Nuclear condensates of the Polycomb protein chromobox 2 (CBX2) assemble through phase separation

Received for publication, November 7, 2018, and in revised form, November 28, 2018. Published, Papers in Press, December 4, 2018, DOI 10.1074/jbc.RA118.006620

Roubina Tatavosian‡, Samantha Kent‡1, Kyle Brown‡1, Tingting Yao‡, Huy Nguyen Duc‡, Thao Ngoc Huynh‡, Chao Yu Zhen‡, Brian Ma‡, Haobin Wang‡, and @ Xiaojun Ren‡2

From the ‡Department of Chemistry, University of Colorado, Denver, Colorado 80217-3364 and §Department of Biochemistry and Molecular Biology, Colorado State University, Fort Collins, Colorado 80523

Edited by Ronald C. Wek

Polycomb group (PcG) proteins repress master regulators of development and differentiation through organization of chromatin structure. Mutation and dysregulation of PcG genes cause developmental defects and cancer. PcG proteins form condensates in the cell nucleus, and these condensates are the physical sites of PcG-targeted gene silencing via formation of facultative heterochromatin. However, the physiochemical principles underlying the formation of PcG condensates remain unknown, and their determination could shed light on how these condensates compact chromatin. Using fluorescence live-cell imaging, we observed that the Polycomb repressive complex 1 (PRC1) protein chromobox 2 (CBX2), a member of the CBX protein family, undergoes phase separation to form condensates and that the CBX2 condensates exhibit liquid-like properties. Using site-directed mutagenesis, we demonstrated that the conserved residues of CBX2 within the intrinsically disordered region (IDR), which is the region for compaction of chromatin in vitro, promote the condensate formation both in vitro and in vivo. We showed that the CBX2 condensates concentrate DNA and nucleosomes. Using genetic engineering, we report that trimethylation of Lys-27 at histone H3 (H3K27me3), which is the catalytic product of Polycomb repressive complex 2 (PRC2), one complex of Polycomb group (PcG) proteins (3, 4), H3K27me3 is the binding site for PRC1 (another complex of PcG proteins) that can compact chromatin, forming particular chromatin compartments (3, 4). The repressed, compacted chromatin domains are preserved during cell division (3, 4). These functions and behaviors of facultative heterochromatin have raised several fundamental questions. For example, what are the properties of facultative heterochromatin? How is chromatin compaction achieved? How do PcG proteins contribute to establishing facultative heterochromatin compartments? These questions are essential for understanding how facultative heterochromatin functions and preserves cell identity.

Biochemical and genetic studies of PcG proteins have begun addressing some of the questions raised above. For example, in terms of understanding the properties of facultative heterochromatin, it has been shown that PcG proteins directly regulate chromatin structure and chemically modify the histones of facultative heterochromatin (7). PRC2 methylates histone H3 on lysine 27 by the catalytic subunit enhancer of zeste homolog 2 (8–12). Embryonic ectoderm development (EED), one core subunit of PRC2, binds H3K27me3, which allosterically stimu-

The genome in eukaryotic cells can be broadly classified as euchromatin (active transcription) and heterochromatin (repression and silencing) (1, 2). Heterochromatin can be further described broadly as constitutive and facultative heterochromatin (3–5). Constitutive heterochromatin is observed at and near centromeres and telomeres (5). Facultative heterochromatin is found at a specific subset of genes encoding regulators of development and differentiation (3, 4). Another example of facultative heterochromatin is X chromosome inactivation in female mammals (6). Facultative heterochromatin represses gene expression in part through compacting chromatin to reduce the accessibility of DNA (3–5). Facultative heterochromatin is decorated by the trimethylation of Lys-27 at histone H3 (H3K27me3), which is the catalytic product of Polycomb repressive complex 2 (PRC2), one complex of Polycomb group (PcG) proteins (3, 4). H3K27me3 is the binding site for PRC1 (another complex of PcG proteins) that can compact chromatin, forming particular chromatin compartments (3, 4). The repressed, compacted chromatin domains are preserved during cell division (3, 4). These functions and behaviors of facultative heterochromatin have raised several fundamental questions. For example, what are the properties of facultative heterochromatin? How is chromatin compaction achieved? How do PcG proteins contribute to establishing facultative heterochromatin compartments? These questions are essential for understanding how facultative heterochromatin functions and preserves cell identity.

Biochemical and genetic studies of PcG proteins have begun addressing some of the questions raised above. For example, in terms of understanding the properties of facultative heterochromatin, it has been shown that PcG proteins directly regulate chromatin structure and chemically modify the histones of facultative heterochromatin (7). PRC2 methylates histone H3 on lysine 27 by the catalytic subunit enhancer of zeste homolog 2 (8–12). Embryonic ectoderm development (EED), one core subunit of PRC2, binds H3K27me3, which allosterically stimu-

The genome in eukaryotic cells can be broadly classified as euchromatin (active transcription) and heterochromatin (repression and silencing) (1, 2). Heterochromatin can be further described broadly as constitutive and facultative heterochromatin (3–5). Constitutive heterochromatin is observed at and near centromeres and telomeres (5). Facultative heterochromatin is found at a specific subset of genes encoding regulators of development and differentiation (3, 4). Another example of facultative heterochromatin is X chromosome inactivation in female mammals (6). Facultative heterochromatin represses gene expression in part through compacting chromatin to reduce the accessibility of DNA (3–5). Facultative heterochromatin is decorated by the trimethylation of Lys-27 at histone H3 (H3K27me3), which is the catalytic product of Polycomb repressive complex 2 (PRC2), one complex of Polycomb group (PcG) proteins (3, 4). H3K27me3 is the binding site for PRC1 (another complex of PcG proteins) that can compact chromatin, forming particular chromatin compartments (3, 4). The repressed, compacted chromatin domains are preserved during cell division (3, 4). These functions and behaviors of facultative heterochromatin have raised several fundamental questions. For example, what are the properties of facultative heterochromatin? How is chromatin compaction achieved? How do PcG proteins contribute to establishing facultative heterochromatin compartments? These questions are essential for understanding how facultative heterochromatin functions and preserves cell identity.

Biochemical and genetic studies of PcG proteins have begun addressing some of the questions raised above. For example, in terms of understanding the properties of facultative heterochromatin, it has been shown that PcG proteins directly regulate chromatin structure and chemically modify the histones of facultative heterochromatin (7). PRC2 methylates histone H3 on lysine 27 by the catalytic subunit enhancer of zeste homolog 2 (8–12). Embryonic ectoderm development (EED), one core subunit of PRC2, binds H3K27me3, which allosterically stimu-

This work was supported by NCI, National Institutes of Health Grant R03CA191943 (to X. R.), National Science Foundation Grant CHE-1500285 (to H. W.), NIGMS, National Institutes of Health Grant R01GM098401 (to T. Y.), and the Office of Research Services (ORS) at the University of Colorado Denver. The authors declare that they have no conflicts of interest with the contents of this article. The content is solely the responsibility of the authors and does not necessarily represent the official views of the National Institutes of Health.

@ Both authors contributed equally to this work.

2 To whom correspondence should be addressed. Tel.: 303-315-7641; Fax: 303-556-4776; E-mail: xiaojun.ren@ucdenver.edu.

3 The abbreviations used are: H3K27me3, trimethylation of Lys-27 at histone H3; CBX, chromobox; EED, embryonic ectoderm development; IDR, intrinsically disordered region; LLPS, liquid–liquid phase separation; PcG, Polycomb group; PHC, polyhomeotic homolog; PRC, Polycomb repressive complex; RING1B, ring finger protein 2; mES, mouse embryonic stem; HT, HaloTag; YFP, yellow fluorescent protein; FRAP, fluorescence recovery after photobleaching; GST, glutathione S-transferase; DIC, differential interference contrast; ATH, AT-hook; ATHL, ATH-like; SRR, serine-rich region; CD, chromodomains; DMEM, Dulbecco’s modified Eagle’s medium; FBS, fetal bovine serum; TMR, tetramethylrhodamine.

1451

J. Biol. Chem. (2019) 294(5) 1451–1463

October 8, 2020

Downloaded From http://www.jbc.org/ at AIRA LIBRARY on September 8, 2020
lates PRC2 activity, and this stimulation can increase the local H3K27me3 level (13, 14). H3K27me3 provides a binding site for chromobox 7 (CBX7) and CBX8 of the CBX family proteins (CBX2/4/6/7/8) (15), which recruit canonical CBX7–PRC1 and CBX8–PRC1 to chromatin. PRC1 complexes can ubiquitinate histone H2A at lysine 119 by the catalytic subunit ring finger protein 2 (RING1B) (16, 17), which can stimulate PRC2 activity (18, 19). Thus, a feedback loop between PRC1 and PRC2 is created to reinforce the epigenetic modifications of facultative heterochromatin. In terms of understanding compaction, it has been demonstrated that the compaction function of PRC1 in mammals is facilitated by CBX2 of the CBX protein family (20, 21). In Drosophila, the protein posterior sex combs exerts chromatin compaction (22, 23). Mutation of the CBX2 residues that are required for the compaction leads to homeotic transformations that are similar to those observed with PcG loss-of-function mutations (21). In terms of understanding facultative heterochromatin compartments, studies have demonstrated that, in the cell nucleus, PcG proteins form microscopically visible condensates (24–28). PcG condensates function as specific nuclear compartments for target gene silencing (24–28). Overall, these biochemical and genetic studies suggest that PRC1 and PRC2 coordinate to establish and maintain facultative heterochromatin. Despite these exciting advances, the fundamental physicochemical principles that underpin how PcG proteins establish, maintain, and regulate facultative heterochromatin remain incompletely understood. Understanding these questions is critical for appreciating how PcG proteins control development and differentiation.

