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On the role of H3.3 in retroviral silencing

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Epigenetic silencing of endogenous retroviruses (ERVs) is initiated by KRAB-ZFP-KAP1-SETDB1 repression complexes during early mammalian development. On the basis of biochemical evidence from histone H3.3-knockout embryonic stem (ES) cells, Elsässer *et al.*¹ reported that histone H3.3 is deposited at KAP1-SETDB1-targeted ERVs by the chaperone DAXX-ATRX complex and that this deposition is required to repress ERV transcription and retrotransposition. However, our re-analysis of the published data revealed little evidence of genome-wide ERV upregulation in H3.3-knockout ES cells, and, more importantly, that the ES cells used for the analysis include polymorphic ERV insertions, which probably reflect a mixed genetic background and compromises their use for ERV expression and re-integration analysis. Thus, despite the strong evidence for H3.3 deposition at KAP1-SETDB1-targeted ERV elements, it remains to be determined whether this deposition plays a major role in preventing ERV reactivation. There is a Reply to this Comment by Elsässer, S. J. *et al.* Nature 548, <http://dx.doi.org/10.1038/nature23278> (2017).

Elsässer *et al.*¹ reported that H3.3 is deposited at ERVs by the H3.3 histone chaperone DAXX in mouse ES cells¹. While we were able to confirm DAXX-dependent H3.3 deposition at ERVs in a newly generated conditional *Daxx*-knockout ES cell line (Extended Data Fig. 1a, b), we have concerns about the authors' conclusion that H3.3 knockout leads to increased ERV expression¹. Intracisternal A-type particle (IAP) ERVs were upregulated less

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Author Contributions G.W., T.S.M., M.C.L., D.L.M., K.O. and G.J.F. planned and designed the study. G.W., R.R., W.W., B.W., M.B. and R.K. performed experiments. G.W., M.M.K. and A.D.E. analysed the data. G.W. and T.S.M. wrote the manuscript with contributions from M.C.L., D.L.M. and G.J.F.

Supplementary Information accompanies this Comment.

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and H3.3-knockout' IAPs reported¹ were present in 129 mice in one or both datasets. For the 80 'H3.3-knockout only' IAP copies, 16 were found in 129 strains and therefore excluded from further analysis.

However, we found 47 of the remaining 64 'H3.3-knockout only' copies in various other mouse strains, but not in 129 or C57BL/6, and only 17 were not present in the published datasets⁷ (Supplementary Table 1). This indicates that H3.3-knockout ES cells have a complex genetic background that includes additional laboratory mouse strains. To verify these insertions, we performed a more stringent re-analysis of the relevant whole-genome sequencing data using our own pipeline (described in the Supplementary Methods). We identified 37 out of the 47 polymorphic and 13 out of the 17 potentially new IAP copies (Supplementary Table 1). The majority of these IAP copies were found in multiple mouse strains, indicating that they were misannotated polymorphic IAPs and not *de novo* retrotransposition events. Notably, we also found 8 of the IAP copies reported as H3.3-knockout only¹ to be present in both wild-type and H3.3-knockout ES cells, as well as in various other strains, including 129 (Supplementary Table 2). Moreover, 6 out of 10 randomly chosen polymorphic and putative new IAP copies could be readily detected in CD-1 feeder DNA (Fig. 2 and Supplementary Table 1), which share considerable genetic similarity to NOD/LtJ mice⁸. Altogether, these analyses show that the vast majority of the reported 'new' IAP insertions¹ are polymorphic IAP elements already present in various laboratory mouse strains.

In summary, our data and the data from Elsässer *et al.*¹ confirm the presence of H3.3-containing heterochromatin at ERV sequences in ES cells, but do not support a vital role for this H3 variant in suppressing ERV expression or mobilization. We speculate that in ES cells, SETDB1-dependent methylation of canonical H3 is generally sufficient to compensate for the loss of H3K9me3-marked H3.3 at ERVs.

