Transcriptional repression by PRC1 in the absence of H2A monoubiquitylation

Ana Raquel Pengelly, Reinhard Kalb, Katja Finkl, and Jürg Müller

Laboratory of Chromatin Biology, Max Planck Institute of Biochemistry, 82152 Martinsried, Germany

Histone H2A monoubiquitylation (H2Aub) is considered to be a key effector in transcriptional repression by Polycomb-repressive complex 1 (PRC1). We analyzed Drosophila with a point mutation in the PRC1 subunit Sce that abolishes its H2A ubiquitylase activity or with point mutations in the H2A and H2Av residues ubiquitylated by PRC1. H2Aub is essential for viability and required for efficient histone H3 Lys27 trimethylation by PRC2 early in embryogenesis. However, H2Aub-deficient animals fully maintain repression of PRC1 target genes and do not show phenotypes characteristic of Polycomb group mutants. PRC1 thus represses canonical target genes independently of H2Aub.

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Polycomb [Pc] group [PcG] proteins form conserved multigene complexes that repress transcription of developmental regulator genes in cells where they should be inactive (Beisel and Paro 2011; Simon and Kingston 2013). PcG protein complexes are thought to silence gene transcription through covalent and noncovalent modification of chromatin at target genes. Polycomb-repressive complex 2 (PRC2) trimethylates Lys27 in histone H3 (H3-K27me3), and, in Drosophila, cells with an H3-K27R point mutation fail to repress PcG target genes, demonstrating the importance of this modification for repression (Pengelly et al. 2013; McKay et al. 2015). The H3-K27me3 modification is recognized by Pc, a subunit of PRC1, and thus is thought to mark nucleosomes for interaction with PRC1 (Fischle et al. 2003; Min et al. 2003). PRC1 compacts nucleosome arrays in vitro by a nonenzymatic mechanism (Shao et al. 1999; Francis et al. 2004), and domains of PRC1 subunits important for this activity are critical for gene repression in vivo (King et al. 2005; Isono et al. 2013; Gambetta and Müller 2014).

PRC1 also contains the E3 ligase activity for monoubiquitylation of H2A at Lys19 in mammals or at the corresponding Lys118 in Drosophila (Wang et al. 2004; Lagarou et al. 2008). The mechanism and physiological role of monoubiquitylated histone H2A [H2Aub] is poorly understood. Early studies proposed that H2Aub directly blocks transcription elongation by RNA polymerase II (Stock et al. 2007). More recent studies found that H2Aub is bound by a specific form of PRC2 and promotes H3-K27me3 by this complex on H2Aub nucleosomes in vitro (Kalb et al. 2014). In mouse embryonic stem (ES) cells, tethering of PRC1 E3 ligase activity to a chromosomal site was found to result in H2A monoubiquitylation, binding of PRC2, and H3-K27me3 formation at this site, leading to the proposal that a primary function of H2Aub might be to recruit PRC2 (Blackledge et al. 2014). The E3 ligase activity for H2Aub formation is provided by the PRC1 subunit Ring1b and its paralog, Ring1a, in vertebrates and by Sce in Drosophila (de Naples et al. 2004; Wang et al. 2004). Drosophila lacking Sce protein show loss of H2Aub (Lagarou et al. 2008; Gutiérrez et al. 2012), whereas, in vertebrates, only removal of both Ring1a and Ring1b proteins results in complete loss of H2Aub (de Naples et al. 2004). Sce and Ring1a/b also play an architectural role in PRC1. The N-terminal Ring finger domain of Ring1a/b (or Sce) associates with the Ring finger domains of the different PCGF family members (Buchwald et al. 2006; Li et al. 2006) to form the core of the various PRC1-type assemblies (Gao et al. 2012), while the C terminus binds to the Cbx/Pc or Rybp subunits (Scholerammer et al. 1997; Wang et al. 2010). The reduced levels of diverse PRC1 subunits in Ring1b mutant mouse ES cells (Leeb and Wutz 2007) supports the idea that Ring1a/b is critical for integrity of PRC1 assemblies. In turn, this raises the question of to what extent the severe phenotypes of mouse embryos lacking Ring1b protein (Vonecken et al. 2003) or Drosophila lacking Sce protein (Gutiérrez et al. 2012) can be attributed to the loss of H2Aub.