Spatial and temporal compartmentalization of intracellular components into organelles in eukaryotic cells is a generic theme for organizing biochemical reactions (29–34). These organelles can be membrane-bound or membraneless. A large number of membraneless compartments, including the nucleolus, stress granules, Cajal bodies, promyelocytic leukemia nuclear bodies, and others, are condensates formed by condensation of cellular components through liquid–liquid phase separation (LLPS) (29–33). The forces that drive LLPS are multivalent interactions among proteins and other macromolecular polymers such as RNA and DNA (29–31). Phase-separated condensates have been shown to be involved in multiple cellular processes and functions (29–31). Over the past year, phase-separated membraneless condensates have been suggested to be implicated in transcriptional activation and repression (35–41). A phase-separated model has been emerging to explain transcriptional activation: transcription factors and coactivators phase separate to form condensates that interact with condensates of RNA polymerase II to efficiently activate transcription (35–39). Phase-separated condensates also function in transcriptional repression. Heterochromatin protein 1α phase separates to form condensates that compartmentalize constitutive heterochromatin (40, 41). Facultative heterochromatin represents one major class of chromatin structures. Whether the PcG proteins that are responsible for the formation of facultative heterochromatin phase separate to form condensates remains unknown.

Here, we provide the first experimental evidence that the PRC1 protein CBX2 phase separates to form condensates that can concentrate DNA and nucleosomes. The conserved residues within the CBX2 intrinsically disordered region (IDR) are required for the CBX2 condensate formation both in vitro and in vivo. We show that H3K27me3 contributes little to the CBX2 condensate formation. We demonstrate that CBX2 determines the condensate formation of CBX2–PRC1; however, the CBX2–PRC1 subunits are not required for the formation of CBX2 condensates. Thus, we provide a general experimental framework that can explain how PcG condensates assemble and organize PcG-bound chromatin. Our results provide a starting point for further exploring how phase separation facilitates efficient and specific control of transcription.

**Results**

**CBX2 forms condensates in cells**

We first investigated whether CBX2 forms condensates in living cells. We integrated YFP-Cbx2 and HaloTag (HT)-Cbx2 into the genome of PKG12.1 mouse embryonic stem (wildtype WT) mES cells, respectively. The expression of these fusion genes is under the control of an inducible, tetracycline response element–tight promoter. To observe the cellular distribution of HT-CBX2, we labeled the fusion protein by HaloTag® TMR ligand in living cells. Both YFP-CBX2 and HT-CBX2 formed condensates in living WT mES cells (Fig. 1, a and b), which is consistent with the previous observations reporting that both exogenous and endogenous CBX2 forms condensates (42–44). About half of cells contained microscopically visible CBX2 condensates. The average area of condensates was 0.19 μm² (~250 nm in radius), and their fluorescence intensity was ~1.5-fold higher than the average intensity. It is possible that the formation of CBX2 fusion condensates in mES cells is due to their overexpression. To resolve this possibility, we integrated YFP-CBX2 and HT-CBX2 into the genome of Cbx2−/− mES cells. Under the basal expression, the protein level of YFP-CBX2 was similar to that of the endogenous protein CBX2 (45). Without doxycycline induction, the distribution of YFP-CBX2 and HT-CBX2 in Cbx2−/− mES cells was similar to that in WT mES cells (Fig. 1, a and b). These data indicate that CBX2 forms microscopically visible condensates in living cells.

CBX2 forms a stable PRC1 complex (CBX2–PRC1), including polyhomoetic homolog 1 (PHC1) and RING1B (Fig. 1c) (7, 46), so we investigated whether YFP-CBX2 condensates colocalize with CBX2–PRC1 subunits. We stained endogenous RING1B and PHC1 as well as YFP-CBX2, RING1B and PHC1 formed condensates in cells (Fig. 1d), consistent with the previous reports (25, 43, 47, 48). YFP-CBX2 condensates colocalize with condensates of RING1B and PHC1 (Fig. 1d). Because H3K27me3 marks PcG-targeted genes, we investigated whether CBX2 condensates colocalize with H3K27me3. Immunofluorescence of H3K27me3 and YFP-CBX2 showed that CBX2 condensates colocalize with chromatin with the dense H3K27me3 mark (Fig. 1d), suggesting that PcG-targeted genes are recruited to CBX2 condensates or vice versa. Thus, our results show that CBX2 condensates colocalize with CBX2–PRC1 subunits and H3K27me3-marked chromatin regions.

To determine whether the condensate formation of RING1B and PHC1 depends on CBX2, we first stained RING1B and PHC1

---

*Polycomb CBX2 condensates by phase separation*

---

1.452 J. Biol. Chem. (2019) 294(5) 1451–1463

Downloaded From: http://www.jbc.org/ at ATRARIA LIBR-ESS on September 8, 2020
in Cbx2<sup>−/−</sup> mES cells. In contrast to WT mES cells, RING1B and PHC1 exhibited a granular distribution in Cbx2<sup>−/−</sup> mES cells, and their large condensates disappeared (Fig. 1e). We then stained RING1B and PHC1 in YFP-Cbx2/Cbx2<sup>−/−</sup> mES cells. RING1B and PHC1 formed condensates that colocalize with YFP-CBX2 condensates (Fig. 1e). Our results indicate that the formation of RING1B and PHC1 condensates depends on CBX2.

Next, we interrogated whether CBX2 condensates exhibit liquid-like features that are characterized with rapid exchange kinetics, which can be studied by measuring the recovery rate using fluorescence recovery after photobleaching (FRAP). We performed FRAP experiments on condensates of YFP-CBX2 stably expressed in mES cells. FRAP showed that 80% of YFP-CBX2 within condensates is recovered within 3 min (Fig. 1, f and g), consistent with our previous reports (42, 49). These results indicate that CBX2 within condensates dynamically exchanges with surrounding environments and has liquid-like properties in cells.

If CBX2 condensates were liquid-like, reducing the concentration of YFP-CBX2 would dissolve these condensates. We lysed cells stably expressing YFP-CBX2 in lysis buffer to cause a local decrease of YFP-CBX2 through diffusion. We did not detect YFP-CBX2 condensates in the lysate. We expected that formaldehyde cross-linking would preserve CBX2 condensates.
After sonication and centrifugation, YFP-CBX2 condensates would be within the pellets. To test this speculation, prior to lysis, we cross-linked cells with formaldehyde and prepared lysates from the cross-linked cells. These lysates were subjected to sonication. Using fluorescence microscopy, we observed CBX2 condensates within the sonicated lysate (Fig. 1h). After centrifugation, we did not observe CBX2 condensates in the supernatant but instead observed them in the resuspended pellets (Fig. 1h). These data further support that CBX2 condensates form in the cells and possess liquid-like properties.

**CBX2 phase separates to form condensates in vitro**

Proteins can undergo LLPS through IDRs, leading to formation of condensate in aqueous solution (29–31). We analyzed the properties of the primary sequence of CBX2 and found that 59% of the CBX2 sequence is intrinsically disordered as predicted by MobiDB 3 (50). CBX2 is the protein that condenses chromatin through the highly basic residues within the IDR (20, 21). Thus, we reasoned that CBX2 could form condensates in vitro through LLPS. To test this hypothesis, we expressed and purified recombinant GST-CBX2-FLAG (GST-CBX2) from *Escherichia coli* at high salt concentration or in the presence of glutathione (GSH) (Fig. 2b). We found that both high salt and GSH prevent aggregation of CBX2. We placed the tags at the respective N-terminal and C-terminal ends of CBX2 to remove truncated CBX2 during the purification. We dialedyzed the high salt of GST-CBX2 fusion to 140 mM NaCl at 4 °C overnight and transferred 10 μl of sample to a coverslip. After condensates settled down on the surface of the coverslip, we performed differential interference contrast (DIC) imaging and observed CBX2 condensates with a size of a few hundred nanometers (Fig. 2c). Next, we generated CBX2-FLAG (CBX2) without GST fusion (Fig. 2b). CBX2 also underwent LLPS to form condensates (Fig. 2c), suggesting that the condensate formation is not driven by GST. To determine the identity of these condensates, we produced GST-YFP-CBX2-FLAG (GST-YFP-CBX2) (Fig. 2b). Fluorescence imaging showed that GST-YFP-CBX2 assembles into condensates (Fig. 2c). Under the same conditions, GST and BSA did not form condensates (Fig. 2c).

