinery upon DNA damage. Key to controlling the activity of these ATPases are chromatin targeting mechanisms and regulatory interactions (Dann et al., 2017). Such mechanisms help ensure that remodelers are only active where and when needed. While the mechanisms through which the Chd1 and ISWI remodelers are regulated by nucleosomes have been explored (Clapier and Cairns, 2012; Hauk et al., 2010; Ludwigsen et al., 2017; Yan et al., 2016), less is known about how DNA damage triggers the activity of remodelers such as the PARP1-dependent ALC1 (Ahel et al., 2009; Gottschalk et al., 2012; 2009), which massively decompacts chromatin upon DNA damage (Movie S1; Sellou et al., 2016). Considering ALC1's validated roles as an oncogene (ALC1 is amplified in several cancers and promotes metastases, proliferation, and pluripotency; Chen et al., 2010; Jiang et al., 2015; Kulkarni et al., 2013; Ma et al., 2008), understanding how PAR triggers ALC1 activity would advance our molecular understanding of how DNA damage impacts our genome, shed light on how a NAD⁺ metabolite and nucleic acid triggers the activation of an oncogene, and reveal approaches that might allow us to target ALC1 therapeutically.

Single-strand DNA breaks rapidly induce the activity of PARP1, PARP2, and PARP3, enzymes that use NAD⁺ to ADP-ribosylate chromatin and other cellular targets (Carter-O'Connell



et al., 2016; Gibson et al., 2016; Grundy et al., 2016). The clinical promise of PARP1 inhibitors in cancer therapy (Lord and Ashworth, 2017) and the identification of domains that recognize ADP-ribosylated proteins, including ADP-ribose binding macrodomains (Karras et al., 2005; Kustatscher et al., 2005), has rekindled interest in NAD⁺ signaling (Cambronne et al., 2016; Kim et al., 2004; Petesch and Lis, 2012; Tulin and Spradling, 2003). While cellular mono-ADP-ribosylation, catalyzed by related PARP enzymes, is thought to act as a reversible, regulatory post-translational modification (PTM) (Jankevicius et al., 2013; Rosenthal et al., 2013; Sharifi et al., 2013), the tightly regulated synthesis of PAR by PARP1 and PARP2 profoundly alters nuclear organization and cellular homeostasis (Altmeyer et al., 2015; Asher et al., 2010; Bai et al., 2011a; 2011b; Wright et al., 2016).

PAR is a nucleic acid with important roles in the stress response to DNA damage. It is as an abundant, transient polymeric anion that can promote phase separation (Altmeyer et al., 2015; Asher et al., 2010; Bai et al., 2011a, 2011b; Wright et al., 2016). Sites of high PARP1 activity *in vivo* recruit ATP-dependent remodelers, including ALC1 (Amplified in Liver Cancer 1; also known as CHD1L), CHD2, CHD4, SMARCA5/SNF2H, and *Drosophila* Mi-2 (Chou et al., 2010; Murawska et al., 2011; Polo et al., 2010; Smeenk et al., 2013). Remodelers such as ALC1 and CHD2 mediate chromatin relaxation through unknown mechanisms at the site of DNA damage (Luijsterburg et al., 2016; Sellou et al., 2016). These are some of the earliest, PARP-dependent changes in chromatin structure upon DNA damage (Kruhlak et al., 2006; Poirier et al., 1982; Strickfaden et al., 2016).

Others and we have shown that ALC1 recruits to DNA damage sites upon PARP1 activation. Recruitment and PAR binding requires its C-terminal, PAR-binding macrodomain module (Ahel et al., 2009; Gottschalk et al., 2009, 2012; Jiang et al., 2015). Interestingly, its ATPase and nucleosome-remodeling activities depend on PARP1 activation. *In vitro* assays reveal that PARylated PARP1 promotes ALC1-dependent nucleosome sliding (Ahel et al., 2009; Gottschalk et al., 2009, 2012). Key to ALC1's activity is the ability of its macrodomain to recognize PARylated PARP1. However, what keeps ALC1 inactive prior to PARP1 activation and how the nuclear metabolite and nucleic acid PAR triggers ALC1 activation are not known.

RESULTS

Modular Auto-Inhibition in the Remodeler ALC1

We set out to investigate what suppresses ALC1 remodeler activity when PARP1 is inactive. Unlike most remodelers, endogenous ALC1 does not purify as a multi-subunit complex, and its remodeling activity can be efficiently reconstituted *in vitro* using recombinant protein and DNA, together with PARP1 and NAD⁺ (Ahel et al., 2009; Gottschalk et al., 2009, 2012). The enzyme consists of a two-lobed catalytic Snf2-like ATPase domain with homology to ATP-dependent DExx-box helicase (ATPase, Figure 1A), which is connected through a linker region of unknown function to a C-terminal macrodomain (macro), which mediates PARP1 activity-dependent chromatin-targeting (Ahel et al., 2009; Gottschalk et al., 2009, 2012).

To establish whether the ALC1 ATPase domain and macrodomain interact, we generated an ATPase fragment (residues 31–615; “ATPase module”) and a fragment containing both linker and C-terminal macrodomain (residues 616–878; “macro module”) (Figure 1A). The domain boundaries were identified using limited proteolysis (Figure S1). The recombinant ATPase and macrodomain modules can be expressed individually in *E. coli* and purified to homogeneity. Multiple lines of biochemical evidence show that the two modules form a stable complex. The two fragments interact with each other in pull-down assays (Figure 1B). Size-exclusion chromatography assays reveal the formation of a stoichiometric 1:1 complex (Figure 1C), which elutes with a molecular size (~138 kDa) close to that of the (near) full-length ALC1 construct (residues 31–878; eluting at ~131 kDa; Figure 1D). Thus, ALC1 behaves as a monomer. To determine the affinity of the two ALC1 modules for each other, we employed isothermal titration calorimetry (ITC) assays, which reveal an equilibrium dissociation constant of 96 ± 22 nM (Figure 1E). These results indicate that the ATPase and macrodomain modules of the ALC1 remodeler engage through a tight, intra-molecular interaction.

To test whether this interaction is observed in living cells, we used fluorescence-two-hybrid (F2H) analysis (Zolghadr et al., 2012). Tethering of a fluorescent mCherry-LacI-ALC1 macrodomain bait to an integrated LacO array in U2OS cells enriches the eYFP-tagged ALC1 ATPase prey (Figures 1F and S1), while unrelated macrodomains do not recognize the ALC1 ATPase. We conclude that in the absence of exogenous DNA damage (i.e., when PARP1 has not been induced), the ALC1 modules specifically interact with each other. To rule out the possibility that ALC1 may form dimers, trimers, or other, higher-order complexes through intermolecular domain swapping, we conducted co-immunoprecipitation and F2H assays with full-length ALC1 (Figure S2). Both assays indicate that ALC1 is a monomer *in vivo* (compared to positive controls). Our data suggest that the C-terminal macrodomain of ALC1 packs against one or both of its ATPase lobes in the context of the full-length ALC1 protein, hinting at an intramolecular “gating” function of the ALC1 macrodomain, as described for the unrelated chromodomain of γ CHD1 and the NTR domain of ISWI (Hauk et al., 2010; Ludwigsen et al., 2017).

To probe the domain topology of ALC1, we used MS cross-linking. We mapped multiple crosslinks within each of the two ALC1 modules, including between the flexible linker region and the canonical ATPase and macrodomain folds (Figure S1; Table S1). The cross-linking pattern complements well the domain boundaries identified by limited proteolysis (Figure S1). The MS-based crosslinks indicate that the ALC1 hinge contacts the macrodomain and ATPase, confirming our limited proteolysis data. Together, crosslinking and limited proteolysis (Figure S1) hint at a compact arrangement of the ALC1 ATPase and macrodomain modules with respect to each other, consistent with a “gated” structure, which may restrict DNA access to the ATPase motor. We conclude that in the absence of activated PARP1, intramolecular interactions between the macrodomain and ATPase modules establish an auto-inhibited ALC1 conformation.

