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Restricted epigenetic inheritance of H3K9 methylation

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Abstract

Post-translational histone modifications are believed to allow the epigenetic transmission of distinct chromatin states, independently of associated DNA sequences. H3K9 methylation is essential for heterochromatin formation, however, a demonstration of its epigenetic heritability is lacking. Fission yeast has a single H3K9 methyltransferase, Clr4, that directs all H3K9 methylation and heterochromatin. Utilizing releasable tethered Clr4 reveals that an active process rapidly erases H3K9 methylation from tethering sites in wild-type cells. However, inactivation of the putative histone demethylase Epe1 allows H3K9 methylation and silent chromatin maintenance at the tethering site through many mitotic divisions, and transgenerationally through meiosis, after release of tethered Clr4. Thus, H3K9 methylation is a heritable epigenetic mark whose transmission is usually countered by its active removal, which prevents the unauthorised inheritance of heterochromatin.
heterochromatin nucleation, spreading and maintenance (13-15). The inheritance of heterochromatin on centromere repeat DNA inserted at ectopic locations also requires RNAi and mating-type locus heterochromatin is dependent on DNA binding factors in the absence of RNAi (13-17). However, constitutive tethering of Clr4 to a euchromatic locus via the Gal4 DNA binding domain (GBD) allows the assembly of an extensive domain of H3K9me-heterochromatin, independently of RNAi (18). Here we tether a regulatable TetR\textsuperscript{off}-Clr4 fusion protein to determine if H3K9me is a persistent histone modification that can be stably copied through mitotic cell divisions and meiosis following release or loss of the TetR\textsuperscript{off}-Clr4 initiator.

TetR\textsuperscript{off}-2xFLAG-Clr4-cdd fusion protein (abbreviated TetR-Clr4\textsuperscript{*}), lacking the Clr4 chromodomain, was stably expressed in cells with an ade6\textsuperscript{+} gene downstream of 4xtetO binding sites at the ura4 locus (4xtetO-ade6\textsuperscript{+}; Fig. 1A) (19). TetR-Clr4\textsuperscript{*} silences 4xtetO-ade6\textsuperscript{+} independently of RNAi (ago1Δ, dcr1Δ), similar to GBD-Clr4 (18), resulting in reduced RNAPII association, and high H3K9me2 levels and silencing over a broad region (Fig. 1C, fig. S1, and fig. S9C-E). TetR-Clr4\textsuperscript{*} is released within 5 minutes from tetO sites by addition of anhydrotetracycline (AHT; Fig. 1B). All strains utilised also express wild-type Clr4 which can interact via its chromodomain with TetR-Clr4-directed H3K9me and thus potentially use its read-write capabilities to methylate newly incorporated H3 and allow heterochromatin transmission to daughter cells following TetR-Clr4\textsuperscript{*} release. However, in a time course, H3K9me2 rapidly declines over 4xtetO-ade6\textsuperscript{+} following AHT addition; >90\% is lost within six hours (Fig. 1C, and fig. S2A). AHT itself does not affect endogenous heterochromatin integrity (fig. S2B). H3 levels do not decline on 4xtetO-ade6\textsuperscript{+} over this period (fig. S2A). Swi6\textsuperscript{HP1} is also lost from 4xtetO-ade6\textsuperscript{+} when cells are grown with AHT (fig. S1F).