**Figure 2. CBX2 phase separates to form condensates in vitro.** a, CBX2 is an intrinsically disordered protein predicted by MobiDB 3 (50). A PONDR score greater than 0.5 indicates intrinsically disordered regions. 59% of CBX2 sequence is intrinsically disordered. b, SDS-PAGE analysis of recombinant CBX2 proteins. Left, recombinant GST-CBX2-FLAG (GST-CBX2). Middle, recombinant CBX2-FLAG (CBX2). Right, recombinant GST-YFP-CBX2-FLAG (GST-YFP-CBX2). The molecular mass ladder is shown at the left of the gel image. c, representative DIC images of GST-CBX2 (4.8 μM) and CBX2 (4.8 μM) condensates as well as the control BSA (10 μM) and GST (10 μM) on the surface of a coverslip. A representative epifluorescence image of GST-YFP-CBX2 (4.8 μM) condensates is shown. Scale bars, 2.0 μm. d, dependence of the formation of CBX2 condensates on its concentrations shown above the image. Representative DIC images of condensates on the surface of coverslip are shown. Scale bars, 2.0 μm. e–g, increasing concentrations of NaCl, Triton X-100, and hexanediol dissolve CBX2 condensates. We incubated CBX2 (4.8 μM) condensates with the indicated concentrations of NaCl, Triton X-100, and hexanediol for 30 min on ice. The mixture was loaded to a coverslip for imaging. We counted condensates using ImageJ. The data were from at least 10 frames for each sample. Error bars represent S.D. p values were calculated based on Student’s t test. h, representative epifluorescence images of WT mES cells stably expressing YFP-CBX2 before and after treatment with 10% hexanediol for 5.0 min. Scale bar, 10 μm.
found that the phase separation of CBX2 is concentration-dependent (Fig. 2d). Thus, our results demonstrate that CBX2 can undergo LLPS in form condensates in vitro.

Classical polymer theory predicts that polymers undergo LLPS through multivalency-driven interactions such as cation–π, electrostatic, dipolar, and hydrophobic interactions (29–31). Thus, we investigated whether NaCl and Triton X-100 dissolve CBX2 condensates. We found that treatment of CBX2 condensates with increasing concentrations of NaCl and Triton X-100 causes a reduction in the number of CBX2 condensates (Fig. 2, e and f). Hexanediol is known to dissolve liquid-like condensates (35, 37), possibly through disruption of hydrophobic interactions. We found that treatment of CBX2 condensates with hexanediol results in a reduction in the number of condensates in vitro (Fig. 2g). Treatment of mES cells expressing YFP-CBX2 with hexanediol caused mild effects on the formation of CBX2 condensates (Fig. 2h), suggesting that other noncovalent interactions are also involved in the phase separation of CBX2. These results indicate that multivalent interactions contribute to the LLPS of CBX2.

**CBX2 condensates concentrate DNA and nucleosomes**

One of the characteristic properties of cellular compartmentalization is that compartments can increase the local concentration of resident biochemical molecules (29–34). PcG condensates are the physical sites of PcG-mediated silencing, involving organization of the PcG-bound chromatin (24–26). Given that CBX2 can directly bind DNA (51), we investigated whether CBX2 condensates can concentrate DNA. We labeled 24-bp dsDNA with fluorescent dye and mixed them with CBX2. Dye-labeled DNA did not form condensates; however, in the presence of CBX2, DNA was concentrated, and CBX2 condensates colocalized with the concentrated DNA (Fig. 3a). Previous studies have shown that CBX2 can compact chromatin on its own (20), so we tested whether CBX2 can concentrate core nucleosome particles. We prepared dye-labeled nucleosomes and mixed them with CBX2. After dialysis, we observed that CBX2 condensates colocalize with the concentrated dye-labeled nucleosomes (Fig. 3a). Under the same conditions, CBX2 condensates could not enrich the dye (Fig. 3a). Thus, these results suggest that CBX2 condensates can concentrate DNA and nucleosomes in vitro.

Given that PcG condensates are the repressive compartments for PcG-targeted genes, we should be able to detect DNA within CBX2 condensates isolated from cells. We cross-linked cells stably expressing YFP-CBX2 with formaldehyde. After sonication and centrifugation, we resuspended the pellet and stained DNA with Hoechst. Fluorescence images showed that YFP-CBX2 condensates contain concentrated DNA labeled by Hoechst (Fig. 3b). Our data indicate that CBX2 condensates can enrich chromatin/DNA within cells.

**Conserved residues within the IDR of CBX2 contribute to LLPS in vitro and in living cells**

The driving forces of phase separation of proteins containing IDRs are noncovalent interactions, particularly cation–π and electrostatic interactions (29–34). The cation–π interactions occur between aromatic residues and Lys or Arg residues (52–57). We found that CBX2 contains a high content of Lys and Arg; however, the frequency of aromatic residues is lower than their respective average frequency in vertebrate proteins. Because proteins whose phase separations are promoted by the cation–π interactions contain a high content of aromatic residues (52), we expect that cation–π interactions may not be the major driving forces for the phase separation of CBX2. Instead, we found that, in CBX2, many positively and negatively charged residues are grouped into a series of clusters across the sequence (Fig. 4). Alternative clusters of positive and negative charges can form an electrostatic interaction, which is one of the major driving forces that promote the phase separation of IDR-containing proteins (41, 56, 58–62). It is interesting to note that the three conserved regions, AT-hook (ATH), AT-like 1 (ATHL1), and ATHL2 (63), are positively charged clusters (Fig. 4, a and b). We substituted residues PRG (amino acids 77–79) for AAA to generate CBX2ATH, PRG (amino acids 134–136) for AAA to generate CBX2ATHL1, and RKKRGKR (amino acids 161–167) for AAAAGAA to generate CBX2ATHL2 (Fig. 4b). The net positive charge per residue of the mutated regions of CBX2ATH and CBX2ATHL1 was slightly reduced compared with CBX2, whereas the net positive charge per residue of the mutated region of CBX2ATHL2 was completely eliminated (Fig. 4b). We generated these mutant proteins and compared their
Polycomb CBX2 condensates by phase separation

Figure 4. Conserved residues promote the LLPS of CBX2. a, schematic representation of CBX2. The IDR was predicted by MobidiB 3 (50). Conserved regions include CD, ATH, SRR, ATHL1, ATHL2, and chromobox (Cbox) (63). b, charge distribution of CBX2 and its variants calculated by EMBOSS charge. The net charge per residue was averaged over a sliding window of eight residues. The three conserved regions, ATH, ATHL1, and ATHL2, are positively charged (top panel). The three conserved regions were mutated to generate CBX2ATHL1 and CBX2ATHL2 (bottom panel). The red asterisks indicate the three conserved regions that are mutated. c, representative DIC images of condensates of CBX2 and its variants (CBX2ATHL1, CBX2ATHL2, and CBX2SRR) on the surface of coverslips. The formation of condensates was carried out at a concentration of 4.8 μM for both CBX2 and its variants. d, quantification of condensates per frame from e. The data were from at least 10 frames for each sample. Error bars represent S.D. The p value was calculated based on Student’s t-test. e, representative epifluorescence images of WT mES cells stably expressing HT-CBX2ATH, HT-CBX2ATHL1, HT-CBX2ATHL2, and HT-CBX2SRR. These CBX2 mutants were labeled with HaloTag TMR ligand and then performed live-cell imaging of these mutants (Fig. 4e). To compare their distribution, we imaged cells with similar fluorescence intensity. Quantitative analysis showed that the size and the number of condensates of these CBX2 mutants are significantly reduced compared with WT CBX2 (Fig. 4, f and g). We also noted that the size and the number of condensates of CBX2ATHL1 and CBX2SRR were slightly smaller than for CBX2ATH and CBX2ATHL2 (Fig. 4, f and g), consistent with in vitro analysis. Thus, our data demonstrate that the conserved residues within the IDR that are critical for the LLPS of CBX2 in vitro are also critical for the formation of CBX2 condensates in vivo.

H3K27me3 has minimal effects on the formation of CBX2 condensates

The PRC2-catalyzed product H3K27me3 is the marker of Pcg-targeted chromatin (7). H3K27me3 has been hypothesized to be the mark for recruiting CBX–PRC1 to chromatin (Fig. 5a) (64). Thus, we asked whether H3K27me3 affects the formation of CBX2 condensates in vitro. To this end, we integrated HT-CBX2 into the genome of Eed−/− mES cells. EED is the core component of PRC2, and Eed knockout results in a complete loss of H3K27me3 (15). Live-cell imaging of HT-CBX2 labeled with HaloTag TMR ligand showed that HT-CBX2 forms condensates in Eed−/− mES cells (Fig. 5c). Quantitative analysis indicated that the size and the number of CBX2 condensates in Eed−/− mES cells are not significantly different from those in WT mES cells (Fig. 5, d and e). Previous studies have shown that the aromatic cage, consisting of three aromatic residues, of the chromodomain (CD) of CBX2 is critical for the H3K27me3 binding in vitro (65). We mutated the cage residue Phe-12 of CBX2 to Ala (CBX2F12A) (Fig. 5b). We stably expressed HT-CBX2F12A in WT mES cells. Live-cell imaging showed that HT-CBX2F12A forms condensates (Fig. 5c). The size and the number of HT-CBX2F12A were similar to HT-CBX2 (Fig. 5, d and e). These data suggest that H3K27me3 contributes little to the formation of CBX2 condensates in living cells.