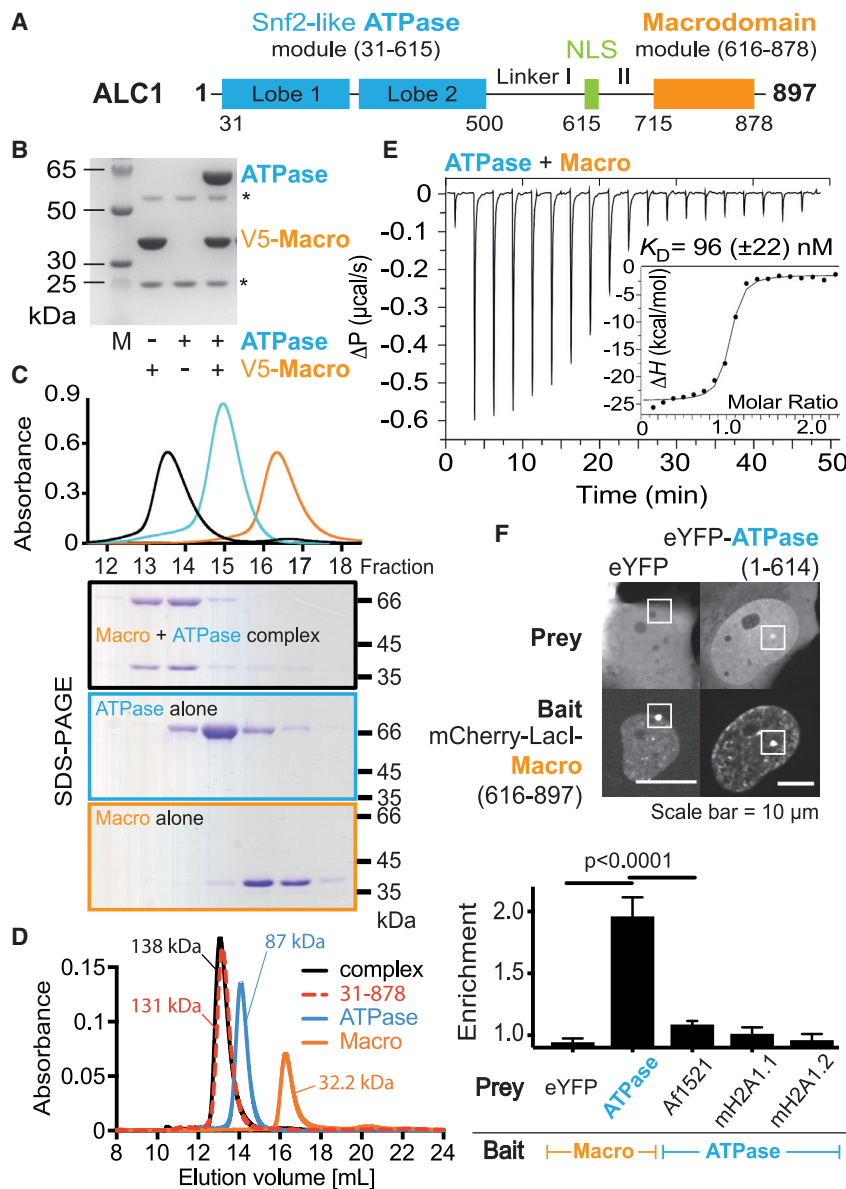


Figure 1. The ALC1 Macrodomain Interacts with the ATPase Module in the Enzyme's Inhibited State

(A) The ALC1 oncogene is composed of two primary modules: an N-terminal Snf2-like ATPase module (residues 31–615) and a C-terminal macrodomain module (616–878). The boundaries were defined using limited proteolysis (Figure S1). NLS, nuclear localization signal.

(B) SDS-PAGE of a V5-based pull-down with recombinant, purified ALC1 macrodomain and ATPase module. The asterisk denotes anti-V5 IgG heavy and light chains.

(C) Size exclusion chromatography (SEC) of recombinant, purified ALC1 macrodomain (residues 636–878, orange), ATPase domain (residues 31–615, cyan), and *in vitro*-reconstituted complex (black), plus SDS-PAGE of the eluted fractions.

(D) Comparison of the elution profiles by gel filtration of the reconstituted ALC1 ATPase–macrodomain complex with purified, near-full-length ALC1 (residues 31–878).

(E) Isothermal titration calorimetry (ITC) assays show that ALC1's two modules bind each other in a high-affinity, exothermic reaction and with 1:1 stoichiometry ($N = 0.87 \pm 0.05$).

(F) Fluorescence-two-hybrid (F2H) analysis in live cells (Zolghadr et al., 2012) reveals that ALC1's ATPase (eYFP-ATPase; prey) readily enriches on a LacO-array tethered mCherry-LacI-macrodomain (bait). Example image (top images), quantitation ($n = 20$), and comparison with unrelated macrodomains reveal a specific ALC1 ATPase and ALC1 macrodomain interaction (bottom chart). Error bar represents the SEM, $n \geq 20$.

See also Figures S1 and S2 and Table S1.

PARP1 Activation Disrupts the Auto-Inhibited State

ALC1 rapidly recruits to DNA damage sites and massively decompacts chromatin in response to PARP1 activation (Ahel et al., 2009; Gottschalk et al., 2009; Sellou et al., 2016). These activities require a functional, PAR-binding ALC1 macrodomain. We hypothesized that PAR binding to ALC1 may promote an active conformation of ALC1. To determine whether the activation of PARP1 in living cells alters the modular, intra-molecular interactions within ALC1, we used the F2H assay to measure the interaction of the ALC1 ATPase module with the macrodomain prior to and following UV-laser induced PARP1 activation. DNA damage leads to a time-dependent decrease of ALC1 ATPase prey from the tethered ALC1 macrodomain (Figure 2A; Movie S2). We conclude that PARP1 activation leads to the loss of interaction between the two ALC1 modules. H_2O_2 -induced DNA damage also leads to a loss of interaction

(Figure S2), and the site of PARP1 activation and ALC1 ATPase–macrodomain dissociation do not need to overlap, since FRAP assays indicate high turnover of our fusion proteins (Figure S2). Importantly, a G750E mutant within the macrodomain, which disrupts binding of the pyrophosphate of ADP-ribose in canonical macrodomains (Kustatscher et al., 2005), retains its binding with the ATPase module, even upon PARP1 induction (Figure S2). These data reveal that the interaction between the two ALC1 modules is regulated by PARP1 activation *in vivo*. PAR binding to the macrodomain is coupled to the loss of interaction with the ATPase module, consistent with a direct, allosteric regulation of ALC1 by PAR.

Next, we sought to determine the minimal ALC1 ligand that is necessary and sufficient to trigger ATPase–macrodomain dissociation and PAR-mediated ALC1 activation. We tested the effect of PARP1 activation *in vitro* on the interaction between ALC1 ATPase and macrodomain. Addition of NAD^+ to a pull-down containing PARP1, DNA, and the two ALC1 modules disrupts ALC1 ATPase–macrodomain interaction (Figure 2B, lanes 4 and 5). Addition of PARP1 inhibitors suppresses the disruptive effect of PARylation on ATPase–macrodomain interaction (lanes

mediates this chromatin plasticity (Sellou et al., 2016), we decided to use an *in vivo* chromatin relaxation assay to test the function of engineered ALC1 macrodomain-deletion fragments.

Since ALC1 does not recruit to chromatin upon DNA damage in the absence of its PAR-binding macrodomain, we tethered full-length, fragment, and mutant LacI-ALC1 fusions to an integrated LacO array in human cells. As expected, wild-type, full-length ALC1 (1–897) does not alter the LacO-array when tethered to the LacO-array (Figure 5A). In sharp contrast, when the macrodomain of ALC1 is deleted, the ALC1 fragment (residues 1–673) decompacts the LacO array (Figure 5A). This chromatin relaxation is seen with other ALC1 C-terminal fragments, but not in a fragment as short as 1–614. This indicates that sequences within linker II of ALC1 (residues 615–673; Figure 1A) promote chromatin remodeling activity, while the macrodomain is inhibitory to ALC1 *in vivo*. Importantly, mutation of conserved residues within the ALC1 helicase that disrupt ATPase activity (Ahel et al., 2009; Gottschalk et al., 2009) abolishes ALC1-induced chromatin relaxation (Figure 5A). Tethered ALC1 fragments lacking the macrodomain thus possess remodeling activity in the absence of DNA damage induction and PARP1 activation.

Next, we tested whether the macrodomain of ALC1 alters the inherent chromatin remodeling activity of the LacI-tethered ALC1 ATPase (1–673) construct when added in *trans*. Addition of the ALC1 macrodomain module to the active, tethered ALC1 ATPase reduces chromatin decompaction (Figure 5A). A PARP1 inhibitor enhances this inhibition. Our data indicate that the ALC1 macrodomain inhibits the ATPase activity of ALC1 at physiological levels of PARP1 activity. In its absence, a tethered ALC1 ATPase module remodels chromatin *in vivo*.

Tri-ADP-Ribose De-represses the ATPase Activity of the ALC1 Remodeler

Our LacO-tethering assay identified a constitutively active ALC1 ATPase fragment (1–673), whose activity can be suppressed by addition of the ALC1 macrodomain module. Since tri-ADP-ribose promotes the dissociation of the ALC1 macrodomain from the ALC1 ATPase module, we sought to determine the relevance of tri-ADP-ribose binding on the catalytic activity of the ALC1 remodeler. We established a robust, DNA-dependent ATPase assay for both the ALC1 ATPase module (31–673) and the (near) full-length ALC1 protein (31–878). We find that the ALC1 ATPase module shows robust, DNA-dependent ATPase activity (Figure S6). Importantly, titration of the ALC1 macrodomain module to the ALC1 ATPase lowers ATPase activity (Figure S6). Once a 2.5 molar excess of ALC1 macrodomain is added to the ALC1 ATPase, the resulting complex is inactive, revealing background ATPase activity similar to that of the ALC1 ATPase without DNA. The ALC1 macrodomain thus represses the inherent ATPase activity present in the ALC1 ATPase module.

