A silencing pathway to induce H3-K9 and H4-K20 trimethylation at constitutive heterochromatin

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Histone lysine methylation is a central modification to mark functionally distinct chromatin regions. In particular, H3-K9 trimethylation has emerged as a hallmark of pericentric heterochromatin in mammals. Here we show that H4-K20 trimethylation is also focally enriched at pericentric heterochromatin. Intriguingly, H3-K9 trimethylation by the Suv39h HMTases is required for the induction of H4-K20 trimethylation, although the H4 Lys 20 position is not an intrinsic substrate for these enzymes. By using a candidate approach, we identified Suv4-20h1 and Suv4-20h2 as two novel SET domain HMTases that localize to pericentric heterochromatin and specifically act as nucleosomal H4-K20 trimethylating enzymes. Interaction of the Suv4-20h enzymes with HP1 isoforms suggests a sequential mechanism to establish H3-K9 and H4-K20 trimethylation at pericentric heterochromatin. Heterochromatic H4-K20 trimethylation is evolutionarily conserved, and in *Drosophila*, the *Suv4-20* homolog is a novel PEV modifier to regulate position-effect variegation. Together, our data indicate a function for H4-K20 trimethylation in gene silencing and further suggest H3-K9 and H4-K20 trimethylation as important components of a repressive pathway that can index pericentric heterochromatin.

[**Keywords:** Histone code; histone H4 Lys 20; mono-, di-, trimethylation; Suv4-20h HMTases; heterochromatin; combinatorial histone methyl marks]

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Chromatin is the physiological template of the genetic information and is composed of the DNA polymer, histones, and other chromosomal proteins. The basic repeating unit of chromatin is the nucleosome octamer, which is built by the core histones H2A, H2B, H3, and H4 wrapping 147 bp of DNA (Luger et al. 1997). Histone N termini (tails) are flexible and protrude from the nucleosome octamer. Histone tails are subject to post-translational modifications, including acetylation, methylation, phosphorylation, ubiquitination, and ADP-ribosylation (van Holde et al. 1988). During the recent years, many histone modifying enzymes and several chromatin-associated proteins that specifically bind to these modifications have been identified. Particular histone modification patterns have been associated with distinct

chromatin states and are proposed to represent an indexing mechanism that could extend the information potential of the genetic code (Strahl and Allis 2000; Turner 2000; Jenuwein and Allis 2001).

Histone modifications can act as either repressive or active marks. For histone lysine methylation, modification of H3-K4, H3-K36, and of H3-K79 have been correlated with transcriptional activation, whereas methylation of H3-K9, H3-K27, and H4-K20 are marks of repressive chromatin states (for review, see Fischle et al. 2003; Lachner et al. 2003; Vaquero et al. 2003). Histone lysine residues can be mono-, di-, or trimethylated (Paik and Kim 1971; DeLange et al. 1973), thereby extending the coding potential of a methylatable lysine position. Indeed, the distinctions between di- and trimethylation of H3-K4 (Santos-Rosa et al. 2002) or of H3-K9 residues (Czernin et al. 2002; Kuzmichev et al. 2002; Tamaru et al. 2003) have been shown to be relevant for transcriptional regulation or Polycomb-mediated gene silencing. Distinct methylation states or combinations between several methylation marks could further discriminate

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different chromatin regions or entire chromosomes. For example, H3-K27 trimethylation in conjunction with H3-K9 dimethylation are considered epigenetic imprints of the inactive X chromosome (Plath et al. 2003; Silva et al. 2003; Okamoto et al. 2004), whereas H3-K9 trimethylation and H3-K27 monomethylation are associated with pericentric heterochromatin (Peters et al. 2003; Rice et al. 2003). The combinatorial nature of histone lysine modifications requires controlled interplay between different histone lysine methylation systems. Loss of a given HMTase may also affect methylation on lysine residues for which the enzyme has no intrinsic activity. This is exemplified by the conversion of pericentric H3-K27 monomethylation to H3-K27 trimethylation in the absence of Suv39h enzymes (Peters et al. 2003).

Methylation of H4-K20 has been described as another mark of repressive chromatin domains (Nishioka et al. 2002; Sims et al. 2003). However, previous studies were focused toward detection of H4-K20 methylation, regardless of the methylation status and characterized PR/SET7 (Nishioka et al. 2002a; Rice et al. 2002) or SET8 (Fang et al. 2002) as the responsible enzyme. Here, we examine all three H4-K20 methylation states in mammalian chromatin with highly selective antibodies that discriminate H4-K20 mono-, di-, and trimethylation. We demonstrate that H4-K20 trimethylation is a novel and evolutionarily conserved mark of pericentric heterochromatin. In addition, we identify two murine SET domain HMTases that localize to pericentric heterochromatin and induce H4-K20 trimethylation in a *Suv39h*-dependent manner. In *Drosophila*, mutants of the corresponding gene locus are dominant suppressors of position-effect variegation. These data establish H4-K20 trimethylation as a repressive mark in gene silencing mechanisms and suggest that the sequential induction of H3-K9 and H4-K20 trimethylation by distinct histone lysine methylation systems can index repressive chromatin domains.