Selective impairment of E3 ligase activity in an otherwise intact Ring1b protein can be achieved by the I53A substitution in the interface that binds the E2 ubiquitin-conjugating enzyme (Buchwald et al. 2006; Bentley et al. 2011). Previous studies used transgenes expressing the Ring1bI53A protein to manipulate H2Aub levels in mouse ES cells lacking Ring1b (Eskeland et al. 2010) or Ring1a and Ring1b (Endoh et al. 2012). Endoh et al. [2012] reported that H2Aub was required for maintenance of ES cell identity and efficient repression of a distinct subset of PcG target genes.

Here, we investigated the role of H2Aub in developing Drosophila. We generated animals in which we replaced wild-type Sce with a catalytically inactive Sce protein and, in parallel, animals in which we mutated the residues in H2A and its variant, H2Av, that are monoubiquitylated by PRC1. We show that H2Aub is not required for repression of canonical PcG target genes and that H2Aub-deficient animals arrest development with surprisingly mild morphological defects. Together, these results suggest that PRC1 represses target genes primarily through an H2Aub-independent mechanism.

Keywords: Polycomb, PRC1, H2A and H2Av monoubiquitylation, transcriptional repression, Drosophila

Corresponding author: mueller@biochem.mpg.de

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Results and Discussion

Sce E3 ligase activity for H2Aub formation is not required for repression of canonical PRC1 target genes

In a first approach, we analyzed Drosophila lacking Sce E3 ligase activity. In an initial set of experiments, we generated \( Sce^{KO} \) mutant embryos that were homozygous for \( Sce^{KO} \) [a small chromosomal deletion that removes the entire \( Sce \) ORF] (Gutiérrez et al. 2012) and lacked not only zygotically expressed \( z^{-} \) but also maternally deposited \( m^{-} \) wild-type Sce protein [Fig. 1A, panel 1, cf. lanes 4–6 and 1–3; see Supplemental Fig. S1 for experimental strategy]. Using an antibody that specifically detects H2Aub, we found that H2Aub was undetectable in \( Sce^{KO} \) mutant embryos but readily detected in wild-type embryos (Fig. 1A, panel 9, cf. lanes 4–6 and 1–3), demonstrating that Sce is responsible for all monoubiquitylation of H2A.

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In the next set of experiments, we therefore generated animals that contain a \( Sce^{I48A} \) mutant protein instead of wild-type Sce. The I48A mutation in Sce corresponds to the I53A mutation in Ring1b that impedes its E3 ligase activity for H2Aub (see above). We first generated \( Sce^{I48A} \) mutant embryos in which both zygotically expressed and maternally deposited Sce protein contained the I48A mutation; the SceI48A protein was expressed at normal levels from a genomic Sce fragment (Fig. 1A, panel 1, cf. lanes 7–9 and 1–3; see Supplemental Fig. S1 for experimental strategy). In \( Sce^{I48A} \) mutant animals, all analyzed PcG proteins, including Pc, were present at wild-type levels (Fig. 1A, panels 2–8), but H2Aub was still >98% reduced (Fig. 1A, panel 9, cf. lanes 7–9 and 1–3). These \( Sce^{I48A} \) mutant embryos thus allowed us to selectively assess the requirement for H2Aub in Drosophila.

We first investigated how loss of H2Aub affects H3-K27me3 by PRC2. H3-K27me3 bulk levels in \( Sce^{KO} \) mutant embryos thus allowed us to selectively assess the requirement for H2Aub in Drosophila.