Because the CD of CBX2 is the binding domain for H3K27me3 in vitro (65, 66) (Fig. 5a), we investigated the effects of the CD on
the formation of CBX2 condensates in living cells. We fused CD with HT, generating HT-CD\textsuperscript{CBX2} (Fig. 5b). We also deleted CD to generate \textit{HT-CD\textsuperscript{CBX2}} (Fig. 5b). We established mES cells that stably express \textit{HT-CD\textsuperscript{CBX2}} and \textit{HT-CD\textsuperscript{CBX2}}, respectively. HT-CD\textsuperscript{CBX2} did not form condensates in living cells (Fig. 5c). HT-CD\textsuperscript{CBX2} phase separated to form condensates (Fig. 5c); however, their size and number were significantly reduced compared with HT-CBX2 (Fig. 5d and e). Given that ATH is adjacent to CD and binds DNA (51) (Fig. 5b), we deleted both CD and ATH to generate \textit{HT-CBX2-ATH} (Fig. 5b), which was then stably integrated into the genome of mES cells. HT-CBX2\textsuperscript{CD-ATH} did not phase separate to form condensates within living cells (Fig. 5c). Thus, these results indicate that the interactions of H3K27me3 and CBX2 are not required for the formation of CBX2 condensates; however, the amino acid residues within CD are required for the condensate formation.

Depletion of CBX2–PRC1 subunits does not prevent the formation of CBX2 condensates

CBX2 phase separates on its own \textit{in vitro}, so we speculated that removal of CBX2–PRC1 subunits would not prevent the formation of CBX2 condensates \textit{in vivo}. To address this, we integrated HT-CBX2 into the genome of \textit{Ring1a\textsuperscript{+/−}/Ring1b\textsuperscript{+/−}, Rosa26::CreERT2 and Bmi1\textsuperscript{−/−}/Mel18\textsuperscript{−/−}} mES cells, respectively. Ring1b in \textit{Ring1a\textsuperscript{+/−}/Ring1b\textsuperscript{+/−}}, Rosa26::CreERT2 mES cells was depleted by administrating hydroxytamoxifen as described previously (15, 49). Live-cell imaging showed that depletion of \textit{Ring1a} and \textit{Ring1b} or \textit{Mel18} does not disperse CBX2 condensates (Fig. 6a). Instead, we found that CBX2 condensates in these double-knockout mES cells are typically more numerous and larger compared with WT mES cells (Fig. 6a). Some CBX2 condensates in the double-knockout mES cells were irregular instead of the usual droplet-like shape, suggesting that these CBX2–PRC1 subunits may influence the material properties of CBX2 condensates. Quantitative analysis demonstrated that both the size and the number of CBX2 condensates in the double-knockout mES cells are significantly larger than those in WT mES cells (Fig. 6, b and c). These data indicate that CBX2–PRC1 subunits contribute less to the formation of CBX2 condensates \textit{in vivo} but may regulate their material properties.

Discussion

Numerous studies have demonstrated that PcG proteins form microscopically visible condensates in primary and transformed cells, both in flies and mammals (24–28). These condensates have been shown to be the physical sites for repressing PcG-targeted genes (24–28). Consistent with these previous observations, we demonstrate that PcG protein CBX2 forms condensates in mES cells that colocalize with H3K27me3–dense chromatin regions and CBX2–PRC1 subunits. We further show that CBX2 can undergo LLPS \textit{in vitro} in the absence of other proteins. It is striking that CBX2 mutants that lack the capacity of LLPS \textit{in vitro} have a similar deficient ability to form condensates \textit{in vivo}. The strong correlation between \textit{in vitro} and \textit{in vivo} data indicates that CBX2 condensates in living cells form through LLPS. This is further supported by our following observations: CBX2 condensates in living cells exhibit rapid...
Polycomb CBX2 condensates by phase separation

exchange dynamics, a hallmark of liquid-like condensates, and our data show that CBX2 condensates concentrate DNA and nucleosomes. Previous studies have shown that CBX2 can compact chromatin (20, 21). Similar to heterochromatin protein 1α condensates compaction of heterochromatin (40, 41), we propose that CBX2 compacts chromatin by forming phase-separated condensates.

The phase behavior of proteins containing IDRs can be described by the theory of associative polymers (52, 67). Associative polymers phase separate through interactions between associative motifs called stickers that are separated from one another by spacers (52, 67). Spacers can impart the material properties of polymers and modulate the phase-separation ability of polymers (52, 67). The stickers can be residues that involve cation–π, electrostatic, hydrophobic, or dipolar interactions (29, 30, 52). In the case of CBX2, the stickers appear to be appositively charged clusters. Perturbation of these charged clusters reduces the phase separation of CBX2 both in vitro and in vivo, which is consistent with the notion that the phase separation of IDR-containing proteins can be promoted by interactions between blocks of appositively charged residues (41, 56, 58–62). The SRR also appears to be a sticker because substitution of the Ser residues of SRR with Ala prevents the phase separation of CBX2 both in vitro and in vivo. Because the content of aromatic residues is low and there is no apparent pattern for aromatic residues across the CBX2 sequence, we hypothesize that these aromatic residues are unlikely to be stickers. However, further experiments are required to test this hypothesis.

Our data suggest a scaffold–client model for the assembly of CBX2–PRC1 condensates (30, 68). Our results indicate that CBX2 is the scaffold, and the other subunits of CBX2–PRC1 are clients. Three lines of evidence support this model. First, the depletion of Ring1a and Ring1b or Bmi1 and Mel18 does not prevent the formation of CBX2 condensates in cells. Second, the depletion of Chx1 dissolves condensates of PHC1 and RING1B in cells. Finally, CBX2 can phase separate to form condensates in the absence of CBX2–PRC1 subunits in vitro. Previous studies have mapped the physical interactions between the subunits within the CBX–PRC1 complexes. It has been suggested that one of RING1A/RING1B, MEL18/BMI1, and PHC1/PHC2/PHC3 combines to form a stable heterotrimeric protein complex (69, 70). We expect that the trimeric protein complex could be recruited into CBX2 condensates through the C-terminal domain of RING1B interactions with the chromobox (Cbox) motif of CBX2 (71, 72). The absence of CBX2–PRC1 subunits, leading to irregular shapes of CBX2 condensates, suggests that the trimeric client has critical roles in regulating the material properties of CBX2–PRC1 condensates as well as the assembly of PcG condensates.

How do CBX2–PRC1 condensates compact and organize PcG-bound chromatin domains? There should be at least two kinds of organization of PcG-bound chromatin domains: condensing PcG-bound chromatin and establishing long-range interactions of distal PcG-bound regions (7). CBX2 is the PRC1 protein that is responsible for compaction of chromatin in vertebrates (20, 21). The highly charged positive residues within the IDR of CBX2 that are required for the compaction (20) are also required for the phase separation, suggesting that the phase separation is coupled to the chromatin compaction. The chromatin compaction by CBX2 depends on direct binding to nucleosomes but is independent of H3K27me3 (20). CBX4/6/7/8 proteins also have a high content of IDRs; however, unlike CBX2, they do not contain a high content of highly charged basic residues and cannot compact chromatin (20). We suggest that CBX4/6/7/8 may have less capacity to phase separate than CBX2. We hypothesize that CBX2–PRC1 condensates exert compaction of chromatin by binding DNA elements underlying PcG-targeted domains. Our data and previous observations support this hypothesis. Our data demonstrate that CBX2 condensates can concentrate DNA and core-unmodified nucleosomes and that H3K27me3 has minimal effects on the formation of CBX2–PRC1 condensates. Our previous live-cell single-molecule tracking showed that H3K27me3 has negligible effects on the chromatin-bound level of CBX2 (15). Previous studies demonstrated that CBX2 directly binds DNA with a $K_d$ value hundreds-fold smaller than its binding to H3K27me3 in vitro (51, 65, 66, 73) and that the compaction function is independent of histone tails (20, 21, 23). Thus, these data suggest a model in which CBX2–PRC1 condensates compact chromatin through CBX2 directly binding chromatin.

What are the molecular factors that bring distal PcG-bound regions into CBX2–PRC1 condensates for compaction? We propose a bridge model in which CBX7–PRC1 or CBX8–PRC1 acts as the molecular bridge that recruits H3K27me3-marked domains into CBX2–PRC1 condensates. CBX7–PRC1 and CBX8–PRC1 bind to H3K27me3-chromatin via CBX7 and CBX8 interactions with H3K27me3 and then recruit H3K27me3-chromatin into CBX2–PRC1 condensates through the polymerization ability of PHC proteins. The model is supported by our previous live-cell single-molecule tracking results that demonstrate that the removal of H3K27me3 greatly reduces the bound level of CBX7 and CBX8 (15) and by the fact that the sterile α motif domain of PHC proteins can form long filaments via head-to-tail intermolecular interactions (74, 75). Thus, our model suggests that the phase separation of CBX2–PRC1, CBX7–PRC1, CBX8–PRC1, and H3K27me3 coordinates to organize chromatin structure, thereby regulating gene activity. The model can explain why the presence of PHC protein, RING1B, and H3K27me3 is required for compaction of chromatin (25, 27, 28, 76, 77) and be tested in the future.