Results

H4-K20 trimethylation is a novel mark of pericentric heterochromatin

Pericentric heterochromatin in mammals mainly consists of major satellite repeats and can easily be visualized by the fluorochrome DAPI, which preferentially intercalates with A/T-rich repeat sequences. Differential association of H3-K9 and H3-K27 methylation states in euchromatin versus heterochromatin (Peters et al. 2003) prompted us to analyze other potentially repressive marks, such as histone H4-K20 methylation. Highly specific antibodies that discriminate H4-K20 methylation states were developed by Upstate Biotechnology (UBI) and quality controlled by dot blots presenting a panel of 23 histone tail peptides (Supplementary Fig. S1) and by Western blots of mouse nuclear extracts (Supplementary Fig. S2). Antibody specificity was further confirmed by peptide competition assays in immunofluorescence anal-

yses (Supplementary Fig. S3). By using these and our previously characterized H3-K9 methyl antibodies, we performed comparative immunofluorescence in wild type and *Suv39h* double-null (dn) female mouse embryonic fibroblasts (MEFs). In wild-type MEFs, H4-K20 monomethylation is dispersed in euchromatin but shows focal enrichment at the inactive X chromosome (Fig. 1A, arrow; Kohlmaier et al. 2004). H4-K20 dimethylation is also broadly distributed over euchromatic regions but shows a more speckled pattern, which resembles H3-K9 dimethylation (Fig. 1A). In contrast, H4-K20 trimethylation is strongly enriched at DAPI dense regions and reflects the characteristic accumulation of H3-K9 trimethylation at pericentric heterochromatin (Fig. 1A). We also analyzed H4-K20 methylation states in interphase chromatin of HeLa cells. H4-K20 mono- and dimethylation are uniformly distributed throughout the nuclei (Supplementary Fig. S4). Notably, H4-K20 monomethylation displays variable signal intensities, suggesting a potential cell-cycle regulation for this mark. In contrast, H4-K20 trimethylation is enriched at several nuclear foci, which represent pericentric heterochromatin as demonstrated by immunofluorescence analysis of metaphase spreads (Supplementary Fig. S4).

An interplay between distinct methylation systems at pericentric heterochromatin has been described previously (Peters et al. 2003). For example, disruption of the *Suv39h* enzymes results in the loss of H3-K9 trimethylation by also converting H3-K27 monomethylation to H3-K27 trimethylation. Because of the strikingly similar accumulation of H4-K20 trimethylation and H3-K9 trimethylation at pericentric heterochromatin, we analyzed whether H4-K20 methylation states may also depend on the presence of the *Suv39h* enzymes. In *Suv39h* dn female MEFs, H4-K20 mono- and dimethylation are not altered, but H4-K20 trimethylation is entirely lost from pericentric heterochromatin (Fig. 1A). These data were confirmed by analyzing metaphase chromosomes of wild-type and *Suv39h* dn MEFs. In wild-type mitotic spreads, H4-K20 trimethylation is strongly enriched at pericentric regions and shows a diffuse staining pattern along the chromosomal arms (Fig. 1B). In contrast, in mitotic spreads from *Suv39h* dn MEFs, pericentric H4-K20 trimethylation is lost, whereas the chromosomal arms display enriched signals (Fig. 1B).

From these data, we conclude that the presence of the *Suv39h* enzymes can direct pericentric H4-K20 trimethylation.

Identification of novel heterochromatic SET domain proteins

The requirement of *Suv39h* enzymes for pericentric H4-K20 trimethylation raised the question whether *Suv39h* enzymes might contain an intrinsic activity toward the H4-K20 position. However, in previous and extended *in vitro* HMTase assays (data not shown), the recombinant *Suv39h* enzymes only target the H3-K9 position, with a weak activity also toward histone H1 (Rea et al. 2000;

Figure 1. H3-K9 and H4-K20 methylation states in wild-type and *Suv39h* double-null (dn) MEFs. (A) Female wild-type and *Suv39h* dn MEFs were stained with antibodies directed against H3-K9 mono-, di-, and trimethylation (upper panel) or H4-K20 mono-, di-, and trimethylation (lower panel). DAPI dense foci represent pericentric regions. The inactive X chromosome is enriched for H4-K20 monomethylation and indicated by an arrow. (B) Metaphase spreads of wild-type and *Suv39h* dn MEFs were stained with the H4-K20 trimethylation antibody. Pericentric enrichment of this methylation mark is lost in *Suv39h* dn cells.

Peters et al. 2003). Therefore, other enzymes must exist that can trimethylate the H4-K20 position.

H4-K20 methylation is conserved among eukaryotes, such as *Schizosaccharomyces pombe*, *Caenorhabditis elegans*, *Drosophila*, and mammals, but is not present in *Saccharomyces cerevisiae* (Fang et al. 2002; data not shown). We therefore chose a candidate approach to identify H4-K20-specific HMTases by comparing all SET domain proteins that are shared among mouse, *Drosophila*, and *S. pombe*. We reasoned that HMTases with similar enzymatic specificities could be classified by their relative homology within the SET domain (Fig. 2). Within each of the resulting subgroups, the catalytic specificity of known SET domain HMTases is indicated (Fig. 2, gray boxes). Candidate proteins that are conserved between *Drosophila* and mouse were selected for further analyses and are shown by an asterisk (Fig. 2). Full-length IMAGE clones (RZPD) were used to express cDNAs as EGFP fusion proteins in MEFs under control of a cytomegalo virus (CMV) promoter. Of the 12 candidate cDNAs tested, Cgi-85 (*Suv4-20h1*, see below) and Mgc2705 (*Suv4-20h2*, see below) show a heterochromatic accumulation in wild-type MEFs, which is lost in *Suv39h* dn cells (Fig. 3B). In contrast, all other candidate SET domain proteins, including a previously described H4-K20 HMTase (Fang et al. 2002; Nishioka et al. 2002a;

Rice et al. 2002), displayed broad nuclear staining patterns (data not shown).