We also tethered TetR-Clr4\textsuperscript{*} within two non-essential genes with long open reading frames, which are less likely to contain unannotated features that might interfere with heterochromatin integrity. Moreover, both sib1\textsuperscript{+} (15,005 bp) and vps1302\textsuperscript{+} (9,200 bp) exhibit expression levels and rates of H3 turnover that are ~3-fold lower than those of ade6\textsuperscript{+} (Fig. 2A-C, and fig. S3). 4xtetO and 1xtetO sites were placed within sib1 and vps1203, respectively (Fig. 2D and E). sib1:4xtetO and vps1302:1xtetO were also placed under the control of low, medium (med), and high versions of the constitutive adh1 promoter (20). We also generated sib1:4xtetO and vps1302:1xtetO without promoters (no). All strains expressed wild-type Clr4 and TetR-Clr4\textsuperscript{*}. Both sib1\textsuperscript{+} and vps1302\textsuperscript{+} were expressed at low levels when their promoters were removed and at much higher levels from med-adh1 or high-adh1 compared to their own, or low-adh1 promoters (fig. S4). TetR-Clr4\textsuperscript{*} was unable to establish significant levels of H3K9me2 when tethered to sib1:4xtetO or vps1302:1xtetO expressed from hi-adh1 and relatively low levels when expressed from med-adh1, but substantial H3K9me2 occurred when either gene had no, its own or, the low-adh1 promoter (fig. S4). However, as with 4xtetO-ade6\textsuperscript{+}, rapid loss of H3K9me2 followed TetR-Clr4\textsuperscript{*} release from even no and own promoter constructs; again >90\% was lost within 6 hours (Fig. 2, D and E). Although high levels of transcription across tethering sites prevents the establishment of H3K9me by TetR-Clr4\textsuperscript{*}, neither low promoter strength nor low H3 turnover renders H3K9me more persistent upon methyltransferase release. Thus the inability
to maintain H3K9 methylation upon removal of the initiating tethered Clr4 methyltransferase is likely a general feature of euchromatic loci.

To determine if the loss of H3K9 methylation from the tethering site is coupled to replication or passage through the cell cycle we released TetR-Clr4* from 4xtetO-ade6+ in cdc25-22 synchronized cultures (Fig. 3A). H3K9me2 levels on 4xtetO-ade6+ dropped by 70% within one cell cycle following the addition of AHT to these synchronized cultures and no accelerated H3K9me2 loss was evident during S phase which is coincident with septation (21). We also released TetR-Clr4* from 4xtetO-ade6+ in non-cycling G2 blocked cdc25-22 cells (Fig. 3B). TetR-Clr4* was lost from 4xtetO-ade6 within an hour and H3K9me declined to less than 25% of initial levels within 4 hours. Thus, after release of the initiating methyltransferase, rather than being passively diluted through chromatin replication, H3K9 methylation must be removed by an active process.

Known and putative histone demethylases might act to remove H3K9me and thus disassemble heterochromatin from TetR-Clr4* tethering sites. We therefore tested if mutation of genes for six JmjC domain (Epe1, Jmj1, Jmj2, Jmj4, Lid2, Msc1) (22) or two SWIRM/Amino-oxidase domain proteins (Lsd1 or Lsd2; 23) allowed long-term 4xtetO-ade6+ silencing after tethered TetR-Clr4* release. Wild-type 4xtetO-ade6+ TetR-Clr4* cells form red/ade6-repressed colonies on indicator plates lacking AHT, white/ade6-expressing colonies appear on +AHT plates due to loss of H3K9me-dependent heterochromatin over 4xtetO-ade6+. Of the eight tested mutants only epe1Δ consistently formed red/pink colonies on +AHT plates, indicating that 4xtetO-ade6+ can remain repressed without bound TetR-Clr4* (Fig. 4A, fig. S5 and fig. S6). Catalytically inactivating mutations in the Fe(II) or 2-oxoglutarate binding sites of the Epe1 putative demethylase (epe1-H297A and epe1-K314A) had a similar phenotype (Fig. 4A, fig. S6 and Table S3). The variable silencing and colony colour most likely reflects stochastic events at the 4xtetO-ade6+ locus in epe1Δ cells in which H3K9me domains are known to expand and additional heterochromatin islands also appear, potentially titrating and redistributing heterochromatin proteins between various loci in individual cells (24-27). Maintenance of the silenced state in epe1Δ cells is not dependent on the RNAi component Ago1 as ago1Δepe1Δ cells form red/ade6-silent colonies on +AHT plates (fig. S7A) but it does require untethered wild-type Clr4 with an intact Clr4 chromodomain and Swi6 (fig. S8). This reliance on untethered, intact Clr4 and Swi6 is consistent with a simple read-write propagation mechanism (fig. S10).