In summary, our results demonstrate, for the first time, that PcG condensates form through LLPS, and these condensates can concentrate nucleosomes and DNA. Our data show that the charged basic residues of CBX2 that are responsible for compaction of chromatin promote LLPS of CBX2. We further show that H3K27me3 has minimal effects on the formation of CBX2–PRC1 condensates. Our data suggest a scaffold–client model that underpins how CBX2–PRC1 condensates assemble and compact chromatin. Together, these results provide a starting point for conceptualizing the roles of PcG proteins in the assembly, structure, and functions of facultative heterochromatin.
Materials and methods

Cell culture

PGK12.1 mES cells (78) were provided by Dr. Neil Brockdorff (University of Oxford, UK). Cbx2−/− mES cells (79), Eed−/− mES cells (80), Ring1a−/−/Ring1bβ/β, Rosa26::CreERT2 mES cells (80), and Bmi1−/−/Mel18−/− mES cells (80) were provided by Dr. Haruhiko Koseki (RIKEN Center for Integrative Medical Sciences, Japan). To deplete Ring1b, Ring1a−/−/Ring1bβ/β; Rosa26::CreERT2 mES cells were treated with 4-hydroxytamoxifen (H7904, Sigma-Aldrich) for 2.0 days under a concentration of 1.0 μM as described previously (15, 49). Hereafter, we refer to Ring1a−/−/Ring1bβ/β; Rosa26::CreERT2 mES cells as Ring1a−/−/Ring1bβ/β−/− mES cells. Dr. Tom Kerppola (University of Michigan, Ann Arbor, MI) provided HEK293T cells. mES cells were grown in the mES cell medium (DMEM (D5796, Sigma-Aldrich) supplemented with 15% FBS, 0.1 mM nonessential amino acids (11140050, Life Technologies), 100 units/ml penicillin-streptomycin (15140-122, Life Technologies), 55 μM β-mercaptoethanol (21985-023, Life Technologies), 2 mM glutamine (G7513, Life Technologies), and 10° 3 units/ml leukemia inhibitor factor at 37 °C in 5% CO2. HEK293T cells were maintained in the HEK293T cell medium (DMEM (D5796, Sigma-Aldrich) supplemented with 15% FBS, 0.1 mM nonessential amino acids (11140050, Life Technologies), 100 units/ml penicillin-streptomycin, 2 mM glutamine, and 55 μM β-mercaptoethanol at 37 °C in 5% CO2.

Plasmids

The pTRIPZ (M1)-HT-Cbx2 plasmid harbors a puromycin resistance gene (15). To generate Cbx2 variants fused with HT, we replaced the Cbx2 sequence in the plasmid pTRIPZ (M1)-HT-Cbx2 with the Cbx2 variant sequence. We generated the following Cbx2 variants: 1) Cbx2ATH, substitution of PRG with AAA (amino acids 77–79); 2) Cbx2ATHL1, substitution of PRG with AAA (amino acids 134–136); 3) Cbx2ATHL2, substitution of PRG with AAA (amino acids 102–120); 4) Cbx2SRR, substitution of SKSKSSSSSSTSSSSSS with SKSKASASASTASASAA (amino acids 102–120); 5) Cbx2F12A, substitution of Phe-12 with Ala; 6) CDCBX2, amino acids 161–167; 7) HT-Cbx2 with the variant sequence. We generated the following CBX2 variants by PCR and used them to replace the Cbx2 sequence in the plasmid pGEX-6P-1-GST-CBX2-FLAG. We amplified the sequence encoding the Cbx2 variant sequence in the plasmid pGEX-6P-1-GST-YFP-CBX2-FLAG. To generate recombinant CBX2 in E. coli, we amplified the Cbx2 sequence by PCR and inserted it into the downstream GST sequence within the pGEX-6P-1-GST vector (GE Healthcare) to generate pGEX-6P-1-GST-CBX2. To facilitate double-affinity purification, we added a FLAG tag downstream of the Cbx2 sequence to generate pGEX-6P-1-GST-CBX2-FLAG. We amplified the YFP sequence to insert it upstream of the Cbx2 sequence to generate pGEX-6P-1-GST-YFP-CBX2-FLAG. To generate plasmids for expressing CBX2 variants in E. coli, we amplified the sequence encoding the Cbx2 variant sequence by PCR and used them to replace the Cbx2 sequence in the plasmid pGEX-6P-1-GST-CBX2-FLAG. We generated the following CBX2 variants: 1) Cbx2ATH, substitution of PRG with AAA (amino acids 77–79); 2) Cbx2ATHL1, substitution of PRG with AAA (amino acids 134–136); 3) Cbx2ATHL2, substitution of PRG with AAA (amino acids 102–120); 4) CBX2SRR, substitution of SKSKSSSSSSTSSSSSS with SKSKASASASTASASAA (amino acids 102–120).

Establishing cell lines

24 h before transfection, HEK293T cells were seeded into a 100-mm dish to reach 85–90% confluence at the time of transfection. Cells were cotransfected with 21 μg of pTRIPZ (M) containing the fusion gene, 21 μg of psPAX2, and 10.5 μg of pMD2.G using calcium phosphate precipitation. 12 h after transfection, the medium was replaced with fresh medium. 50 h after the medium change, the medium was harvested and centrifuged at 1000 × g to remove cell debris. Cells were mixed with the harvested medium in the presence of 8.0 μg/ml Polybrene (H9268, Sigma-Aldrich). 48–72 h after transfection, 1.0–2.0 μg/ml puromycin (P8833, Sigma-Aldrich) was added to cells. Cells were selected in the presence of puromycin for at least 1 week.

Generating recombinant protein of CBX2 and its variants

Recombinant CBX2 and its variants were generated and purified according to previous reports with modifications (81). The pGEX-6P-1-GST-FLAG vector containing the Cbx2 fusion gene was transformed into RosettaTM 2 (pLysS) host strains (71403, Novagen). A single colony was used to inoculate 5.0 ml of LB medium. Following overnight culture at 37 °C, 1.0 ml of the overnight culture was transferred into 1.0 liter of LB medium. After 6-h culture at 37 °C, the protein expression was induced overnight at 18 °C in the presence of 1.0 mM isopropyl 1-thio-β-D-galactopyranoside (IB02105, IBI Scientific). After centrifugation, cell pellets were resuspended in 25 ml of lysis buffer (50 mM HEPES, pH 7.5, 1.6 M KCl, 0.2 mM EDTA, 0.5 mM MgCl2, 0.5 mM DTT, and 0.2 mM phenylmethylsulfonyl fluoride). After three freeze-thaw cycles using liquid nitrogen, cells were disrupted using a sonicator (VCX130, Vibra-CellTM) for 3.0 min at 45% amplitude using 15-s on and 45-s off cycles. Cell debris was removed by centrifugation at 10,000 × g for 20 min at 4 °C. To precipitate nucleic acids, 10% polyethyleneimine in 20 ml of HEPES, pH 7.5, was added to the lysate to achieve a final concentration of 0.15%. The mixture was incubated for 30 min at 4 °C. After centrifugation at 20,000 × g for 20 min, the supernatant was incubated with 0.5 ml of prewashed GSH-Sepharose 4B beads (17-0756-01, GE Healthcare) for 1.0 h at 4 °C. After washing four times with washing buffer (20 ml HEPES, pH 7.5, 500 mM KCl, 0.2 mM EDTA, 1.0 mM DTT, and 0.2 mM phenylmethylsulfonyl fluoride), recombinant protein was eluted with 1.0 ml of 40 mM reduced L-GSH (G4251, Sigma-Aldrich). Alternatively, recombinant protein was eluted by incubating with 80 units of PreScission protease (27-0843-01, GE Healthcare), which cleaves the GST tag, at 4 °C overnight. Eluted protein was incubated with 100 μl of anti-FLAG-M2 affinity resin (A2220, Sigma) for 2.0 h at 4 °C. After washing four times with washing buffer supplemented with 1.0 M KCl, recombinant protein was eluted with 0.4 mg/ml FLAG peptide (F3290, Sigma-Aldrich) dissolved in washing buffer supplemented with 1.0 M KCl. Recombinant protein was resolved by SDS-PAGE to determine its purity and identity.
Polycomb CBX2 condensates by phase separation

In vitro condensate formation

After purification, KCl was added to the purified protein to reach a final concentration of 2.0 M, and then the mixture was concentrated to 30–50 μL by using an Amicon centrifugal tube (UFC500324, Millipore). Protein concentration was quantified by the Bradford (Bradford) protein assay (1856209, Thermo Scientific). 30 μL of protein samples were dialyzed with Spectra/Pro 1 dialysis tubing (132645, Spectrum Labs) in 1.0 liter of dialysis buffer (10 mM phosphate buffer, pH 7.4, containing 2.7 mM KCl and 137 mM NaCl) or dialysis buffer supplemented with 1.0 mM DTT at 4 °C. After changing buffer once, dialysis was performed overnight. 10 μL of the dialyzed sample were added to a coverslip dish made as described previously (82). After all condensates had settled down to the surface of the coverslip, DIC or fluorescence images of condensates were acquired using an Axio Observer D1 microscope (Zeiss, Germany) equipped with a 100×/1.40 numerical aperture oil immersion objective with additional 2.5× magnification and an Evolve electron-multiplying charge-coupled density camera (512 × 512; Photometrics, Tucson, AZ). For the excitation and emission of YFP, a Brightline® single-band laser filter set (Semrock; excitation filter, FF02-482/18-25; emission filter, FF01-525/25-25; dichroic mirror, Di02-R488-25) was used. The number of condensates per frame was counted by using ImageJ.