Cgi-85 (876 amino acids) and Mgc2705 (468 amino acids) are two closely related proteins present in mammals, whereas only one corresponding gene product is found in *Drosophila* and *S. pombe* (Supplementary Fig. S2). As we will show below, the *Drosophila* homolog, CG13363, is a dominant suppressor of position-effect variegation. We therefore classify Cgi-85 and Mgc2705 as novel Su(var) genes and suggest to rename these loci as *Suv4-20h1* and *Suv4-20h2*.

With the exception of the SET domain, which is located close to the N terminus, no other conspicuous domains could be identified in *Suv4-20h* proteins. However, the SET domain is surrounded by regions that are highly conserved within these orthologs. In addition, there is a small region (~10 amino acids) in the C termini that is shared between the various *Suv4-20h* proteins (Fig. 3A; Supplementary Fig. S2).

Suv4-20h proteins mediate H4-K20 trimethylation at pericentric heterochromatin

Next, we investigated the function of the putative *Suv4-20h* enzymes by RNA interference (RNAi) in MEF cells.

Figure 2. Neighbor-joining tree of mouse SET domain proteins. Sequences of mouse SET domain proteins were identified from public databases. SET domains sequences were aligned, and a neighbor-joining tree showing related SET domain proteins was constructed. Homologous protein sequences in *Drosophila* and *S. pombe* were identified by blast searches. HMTase specificity of subtrees was assigned according to the described HMTase activity of representative enzymes, such as H3-K4 [Ash1 [Beisel et al. 2002], Mil [Milne et al. 2002], Set7/9 [Nishioka et al. 2002b; Wang et al. 2001]]; H3-K9 [Suv39h1 [Rea et al. 2000], ESET [Yang et al. 2002], G9a [Tachibana et al. 2001]]; H3-K27 [Ezh2 [Kuzmichev et al. 2002; Müller et al. 2002]]; H3-K36 [Nsd1 [Rayasam et al. 2003]]; and H4-K20 [Pr-Set7 [Nishioka et al. 2002a; Fang et al. 2002]]. Proteins that were selected for the candidate approach are indicated by an asterisk.

MEFs were cotransfected with a pSUPER vector expressing hairpin RNAs and pEGFP-N1 conferring G418 resistance. Transfected cells were cultured for 5 d under selection medium and then analyzed for H4-K20 and H3-K9 methylation patterns by indirect immunofluorescence. As controls, we included RNAi for both Suv39h enzymes.

RNA hairpins directed against both Suv39h1 and Suv39h2 result in the loss of H3-K9 and H4-K20 trimethylation at pericentric heterochromatin (Fig. 3C). Thus, the Suv39h-specific RNAi knock-down accurately reflects the histone methylation defects observed in *Suv39h* dn cells (Fig. 1). In contrast, RNAi knock-down of both Suv4-20h1 and Suv4-20h2 selectively impair pericentric accumulation of H4-K20 trimethylation, but do not perturb H3-K9 trimethylation (Fig. 3C).

Suv4-20h proteins are nucleosomal HMTases with specificity for H4-K20 trimethylation

Next, we characterized the activity and substrate specificity of Suv4-20h proteins. N-terminal fragments of

Suv4-20h1 (amino acids 1–387) and Suv4-20h2 (amino acids 1–280) containing the SET domain were bacterially expressed as GST fusion proteins. Increasing amounts of fusion protein were used in a reaction mix containing ³H-labeled SAM as methyl donor, and recombinant histone H4, histone octamers, or nucleosomes. Reaction products were separated by SDS-PAGE, and methyl transfer to histones was detected by fluorography. Both Suv4-20h1 and Suv4-20h2 show some HMTase activity toward recombinant octamers but display a significantly higher transfer reaction toward nucleosomes. Peptides and free histones were not accepted as substrates (Fig. 4A; data not shown). The HMTase activity appears to be specific for H4-K20, because no methyl transfer is detected when nucleosomes containing a mutated histone H4 (K20A) were used.

Because histone lysine positions can be mono-, di-, or trimethylated, we next investigated the methylation state of the reaction products. Bacterially expressed N termini of Suv4-20h1 and Suv4-20h2 were used in an HMTase assay with nonradioactive SAM and recombinant substrates. Reaction products were separated by SDS-PAGE, and the methylation state of histone H4 was

Figure 3. Identification of Suv4-20h1 and Suv4-20h2 as heterochromatic H4-K20 trimethylating HMTases. (A) Protein structure of Suv4-20h proteins. The SET domain is located in the N terminus and flanked by N- as well as C-terminal extensions of high sequence homology within Suv4-20h orthologous proteins. In addition, there is a conserved region in the C terminus of these proteins. (B) EGFP-tagged Suv39h1, Suv4-20h1, and Suv4-20h2 were expressed in wild-type and *Suv39h* double-null (dn) female MEFs. After fixation, distribution of EGFP-tagged proteins was analyzed by fluorescence microscopy. DAPI dense foci represent pericentric heterochromatin. (C) RNAi mediated knock-down of Suv39h and Suv4-20h proteins. Female wild-type MEFs were cotransfected with pEGFP-N1 as transfection marker and pSUPER vector expressing hairpin oligos directed against Suv39h1/Suv39h2 or Suv4-20h1/Suv4-20h2. After 5 d in selection medium, cells were stained with α -trimethyl H3-K9 and α -trimethyl H4-K20 antibodies. Immunostaining was examined in EGFP-positive cells.