We next compared the phenotype of \( Sce^{I48A} \) mutant embryos with that of \( Sce^{KO} \) mutant embryos. The embryonic cuticle of \( Sce^{KO} \) mutant embryos shows extensive homeotic transformations (Fig. 1B) caused by widespread misexpression of HOX genes such as Antennapedia (Antp), Ultrabithorax (Ubx), and Abdominal-B (Abd-B) [Fig. 1B]. In contrast, \( Sce^{I48A} \) mutant embryos looked indistinguishable from wild-type embryos: They did not show any detectable homeotic transformations or any other defects in the embryonic cuticle, and expression of Antp, Ubx, and Abd-B was confined to their normal expression domains (Fig. 1B). Moreover, engrailed (en), another classic PcG target gene that is misexpressed in \( Sce^{KO} \) mutant embryos (Fig. 1B), also showed a wild-type expression pattern in \( Sce^{I48A} \) mutant embryos (Fig. 1B). Polycomb repression of the HOX and \( en \) genes is thus unaffected in embryos lacking H2Aub.

Even though \( Sce^{I48A} \) mutant animals showed overall normal morphology (Fig. 1B), they nevertheless...
arrested development at the end of embryogenesis. To explore whether morphogenesis or tissue patterning might be affected in more subtle ways, we stained late-stage *Sce* 

mutant embryos with anti-Futsch [22C10] antibody to visualize the cells of the peripheral nervous system (PNS) or with anti-myosin heavy chain (Mhc) antibody to visualize muscle cells. Development and patterning of PNS and muscle tissues in *Sce* 

mutant embryos was comparable with that of wild-type embryos (Supplemental Fig. S3). However, *Sce* 

mutant animals occasionally showed aberrant neuronal or muscle fiber connections (Supplemental Fig. S3). These defects were detected in only a fraction of the *Sce* 

mutant embryos and occurred in random locations rather than a stereotype pattern in the body, making it difficult to quantify the defects. Together, these data suggest that overall cell type specification and patterning of epidermal, mesodermal, and PNS tissues occurs normally in embryos lacking H2Aub but that random cells or small groups of cells display morphogenetic defects.

**Post-embryonic development, metamorphosis, and differentiation in the absence of H2Aub**

We next generated *Sce* 

mutant animals that expressed *Sce* 

mutant protein from their zygotic genome but contained a supply of maternally deposited wild-type Sce protein during the early stages of embryogenesis. To investigate perdurance of maternally deposited Sce protein, we analyzed extracts from *Sce* 

mutant embryos (i.e., lacking zygotic expression of Sce) and found that residual maternally supplied wild-type Sce protein and H2Aub generated by this protein are present in late stage embryos (Supplemental Fig. S4A). *Sce* 

mutant animals developed beyond the end of embryogenesis, completed all stages of larval development, and formed pupae, and a small fraction developed into pharate adults (Supplemental Fig. S4B). As expected, because of dilution of maternally deposited Sce and H2Aub by cell division and turnover during larval growth, H2Aub was no longer detectable in *Sce* 

mutant third instar larvae (Fig. 2A). Of note, bulk levels of H3-K27me3 in these animals were comparable with those in wild-type larvae (Fig. 2A, discussed below). The exoskeleton of *Sce* 

pharate adults showed all of the features of wild-type Drosophila (Fig. 2B) with variable morphological defects (see below). However, these animals showed no homeotic transformations in head, thoracic, or abdominal segments, whereas such transformations were readily observed in clones of *Sce* 

mutant cells (Fig. 2B,C). Because the majority of *Sce* 

mutant arrests development as late pupae without forming an adult exoskeleton (Supplemental Fig. S4B), we also analyzed HOX gene expression in these animals. The HOX genes *Ubx* and *Abd-B* remained fully silenced in imaginal discs of *Sce* 

mutant larvae but were widely misexpressed in cell clones lacking Sce protein (Fig. 2C). Taken together, these data suggest that H2Aub is not required for repression of HOX genes during post-embryonic development.