Next, we tested whether the addition of the nanomolar tri-ADP-ribose ligand of ALC1 alters the ATPase activity of the enzyme *in vitro*. Addition of a 2-fold molar excess of tri-ADP-ribose to the inactive ALC1 ATPase-macrodomain complex robustly de-represses the ALC1 ATPase, going from <2% without tri-ADP-ribose to ~60% of the activity of the free ALC1 ATPase module (Figure 5B). Importantly, addition of a 6-fold molar excess of mono-ADP-ribose to the ATPase-macrodomain

complex fails to rescue ATPase activity (Figure 5B), consistent with the lack of binding of monomeric ADP-ribose for the ALC1 macrodomain (Figure 3A). Thus, tri-ADP-ribose binding to the ALC1 ATPase-macrodomain complex strongly activates the ATPase activity of the ALC1 remodeler.

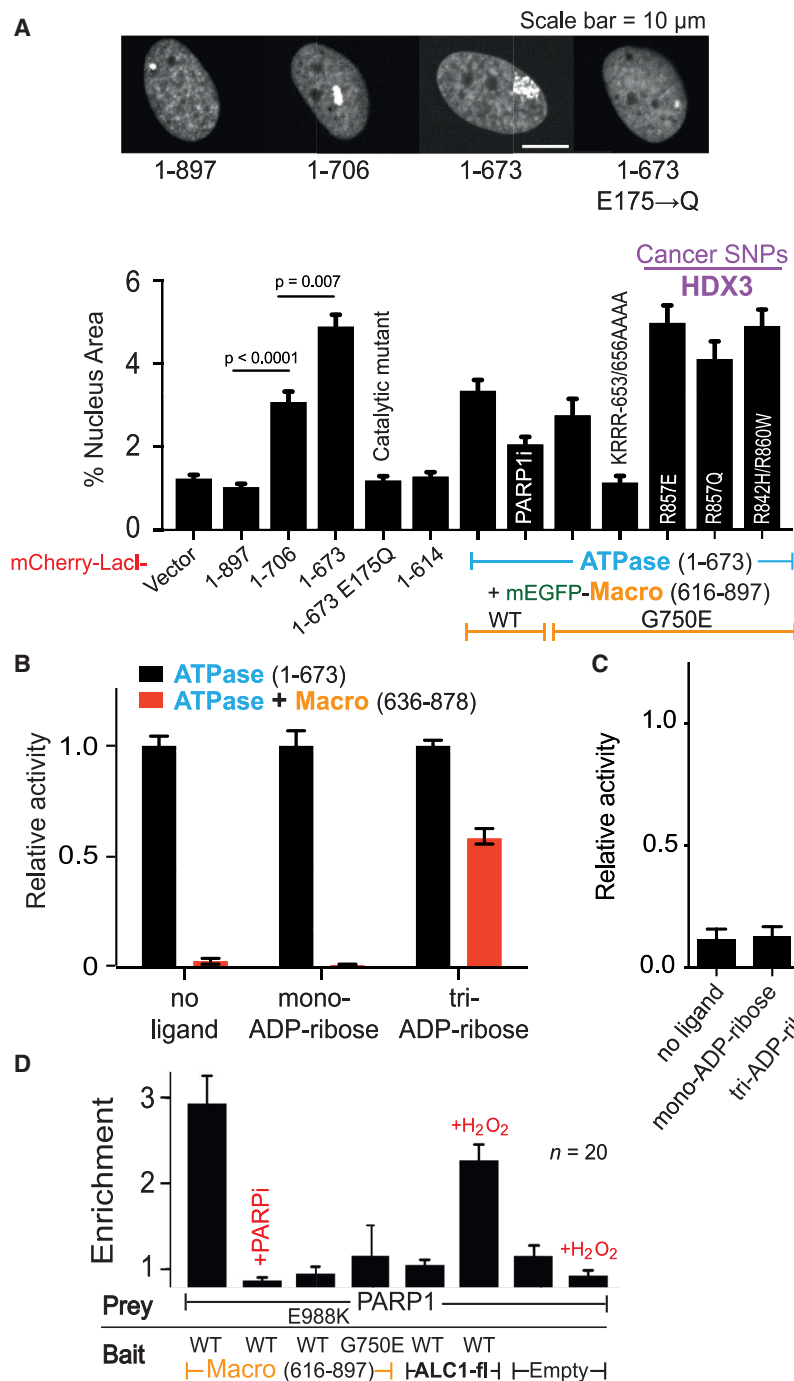
In addition, we tested whether tri-ADP-ribose alters the activity of (near) full-length ALC1. As expected, the ATPase activity of this construct is low, including in the presence of mono-ADP-ribose. However, tri-ADP-ribose strongly activates the ATPase activity in the ALC1 remodeler (Figure 5C). The level of activation (fold induction) induced by tri-ADP-ribose is lower than in our assays using the reconstituted ALC1 complex. However, this is likely the result of degradation products present in our recombinant ALC1 (31–878) construct (Figure S6). Indeed, some of the proteolytic fragments observed in our ALC1 construct (31–878) likely lack (parts of) the inhibitory macrodomain and may thus display catalytic ATPase activity independently of the tri-ADP-ribose trigger. Our assays show that tri-ADP-ribose is a potent activator of the DNA-dependent ATPase activity of the ALC1 remodeler.

Somatic Cancer Mutants in ALC1 Drive Chromatin Remodeling

We identified cancer SNPs located within the HDX3 region of the ALC1 macrodomain, which lead to a loss-of-interaction phenotype between the ALC1 ATPase and macrodomain (Figure 4C). We thus tested the effect of these mutant macrodomains on the activity of the constitutively active ALC1 ATPase tethered to the LacO array when expressed in *trans*. Interestingly, the point mutants R857Q and R842H/R860W show a decompaction of the LacO similar to that of the constitutively active ALC1 ATPase module without any macrodomain expressed in *trans*. In sharp contrast, co-expression of the wild-type macrodomain, or of the G750E macrodomain mutant, strongly reduces the decompacted area, indicating that these macrodomain constructs inhibit the remodeler activity in *trans* and *in vivo* (Figure 5A). Furthermore, FCS assays in the context of the full-length ALC1 protein show a decreased diffusion behavior of the HDX1 and HDX3 mutants compared to wild-type ALC1 (Figure S6). This suggests that HDX mutants that disrupt the intramolecular ATPase-macrodomain interactions promote an ungated structure in ALC1, which may potentially lead to increased DNA binding. Taken together, our data show that somatic cancer mutations phenocopy the activity of the constitutively active ALC1 ATPase fragment. While the relevance for this in cancer remains to be established, our tethering assay indicates that mutations in ALC1 that disrupt its inhibitory intra-molecular interactions (Figure 4C) promote the deregulated, constitutive activity of this chromatin remodeler (Figure 5A).

Modular Allosteric in ALC1 Regulates Interaction with PARP1

To further probe the PAR-regulated modular allostery in ALC1, we tested whether, in turn, the ATPase module affects the ability of the ALC1 macrodomain to recognize its effector molecule, PARylated PARP1, in living cells. While full-length ALC1 does not readily interact with full-length PARP1 in untreated human cells using the F2H assay (Figure 5D; Movie S3), DNA damage induction with H₂O₂ promotes the interaction between



these two proteins, consistent with the recognition of activated, PARylated PARP1 by ALC1. Interestingly, an ALC1 fragment lacking the catalytic ATPase domain readily interacts with PARP1, even in the absence of exogenous DNA damage (Figure 5D). Treatment of cells with a PARP1 inhibitor abrogates this interaction. This indicates that the isolated macrodomain of ALC1 recognizes ADP-ribosylated forms of PARP1 under “non-DNA-damage” conditions (Figure 5D), likely reflecting

Figure 5. Release from Auto-Inhibition Drives ALC1's Chromatin Remodeling Activity

(A) Tethering of engineered mCherry-LacI-ALC1 to an integrated LacO array decompacts chromatin in U2OS cells (representative images; top). The decompaction of the LacO array is calculated as percent of the nucleus area (bottom). The deletion of ALC1's macrodomain generates a constitutively active ALC1 that decompacts the LacO-array *in vivo*. Constructs assayed: full-length ALC1 (1–897), macrodomain deletion (1–707), a hyperactive construct (1–673), ATPase-dead point mutation (1–673, E175Q), plus ALC1 (1–614), which represents the ATPase module identified in our limited proteolysis (Figure S1). Importantly, co-transfection of the ALC1 macrodomain (mEGFP-616–897) with the constitutively active ALC1 fragment (1–673) reduces the decompacted area. Further, cancer SNPs within HDX3 that disrupt interaction with the ATPase module (Figure 4C) do not decrease the chromatin decompaction catalyzed by the ALC1 ATPase (1–673) module. Error bar represents the SEM, $n \geq 20$.