Silencing of 4xtetO-ade6+ can be propagated through multiple cell divisions in epe1 mutants (lost in 4% of cells/division), and a high proportion of descendant cells retain silencing of, and 30-70% of H3K9me2 on, 4xtetO-ade6+ after TetR-Clr4* release by AHT. In contrast, 4xtetO-ade6+ silencing and H3K9me2 are completely lost in wild-type cells (Fig. 4A, and fig. S7B-E). The relative levels of H3K9me2 and H3K9me3 detected on 4xtetO-ade6+ are similar in wild-type and epe1Δ cells and surrounding genes are silenced by H3K9me2 in both wild-type and epe1Δ. (fig. S9). To determine if H3K9me on 4xtetO-ade6+ in epe1Δ cells is maintained through meiosis in the absence of TetR-Clr4*, epe1Δ 4xtetO-ade6+ tetR-clr4* cells (F0) were crossed to epe1Δ cells devoid of both 4xtetO-ade6+ and TetR-Clr4* and then F1 epe1Δ 4xtetO-ade6+ progeny lacking TetR-Clr4* were again crossed to epe1Δ cells. A high proportion of resulting F2 epe1Δ 4xtetO-ade6+ progeny
formed red-pink/ade6-repressed colonies and H3K9me2 was retained (Fig. 4B, and fig. S6B). Thus, epe1Δ allows silencing and H3K9me to persist through multiple mitotic divisions, and meiosis, in the complete absence of the tethered TetR-Clr4* that initiated H3K9me-dependent heterochromatin on 4xtetO-ade6+. Crossing of red F2 epe1Δ 4xtetO-ade6+ cells to wild-type epe1+ cells resulted in loss of silencing (white colonies only) and H3K9me2 from the 4xtetO-ade6+ locus. Thus, provision of epe1+ results in removal of persistent H3K9me and loss of silencing (Fig. 4B). Genetically identical naïve epe1Δ 4xtetO-ade6+ cells, that were never exposed to the TetR-Clr4* initiator, formed only white/ade6-expressing colonies and H3K9me2 was absent (Fig. 4B). We conclude that the transient tethering of TetR-Clr4* adjacent to 4xtetO-ade6+ allows establishment of H3K9me-dependent heterochromatin which can be propagated epigenetically through mitotic cell divisions and meiosis using endogenous read-write copying mechanisms, provided Epe1 is rendered non-functional (Model: fig. S10).

Propagation of heterochromatin on 4xtetO-ade6+ in epe1 mutants requires recognition of TetR-Clr4*-mediated H3K9me by the chromodomain of Clr4, and also Swi6 (fig. S8). Epe1 associates with Swi6HP1 and clearly opposes heterochromatin formation (24-28). Indeed Epe1 associates with TetR-Clr4*-mediated heterochromatin (fig. S7C). Although Epe1 contains a JmjC domain, its Fe(II) binding site is unusual and histone demethylase activity has not been detected (22). However, the human PHF2 JmjC domain bears a similar anomaly but phosphorylation activates its latent H3K9 demethylase activity (29). The analyses presented here are consistent with Epe1 normally acting as an H3K9 demethylase that removes H3K9 methylation from ectopic sites of heterochromatin formation. Moreover, additional heterochromatin islands and domain expansion in epe1 mutants are best explained by loss of an H3K9 demethylase that prevents excessive H3K9me-dependent heterochromatin formation. Epe1-dependent removal of H3K9me ensures regulation of centromeric heterochromatin and makes the RNAi pathway essential for the systematic replenishment of H3K9me every cell cycle (30, 31). Epe1 itself may be regulated in response to environmental cues in order to retain or eliminate H3K9 methylation at specific locations (26). Indeed Epe1 levels are regulated and this may aid the persistence of centromeric H3K9me-dependent heterochromatin (28). Thus opposing H3K9 methyltransferase and demethylase activities must be finely tuned to allow controlled heterochromatin formation and prevent its inappropriate mitotic and transgenerational inheritance. It seems counterintuitive for heterochromatin to carry a means of self-destruction, however, such an inbuilt safety mechanism averts the inappropriate, and potentially deleterious, silencing of genes by removing repressive heterochromatin and preventing its propagation.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgments

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References and Notes


19. Materials and methods are available as supporting material online.


Fig. 1. H3K9 methylation is rapidly lost upon release of tethered TetR-Clr4*

(A) Positions of 4xtetO, tethered TetR-Clr4* beside ade6+ at ura4, and surrounding S. pombe chromosome III genes. Dumbbells indicate primer pairs.