analyzed by probing Western blots with H4-K20 mono-, di-, and trimethyl-specific antibodies. No H4-K20 monomethylated product could be observed (Fig. 4B). The weak signal visualized by the H4-K20 monomethyl antibody is a cross-reactivity with unmodified histone H4 (shown by an asterisk). In contrast, reaction products of Suv4-20h enzymes display some H4-K20 dimethylation but are strongly enriched for H4-K20 trimethylation (Fig. 4B). The absence of H4-K20 monomethylation in reaction products of Suv4-20h2 was further confirmed by a time-course experiment. Reactions of Suv4-20h2 with recombinant nucleosomes were stopped after sequential incubation times and analyzed by Western blotting. In the reaction products, H4-K20 di- and trimethylation increased over time, but even at very early time points (2 min), no H4-K20 monomethylation is detected (Supplementary Fig. S3). The collective interpretation of the results presented above allow us to conclude that Suv4-20h enzymes are nucleosomal HMTases with a preferential activity to direct H4-K20 trimethylation.

Targeting of Suv4-20h2 by HP1 interactions

To identify the targeting mechanism(s) of Suv4-20h enzymes to heterochromatic regions, we expressed various protein truncations as EGFP fusions in MEFs. For this analysis we focused on Suv4-20h2 (468 amino acids). The N terminus of Suv4-20h2 (S#1: amino acids 1–280) comprising the SET domain is broadly dispersed in the nucleus, whereas the C terminus (S#2: amino acids 280–468) shows prominent association with pericentric heterochromatin (Fig. 5A). To further define the domain required for heterochromatic targeting, we also analyzed subfragments of the C terminus. One subfragment (S#4: amino acids 280–349) shows aberrant nuclear distribution. Although there are EGFP speckles in the nucleus, these do not correspond to DAPI dense heterochromatin. All other protein fragments display heterochromatic associations. Thus, the minimal heterochromatic targeting module of Suv4-20h2 is localized within amino acids 348–441.

A possible mechanism for Suv39h-dependent targeting of Suv4-20h2 could be an interaction with other heterochromatic proteins, such as HP1, the localization of which also depends on Suv39h function (Bannister et al. 2001; Lachner et al. 2001). Therefore, we investigated whether Suv4-20h2 could directly interact with HP1 isoforms. Protein fragments of the above-mentioned localization assay were generated as recombinant GST fusion proteins and bound to glutathione-Sepharose beads. Full-length myc-tagged HP1 α , HP1 β , and HP1 γ were expressed by *in vitro* translation and incubated with GST-Suv4-20h2 protein variants. After extensive washing steps, bound proteins were separated on SDS-PAGE, and Western blots were probed with α -myc antibody for detection of HP1 isoforms. The N terminus of Suv4-20h2 cannot bind HP1 isoforms, but the C terminus interacts with HP1 α , HP1 β , and HP1 γ (Fig. 5B). Recombinant GST does not interact with HP1, excluding unspecific binding to the GST tag (Fig. 5B). Subfragments of the C terminus show a differential interaction pattern with HP1 isoforms. Suv4-20h2 S#5, which was found to be the minimal heterochromatic targeting module (Fig. 5A), is also the minimal subfragment that interacts with HP1 α , HP1 β , and HP1 γ . However, the aberrantly localized subfragment S#4 does not interact with HP1 α or HP1 β but can still associate with HP1 γ . Thus, although there are several distinct HP1 binding sites in amino acid region 280–441 (which are different from the conserved C-terminal domain), it is the interactions with HP1 α and HP1 β that appear to localize Suv4-20h2 to pericentric heterochromatin (Fig. 5C).

The *in vitro* binding data together with the heterochromatic localization of Suv4-20h2 suggest a directed targeting of this HMTase by interaction with HP1. If this model is correct, Suv4-20h enzymes and H4-K20 trimethylation should be lost in HP1-deficient cells. This question cannot easily be addressed in mammals, because of the lack of available HP1 mutants. Moreover, our attempts using RNAi-mediated knock-down of all three HP1 isoforms in MEFs failed to deplete HP1 func-

Figure 4. Substrate and product specificity of Suv4-20h HMTases. (A) Recombinant N termini of Suv4-20h1 (amino acids 1–387) and Suv4-20h2 (amino acids 1–280) were tested in HMTase assays with recombinant histone H4 (rH4), recombinant histone octamers (rOct), recombinant nucleosomes (rNuc), and recombinant nucleosomes containing a point mutation in H4 (K20A). The *top* panel shows HMTase activity; the *bottom* panel shows coomassie staining of the membrane used for the HMTase assay. (B) Reaction products of HMTase assays, as in A, were Western blotted and probed with α -monomethyl, α -dimethyl, and α -trimethyl H4-K20 antibodies (*top* panels, antibody; *bottom* panels, Ponceau staining). As a control, recombinant and native nucleosomes that were not processed by the enzyme were loaded (asterisks indicate slight cross-reactivities, observed with long exposure times).

tion. We therefore extended our studies to *Drosophila*, in which H4-K20 trimethylation is present at pericentric heterochromatin and in a high number of bands (as judged by comparison with propidium iodide labeling of DNA) on salivary gland chromosomes (Fig. 6). In *Dro-*

sophila, pericentric H4-K20 trimethylation is severely reduced in *Su(var)3-9* mutants (Fig. 6A), indicating that cross-talk between H3-K9 and H4-K20 trimethylation systems is evolutionarily conserved. In *Su(var)2-5* (HP1) mutants, H3-K9 methylation and pericentric accumula-