As in *Sce* 

mutant embryos (Supplemental Fig. S3), the morphological defects in *Sce* 

pharate adults were variable and showed no consistent pattern (Fig. 2B, cf. the two *Sce* 

individuals). The most conspicuous and consistent phenotype in these *Sce* 

animals was defective fusion of left and right hemisegments in thoracic and abdominal segments, accompanied by a loss of macrochaete and microchaete in the affected area (Fig. 2B, Supplemental Fig. S4B).
Cells containing H2A and H2Av that cannot be ubiquitylated maintain repression of PcG target genes

To complement the analysis of mutants lacking Sce E3 ligase activity, we generated animals in which wild-type H2A was replaced with a mutant form of H2A that cannot be ubiquitylated. On vertebrate nucleosomes, PRC1-type complexes ubiquitylate H2A at K119 and K121 but are unable to add ubiquitin to the more C-terminally located K125, K127, or K129 (Wang et al. 2004; Elderkin et al. 2007; McGinty et al. 2014). In Drosophila, the C terminus of H2A contains K117 and K118 (which correspond to K118 and K119 in mammals) and two further lysines nearby—K121 and K122 (Supplemental Fig. S5A). In vitro ubiquitylation reactions with the Ring1b/Bmi1 Ring finger module of mammalian PRC1 on reconstituted recombinant Drosophila nucleosomes revealed that only mutation of K117, K118, K121, and K122 completely abolished monoubiquitylation of H2A [Supplemental Fig. S5A]. For in vivo investigation of an H2A mutant that cannot be ubiquitylated, we therefore used a H2A<sup>K117R/K118R/K121R/K122R</sup> mutant (referred to below as H2A<sup>Δ−4R</sup>). In Drosophila, the canonical histone genes are all located in the histone gene cluster (HisC) that consists of 23 repeats of the histone gene unit comprising the H2A, H2B, H3, H4, and H1 genes. Transgene cassettes providing 12 copies of the wild-type histone gene unit rescue animals that are homozygous for a HisC deletion (HisC<sup>Δ−</sup>) into viable adults (Günesdogan et al. 2010). HisC<sup>Δ−</sup> homoyzogotes that carried the same transgene cassettes with a H2A<sup>Δ−4R</sup> mutant instead of wild-type H2A arrested development at the end of embryogenesis, showing an embryonic cuticle indistinguishable from wild-type embryos [data not shown]. We did not investigate the phenotype of these embryos further because their cells still contained maternally deposited wild-type H2A that had been only partially replaced by H2A<sup>Δ−4R</sup> during the few cell divisions that took place between the blastoderm stage and the end of embryogenesis. To analyze cells with a more complete replacement of H2A by H2A<sup>Δ−4R</sup>, we used a previously reported strategy (Pengelly et al. 2013) and generated clones of HisC<sup>Δ−</sup> homozygous cells in imaginal discs of HisC<sup>Δ−</sup> heterozygotes carrying the H2A<sup>Δ−4R</sup> transgene cassette. The HOX genes Ubx and Abd-B remained fully silenced in such H2A<sup>Δ−4R</sup> mutant cells (cf. Figs. 5A and 2). Moreover, clones of H2A<sup>Δ−4R</sup> mutant cells differentiated to form normal epithelial structures in the cuticle of adult flies (cf. Figs. 3B and 2). Repression of HOX genes is thus not impaired in cells in which H2A can no longer be ubiquitylated by PRC1.