(B and C) qChIP time course of FLAG-TetR-Clr4* (B) and H3K9me2 (C) levels on 4xtetO-ade6+ following AHT addition using indicated primers. Data are mean ±SD (n=3), P<0.05 (t-test).
Fig. 2. Tethering TetR-Clr4* at loci with low expression and histone turnover does not stabilize H3K9 methylation

(A) Read distribution (log$_2$RPKM) from S. pombe polyA RNA-seq relative to gene length. ade6+, sib1+ and vps1302+ indicated.

(B) qRT-PCR of ade6+, sib1+ and vps1302+ RNA levels. Data are mean ±SD (n=3) P<0.005 (t-test)

(C) Recombination-induced tag exchange monitoring incorporation of new H3-T7 on act1+, ade6+, sib1+ vps1302+ and cen-dg repeats. Data are mean ±SD (n=3). H3 turnover on sib1+ and vps1302+ was significantly lower than on act1+ and ade6+ P<0.05 (t-test).

(D and E) sib1+ and vps1302+ lose H3K9me2 after TetR-Clr4* release. Position of tetO sites within sib1 and vps1302. own promoters were replaced with ura4+ (no) or swapped to low, medium (med), or high adh1 promoter versions (20). Dumbbells indicate primers.

qChIP of H3K9me2 levels, at time points relative to AHT addition, on sib1:4xtetO (D) and vps1302:1xtetO (E) with no or indicted promoters. Data are mean ±SD (n=3), P<0.05 (t-test). H3K9me2 level within sib1 carrying its own promoter is decreased with a probability of respectively P=0.068 and P=0.051, 3h and 6h following TetR-Clr4* release.
Fig. 3. H3K9 methylation rapidly declines through the cell cycle and in non-cycling cells
Regimes for release of TetR-Clr4* from 4xtetO-ade6+ following AHT addition to cdc25-22 G2 synchronised cultures (A) or double-blocked cdc25-22 G2 cells (B). Synchrony was assessed by septation index. qChIP time course of H3K9me2 or FLAG-TetR-Clr4* levels on 4xtetO-ade6+ using indicated primers. Data are mean ±SD (n=3).
Fig. 4. *epe1* mutants retain heterochromatin without tethered Clr4 methyltransferase through multiple cell divisions and meiosis

(A) wild-type, *epe1Δ*, *epe1-K314A* and *epe1-H297A* cells carrying 4xtetO-ade6+ and expressing TetR-Clr4*, were grown −/+ AHT. Colony colour assay to assess 4xtetO-ade6+ silencing (red-pink colonies; % of total indicated) and H3K9me2 qChIP on 4xtetO-ade6+ with (−AHT) or without (+AHT) tethered TetR-Clr4*. Data are mean ±SD (n=3), P<0.05 (t-test).

(B) TetR-Clr4* was completely removed from F0 *epe1Δ 4xtetO-ade6+ tetR-Clr4* cells by crossing to *epe1Δ* lacking TetR-Clr4* and 4xtetO-ade6+. F1 progeny were crossed to *epe1Δ* cells, generating *epe1Δ 4xtetO-ade6+ F2* cells. Naïve *epe1Δ 4xtetO-ade6+* cells never expressed TetR-Clr4*. Colony colour, qRT-PCR and qChIP assays to assess silencing and transcription of 4xtetO-ade6+, and H3K9me2 levels on 4xtetO-ade6+ in indicated cell types. Data are mean ±SD (n=3). 4xtetO-ade6+ RNA levels are significantly reduced in F0, F1 and F2 compared to wild-type cells without TetR-Clr4*; P<0.05 (t-test).