Methods

Conditional *Daxx*-knockout ES cells were generated by CRISPR-Cas9-mediated *loxP* insertion into ES cells carrying a 4-hydroxytamoxifen-inducible *Rosa cre-ERT* allele. Native chromatin immunoprecipitation followed by sequencing (ChIP-seq) was performed with *Daxx*-knockout ES cells carrying a retrovirally delivered YFP-tagged H3.3 expression construct as described previously². ERV expression and ChIP-seq analysis were performed as described previously² and in the Supplementary Methods. For IAP integration analysis, genomic DNA from CD-1 feeders (StemCell Technologies) and 129/SvJ mice (Jackson Laboratory) was screened by PCR using primers shown in Supplementary Table 1. Full methods are available in the Supplementary Information.

Data availability.

Daxx-knockout ES cell RNA-seq and ChIP-seq data have been deposited in the Gene Expression Omnibus (GEO) under accession number GSE70967. All other data are available from the previously published accession numbers or from the corresponding author upon reasonable request.

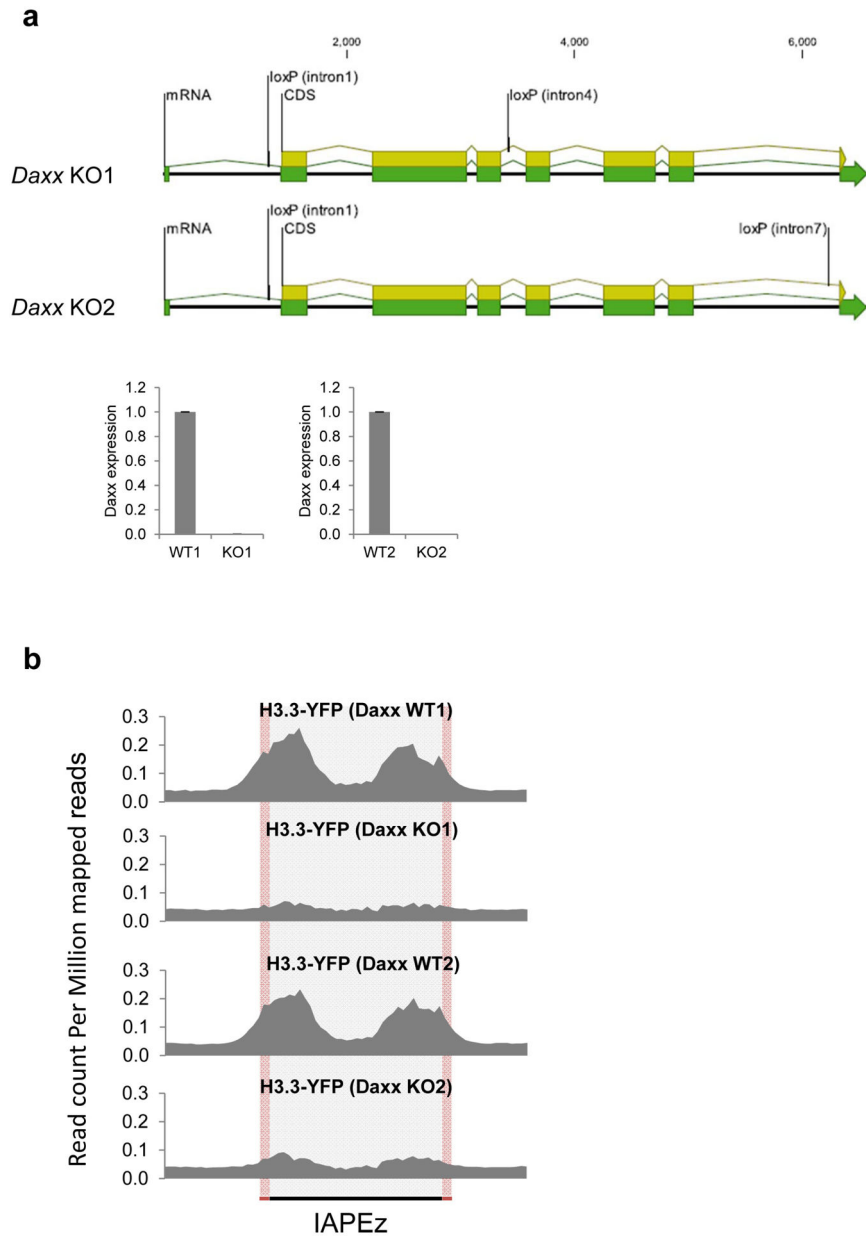
Extended Data

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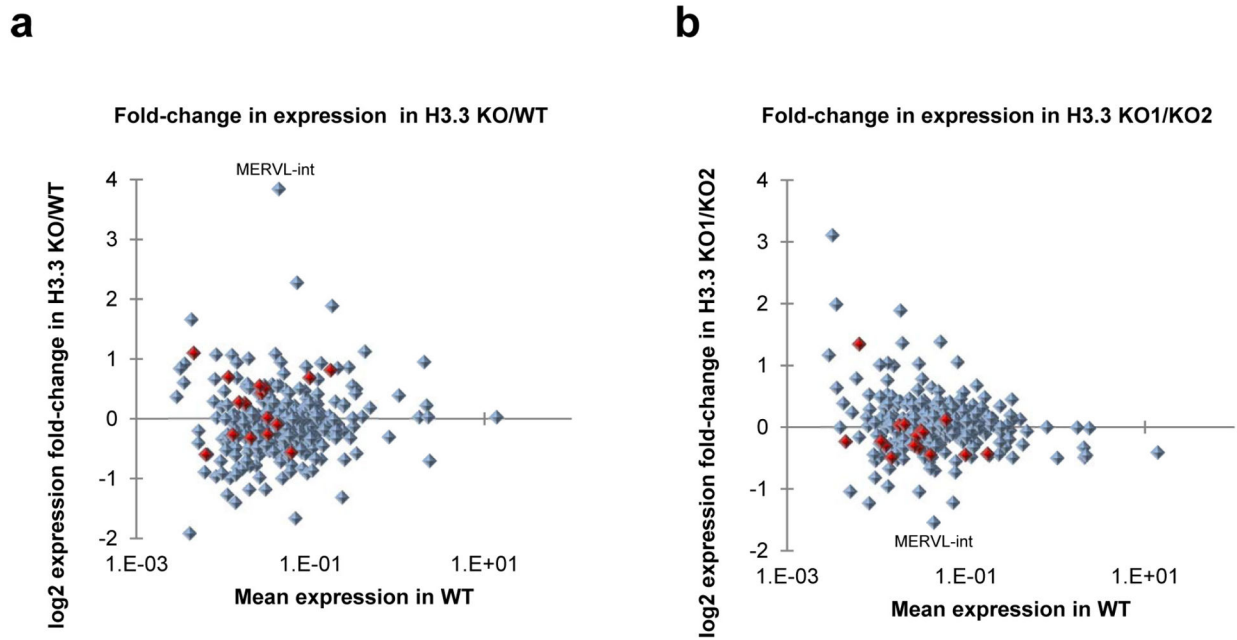
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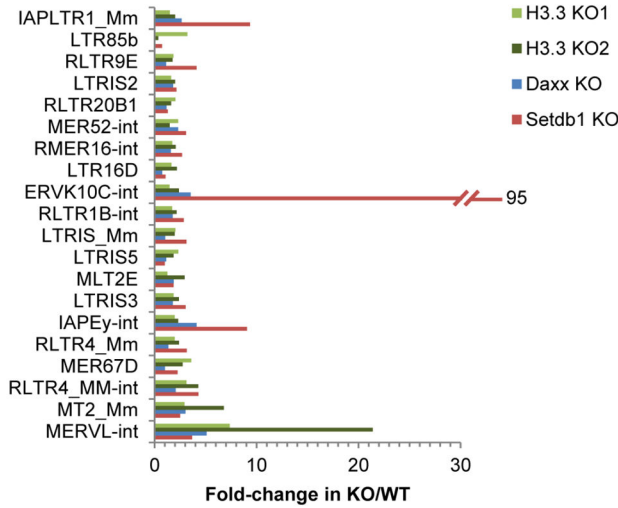
Extended Data Figure 1 | H3.3 is incorporated into ERVs by the chaperone DAXX.
a, Top, representation of the two conditional *Daxx*-knockout alleles generated by CRISPR–Cas9-mediated *loxP* insertion at the indicated introns. Bottom, quantitative RT–PCR analysis of *Daxx* mRNA expression in *Daxx*-knockout ES cells. Data are mean \pm s.d. expression (normalized to *Gapdh*) relative to the corresponding wildtype ES cells ($n = 3$, technical replicates). **b**, H3.3–YFP enrichment at 717 full-length IAP ERVs in two independently derived conditional *Daxx*-knockout ES cell lines.