Figure 5. Definition of a heterochromatic targeting module in Suv4-20h2. (A) Various truncations of Suv4-20h2 (S#1–S#6) were expressed as EGFP fusion proteins in female wild-type MEFs. Cells were fixed, and distribution of fusion proteins was examined with fluorescence microscopy. DAPI dense foci represent pericentric regions. Note, that EGFP foci of subfragment S#4 do not represent pericentric heterochromatin, because they do not colocalize with DAPI dense regions. (B) GST-tagged protein truncations of Suv4-20h2 (S#1–S#6) as well as recombinant GST were bound to glutathione-Sepharose and incubated with *in vitro* translated myc₃-tagged HP1 α , HP1 β , and HP1 γ . After extensive washing steps, bound proteins were separated on SDS-PAGE, and Western blots were probed with α -myc antibodies. (C) Summary table showing structure of Suv4-20h2 protein truncations together with localization and HP1 interaction data. (HET) Heterochromatin; (EU) euchromatin aberrant.

Figure 6. *Drosophila Suv4-20* is a dominant PEV modifier. (A) Polytene chromosomes of wild-type, *Su(var)3-9* null, homozygous *Suv4-20*^{BG00814}, and HP1 null larvae were squashed and stained with α -trimethyl H4-K20 antibody (top panel, DNA staining with propidium iodide; bottom panel, antibody staining; arrow points to chromocenter). (B) *Stubble* variegation in wild-type (top) and *Suv4-20*^{BG00814} mutant flies (bottom). Short bristles (arrow) indicate activation of *Sb*^V in *Suv4-20*^{BG00814} flies. (C) Quantification of *Stubble* position effect variegation in wild-type and *Suv4-20*^{BG00814} mutant flies. Wild-type and *Suv4-20*^{BG00814} females were crossed to *T(2;3)Sb*^V males. The percentage of *Stubble* bristles that were scored in males and females of the progeny is indicated in the table.

tion of *SU(VAR)3-9* remains unaltered (Schotta et al. 2002). Intriguingly, H4-K20 trimethylation is nearly lost from polytene chromatin in these mutants (Fig. 6A). This result indicates that the induction of heterochromatic histone lysine methylation marks occurs in a sequential pathway. At pericentric heterochromatin *Su(var)3-9*-mediated H3-K9 methylation would precede H4-K20 trimethylation by the *Suv4-20* enzymes. Other H3-K9-specific HMTases could direct HP1 and subsequently H4-K20 trimethylation to euchromatic bands.

Drosophila Suv4-20 is a dominant suppressor of position-effect variegation

Our data suggest H4-K20 trimethylation to be a mark of silenced chromatin domains. We therefore investigated whether this modification would indeed be important for gene silencing in well-described PEV models in *Drosophila* (Reuter and Spierer 1992). A single, homozygous-viable P-element insertion (*P{GT1/BG00814}*) into the third exon of *Suv4-20* has been identified in the course of the *Drosophila* gene disruption project (Flybase). H4-K20 trimethylation at polytene chromatin is nearly lost in homozygous mutant larvae (Fig. 6A), demonstrating that the P-element insertion (*Suv4-20*^{BG00814}) represents a strong hypomorphic allele of *Suv4-20*. Because the *Suv4-20* locus maps on the X chromosome, the classical PEV rearrangement *In(1)w^{m4}* (Reuter and Spierer 1992) cannot be used to analyze a potential modifier effect of *Suv4-20*. Therefore, we analyzed another PEV rearrangement that translocates a different marker, *Stubble* (*Sb*), close to pericentric heterochromatin (*T(2;3)Sb*^V; Sinclair

et al. 1983). The dominant mutation *Stubble* induces short bristles, but heterochromatin-induced silencing of *Sb*^V results in wild-type (long) bristles (Fig. 6B). Homozygous *Suv4-20*^{BG00814} as well as control wild-type females were crossed to *T(2;3)Sb*^V males. In the progeny, the extent of *Sb*^V reactivation was determined as the ratio of short bristles (active *Sb*^V) to long bristles (inactive *Sb*^V). In males and females of the wild-type crosses, only 1%–2% of bristles show a *Sb* phenotype, indicating that *Sb*^V is largely inactivated (Fig. 6B). In contrast, *Sb*^V becomes derepressed in the progeny of *Suv4-20*^{BG00814} flies, because now ~25% of the bristles are short (Fig. 6B). This result classifies *Suv4-20* as a dominant PEV modifier and further indicates a functional role for *Suv4-20*-dependent H4-K20 trimethylation in gene silencing.

Discussion

A combinatorial histone methyl code for constitutive heterochromatin

One of the predictions of the “histone code” hypothesis suggests that distinct combinations of histone modifications participate in the formation of different chromatin domains (Strahl and Allis 2000; Turner 2000; Jenuwein and Allis 2001). Of the five lysine positions in the histone N termini that display prominent methylation (Peters et al. 2003), H3-K9, H3-K27, and H4-K20 are supposed to have repressive functions (Fischle et al. 2003; Lachner et al. 2003; Vaquero et al. 2003). A combination of H3-K27 trimethylation together with H3-K9 dimethylation and H4-K20 monomethylation has been described to index the inactive X chromosome (Silva et al.