In a final set of experiments, we explored whether repression of PcG target genes was also maintained in H2A<sup>Δ−4R</sup> mutant cells in which the histone variant H2Av could no longer be ubiquitylated. H2Av, a single-copy gene, encodes the only H2A variant in Drosophila and is not located in the HisC locus. H2Av contains a lysine pair, K120 and K121, at the position corresponding to K118/K119 in H2A and, in vivo, is also ubiquitylated by Sce (Supplemental Fig. S5B, cf. lanes 1–3). First, we constructed a Drosophila strain in which the H2Av gene was replaced by a H2Av<sup>K120R/K121R</sup> mutant [referred to below as H2Av<sup>Δ−2R</sup>]. The strategy to generate H2Av<sup>Δ−2R</sup>-null allele and H2Av<sup>Δ−2R</sup> mutant animals is presented in Supplemental Figure S6A–D and the Materials and Methods. H2Av<sup>Δ−2R</sup> mutants were viable and fertile and showed no obvious morphological defects even though the H2Av<sup>Δ−2R</sup> protein in these animals was no longer monoubiquitylated (Supplemental Fig. S5B, cf. lanes 4–6 and 1–3). Monoubiquitylation of H2Av by PRC1 is thus dispensable for normal development, viability, and fertility in Drosophila. Next, we generated clones of H2A<sup>Δ−4R</sup> and H2Av<sup>Δ−2R</sup> mutant cells in imaginal discs [Supplemental Fig. S6E]. Ubx and Abd-B remained fully silenced in H2A<sup>Δ−4R</sup> and H2Av<sup>Δ−2R</sup> double-mutant cells [Fig. 3C]. Together, these results argue that the ubiquitylation of H2A or H2Av is not critical for PRC1-mediated repression of HOX genes.

**Implications for mechanisms of PcG-mediated gene repression and H2Aub function**

Our analyses of developing Drosophila lacking H2Aub lead to the following main conclusions. First, monoubiquitylation of H2A and H2Av is dispensable for repression of canonical PRC1 targets such as the HOX genes or en. This suggests that nonenzymatic modification of chromatin (Shao et al. 1999; Francis et al. 2004) is the main mechanism by which PRC1 represses these genes. The requirement of Sce protein in canonical PRC1 may primarily reflect its architectural role in physically linking the H3-K27me3-binding activity of Pc [Fischle et al. 2003; Min et al. 2003] to the chromatin-compacting activities of the Psc and Ph subunits [Francis et al. 2004]. A second conclusion can be drawn from the reduction of bulk H3-K27me3 levels in embryos lacking H2Aub. On the one hand, this observation supports our earlier finding that H2Aub promotes H3-K27 trimethylation by PRC2 on nucleosomes in vitro [Kalb et al. 2014]. On the other hand, the reduction of H3-K27me3 is less than twofold, and, even though we do not know where...
it occurs in the genome, a drastic reduction at the HOX or ev genes seems unlikely because high H3-K27me3 levels at these genes are critical for repression [Nekrasov et al. 2007; Pengelly et al. 2013]. We conclude that canonical PcG target genes, PRC2 generates sufficient levels of H3-K27me3 independently of H2Aub. A third point to discuss is the phenotype and the cause of lethality of animals lacking H2Aub. SceKO m–z– embryos and SceKO 2– embryos that develop into pharate adults show remarkably subtle morphological defects that seem to occur stochastically in different locations of the body. Similar, even though the H2A lysine residues monoubiquitylated by Sce are critical for organism viability, H2A 2KR–KR mutant cells can differentiate to form normal epidermal structures. Our transcriptome analyses in SceKOA m–z– embryos failed to identify genes that are significantly misregulated in these embryos [data not shown]. It is possible that lethality in H2A-deficient animals is caused by a combination of multiple minor defects in regulating genes that control morphogenesis or, alternatively, a failure to regulate genes controlling changes in physiology or behavior at specific developmental time points. What these genes are and whether Sce E3 ligase activity regulates them as part of canonical PRC1 or another PRC1-type complex remain to be determined. Finally, we note that the requirement for H2Aub in developing Drosophila appears to be quite different from that in ES cells, where H2Aub was reported to be critical for repression of HOX and other PRC1 target genes and for maintenance of ES cell identity [Endoh et al. 2012]. It will be interesting to investigate the requirement of H2Aub during mouse development and find out how the role of this modification evolved to contribute to PRC1-mediated repression in higher metazoans.

Materials and methods

Drosophila strains and antibodies

Strain genotypes and antibodies used in this study are described in Supplemental Tables S1 and S2.