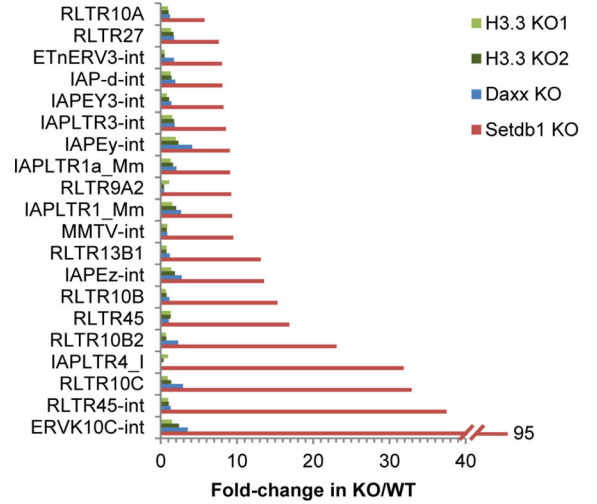
Extended Data Figure 2 | ERV deregulation in H3.3-knockout ES cells.

a, Mean fold change in ERV (LTR elements annotated in UCSC RepeatMasker) expression in H3.3-knockout ES cells (two cell lines) over wild-type ES cells (one cell line). **b**, Fold change in ERV expression comparing two H3.3-knockout ES cell lines. ERVs belonging to the IAP family are marked in red. Only ERV groups with more than 100 family members were considered for analysis.

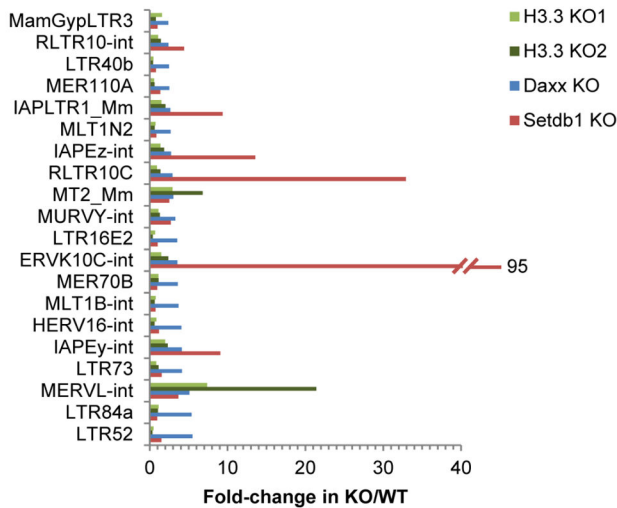
Top 20 ERVs upregulated in H3.3 KO



Top 20 ERVs upregulated in Setdb1 KO



Top 20 ERVs upregulated in Daxx KO



Extended Data Figure 3 | ERV expression in *Setdb1*-, H3.3- and *Daxx*-knockout ES cells.

The fold change in expression over the corresponding wild-type control is shown for the top 20 upregulated ERV annotations in *Setdb1*-, H3.3- and *Daxx*-knockout ES cells.

Annotations including –int represent the internal regions, which are transcribed from the cognate 5' LTR, of the annotated ERV. For example, ERVK10C-int is the internal region of ERVK10C elements, with flanking LTRs: RLTR10A, RLTR10B and RLTR10C (depending on the specific genomic copy), which are also presented among the graphs. Similarly, IAPEz-int is the internal region of IAPEz elements with flanking cognate LTRs:

IAPLTR1_Mm and IAPLTR1a_Mm, which are also represented. As the internal region is much longer and transcribed across its length, this is the most useful annotation to consider for expression analysis. The following published RNA-seq GEO data were re-analysed:

GSM727424 (*Setdb1*-knockout ES cells); GSM1428580 and GSM1428581 (H3.3-knockout

ES cells). Only ERV groups with more than 100 family members were considered for analysis.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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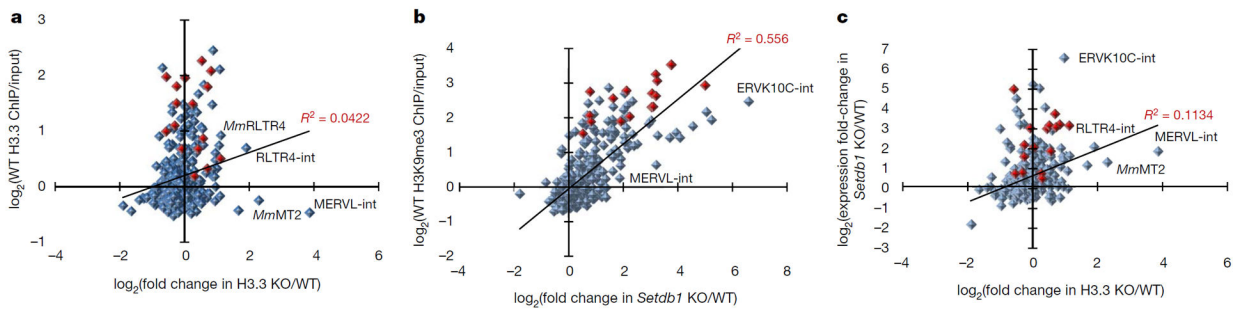


Figure 1 |. Comparative analysis of ERV expression and histone modifications at ERVs in knockout ES cell lines.

a. Mean fold change in ERV (long terminal repeat (LTR) elements annotated in UCSC RepeatMasker) expression in H3.3-knockout (KO) ES cells (two cell lines) over wild-type (WT) ES cells (one cell line) versus H3.3 enrichment in wild-type ES cells. ChIP, chromatin immunoprecipitation. **b.** Fold change in ERV expression in *Setdb1*-knockout ES cells over wild-type ES cells versus H3K9me3 enrichment in wild-type ES cells. **c.** Mean fold change in ERV expression in H3.3-knockout ES cells over wild-type ES cells versus fold change in ERV expression in *Setdb1*-knockout ES cells over wild-type ES cells. ERVs belonging to the IAP family are marked in red. *Mm*, *mus musculus*. Annotations including an '-int' represent the internal regions, which are transcribed from the cognate 5' LTR, of the annotated ERV. Only ERV groups with more than 100 family members were considered for analysis. Linear trend lines and corresponding R^2 values are shown.

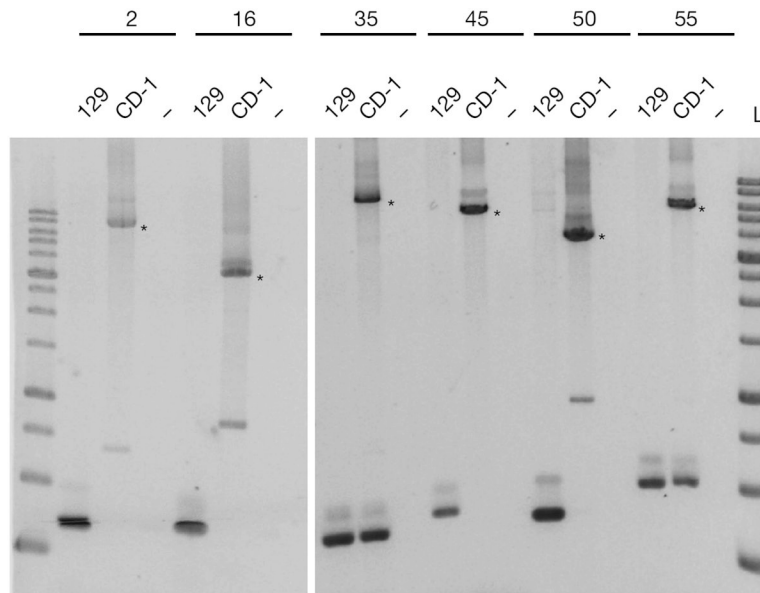


Figure 2 | PCR analysis of polymorphic IAP insertions.

PCR assay of 129/SvJ and genomic DNA of the CD-1 feeders using primers flanking six ‘new’ copies described in H3.3-knockout ES cells in ref. 1. Asterisks indicate larger bands corresponding to the size expected for a typical IAP insertion. Non-template controls are shown (–) and 10-kb size ladders (L) are included on either side of the six copies. IAP copy indexes (2, 16, 35, 45, 50 and 55) are shown above the lanes and the coordinates can be found in Supplementary Table 1 along with the primer sequences.