2003; Plath et al. 2003; Kohlmaier et al. 2004). Prominent marks of constitutive heterochromatin are H3-K9 trimethylation and H3-K27 monomethylation (Peters et al. 2003; Rice et al. 2003). However, the contribution of the three distinct H4-K20 methylation states in the formation of chromosomal subdomains has not been analyzed. Previous studies indicated H4-K20 methylation is broadly enriched at mitotic chromosomes (Fang et al. 2002; Nishioka et al. 2002a; Rice et al. 2002). By using highly selective antibodies, developed by UBI, which discriminate H4-K20 mono-, di-, and trimethylation, we now describe H4-K20 trimethylation as a third component of repressive histone lysine methyl marks in pericentric heterochromatin.

Based on our novel data, we propose the following model for sequential induction of H3-K9 and H4-K20 trimethylation at constitutive heterochromatin in mammals (Fig. 7). In a first step, Suv39h enzymes would be targeted to repeat-rich sequences at pericentric heterochromatin, presumably by interacting with protein complexes containing components of the RNAi machinery (Hall et al. 2002; Jenuwein 2002; Volpe et al. 2002; Schramke and Allshire 2003; Pal-Bhadra et al. 2004; Verdell et al. 2004). Because pericentric heterochromatin is also enriched for H3-K27 monomethylation, and the Suv39h enzymes prefer an H3-K9 monomethylated substrate (Peters et al. 2003), this first step most likely also requires the activity of currently unknown H3-K27 and

H3-K9 monomethylases. Following stabilized binding of HP1 α and HP1 β to H3-K9 trimethylated nucleosomes (Bannister et al. 2001; Lachner et al. 2001), HP1 molecules, probably in conjunction with nucleosomal surfaces, would then recruit the Suv4-20h enzymes, which in turn trimethylate H4-K20. This tentative model would involve collaboration of at least four distinct HMTases to induce the observed combination of H3-K9 trimethyl, H4-K20 trimethyl, and H3-K27 monomethyl marks at pericentric heterochromatin. Future studies are aimed to identify whether HP1 binding is stabilized by a combination of H3-K9 and H4-K20 trimethylation or if other heterochromatin-specific components may exist that could selectively be recruited by this combinatorial histone lysine methylation pattern.

Suv4-20h enzymes are nucleosomal-specific HMTases

Recombinant Suv4-20h HMTases possess selective activity toward the H4-K20 position in a nucleosomal context. Recombinant histone H4 is not used as a substrate, and only poor activity is detected with histone octamers (Fig. 4). In contrast, Suv39h enzymes preferably methylate free histone H3 and recombinant octamers, whereas nucleosomes are only very poor in vitro substrates (data not shown). Thus, selective generation of H3-K9 and H4-K20 methylation marks could occur in a stepwise mechanism (Vermaak et al. 2003), in which histone H3-H4 dimers (Tagami et al. 2004) or tetramers might be methylated by Suv39h and, after nucleosome assembly, Suv4-20h enzymes would trimethylate the H4-K20 position.

In vitro, Suv4-20h enzymes can directly trimethylate unmodified H4-K20 positions (Fig. 4), without requirement for a monomethyl substrate. This is in contrast to Suv39h function, which prefers an H3-K9 monomethylated position (Peters et al. 2003). In *Suv39h* dn cells, H3-K9 monomethylation becomes enriched at pericentric heterochromatin (Peters et al. 2003); however, we did not detect an increase for H4-K20 monomethylation in *Suv4-20h* RNAi knockdown experiments (data not shown). In addition, H3-K9 trimethylation is not required for Suv4-20h activity in vitro (Fig. 4), although cross-tail interactions between nucleosomal H3-K9 and H4-K20 positions may exist in vivo that could affect enzymatic activity of Suv4-20h HMTases.

PR/SET7 is another HMTase with selective activity toward nucleosomal H4-K20 (Nishioka et al. 2002a; Fang et al. 2002), but in contrast to Suv4-20h enzymes, it appears to be an exclusively monomethylating enzyme (K. Sarma, D. Reinberg, and S. Gamblin, unpubl.). Furthermore, PR/SET7 fails to localize to pericentric heterochromatin when expressed as an EGFP fusion protein (data not shown). N- and C-terminal extensions of the SET domain, such as pre- and post-SET domains of Suv39h enzymes, are involved in substrate binding and in conferring enzyme selectivity toward mono-, di-, and trimethylation (Xiao et al. 2003). Suv4-20h enzymes contain extensions to the SET domain that are distinct from pre- and post-SET domains, and that are only conserved

Figure 7. A conserved pathway for H3-K9 and H4-K20 trimethylation at pericentric heterochromatin. Proposed model indicating sequential induction of H3-K9 and H4-K20 trimethylation at pericentric heterochromatin. See text for more detailed explanation. H3-K9 monomethylation is presumably an intermediate state (in parentheses), which is converted to H3-K9 trimethylation by Suv39h enzymes. Hexagons indicate histone lysine methylation states (H3-K9, red; H3-K27, pink; H4-K20, blue).

within their subgroup of orthologous proteins. In addition, there is no significant homology to PR/SET7 within these conserved regions of Suv4-20h enzymes.

Suv4-20h HMTases and PR/SET7 possibly recognize the H4-K20 position in a different nucleosomal context. Indeed, the two H4 tails of one nucleosome are presented in different molecular projections, such that one H4 tail is directed toward the DNA backbone, whereas amino acids 16–25 of the second histone H4 molecule contact the H2A/H2B dimer of the adjacent nucleosome (Luger et al. 1997). It is according to these structural definitions that the H4 tail may be particularly suited for directing transitions between 11- and 30-nm chromatin fibers, and H4-K20 trimethylation appears as a good candidate mark to contribute in facilitating these higher order nucleosomal organizations.