(B) Tri-ADP-ribose de-represses the ATPase activity of the inactive ALC1 ATPase-macrodomain complex. The DNA-dependent ATPase activity of the ALC1 ATPase module was measured using a malachite green assay in the presence and absence of a 2.5 molar excess of ALC1 macrodomain module, as well as in the absence (left) or presence of either a 6-fold molar excess of ADP-ribose (middle) or a 2-fold molar excess of tri-ADP-ribose (right). The data are normalized to the respective mean activity of the ATPase module alone (black bars; $n = 3$; mean \pm SEM).

(C) Tri-ADP-ribose promotes the activation of the ALC1 chromatin remodeling enzyme. The (near) full-length ALC1 construct (31–878) shows only basal ATPase activity in the absence or presence of a 15-fold molar excess of ADP-ribose. In contrast, a 5-fold molar excess of tri-ADP-ribose greatly stimulates ALC1-catalyzed and dsDNA-dependent ATP hydrolysis. The data are normalized to the mean value of ALC1 activity in the presence of tri-ADP-ribose ($n = 3$; mean \pm SEM).

(D) F2H assay testing the interaction of tethered ALC1 macrodomain (wild-type, WT; G750E mutant) with fluorescently tagged PARP1 (wild-type and E988K PAR elongation mutant). Indicated experiments were done in the presence of a PARP1 inhibitor (+PARPi) or H₂O₂. Error bar represents the SEM, $n \geq 20$.

See also Figure S6.

background ADP-ribosylation. Consistently, a point mutant in the ALC1 macrodomain that reduces PAR binding (G750E), or mutation of a key residue in PARP1 (E988K) that is responsible for the elongation of mono-ADP-ribosyl-PARP1 to poly-ADP-ribosyl-PARP1, disrupts the interaction between the ALC1 macrodomain and PARP1 (Figure 5D). We conclude that the isolated ALC1 macrodomain interacts with ADP-ribosylated PARP1 under physiological conditions, while full-length ALC1 requires a

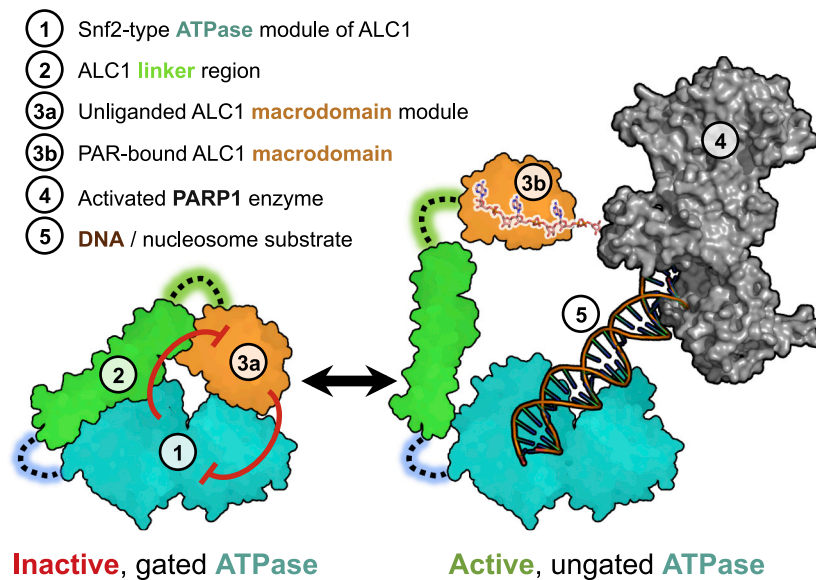


Figure 6. Modular Allostery Sets a Threshold for PARP1-Induced ALC1 Activation

Modular allostery in the chromatin remodeler ALC1 regulates auto-inhibition through the reciprocal interaction of ALC1's ATPase and macrodomain modules. This helps to ensure that the PARP1 product PAR acts as an allosteric activator and potent trigger of ALC1-promoted chromatin relaxation only once acute DNA damage has induced PARP1 activity.

In vivo, binding of the PAR effector to the macrodomain occurs when ALC1 recruits to DNA damage sites, which tethers the remodeler to chromatin and allows ALC1 to remodel chromatin. Our study does not identify the specific, physiological substrate that ALC1 remodels on chromatin. Swi2/Snf2 remodelers such as Mot1 remodel non-nucleosomal substrates (Wollmann et al., 2011). *In vivo*, the PARylation of histones, PARP1, ALC1, and/or other chromatin factors may thus contribute to how ALC1 catalyzes chromatin relaxation.

high threshold of DNA damage and PARP1 activation in order to interact with PARylated PARP1. Thus, the ATPase of ALC1 lowers the affinity of the macrodomain for PARylated-PARP1, consistent with modular allostery. We suggest that the modularity of ALC1 allows the remodeler to be activated only once a threshold of PARP1 activation has been reached.

DISCUSSION

Auto-inhibitory interactions play important roles in signaling and in the regulation of chromatin and repair factors (DaRosa et al., 2015; Guo et al., 2015). Considering the emergent role of remodelers in cancer (St Pierre and Kadoch, 2017; Zhao et al., 2017), a better understanding of how DNA damage alters chromatin is important. While the mechanisms that ALC1 and CHD2 employ to relax chromatin upon DNA damage *in vivo* (Movie S1) are not known, and the remodelers' substrate(s) *in vivo* remains to be identified, here we have identified and dissected the mechanisms that allow the oncogene ALC1 to be tightly regulated by the NAD⁺ metabolite PAR (Figure 6). We show that reciprocal interactions between the ALC1 ATPase and its macrodomain allow ALC1 activity to be controlled by PARP1 activation. The binding of an oligomer of at least three ADP-ribose units to ALC1's macrodomain triggers conformational changes that disrupt auto-inhibitory interactions. This "ungates" the ATPase module, promoting DNA-dependent ATPase activity *in vitro* and remodeling *in vivo* (Figures 3, 4, and 5). Modular allostery thus ensures that ALC1 is exquisitely sensitive to and selective for oligomeric forms of ADP-ribose. We infer that PAR acts as a catalytic trigger only once a threshold of PARP1 induction has been reached. Most "reader" modules in chromatin biology are thought to play a recruitment and tethering function. Our identification of a reciprocal interaction between a PAR-binding macrodomain ("reader" module) and the catalytic ATPase module of ALC1 adds to the allostery described for DNA methyltransferases (Guo et al., 2015; Jeltsch and Jurkowska, 2016).

or other chromatin factors may thus contribute to how ALC1 catalyzes chromatin relaxation.

The selectivity of ALC1 toward oligo-ADP-ribose, and the fact that the enzyme's ATPase impairs the ability of ALC1's macrodomain to bind PARP1 under non-DNA damage conditions (Figure 5D), likely helps to ensure that PARP1-dependent chromatin relaxation is only catalyzed once PARP1 has been activated, such as during DNA damage. Our data reveal how the ATPase activity of a remodeler is gated by the PARP1-product and nucleic acid PAR through regulatory interactions mediated by ALC1's macrodomain. This adds to our mechanistic understanding of how the ATPase activity of chromatin remodelers is regulated. Further work will be necessary to dissect how poly-ADP-ribosylated PARP1 promotes the efficient remodeling of nucleosomes. Our data show that in the special case of ALC1, tethering and activation occur through a NAD⁺ metabolite, which acts as an allosteric ATPase trigger, in a mechanism mediated by the remodeler's core macrodomain fold and additional contacts with the PAR ligand.

Our analysis also identifies how the oncogene ALC1 might be targeted in cancer. Small molecules that inhibit its allostery or activity should reproduce ALC1 knockdown phenotypes, such as reduced tumor growth, reduced reprogramming, and increased sensitivity to chemotherapy (Jiang et al., 2015; Liu et al., 2016). Compounds that stabilize the inactive, gated conformation of ALC1, which lower its catalytic activity, or that disrupt its ability to recognize PAR should suppress the potent chromatin relaxation activity of this oncogene.