To determine the critical/saturation concentration of phase separation of CBX2, a series of concentrations of CBX2 (1.2, 2.4, 4.8, and 12 μM) was dialyzed under the same conditions, and the number of condensates per frame was counted as described above. To investigate driving forces that contribute to the formation of CBX2 condensates, 10 μL of the dialyzed sample, NaCl, Triton X-100, and 1,6-hexanediol (240117, Sigma-Aldrich) were added, respectively. The mixture was incubated at 4 °C for 30 min. Condensates were imaged and analyzed as described above.

To prepare CBX2 condensates that concentrate DNA, 4.8 μM CBX2 was mixed with Alexa Fluor 488–labeled DNA (0.5 μM) or Alexa Fluor 488 (1.0 μM), respectively. To prepare CBX2 condensates that concentrate nucleosomes, 2.4 μM CBX2 was mixed with Cy5-labeled mononucleosomes (40 nM). The mixture was dialyzed as described above. DIC and fluorescence images were taken using an Axio Observer D1 microscope as described above. For the excitation and emission of Alexa Fluor 488, a Brightline single-band laser filter set (Semrock; excitation filter, FF02-482/18-25; emission filter, FF01-525/25-25; dichroic mirror, Di02-R488-25) was used. For the excitation and emission of Cy5, a Brightline long-pass laser filter set (Semrock; excitation filter, FF01-640/14-25; emission filter, BLP01-609/14; dichroic mirror, Di02-R561-25) was used. Visible condensates were counted using ImageJ. Images were presented using Photoshop.

Live-cell imaging of YFP-CBX2 treated with 1,6-hexanediol

We seeded mES cells stably expressing YFP-CBX2 to a gelatin-coated coverslip-bottom dish 24 h before the imaging. Cell culture medium was replaced with the live-cell imaging medium and maintained at 37 °C using a heater controller. Hexanediol was added to the medium to reach a final concentration of 10%. An image stack was taken at every 2-min interval for 20 min using an Axio Observer D1 microscope as described above. For the excitation and emission of YFP, a YFP-2427B filter set (Semrock; excitation filter, FF01-500/24; emission filter, FF01-542/25; dichroic mirror, FF520-Di02) was used.

Immunofluorescence

Cells stably expressing YFP-CBX2 were seeded to coverslips and cultured for 24 h. Cells were fixed with 1.0% paraformaldehyde for 10 min at room temperature. After treatment with 0.2% Triton X-100 for 10 min, cells were washed with basic buffer (10 mM PBS, pH 7.2, 0.05% Tween 20, and 0.1% Triton X-100) and incubated with basic blocking buffer (basic buffer supplemented with 3% goat serum and 3% BSA) overnight. Primary antibodies anti-PHC1 (39723, Active Motif; 1:200 dilution), anti RING1B (D-319, MBL; 1:200 dilution), anti-H3K27me3 (9733, Cell Signaling Technology; 1:200 dilutions), and anti-GFP (A11122, Life Technology; 1:500 dilution) were added to cells and incubated for 2.0 h at room temperature. After washing with basic blocking buffer, cells were incubated with secondary antibodies Alexa Fluor 488–labeled anti-rabbit (A-11008, Life Technologies; 1:1000 dilutions) and/or Alexa Fluor 647–labeled anti-mouse (A32728, Invitrogen; 1:1000 dilution) for 2.0 h at room temperature. Cells were washed and mounted with ProLong Antifade reagents (P7481, Life Technologies) and imaged using a Zeiss LSM 700 Observer Z1 equipped with a 100× oil objective (numerical aperture, 1.4) and an electron-multiplying charge-coupled density camera. For Alexa Fluor 488, 514-nm excitation and 527-nm emission filters were used. For Alexa Fluor 647, 639-nm excitation and 665-nm emission filters were used.

FRAP

Cells stably expressing YFP-CBX2 were seeded to a 35-mm gelatin-coated coverslip-bottom dish. Cells were maintained as described for live-cell imaging. FRAP imaging was performed using a Zeiss LSM 700 Observer with the following parameters: pinhole, full open; scan speed, 177.32 μs/pixel. Before photobleaching, two images were taken. Immediately after photobleaching, 20 images were taken with 15-s intervals.
The images were analyzed using ImageJ. We used TurboReg to correct images for movement in the xy plane. After correcting fluctuations in background and total signal, the fluorescence intensities were normalized to the signal before photobleaching to obtain the fluorescence recovery as described previously (42, 49).

**Author contributions**—R. T., S. K., K. B., H. N. D., T. N. H., H. W., and X. R. formal analysis; R. T., S. K., K. B., T. Y., H. N. D., T. N. H., C. Y. Z., B. M., and X. R. investigation; R. T., S. K., K. B., T. Y., H. N. D., T. N. H., C. Y. Z., B. M., and X. R. methodology; T. Y., H. W., and X. R. resources; H. W. and X. R. supervision; X. R. conceptualization; X. R. data curation; X. R. funding acquisition; X. R. validation; X. R. visualization; X. R. writing—original draft; X. R. project administration; X. R. writing—review and editing.

**Acknowledgments**—We thank Dr. Haruhiko Koseki for providing PGK12.1 mES cells. Dr. Neil Brockdorff for providing PGK12.1 mES cells. Xu, C., Bian, C., Yang, W., Galka, M., Ouyang, H., Chen, C., Qiu, W., Liu, R., Jones, A. E., MacKenzie, F., Pan, P., Li, S. S., Wang, H., and Min, J. (2010) Binding of different histone marks differentially regulates the activity and specificity of polycomb repressive complex 2 (PRC2). Proc. Natl. Acad. Sci. U.S.A. 107, 19266–19271 CrossRef Medline

1. Brown, S. W. (1966) Heterochromatin. *Science* 151, 417–425 Medline
2. Larson, A. G., and Narlikar, G. J. (2018) The role of phase separation in heterochromatin formation, function, and regulation. *Biochemistry* 57, 2540–2548 CrossRef Medline
3. Trojer, P., and Reinberg, D. (2007) Facultative heterochromatin: is there a distinctive molecular signature? *Mol. Cell* 28, 1–13 CrossRef Medline
4. Heard, E. (2005) Delving into the diversity of facultative heterochromatin: the epigenetics of the inactive X chromosome. *Curr. Opin. Genet. Dev.* 15, 482–489 CrossRef Medline
5. Kondo, T., Isono, K., Kondo, K., Endo, T. A., Itohara, S., Vidal, M., and Brockdorff, N. (2004) Polycomb group proteins Ring1A/B link ubiquitylation of histone H2A to heritable gene silencing and X inactivation. *Dev. Cell* 7, 663–676 CrossRef Medline
6. Kalb, R., Latwiel, S., Baymaz, H. I., Jansen, P. W., Müller, C. W., Vermeulen, M., and Müller, J. (2014) Histone H2A monoubiquitination promotes histone H3 methylation in Polycomb repression. *Nat. Struct. Mol. Biol.* 21, 569–571 CrossRef Medline
7. Schuettengruber, B., Bourbon, H. M., Di Croce, L., and Cavalli, G. (2017) Epigenetic consequences of disruptions to the Polycomb repression complex. *Cell* 168, 1039–1043 CrossRef Medline
8. Cao, R., Wang, L., Wang, H., Xia, L., Erdjument-Bromage, H., Tempst, P., Jones, R. S., and Zhang, Y. (2002) Role of histone H3 lysine 27 methylation in Polycomb-group silencing. *Science* 298, 1039–1043 CrossRef Medline
9. Czermin, B., Melfi, R., McCabe, D., Seitz, V., Imhof, A., and Pirrotta, V. (2002) *Drosophila* enhancer of Zeste/ESC complexes have a histone H3 methyltransferase activity that marks chromosomal Polycomb sites. *Cell* 111, 185–196 CrossRef Medline
10. Kuzmichev, A., Nishioka, K., Erdjument-Bromage, H., Tempst, P., and Reinberg, D. (2002) Histone methyltransferase activity associated with a human multiprotein complex containing the Enhancer of Zeste protein. *Genes Dev.* 16, 2893–2905 CrossRef Medline
11. Müller, J., Hart, C. M., Francis, N. J., Vargas, M. L., Sengupta, A., Wild, B., Miller, E. L., O’Connor, M. B., Kingston, R. E., and Simon, J. A. (2002) Histone methyltransferase activity of a *Drosophila* Polycomb group repressor complex. *Cell* 111, 197–208 CrossRef Medline
12. Shen, X., Liu, Y., Hsu, Y. I., Fujiwara, Y., Kim, I., Mao, X., Yuan, G. C., and Orkin, S. H. (2008) EZH1 mediates methylation on histone H3 lysine 27 and complements EZH2 in maintaining stem cell identity and executing pluripotency. *Mol. Cell* 32, 491–502 CrossRef Medline
13. Margueron, R.,ustin, N., Ohno, K., Sharpe, M. L., Son, J., Drury WJ 3rd, Voigt, P., Martin, S. R., Taylor, W. R., De Marco, V., Pirrotta, V., Reinberg, D., and Gamblin, S. J. (2009) Role of the polycomb protein EED in the propagation of repressive histone marks. *Nature* 461, 762–767 CrossRef Medline
14. Xu, C., Bian, C., Yang, W., Galka, M., Ouyang, H., Chen, C., Qiu, W., Liu, R., Jones, A. E., MacKenzie, F., Pan, P., Li, S. S., Wang, H., and Min, J. (2014) Role of histone H2A ubiquitination in Polycomb silencing. *Nature* 431, 873–878 CrossRef Medline
15. de Napoles, M., Mermoud, J. E., Wakao, R., Tang, Y. A., Endoh, M., Ap­panah, R., Nesterova, T. B., Silva, J., Otte, A. P., Vidal, M., Koseki, H., and Brockdorff, N. (2004) Polycomb group proteins Ring1A/B link ubiquitylation of histone H2A to heritable gene silencing and X inactivation. *Dev. Cell* 7, 663–676 CrossRef Medline
16. Wang, H., Wang, L., Erdjument-Bromage, H., Vidal, M., Tempst, P., Jones, R. S., and Zhang, Y. (2004) Role of histone H2A ubiquitination in Polycomb silencing. *Nature* 431, 873–878 CrossRef Medline
17. Kalb, R., Latwiel, S., Baymaz, H. I., Jansen, P. W., Müller, C. W., Ver­meulen, M., and Müller, J. (2014) Histone H2A monoubiquitination promotes histone H3 methylation in Polycomb repression. *Nat. Struct. Mol. Biol.* 21, 1029–1034 CrossRef Medline
18. Cao, R., Wang, L., Wang, H., Xia, L., Erdjument-Bromage, H., Tempst, P., Jones, R. S., and Zhang, Y. (2002) Role of histone H3 lysine 27 methylation in Polycomb-group silencing. *Science* 298, 1039–1043 CrossRef Medline
Polycomb CBX2 condensates by phase separation