H4-K20 trimethylation is evolutionarily conserved

In mammals, ~50 SET domain genes have been identified (Lachner and Jenuwein 2002; Kouzarides 2002), of which a subset is evolutionarily conserved in *Drosophila* and *S. pombe*. Intriguingly, nearly all putative SET domain HMTases that target repressive lysine positions (H3-K9, H3-K27, H4-K20) are absent in *S. cerevisiae*. In agreement, H3-K9, H3-K27, and H4-K20 methylation cannot be detected in *S. cerevisiae* (R. Sengupta and T. Jenuwein, unpubl.), indicating that this organism largely lacks extensive heterochromatic domains and uses other repressive systems (e.g., SIR proteins) to silence gene activity (Kurdistan and Grunstein 2003).

H3-K9 methylation has emerged as an important and evolutionarily conserved mark to silence gene activity at repressive chromatin domains. The concerted function of Suv39h and Suv4-20h enzymes described here raises the question as to the functional role of H4-K20 methylation in well-described epigenetic paradigms. Homologs of the *Suv39h/Suv4-20h* methylation systems are also conserved in *Drosophila* (*Su(var)3-9* and *Suv4-20*) and *S. pombe* (*Clr4* and *SET9*). Surprisingly, H3-K9 methylation appears severely underrepresented (and can hardly be detected) in bulk histone preparations from *S. pombe* extracts, which instead display robust signals for all three H4-K20 methylation states (R. Sengupta and T. Jenuwein, unpubl.). Because HP1 is also conserved in *S. pombe* (as the Swi6 protein), a functional synergy between H3-K9 and H4-K20 methylation systems might exist in *S. pombe* that could be operative to safeguard gene silencing and/or to index distinct chromosomal regions. Together, our definition of novel H4-K20 trimethylating HMTases and the presence of H4-K20 trimethylation from *S. pombe* to mammals provides evidence for an evolutionary conservation of the proposed silencing pathway between H3-K9 and H4-K20 methylation.

Suv4-20h enzymes are novel Su(var) genes

H4-K20 trimethylation is a novel hallmark of pericentric heterochromatin in mammals, but the current lack of

mutants for Suv4-20h enzymes prevents the analysis of its regulatory function during mouse development. However, the identification of a P-element insertion in *Drosophila Suv4-20*, which results in nearly complete loss of H4-K20 trimethylation (Fig. 6A), allowed a first functional analysis. In *Drosophila*, there are ≥ 50 *Su(var)* loci (Reuter and Spierer 1992; Schotta et al. 2003), which encode products involved in heterochromatin-induced gene silencing. Only ~15 of these *Su(var)* genes have been identified to date, the most revealing of which include *Su(var)2-5* (HP1; Eissenberg et al. 1990), *Su(var)3-7* (Cleard et al. 1997), and *Su(var)3-9* (Tschiersch et al. 1994). Here we show that *Drosophila Suv4-20* is a novel, dominant suppressor of position-effect variegation (Fig. 6B). The *Su(var)* effect of *Suv4-20* is about as strong as that of *Su(var)3-9* (data not shown), suggesting that H3-K9 and H4-K20 trimethylation may have a similar potential to establish heterochromatin-induced gene silencing. In addition, both the hypomorphic *Suv4-20* and the *Su(var)3-9* null mutants are homozygous viable. Based on the observed synergy, it is conceivable that the combined disruption of *Suv4-20* and *Su(var)3-9* genes might have a much more pronounced phenotype that could compromise establishment and stability of heterochromatin.

Major functions of heterochromatin include to safeguard a specialized chromatin structure around centromeres (Karpen and Allshire 1997; Bernard and Allshire 2002) and to protect genome stability by silencing of transposable elements (Birchler et al. 2000; Pal-Bhadra et al. 2004). The pericentric accumulation and the silencing defect observed in *Suv4-20* mutants predict H4-K20 trimethylation to participate in these mechanisms, but H4-K20 trimethylation may also be involved in gene repression at euchromatic targets or other functions of chromatin control, in which it could even be independent of Suv39h activity. As histone lysine methylation is implicated in almost all epigenetic paradigms, and H4-K20 trimethylation has been shown to be enriched in aged organs (Sarg et al. 2002), future analyses into the physiological roles of H4-K20 methylation promise not only to provide a better understanding of the interplay between distinct histone lysine methylation systems but may even offer novel insights into fundamental biological processes, such as normal and perturbed development, aging, and regeneration.

Materials and methods

Plasmid construction

Full-length open reading frames of all candidate SET domain proteins as well as protein truncations of Suv4-20h1 and Suv4-20h2 were PCR amplified from IMAGE cDNA clones (purchased from Deutsches Ressourcenzentrum für Genomforschung GmbH (RZPD)) and cloned into the GATEWAY entry vector pDONR207 (Invitrogen). A list of clones used as well as primer sequences is available on request. The reference sequences of Suv4-20h1 and Suv4-20h2 have been deposited at GenBank (AY555192, AY555193).

The two GATEWAY compatible target vectors for expression of EGFP-tagged proteins in mammalian cells (pEGFP-GW) and bacterial expression of GST-tagged proteins (pGEX-GW) were constructed by inserting GATEWAY cassettes into SmaI sites of pEGFP-N1 (Clontech) and pGEX-6P1 (Amersham), respectively. Entry clones were recombined into respective target vectors by GATEWAY LR Reaction.