STAR★METHODS

Detailed methods are provided in the online version of this paper and include the following:

- KEY RESOURCES TABLE
- CONTACT FOR REAGENT AND RESOURCE SHARING

METHOD DETAILS

- Cloning, protein expression and purification
- Limited trypsin proteolysis
- Full-length ALC1 baculovirus cloning expression and purification
- Hydrogen deuterium exchange mass-spectrometry (HDX-MS)
- Chemical cross-linking coupled to mass spectrometry (XL-MS)
- Isothermal titration calorimetry (ITC) assays
- Plasmids for cellular assays
- Cell culture and transfections
- Microscopy experiments
- UV-laser micro-irradiation assays
- PARP1 inhibitor treatment and H₂O₂ treatments
- Fluorescence two-hybrid (F2H) assay
- Co-immunoprecipitation assays
- LacO array remodelling assays
- V5-Macrodomein pull-down assays
- Fluorescence correlation spectroscopy (FCS)
- Thermal shift assays
- Chromatin remodeler ATPase assays
- Analytical size exclusion chromatography (aSEC)

QUANTIFICATION AND STATISTICAL ANALYSIS

- For *in vitro* ATPase and Thermofluor assays
- For the F2H and lacO array assays
- For HDX-MS measurements

SUPPLEMENTAL INFORMATION

Supplemental Information includes six figures, three tables, and three movies and can be found with this article at <https://doi.org/10.1016/j.molcel.2017.11.019>.

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Conceptualization, H.R.S., M.H., G.T., and A.G.L.; Methodology, H.R.S., A.P.N., I.R.M., G.K., F.M.P., G.T., K.D.R., and A.G.L.; Investigation, H.R.S., A.P.N., I.R.M., G.K., M.H., N.H., C.B., C.K., and S.H.; Formal Analysis, H.R.S., A.P.N., I.R.M., G.K., M.H., N.H., C.B., F.M.P., G.T., K.D.R., and A.G.L.; Writing – Original Draft, H.R.S. and A.G.L.; Writing – Review & Editing, H.R.S., G.K., and A.G.L.; Funding Acquisition, H.A.V.K., D.V.F., F.M.P., and A.G.L.; Resources, H.A.V.K., D.V.F., S.E., S.H., and C.K.; Supervision, S.H., F.M.P., G.T., K.D.R., and A.G.L.

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STAR★METHODS

KEY RESOURCES TABLE

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Antibodies		
Rabbit anti-mCherry-antibody	Novus Bio	NBP2-25157
Bacterial and Virus Strains		
<i>Escherichia coli</i> DH5 alpha	Thermo-Fisher Scientific	18265017
<i>Escherichia coli</i> Rosetta (DE3) pLysS Competent Cells - Novagen	Merck Millipore	70956
<i>Escherichia coli</i> DH10MultiBac	Geneva Biotech	DH10MultiBac
<i>Spodoptera frugiperda</i> Sf21 insect cells	ThermoFisher Scientific (Invitrogen)	11497013
<i>Trichoplusia ni</i> High Five insect cells	ThermoFisher Scientific (Invitrogen)	B85502
Chemicals, Peptides, and Recombinant Proteins		
Immobilized pepsin (agarose resin) for online digestion (HDX-MS)	ThermoFisher Scientific	20343
AG14361	Selleckchem	S2178
Homo bi-functional cross-linker BS3	ProteoChem	c1103
Ammonium bicarbonate (NH ₄ HCO ₃)	Sigma	09830
DMSO	Life Technologies	D12345
Trypsin	Promega	V511B
Sep-Pak tC18 Cartridges	Waters	WAT043410
PARP1inhibitor AG14361	Selleck Chemicals	S2178
Di- and tri-ADP-ribose	(Kistemaker et al., 2015)	N/A
Adenosine 5'-diphosphoribose (ADPr)	Sigma	A0752
DMEM Dulbecco's	Sigma	S5796
FBS	GIBCO	10270
Sodium pyruvate	Sigma	S8636
L-glutamine	Sigma	G7513
Penicillin	Sigma	P3032
Streptomycin	Sigma	S9137
Hygromycin B	Sigma	H3274
CO ₂ -independent imaging medium	GIBCO by Life technologies	18045-054
Cell culture Dulbecco's PBS	Sigma	D8537
Critical Commercial Assays		
SYPRO Orange, 5000x in DMSO	Sigma	S5692
Biomol Green	Enzo Life Sciences	BML-AK111
Gel filtration calibraton kit, Low Molecular Weight	GE Healthcare	17-0442-01
Gel filtration standards	BioRad	151-1901
384-well Microplates; ATPase assay	Greiner Bio One	781101
Borosilicate 8-well LabTeks	Thermo-Scientific	155411
X-fect transfection reagent	Clontech	631317
GFP-Trap_A	Chromotek	gta-20
Experimental Models: Cell Lines		
Human U2OS cells harboring the stably integrated lacO (256x) array	(Soutoglou and Misteli, 2008)	N/A

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REAGENT or RESOURCE	SOURCE	IDENTIFIER
Recombinant DNA		
pmCherry-C1 Vector	This study	See Table S3
pmCherry-Lacl-C1	This study	See Table S3
pmCherry-N1Human SSRP1	This study	See Table S3 ; digested with XhoI and HindIII and cloned into mCherry-N1 vector
pEYFP-ALC1-1-897-C1	(Gottschalk et al., 2009)	N/A
pmCherry-Lacl-ALC1-1-897-C1	This study	See Table S3
pmCherry-Lacl-ALC1-1-706-C1	This study	See Table S3
pmCherry-Lacl-ALC1-1-673-C1	This study	See Table S3
pmCherry-Lacl-ALC1-1-614-C1	This study	See Table S3
pmEGFP-Lacl-ALC1-616-897-C1	This study	See Table S3 ; subcloned from pEYFP-ALC1-1-897-C1 between BglIII-EcoR1
pmCherry-Lacl-ALC1-616-897-C1	This study	See Table S3 ; subcloned from pmEGFP-Lacl-616-897-C1; BglIII and EcoR1
pEYFP-ALC1-1-614-C1	This study	See Table S3
pmEGFP- ALC1-1-897-C1	This study	See Table S3 ; subcloned from pEYFP-ALC1-1-897-C1 (Restriction sites BglIII and EcoR1)
pmEGFP-ALC1-616-897-C1	This study	See Table S3 ; subcloned from pmEGFP-Lacl-616-897-C1; Restriction sites BglIII and EcoR1
pET-MCN-ALC1-31-878	This study	See Table S3
pET-MCN-ALC1-31-615	This study	See Table S3
pET-MCN-ALC1-31-605	This study	See Table S3
pET-MCN-ALC1-616-876	This study	See Table S3
pET-MCN-V5-ALC-616-876	This study	See Table S3
pET-MCN-ALC-636-876	This study	See Table S3
pETM-11: ALC1 31-673	This study	See Table S3
pmEGFP-C1Human Spt16	This study	See Table S3
pFBDM-TWINStrep-ALC1-1-897	This study	See Table S3
pcDNA3eYFP-Af1521-macrodomein	(Timinszky et al., 2009)	N/A
pcDNA3eYFP-macroH2A1.1-macrodomein	(Timinszky et al., 2009)	N/A
pcDNA3eYFP-macroH2A1.2-macrodomein	(Timinszky et al., 2009)	N/A
pET-MCN-V5-ALC-616-876: G750E	This study	See Table S3
pmEGFP-ALC1-616-897-C1: G750E	This study	See Table S3
pmEGFP-ALC1-1-897-C1: G750E	This study	See Table S3
pmCherry-Lacl-ALC1-1-614-C1: E175Q	This study	See Table S3
pmCherry-Lacl-ALC1-1-614-C1: EPEPFE/APAPAA	This study	See Table S3
pmEGFP-ALC1-1-897-C1: EPEPFE/APAPAA	This study	See Table S3
pmCherry-Lacl-ALC1-1-614-C1: PEPFE/PAPAA	This study	See Table S3
pmCherry-Lacl-ALC1-1-614-C1: RK319/320EE-KR407/422DE	This study	See Table S3
pmCherry-Lacl-ALC1-1-614-C1: KR407/422DE	This study	See Table S3
pmCherry-Lacl-ALC1-1-614-C1: KK307/308EE-K398E	This study	See Table S3
pmCherry-Lacl-ALC1-1-614-C1: S420A	This study	See Table S3
pmEGFP-ALC1-616-897-C1: 653-KRRR-656/AAAA-G750E	This study	See Table S3
pmEGFP-ALC1-616-897-C1: R857E/G750E	This study	See Table S3
pmEGFP-ALC1-616-897-C1: R857Q/G750E	This study	See Table S3
pmEGFP-ALC1-1-897-C1: R857Q	This study	See Table S3