Tompa, P., and Fuxreiter, M. (2018) Protein phase separation: a new phase in cell biology. Trends Cell Biol. 28, 420–435 CrossRef Medline

Woodyruff, J. B., Hyman, A. A., and Boke, E. (2018) Organization and function of non-dynamic biomolecular condensates. Trends Biochem. Sci. 43, 81–94 CrossRef Medline

Meldi, L., and Brickner, J. H. (2011) Compartmentalization of the nucleus. Trends Cell Biol. 21, 701–708 CrossRef Medline

Chong, S., Dugast-Darzacq, C., Liu, Z., Dong, P., Dailey, G. M., Cattoglio, C., Heckert, A., Banala, S., Lavis, L., Darzacq, X., and Tijan, R. (2018) Imaging dynamic and selective low-complexity domain interactions that control gene transcription. Science 361, eaar2555 CrossRef Medline

Alkema, M. J., Bronk, M., Verhoeven, E., Otte, A., van 't Veer, L. T., Berns, M. G. W., van Oudenaarden, A., and van Dieren, R. P. (2018) RNA polymerase II clustering through carboxy-terminal domain phase separation. Nat. Struct. Mol. Biol. 25, 833–840 CrossRef Medline

Chen, S., Dugast-Darzacq, C., Liu, Z., Dong, P., Dailey, G. M., Cattoglio, C., Heckert, A., Banala, S., Lavis, L., Darzacq, X., and Tijan, R. (2018) Imaging dynamic and selective low-complexity domain interactions that control gene transcription. Science 361, eaar2555 CrossRef Medline

Rau, M. K., Weidinger, C., and Klose, R. J. (2015) Targeting Polycomb CBX2 complexes by phase separation links phase separation and gene control. J. Biol. Chem. 290, 28038–28054 CrossRef Medline

Levine, S. S., Weiss, A., Erdjument-Bromage, H., Shao, Z., Tempst, P., and Kingston, R. E. (2002) The core of the polycomb repressive complex is compositionally and functionally conserved in flies and humans. Mol. Cell. Biol. 22, 6070–6078 CrossRef Medline

Satijn, D. P., Gunster, M. J., van der Vlag, J., Hamer, K. M., Schul, W., Alkema, M. J., Saurin, A. J., Freemont, P. S., van Driel, R., and Otte, A. P. (1997) Ring1A is associated with the polycomb group protein complex and acts as a transcriptional repressor. Mol. Cell. Biol. 17, 4105–4113 CrossRef Medline

Gunter, M. J., Satijn, D. P., Hamer, K. M., de Blauwe, J., de Brujin, D., Alkema, M. J., van Lohuizen, M., van Driel, R., and Otte, A. P. (1997) Identification and characterization of interactions between the vertebrate polycomb-group protein BM11 and human homologs of polycombotic. Mol. Cell. Biol. 17, 2326–2335 CrossRef Medline

Zhen, C. Y., Duc, H. N., Sokotovic, M., Phiel, C. J., and Ren, X. (2014) Cbx2 stably associates with mitotic chromosomes via a PRC2- or PRC1-independent mechanism and is needed for recruiting PRC1 complex to mitotic chromosomes. Mol. Biol. Cell 25, 3726–3739 CrossRef Medline

Piovesan, D., Tabaro, F., Paladini, L., Necci, M., Micetic, I., Camilloni, C., Davey, N., Dosztányi, Z., Mészáros, B., Monzon, A. M., Parisi, G., Schad, E., Sormanni, P., Tompa, P., Vendruscolo, M., et al. (2018) MobiDB 3.0: more annotations for intrinsic disorder, conformational diversity and interactions in proteins. Nucleic Acids Res. 46, D471–D476 CrossRef Medline

Kawaguchi, T., Machida, S., Kurumizaka, H., Tagami, H., and Nakayama, J. I. (2017) Phosphorylation of CBX2 controls its nucleosome-binding specificity. J. Biochem. 162, 343–355 CrossRef Medline

Wang, J., Choi, J. M., Holehouse, A. S., Lee, H. O., Zhang, X., Jahnel, M., Maharana, S., Lemaître, R., Pozniakovsky, A., Drechsel, D., Poser, I., Pappu, R. V., Alberti, S., and Hyman, A. A. (2018) A molecular grammar governing the driving forces for phase separation of prion-like RNA binding proteins. Cell 174, 688 – 699.e16 CrossRef Medline

Boeynaeens, S., Bogaert, E., Kovacs, D., Komijnberg, A., Timmerman, E., Volkov, A., Guharoy, M., De Decker, M., Jaspers, T., Ryan, V. H., Janke, A. M., Baatsen, P., Vercruyssen, T., Kolaïtas, R. M., Daelemans, D., et al. (2017) Phase separation of C9orf72 dipeptide repeats perturbs stress granule dynamics. Mol. Cell 65, 1044–1055.e5 CrossRef Medline

Lee, K. H., Zhang, P., Kim, H. I., Mitrea, D. M., Sarkar, M., Freibau, B. D., Cika, C., Coughlin, M., Messing, I., Moliex, A., Maxwell, B. A., Kim, N. C., Temirov, J., Moore, J., Kolaïtas, R. M., et al. (2016) C9orf72 dipeptide repeats impair the assembly, dynamics, and function of membrane-less organelles. Cell 167, 774–788.e17 CrossRef Medline

Jiang, H., Wang, S., Huang, Y., He, X., Cui, H., Zhu, X., and Zheng, Y. (2015) Phase transition of spindel-associated protein regulate spindle apparatus assembly. Cell 163, 108–122 CrossRef Medline

Nott, T. J., Petsalaki, E., Farber, P., Jervis, D., Fussner, E., Plochowietz, A., Craggs, T. D., Bazett-Jones, D. P., Watson, T., Forman-Kay, J. D., and Baldwin, A. J. (2015) Phase transition of a disordered nuce protein generates environmentally responsive membraneless organelles. Mol. Cell 57, 936–947 CrossRef Medline

Pak, C. W., Kosmo, M., Holehouse, A. S., Padrick, S. B., Mittal, A., Ali, R., Yunus, A. A., Liu, D. R., Pappu, R. V., and Rosen, M. K. (2016) Sequence determinants of intracellular phase separation by complex coacervation of a disordered protein. Mol. Cell 63, 72–85 CrossRef Medline

Mitrea, D. M., Cika, I. A., Stanley, C. B., Nourse, A., Onuchic, P. L., Banerjee, P. R., Phillips, A. H., Park, C. G., Deniz, A. A., and Krivacki, R. W. (2018) Self-interaction of NPM1 modulates multiple mechanisms of liquid-liquid phase separation. Nat. Commun. 9, 842 CrossRef Medline

Feric, M., Vaidya, N., Harmon, T. S., Mitrea, D. M., Zhu, L., Richardson, T. M., Krivacki, R. W., Pappu, R. V., and Brangwynne, C. P. (2016) Coexisting liquid phases underlie nucleolar subcompartments. Cell 165, 1686–1697 CrossRef Medline

Elbaum-Garfinkle, S., Kim, Y., Szczepaniak, K., Chen, C. C., Eckmann, C. R., Myong, S., and Brangwynne, C. P. (2015) The disordered P granule protein LAF-1 drives phase separation into droplets with tunable viscosity and dynamics. Proc. Natl. Acad. Sci. U.S.A. 112, 7189–7194 CrossRef Medline