Transient transfection and immunofluorescence on interphase cells

Candidate SET domain proteins were expressed in female wild-type and *Suv39h* dn cells. In each chamber of a multiwell slide, 5000 cells were plated 1 d before transfection. Cells were transfected with 1 µg plasmid DNA by using the Lipofectamine protocol (Invitrogen). One day posttransfection, cells were fixed in 2% PFA/PBS, extracted with 0.1% sodium citrate/0.1% Triton X-100, washed with PBS, and mounted with Vectashield (Vector Laboratories). EGFP fluorescence was analyzed by fluorescence microscopy. Indirect immunofluorescence was performed according to the method of Peters et al. (2003), using antibodies against H4-K20 mono-, di-, and trimethylation (Upstate Biotech; nos. 07-440, 07-367, 07-463) and H3-K9 mono-, di-, and trimethylation (Peters et al. 2003).

RNAi knock-down

DNA oligos containing the 19mer targeting sequence, a loop region, and HindIII/BglII restriction sites were cloned into pSUPER (Brummelkamp et al. 2002). For each target gene, two different 19mer sequences were used. Fifty thousand wild-type MEFs were plated on a 10-cm dish 1 d before transfection. Cells were transfected with a mixture of 2 µg pSUPER plasmids containing the 19mer targeting sequences and 0.2 µg pEGFP-N1 using Lipofectamine (Invitrogen). One day after transfection selection, medium containing 600 µg/mL G418 was applied to the cells. Five days after transfection, cells were transferred on chamber slides and analyzed by indirect immunofluorescence as described (Peters et al. 2003). Because pEGFP-N1 vector was underrepresented in the transfection mix, cells showing EGFP fluorescence were assumed to also contain pSUPER vector.

HMTase assays

GST fusion proteins of *Suv4-20h1* (amino acids 1–387) and *Suv4-20h2* (1–280) containing the SET domain and N- as well as C-terminal associated regions (Fig. 3A) were expressed in *Escherichia coli*. HMTase assays were performed as described (Nishioka et al. 2002b). In particular, samples of the *Suv4-20h* enzymes were incubate for 1 h at 30°C in reaction buffer (50 mM Tris-HCl at pH 8.5, 5 mM MgCl₂, 4 mM DTT) with 1 µM ³H-labeled SAM (Amersham Pharmacia Biotech). Fifty nanograms of recombinant histone H4 and 2 µg of recombinant octamers/nucleosomes were used as substrates.

In vitro binding assays

GST fusion proteins of *Suv4-20h2* were expressed in *E. coli* and purified on glutathione-S-Sepharose (Pharmacia). Equal amounts of GST fusion proteins were incubated with in vitro translated myc₃-tagged HP1 isoforms (HP1α, amino acids 5–191; HP1β, amino acids 5–185; HP1γ, amino acids 5–173; Lachner et al. 2001) in IP buffer (50 mM Tris-HCl at pH 7.5, 500 mM NaCl, 1 mM EDTA, 0.5% NP-40, 10% glycerol, proteinase inhibitor cocktail). Incubation was carried out overnight at 4°C with gentle agitation. The beads were washed six times with IP

buffer and finally resuspended in 50 µL loading buffer. Bound proteins were separated on SDS-polyacrylamide gels and detected by immunoblotting using α-myc antibody (Upstate Biotech; 05-419).

Immunostaining of polytene chromosomes

Preparation of polytene chromosomes was performed as described (Schotta et al. 2002) with the following modifications: salivary glands of third instar larvae were dissected in 0.7% NaCl, fixed in 4% formaldehyde and squashed in 45% acetic acid/2% formaldehyde. Chromosomes were incubated with α-trimethyl H3-K9 (Peters et al. 2003) as well as α-trimethyl H4-K20 antibodies overnight at 4°C, followed by incubation with secondary alexA 488-conjugated antibodies (Molecular Probes) for 2 h at 37°C. DNA was stained with propidium iodide (Molecular Probes). Preparations were mounted in Vectashield medium and examined with a confocal laser scanning microscope (LSM510, Zeiss).

Phenotypic characterization of Stubble variegation

The P-element P[GT1]BG00814 is inserted into the third exon of *Suv4-20* (CG13363) and results in a strong hypomorphic allele (*Suv4-20^{BG00814}*). The PEV modifier effect of *Suv4-20^{BG00814}* was evaluated by using *T(2;3)Sb^V* (Sinclair et al. 1983). This translocation brings the dominant mutation *Sb¹* (short-bristles phenotype) near to chromosome 2R heterochromatin. Because *Sb^V* is dominant, Su(var)s increase the stubble phenotype. *Suv4-20^{BG00814}/Suv4-20^{BG00814}* as well as control wild-type females were crossed to *T(2;3)Sb^V*, *In(3R)Mo*, *Sb¹*, *sr¹/TM3* males, and 14 defined bristles were scored as being wild type or *Sb*. The ratio between the total numbers of *Stubble* to wild-type bristles is a mark for the extent of *Sb* inactivation and has been evaluated for >25 flies of each genotype.

Bioinformatic methods

SET domain proteins in mouse were identified by using a hidden Markov model (HMM) profile for the SET domain (Pfam, PF00856). IPI database (<http://www.ebi.ac.uk/IPI>) was searched by using HMMER software (<http://hmmer.wustl.edu>). An alignment of all SET domains and the neighbor-joining tree was created by using the t-coffee alignment program (Notredame et al. 2000). Orthologs of mouse SET domain proteins in *S. pombe* and *Drosophila* were identified by using protein blast searches (<http://www.ncbi.nlm.nih.gov/blast>).

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A silencing pathway to induce H3-K9 and H4-K20 trimethylation at constitutive heterochromatin

Gunnar Schotta, Monika Lachner, Kavitha Sarma, et al.

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