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REAGENT or RESOURCE	SOURCE	IDENTIFIER
pmEGFP-ALC1-616-897-C1: R842H/R860W/G750E	This study	See Table S3
pmEGFP-ALC1-1-897-C1: R842H/R860W	This study	See Table S3
pmEGFP-ALC1-1-897-C1: R842H	This study	See Table S3
pmEGFP-ALC1-1-897-C1: R860W	This study	See Table S3
Software and Algorithms		
ProteinLynx Global Server 3.0	http://www.waters.com/waters/home.htm (Waters)	N/A
DynamX 3.0	http://www.waters.com/waters/home.htm (Waters)	N/A
MS Convert Proteowizard Tools	http://proteowizard.sourceforge.net/tools.shtml	N/A
Crossfinder	(Mueller-Planitz, 2015)	N/A
PEAQ-ITC Analysis Software	https://www.malvern.com/en/ (Malvern)	N/A
Graphpad-Prism	https://www.graphpad.com/scientific-software/prism/	N/A
Fiji(ImageJ)	(Schindelin et al., 2012)	N/A
Fluctuation Analyzer 4G software	(Wachsmuth et al., 2015)	N/A
Other		
Superdex 200 Increase 10/300 GL	GE Healthcare	28990944
15-cm analytical column C18 nanocolumn (75 μ m ID homepacked with ReproSil-Pur C18-AQ 2.4 μ m)	N/A	N/A
Ultimate 3000 HPLC system	ThermoFisher Scientific	IQLAAAGABHFAPBMBFB

CONTACT FOR REAGENT AND RESOURCE SHARING

Further information and requests for resources and reagents should be directed to and will be fulfilled by the Lead Contact, Andreas G. Ladurner (andreas.ladurner@bmc.med.lmu.de).

METHOD DETAILS**Cloning, protein expression and purification**

All human ALC1 fragments and mutants were engineered by sub-cloning and site-directed mutagenesis, using oligonucleotides defined in [Table S3](#). Recombinant proteins were expressed and purified as N-terminally 6 \times His-tagged fusion proteins from *E. coli* Rosetta (DE3) pLys using pETM-CN or pETM-11 (only for ALC1 31-674) vectors. Starting cultures were grown in LB-medium supplemented with the appropriate antibiotics over night at 30°C and used at a 1:50 dilution to inoculate the expression cultures. Expression cultures were then grown in rich medium (supplemented with antibiotics) at 37°C and 160-200 rpm until reaching an OD₆₀₀ of 0.6 – 0.8 before protein expression was induced with 0.2 mM isopropyl β -D-1 thiogalactopyranoside (IPTG). After induction, protein expression was allowed to proceed for 18 hr at 18°C (ALC1 570-897, 31-605, 31-615, 615-876, V5-615-876, 636-878) or 16°C (ALC1 31-674, 3-897, 31-878) until the cells were harvested by centrifugation at 8,000 \times g for 10 min. Bacterial pellets were either directly used for protein purification or stored at –80°C. The cells were lysed by sonication at 4°C in lysis buffer (50 mM Tris-HCl pH 7.5, 500 mM NaCl, 20 mM imidazole, 5 mM dithiothreitol (DTT) and Complete EDTA free protease inhibitor tablets (Roche). The lysate was cleared by centrifugation at 45,000 \times g and loaded onto Ni-Sepharose 6FF beads for batch purifications or a HisTrap HP 1mL column operated on a ÄKTA pure FPLC system (GE Healthcare). After washing with several (15-30) column volumes (CV) lysis buffer, proteins were eluted either stepwise (batch purification) or in a 15 CV gradient of elution buffer (lysis buffer supplemented with 500 mM imidazole). The eluate was concentrated by ultrafiltration (Vivaspin 30,000 MWCO, Sartorius) and loaded onto a Superdex 200 16/60 column (GE Healthcare) equilibrated in SEC-buffer (25 mM Tris-HCl pH 7.5, 200 mM NaCl, 1 mM DTT). Peak fractions were pooled and diluted with dilution buffer (25 mM Tris-HCl pH 7.5, 2 mM DTT) to a final salt concentration of 50 mM NaCl (low-salt buffer) and loaded onto a cation exchange column (MonoS 5/5 HR or Resource S 6mL (GE Healthcare), full length and Macro domain constructs), or an anion exchange column (MonoQ 5/5 HR or Resource Q 6mL (GE Healthcare), ATPase constructs) pre-equilibrated with low-salt buffer. After washing with 5-10 CV low-salt buffer, proteins were eluted by increasing the NaCl concentrations to

500 mM in a linear gradient of 10-20 CV. Peak fractions were again pooled and concentrated by ultrafiltration (Vivaspin 30,000 MWCO, Sartorius). The purified proteins were either used directly or snap-frozen in liquid nitrogen for storage at -80°C . For the ITC assays, the macrodomain of macroH2A1.1 was expressed and purified, as published (Kustatscher et al., 2005; Timinsky et al., 2009). Briefly, N-terminally 6 \times His-tagged of MacroH2A1.1 (residues 162-369) was expressed for 3 hr at 37°C after induction with 1 mM IPTG. The cells were lysed in 50 mM Tris-HCl pH 7.4, 500 mM NaCl, 1 mM MgCl_2 , 5 mM β -ME and 10% (v/v) glycerol. The protein was bound to Ni-Sepharose 6FF beads, washed with lysis buffer containing 1 M NaCl and 40 mM imidazole, and eluted in lysis buffer supplemented with 500 mM imidazole. The ALC1 macrodomain and the ALC1 ATPase module were purified as described above.

Limited trypsin proteolysis

500 μl of purified Hs ALC1 (3-897; full-length) or Hs ALC1 (570-897; macrodomain) at a concentration of 1 mg/ml were incubated at room temperature (RT) with trypsin (Sigma Aldrich) at a ratio of 1:100 (full-length) and 1:10 (macrodomain). Reactions were stopped by addition of protein gel loading buffer and boiling (for SDS-PAGE analysis) or 2 mM PMSF (final; for analytical SEC analysis). The resulting major bands on the SDS-PAGE gels were used for MALDI peptide mass fingerprinting to identify the peptides corresponding to the particular band.

Full-length ALC1 baculovirus cloning expression and purification

Full length human ALC1 (1-897) was expressed as N-terminally Twin-Strep-tagged fusion protein using MultiBac technology (Fitzgerald et al., 2006). Briefly, cDNA was cloned into pFBDM vector for expression under control of a polyhedrin promoter. Transposition into MultiBac baculoviral DNA was performed in *E. coli* DH10MultiBac cells (Geneva Biotech). Isolated bacmid DNA was transfected into *Spodoptera frugiperda* Sf21 insect cells (Invitrogen) in order to generate baculovirus, while large-scale protein expression was carried out using *Trichoplusia ni* High Five insect cells (Invitrogen). High Five cells were infected with 1/100 (v/v) with baculovirus. Cells were cultured for 60 hr at 27°C until they were harvested by centrifugation. Cells were lysed by gentle sonication in lysis buffer (20 mM HEPES pH 8.0, 500 mM NaCl, 10% [v/v] Glycerol, 2 mM MgCl_2 , 1 mM DTT) and complete EDTA free protease inhibitor tablets (Roche). The lysate was cleared by centrifugation and loaded onto Streptactin-Sepharose (IBA) for batch purification. After washing beads with lysis buffer, protein was eluted using 3.5 mM d-Desthiobiotin (Sigma). The eluate was concentrated by ultrafiltration (Vivaspin 30,000 MWCO, Sartorius) and loaded onto a Superdex 200 16/60 column (GE Healthcare) equilibrated in SEC-buffer (20 mM HEPES pH 8.0, 500 mM NaCl, 10% [v/v] Glycerol, 2 mM MgCl_2 , 1 mM DTT). Purified, monodisperse ALC1 protein was snap-frozen in liquid nitrogen for storage at -80°C .