Patel, A., Lee, H. O., Jawerth, L., Maharana, S., Jahnel, M., Hein, M. Y., Stoynov, S., Mahmadi, J., Saha, S., Franzmann, T. M., Pozniakovsky, A., Poser, I., Maghelli, N., Ro yer, L. A., Weigert, M., et al. (2015) A liquid-to-solid phase transition of the ALS protein FUS accelerated by disease mutation. Cell 162, 1066–1077 CrossRef Medline

Altmeier, M., Nelesen, K. J., Teloni, F., Pozniakovsky, I., Pellegrino, S., Grafte, M., Rask, M. B., Streicher, W., Jungmichel, S., Nielsen, M. L., and Lukas, J. (2015) Liquid demixing of intrinsically disordered proteins is seeded by poly(ADP-ribose). Nat. Commun. 6, 8088 CrossRef Medline

Senthilkumar, R., and Mishra, R. K. (2009) Novel motifs distinguish multiple homologues of Polycomb in vertebrates: expansion and diversification of the epigenetic toolkit. BMC Genomics 10, 549 CrossRef Medline

Blackledge, N. P., Rose, N. R., and Klose, R. J. (2015) Targeting Polycomb systems to regulate gene expression: modifications to a complex story. Nat. Rev. Mol. Cell Biol. 16, 643–649 CrossRef Medline

Kaustov, L., Ouyang, H., Amaya, M., Lemak, A., Nady, N., Duan, S., Wasney, G. A., Li, Z., Vedadi, M., Schapira, M., Min, J., and Arrowsmith, C. H.
Recognition and specificity determinants of the human CBX chromodomains. J. Biol. Chem. 286, 521–529 CrossRef Medline

Bernstein, E., Duncan, E. M., Masui, O., Gil, J., Heard, E., and Allis, C. D. (2006) Mouse polycomb proteins bind differentially to methylated histone H3 and RNA and are enriched in facultative heterochromatin. Mol. Cell. Biol. 26, 2560–2569 CrossRef Medline

Semerenko, A. N., and Rubinstein, M. (1998) Thermoreversible gelation in solutions of associative polymers. 1. Statics. Macromolecules 31, 1373–1385 CrossRef Medline

Banani, S. F., Rice, A. M., Peeples, W. B., Lin, Y., Jain, S., Parker, R., and Rosen, M. K. (2016) Compositional control of phase-separated cellular bodies. Cell 166, 651–663 CrossRef Medline

Hemenway, C. S., Halligan, B. W., and Levy, L. S. (1998) The Bmi-1 oncogene interacts with dinG and MP12: the role of RING finger domains. Oncogene 16, 2541–2547 CrossRef Medline

Satijn, D. P., and Otte, A. P. (1999) RING1 interacts with multiple Polycomb-group proteins and displays tumorigenic activity. Mol. Cell. Biol. 19, 57–68 CrossRef Medline

Wang, R., Ilango, U., Robinson, A. K., Schirf, V., Schwarz, P. M., Lafer, E. M., Demeler, B., Hinck, A. P., and Kim, C. A. (2008) Structural transitions of the RING1B C-terminal region upon binding the polycomb box domain. Biochemistry 47, 8007–8015 CrossRef Medline

Tardat, M., Albert, M., Kunzmann, R., Liu, Z., Kaustov, L., Thierry, R., Duan, S., Brykczynska, U., Arrowsmith, C. H., and Peters, A. H. (2015) Cbx2 targets PRC1 to constitutive heterochromatin in mouse zygotes in a parent-of-origin-dependent manner. Mol. Cell 58, 157–171 CrossRef Medline

Kim, C. A., Gingery, M., Palpa, R. M., and Bowie, J. U. (2002) The SAM domain of polyhomeotic forms a helical polymer. Nat. Struct. Biol. 9, 453–457 CrossRef Medline

Kim, C. A., and Bowie, J. U. (2003) SAM domains: uniform structure, diversity of function. Trends Biochem. Sci. 28, 625–628 CrossRef Medline

Eskeland, R., Leeb, M., Grimes, G. R., Kress, C., Boyle, S., Sproul, D., Gilbert, N., Fan, Y., Skoultchi, A. I., Wutz, A., and Bickmore, W. A. (2010) Ring1B compacts chromatin structure and represses gene expression independent of histone ubiquitination. Mol. Cell 38, 452–464 CrossRef Medline

Schoenfelder, S., Sugar, R., Dimond, A., Javiera, B. M., Armstrong, H., Miou, B., Dimitrova, E., Matheson, L., Tavares-Cadete, F., Furlan-Magari, M., Segonds-Pichon, A., Jurkowski, W., Tabbara, K., Andrews, S., et al. (2015) Polycomb repressive complex PRC1 spatially constrains the mouse embryonic stem cell genome. Nat. Genet. 47, 1179–1186 CrossRef Medline

Penny, G. D., Kay, G. F., Sheardown, S. A., Rastan, S., and Brockdorff, N. (1996) Requirement for Xist in X chromosome inactivation. Nature 379, 131–137 CrossRef Medline

Katoh-Fukui, Y., Tsuchiya, R., Shiroishi, T., Nakahara, Y., Hashimoto, N., Noguchi, K., and Higashinakagawa, T. (1998) Male-to-female sex reversal in M33 mutant mice. Nature 393, 688–692 CrossRef Medline

Endoh, M., Endo, T. A., Endo, T., Fujimura, Y., Ohara, O., Toyoda, T., Otte, A. P., Okano, M., Brockdorff, N., Vidal, M., and Koseki, H. (2008) Polycomb group proteins Ring1A/B are functionally linked to the core transcriptional regulatory circuitry to maintain ES cell identity. Development 135, 1513–1524 CrossRef Medline

Huynh, T. N., and Ren, X. (2017) Producing GST-Cbx7 fusion proteins from Escherichia coli. Bio Protoc. 7, e2333 CrossRef Medline

Duc, H. N., and Ren, X. (2017) Labelling HaloTag fusion proteins with HaloTag ligand in living cells. Bio Protoc. 7, e2526 CrossRef Medline

Polycomb CBX2 condensates by phase separation

Kim, C. A., Gingery, M., Palpa, R. M., and Bowie, J. U. (2002) The SAM domain of polyhomeotic forms a helical polymer. Nat. Struct. Biol. 9, 453–457 CrossRef Medline

Kim, C. A., and Bowie, J. U. (2003) SAM domains: uniform structure, diversity of function. Trends Biochem. Sci. 28, 625–628 CrossRef Medline

Eskeland, R., Leeb, M., Grimes, G. R., Kress, C., Boyle, S., Sproul, D., Gilbert, N., Fan, Y., Skoultchi, A. I., Wutz, A., and Bickmore, W. A. (2010) Ring1B compacts chromatin structure and represses gene expression independent of histone ubiquitination. Mol. Cell 38, 452–464 CrossRef Medline

Schoenfelder, S., Sugar, R., Dimond, A., Javiera, B. M., Armstrong, H., Miou, B., Dimitrova, E., Matheson, L., Tavares-Cadete, F., Furlan-Magari, M., Segonds-Pichon, A., Jurkowski, W., Tabbara, K., Andrews, S., et al. (2015) Polycomb repressive complex PRC1 spatially constrains the mouse embryonic stem cell genome. Nat. Genet. 47, 1179–1186 CrossRef Medline

Penny, G. D., Kay, G. F., Sheardown, S. A., Rastan, S., and Brockdorff, N. (1996) Requirement for Xist in X chromosome inactivation. Nature 379, 131–137 CrossRef Medline

Katoh-Fukui, Y., Tsuchiya, R., Shiroishi, T., Nakahara, Y., Hashimoto, N., Noguchi, K., and Higashinakagawa, T. (1998) Male-to-female sex reversal in M33 mutant mice. Nature 393, 688–692 CrossRef Medline

Endoh, M., Endo, T. A., Endo, T., Fujimura, Y., Ohara, O., Toyoda, T., Otte, A. P., Okano, M., Brockdorff, N., Vidal, M., and Koseki, H. (2008) Polycomb group proteins Ring1A/B are functionally linked to the core transcriptional regulatory circuitry to maintain ES cell identity. Development 135, 1513–1524 CrossRef Medline

Huynh, T. N., and Ren, X. (2017) Producing GST-Cbx7 fusion proteins from Escherichia coli. Bio Protoc. 7, e2333 CrossRef Medline

Duc, H. N., and Ren, X. (2017) Labelling HaloTag fusion proteins with HaloTag ligand in living cells. Bio Protoc. 7, e2526 CrossRef Medline
Nuclear condensates of the Polycomb protein chromobox 2 (CBX2) assemble through phase separation
Roubina Tatavosian, Samantha Kent, Kyle Brown, Tingting Yao, Huy Nguyen Duc, Thao Ngoc Huynh, Chao Yu Zhen, Brian Ma, Haobin Wang and Xiaojun Ren

J. Biol. Chem. 2019, 294:1451-1463.
doi: 10.1074/jbc.RA118.006620 originally published online December 4, 2018

Access the most updated version of this article at doi: 10.1074/jbc.RA118.006620

Alerts:
- When this article is cited
- When a correction for this article is posted

Click here to choose from all of JBC's e-mail alerts

This article cites 82 references, 26 of which can be accessed free at http://www.jbc.org/content/294/5/1451.full.html#ref-list-1