Preparation of nuclear and chromatin extracts

Soluble nuclear extracts from embryos were prepared as described [Scheuermann et al. 2010]. For chromatin extracts, the pellet obtained after ultracentrifugation was solubilized in urea buffer [8 M urea, 20 mM Tris at pH 8, 10 mM DTT] and sonicated (Bioruptor).

For small-scale preparations of nuclei, the subcellular protein fractionation kit for tissues (Pierce) was used. Pelleted nuclei were resuspended in pH 8, 10 mM DTT and sonicated (Bioruptor). Soluble nuclear extracts from embryos were prepared as described (Kalb et al. 2014).

Acid extraction of histones

Histones were acid-extracted from embryos or diploid larval imaginal disc and CNS tissues as described [Nekrasov et al. 2007].

Generation of SceKO m–z– embryos

The conditionally removable genomic Sce rescue construct >Sce< was based on a previously described vector [Gambetta and Müller 2014] that was modified to contain an attP site. The genomic Sce< fragment comprised chs3R sequences 27680208–27683747 [Berkeley Drosophila Genome Project [BDGP] R6]. The construct was integrated into the I27 [Bloomington Drosophila Stock Center [BDSC] 24482] attP landing site. One copy of the >Sce< transgene rescued SceKO homozygotes into viable and fertile flies that were indistinguishable from wild type. SceKO m–z– and SceKO 2– embryos were generated as described in Supplemental Figure S1 and were identified by loss of the GFP reporter in the >Sce< cassette. The SceKO transgene on chs2 [Supplemental Fig. S1] was integrated in attP site VK37 [BDSC 24872]; it contained the same genomic Sce fragment but with an I48A point mutation and lacked FRT-LoxP and GFP elements. An >Sce< transgene in VK37, used as a control, fully rescued SceKO homozygotes. Excision efficiency of the >Sce< cassette was 98%, as previously reported for another cassette [Gambetta and Müller 2014]. Plasmid maps are available on request.

Generation of the H2AubKO deletion allele

Ends-out recombination was used to disrupt H2Aub and replace its entire coding region [BDGP R6 chs3R: 26,866,929–26,867,220 by mini-white using a previously described strategy [Gong and Golic 2003]. Three-thousand-eight-hundred-twenty-one base pairs of H2Aub 5′ flanking sequences [BDGP R6 chs3R: 26,863,224–26,867,045] and 4881 bp of 3′ flanking sequences [BDGP R6 chs3R: 26,869,197–26,874,078] were cloned into pwp35 [Gong and Golic 2003]. Several independent targeting events that failed to complement the lethality of H2AubKO were isolated. Successful disruption of H2Aub was confirmed by PCR analysis with primers pairs [5′ to 3′] GACCTTGGACGCACTGTC and CACCAAAGGCTCAAC TACTG, ACTCTGTGACGACCTGAAC and CACATGGTTAGAT GCTCAG, and GCGCAGGTAAGTGCATC and CACCCCGTGGCA GTGCCT (Supplemental Fig. S5). The H2Aub rescue transgene H2Awt containing H2A coding and flanking sequences [BDGP R6 chs3R: 26,866,577–26,869,660] in a modified attB vector and was integrated into the VK37 attP site. The same strategy was used to generate animals carrying the H2AubKO transgene.

Generation of histone transgenes

Site-directed mutagenesis on pENTR223-HisGU.WT, pENTR41R-HisGU.WT, and pENTR2R3-HisGU.WT [Günesdogan et al. 2010] was used to mutate histone H2A specifically at Lys117, Lys118, Lys121, and Lys122 into arginines. The final construct pC31-attB3xHisGU.H2A-K117R/K118R/K121R/K122R, generated by Gateway LR recombination of the above vectors, was integrated at attP sites VK33 [BDSC 9750] and 86Fb [BDSC 130437].

Immunostaining and cuticle preparations of Drosophila

Immunostaining of embryos and imaginal discs and generation of clones in discs and adults were performed following standard protocols.

H2A ubiquitylation assays

In vitro ubiquitylation of recombinant Drosophila mononucleosomes were performed as described [Kalb et al. 2014].

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