Hydrogen deuterium exchange mass-spectrometry (HDX-MS)

For HDX labeling: All proteins were incubated for 30 min at 25°C before initiating the labeling reactions. Deuterated buffer (2.5 mM Tris (pH 7.5), 150 mM NaCl, 0.1 mM DTT) was added in a 10:1 dilution (v/v), resulting in 90% (v/v) D_2O and 0.8 μM protein concentration during labeling. The reactions were incubated for various time intervals ranging from 0.25 to 60 min. The following labeling reactions were prepared in duplicates unless otherwise specified: a) A full time series of the full length ALC1 with and without 5-fold molar excess of tri-ADP-ribose. Single measurements were conducted for the entire time series and from the results, two time points (0.25 and 10 min) were selected for triplicate measurements. b) A 10 min time point of the full length ALC1 with and without 50-fold molar excess of tri-ADP-ribose. c) A 0.25 min time point of the Macrodomain with and without 5-fold molar excess of tri-ADP-ribose or 50-fold molar excess of mono-ADP-ribose. d) A 0.25 min time point of the 1:1 molar ratio mixture of the ATPase domain and the Macrodomain with and without 5-fold molar excess of tri-ADP-ribose. Following the indicated incubation periods, the labeling reactions were quenched by 1:1 (v/v) dilution into ice-cold quench buffer (219 mM potassium phosphate (pH 2.5), 6 M urea). The quenched samples were immediately frozen and stored at -80°C . For LC-MS and LC-MS/MS measurements: Samples were quickly thawed and injected into a Waters nanoAcquity UPLC system with an HDX Manager which allows for online digestion at 20°C and desalting and separation of peptides at 0°C . Following online digestion on column packed with immobilized pepsin on agarose (Thermo Scientific Pierce, Rockford, USA), the peptic peptides were trapped on a C18 Vanguard trap column (Acquity UPLC BEH C18 column, 130 \AA , 1.7 μm , 2.1mm X 5 mm – Waters – Product number: 186003975) and desalted for 3 min with 0.23% formic acid in water (pH 2.5) at a flow rate of 200 $\mu\text{l}/\text{min}$. The peptides were eluted from the trap column to an analytical column (Acquity UPLC BEH C18 column, 130 \AA , 1.7 μm , 1mm X 100 mm – Waters – Product number: 186002346) fitted with a C18 Vanguard column (Acquity UPLC BEH C18 column, 130 \AA , 1.7 μm , 2.1mm X 5 mm – Waters – Product number: 186003975) and separated using a 9 min gradient from 8% to 40% of 0.23% formic acid in acetonitrile at a flow rate of 40 $\mu\text{l}/\text{min}$. The peptides were ionized by electrospray ionization and mass spectra collected using a Waters Synapt G2 HDMS mass spectrometer set in positive ion mode. Peptic peptides of non-deuterated samples were identified in a separate experiment using collision-induced dissociation tandem mass spectrometry with data collected in a data-independent (MS^E) manner. Peptide identification was carried out using ProteinLynx Global Server 3.0 and the HDX-MS data was processed using DynamX 3.0 from Waters.

Chemical cross-linking coupled to mass spectrometry (XL-MS)

Cross-linking reactions were carried out with 1 μM purified ALC1 full-length protein in 25mM HEPES KOH pH 7.6, 0.1 mM EDTA, 10% glycerol, 100 mM KOAc, 1.5 mM MgCl_2 , 1 mM DTT. The homobifunctional cross-linker BS3 (ProteoChem) was dissolved freshly in

DMSO and added to a final concentration of 1 mM. The cross-linking reactions and a mock-treated control were incubated on ice for 20 min before quenching with NH_4HCO_3 to a final concentration of 200 mM for 10 min. The protein sample was loaded onto a pre-equilibrated Superdex 200 (Increase 10/300 GL, GE Healthcare). The elution profile at 280 nm was consistent with ALC1 being monomeric in solution and no evidence for a possible dimer formation was observed. Protein fractions were pooled and concentrated by the addition of 15% (w/v) trichloroacetic acid with pelleting. The pellet was washed twice with acetone and re-dissolved in 8 M urea. In-solution digestion was performed after urea dilution with trypsin (enzyme to substrate ratio 1:50) at 37°C overnight. Digestion was stopped by the addition of 2% (v/v) formic acid. Acidified peptides were purified and concentrated by a C18 resin (Sep-Pak tC18 Cartridges, Waters). The eluate was dried by vacuum centrifugation and re-dissolved in 0.1% trifluoroacetic acid. To map the cross-links by LC-MS/MS, samples were desalted offline using C18 Stagetip and injected in an Ultimate 3000 (Thermo) or RSLCnano system (Thermo). Peptides were separated in a 15 cm analytical column C18 nanocolumn (75 μm ID homepacked with ReproSil-Pur C18-AQ 2.4 μm from Dr. Maisch) with a 40 or 60 min gradient from 5 to 60% acetonitrile in 0.1% formic acid. The effluent from the HPLC was directly electrosprayed into a Q Exactive HF (Thermo). The Q Exactive HF instrument was operated in data dependent mode to automatically switch between full scan MS and MS/MS acquisition. Survey full scan MS spectra (from m/z 375–1600) were acquired with resolution $R = 60,000$ at m/z 400 (AGC target of 3×10^6). The ten most intense peptide ions with charge states between 3 and 5 were sequentially isolated to a target value of 1×10^5 , and fragmented at 27% normalized collision energy. Typical mass spectrometric conditions were: spray voltage, 1.5 kV; no sheath and auxiliary gas flow; heated capillary temperature, 250°C; ion selection threshold, 33,000 counts. Thermo binary raw file were transformed to mzXML files using msconvert (Proteowizard Tools; proteowizard.sourceforge.net/tools.shtml). Cross-linked peptides were identified using Crossfinder (Mueller-Planitz, 2015). An MS1 and MS2 tolerance window of 10 and 12 ppm, respectively, was applied. The number of missed trypsin cleavage sites was set to three. For all peptides, oxidation of methionine was set as variable modification and carbamido methylation as fixed modification. All lysine residues were considered as potential cross-linking sites for BS3. False-discovery rates (FDRs) were estimated by Crossfinder and results were filtered according the following parameters for identification of cross-linking candidates: FDR < 1%, number of fragment ions per spectrum ≥ 6 , number of fragment ions per peptide ≥ 3 , number of fragment ions with cross-linker ≥ 1 , fractional intensity of assigned MS2 peaks ≥ 0.05 , relative filter score: 100. We ensured that cross-linking candidates were not present in the mock-treated sample. MS2 spectra of cross-linking candidates spanning at least 10 amino acids in the primary sequence were manually validated.

Isothermal titration calorimetry (ITC) assays

For ITC binding assays the Peaq-ITC instrument (Malvern) was used. Before the experiment, proteins were dialyzed overnight against 25 mM Tris, 100 mM NaCl buffer (pH 7.5) and 5 mM 2-mercaptoethanol at 4°C. The dialyzed proteins were then centrifuged for 20 min at 35,000 g at 4°C and the protein concentration was determined by absorbance measurements at 280 nm wavelength using calculated molar extinction coefficients. All the ITC experiments were carried out at 25°C. The binding reactions were performed using 5–15 μM macrodomain and 50–150 μM ligand in the syringe. For the ALC1 macrodomain–ATPase interaction studies, all the experiments were carried out using 10 μM ATPase (cell) and 120 μM of ALC1 macrodomain (syringe). For the competition assay, 4.5 μM ALC1 macrodomain was incubated with 2-fold molar excess of tri-ADP ribose (9 μM) at room temperature and then loaded into the cell. The same molar concentration of tri-ADP ribose was added into the ligand-containing solution consisting of 45 μM of ATPase. PEaq-ITC Analysis Software (Malvern) was used for data analysis. GraphPad Prism software was used to plot the data. Adenosine 5'-diphosphoribose was bought from Sigma (A0752). Di- and tri-ADP ribose were synthesized and purified (Kistemaker et al., 2015).

Plasmids for cellular assays

The mammalian expression constructs of N-terminally fluorescent tagged bait and prey were generated by cloning of the relevant DNA sequences in the fluorescently tagged LacR containing mammalian expression vectors and without LacR containing mammalian expression vectors, respectively (Table S3). All plasmids were generated following usual cloning procedure and were validated by Sanger sequencing.

Cell culture and transfections

For all the F2H assays, human U2OS cells harboring the stably integrated lacO (256x) array was used (Soutoglou and Misteli, 2008). Cells were cultured in DMEM (Sigma) supplemented with 10% (v/v) FBS (GIBCO), 1 mM sodium pyruvate (Sigma), 2 mM L-glutamine (Sigma), 100 U ml^{-1} penicillin, 100 mg ml^{-1} streptomycin (Sigma) and 200 mg ml^{-1} hygromycin B (Sigma) with 5% CO_2 at 37°C in a humidified environment.

Microscopy experiments

All microscopy experiments were performed in borosilicate 8-well Lab-Teks (Thermo Scientific). A Zeiss AxioObserver Z1 confocal spinning disk microscope was used for imaging. Images were acquired with an AxioCam HRm CCD/EMCCD/cCMOS camera (Zeiss) using Zeiss C-Apochromat 40x/1.2 water immersion and 63x/oil immersion objective lenses. Cells were imaged 24 hr post-transfection.

UV-laser micro-irradiation assays

Live cells were imaged on a AxioObserver Z1 confocal spinning-disk microscope equipped with an AxioCam HRm CCD camera (Zeiss) through a Zeiss 40x/water objective lens. For laser micro-irradiation, a 355-nm-wavelength diode-pumped solid-state pulsed laser (DPSL-355/14, Rapp OptoElectronics) was used. DNA damage was induced by focusing the 355 nm pulsed laser in the nucleus either as a small circle or as a line through the entire diameter of the nucleus. Cells were imaged at room temperature in a CO₂ independent medium (1x; GIBCO).

PARP1 inhibitor treatment and H₂O₂ treatments

For F2H and laser micro-irradiation experiments, cells were treated with 30 μM of the PARP1 inhibitor AG14361 (Selleck Chemicals) for at least 60 min, for the LacO array de-compaction experiments with PARP1 inhibitor cells were treated with the inhibitor for at least 24 hr before imaging (at the time of transfection itself). Cells were imaged immediately after H₂O₂ treatment, which was diluted to 1:2000 from stock H₂O₂ solution (30% v/v) in the imaging medium.

Fluorescence two-hybrid (F2H) assay

F2H assays were performed as previously described (Bowman et al., 2016; Czarna et al., 2013). Briefly, assays were performed in borosilicate 8-well Lab-Teks (Thermo Scientific). The fluorescently tagged bait and prey proteins were transiently co-transfected, using Xfect reagent (Clontech) according to the manufacturer's recommendations. Cells were imaged at 25°C in a CO₂-independent imaging medium (GIBCO) supplemented with 10% (v/v) FBS (GIBCO), 1 mM Sodium pyruvate (Sigma), 2 mM L-glutamine (Sigma), 100 U ml⁻¹ penicillin, 100 mg ml⁻¹ streptomycin (Sigma). Image analysis was manually performed with FIJI (ImageJ) image analysis software (Schindelin et al., 2012). Briefly, the average fluorescence intensity of the prey (mEGFP fusion) over the LacO array was divided by the average fluorescence intensity in the nucleoplasm to get the enrichment ratio. The fluorescent bait spot was identified by the localization of anchored bait protein (Fluorescent LacI fusion) on the LacO array. Only the cells that show a clearly discernible Lac repressor spot in the nuclei were chosen for analysis. The fluorescence intensity of the prey on the array and outside the array (nucleoplasm) was calculated, background subtracted and plotted as the ratio of the two (as enrichment ratio).

Co-immunoprecipitation assays

To test whether ALC1 oligomerizes *in vivo*, EGFP- and mCherry-tagged ALC1, EGFP-SPT16 and SSRP1-mCherry (positive control) or EGFP and mCherry-ALC1 (negative control) were expressed in U2OS cells for 24 hr and treated with 30 μM AG14361 PARP inhibitor (Selleckchem) for 1 hr before lysis. GFP-tagged proteins were immunoprecipitated using GFP-trap (Chromotek) following manufacturer's instructions. AG14361 (30 μM) was further supplied during all incubation steps. Interaction between GFP- and mCherry-tagged proteins was examined by western blotting using GFP- (Eurogentec) and mCherry-antibodies (Novus Bio, NBP2-25157), respectively.

LacO array remodelling assays

The LacO array de-compaction assays was performed in the same way as the F2H assay, with the only difference in the analysis of the results, wherein the area covered by the Lac repressor fusion protein on the LacO array was measured and normalized by the total area of the same nuclei, which was then plotted as % Nucleus area. Only the cells, which show a clearly discernible Lac Repressor decompacted spot in the nuclei, were chosen for analysis.

V5-Macrodomein pull-down assays

For V5-Macrodomein pull-down assays V5-tagged Macrodomein (615-876) of ALC1 was used as bait on the anti-V5 Agarose affinity gel beads in 1x PBS buffer. Other indicated proteins (like ATPase domain, PARP1) and ligands (DNA, NAD⁺, ADP-ribose ligands, PolyA-DNA, PARP1 inhibitor) were added to the pre-incubated V5-macrodomein bound beads as indicated and the complex was incubated on room temperature for 30 min before centrifugation followed by 3 washes in PBS and was finally boiled in Laemmli buffer before running on a 4%–12% gradient SDS-PAGE gel.

Fluorescence correlation spectroscopy (FCS)

FCS experiments were performed on a Leica SP8 confocal microscope equipped with a Plan APO 63x/1.2 N.A. water immersion objective. GFP fluorescence was excited with a 488 nm laser and emission selected by a bandpass filter at 500–550 nm. Laser power used for FCS measurements was adjusted to minimize photobleaching. Pinhole was set to one Airy unit. Single photons were detected and counted using a single photon avalanche diode and a PicoHarp module from PicoQuant. To estimate the residence time of GFP-tagged constructs in the focal volume, autocorrelation curves were fitted with a one-species model assuming pure diffusion and neglecting the contribution of the photo-physics of GFP using the Fluctuation Analyzer 4G software (Wachsmuth et al., 2015).

Thermal shift assays

Fluorescence-based thermal shift assays were conducted on an Applied Biosystems 7500 fast real-time PCR system. 2 μM protein solutions supplemented with 5 × SYPRO Orange in the absence or presence of 6.5 μM tri-ADP-ribose or 19.5 μM ADP-ribose were

heated in 25mM Na-PO₄ (pH 7.0), 0.25M NaCl and 1mM DTT from 5°C to 95°C at a ramp rate of 1%. The assays were conducted in MicroAmp Fast 96-well reaction plates sealed with MicroAmp Optical Adhesive Film (Applied Biosystems). The fluorescence measurements at 554nm were normalized to the lowest value before the transition and the maximum fluorescence.

Chromatin remodeler ATPase assays

The ALC1 catalyzed ATP hydrolysis was performed in 384-well using a malachite green assay (Greiner Bio One). For this, 1μM recombinant (near) full length ALC1 protein (residues 31-878) or the constitutively active ATPase fragment (residues 31-674) were pre-incubated for 10 min at RT with 1 μM dsDNA (75 bp with 4 nt overhangs on both ends; see [Key Resources Table](#) for their sequence) in 25 mM Bis-Tris pH 7.0, 0.15 M NaCl, 1 mM MgCl₂, 50 μM ZnCl₂, 1 mM DTT and 5% (w/v) glycerol in the absence or presence of ligand (5 μM tri-ADP-ribose or 15μM monomeric ADP-ribose). In case of the constitutively active ATPase fragment, the reactions were additionally performed in the absence or presence of 2.5μM ALC1 macrodomain (residues 636-878). The reactions were started by the addition of 1 μL 1 mM ATP, giving a final volume of 10 μL. After 5 min at RT, the reactions were stopped by the addition of 20 μL Biomol Green (Enzo Life Sciences), and the color was allowed to develop for 15 min. The absorbance of the resulting phosphomolybdate complex was read at 640 nm on a Tecan Infinite M1000 Pro microplate reader.

Analytical size exclusion chromatography (aSEC)

Globular proteins of known molecular weight from the Gel Filtration Calibration kit (GE Healthcare) and a Gel Filtration Standard (BioRad) were used to calibrate a Superdex 200 Increase 10/300 column (GE Healthcare). Blue dextran was used to determine the column void volume. Protein elution volumes were measured by monitoring the absorption at 280 nm. The elution volumes were used to calculate the partition coefficient (K_{av}) with the formula $K_{av} = (V_e - V_0)/(V_t - V_0)$ where V_e is the elution volume, V_0 is the void volume, and V_t is the total volume of the column. The apparent molecular weight was then derived from the inverse logarithm of the partition coefficient. All aSEC experiments were performed in 25 mM Na-PO₄ (pH 7.0), 0.25 M NaCl and 1 mM DTT.

QUANTIFICATION AND STATISTICAL ANALYSIS

For *in vitro* ATPase and Thermofluor assays

The experiments were performed three times as technical triplicates. The blank-subtracted values were normalized to the respective mean ATPase activity in the absence of the macrodomain or to the mean activity of the full-length protein. Blank values were determined by measuring the absorbance under identical conditions without ATP. All Thermofluor experiments were performed multiple times as technical replicates.

For the F2H and lacO array assays

The Wilcoxon matched-pairs signed rank test was used since normal distribution could not be assumed to determine the statistical significance in F2H and LacO decompaction assays (biological replicates $n \geq 2$). The column in the graph represents the *Mean*, error bar represents the SEM and number of cells is ≥ 20 . Actual p values are indicated in the figures. GraphPad Prism software was used to plot and analyze the data.

For HDX-MS measurements

For statistical analysis of the HDX-MS data, the standard deviation of measured deuterium content in each peptide based on technical replicates ($n = 2$ or $n = 3$) and using all relevant charge states. A significance threshold for differences in HDX between individual protein states was set to 0.52 D, corresponding to the 98.75% confidence interval (CI) calculated using technical replicates ($n = 3$) data from a single time-point for full-length ALC1 in the presence or absence of tri-ADP-ribose. The CI was calculated according to:

$$CI = \bar{x} \pm t * \frac{\sigma}{\sqrt{n}}$$

where x is the average difference in deuterium content assuming a zero-centered distribution ($x = 0$), t is 8.86 for the 98.75% CI with 2 degrees of freedom, σ is the average propagated standard deviation of differences in deuterium content for all peptide segments, and n is the number of replicate samples ($n = 3$).