Chromatin-modifying genetic interventions suppress age-associated transposable element activation and extend life span in *Drosophila*

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Transposable elements (TEs) are mobile genetic elements, highly enriched in heterochromatin, that constitute a large percentage of the DNA content of eukaryotic genomes. Aging in *Drosophila melanogaster* is characterized by loss of repressive heterochromatin structure and loss of silencing of reporter genes in constitutive heterochromatin regions. Using next-generation sequencing, we found that transcripts of many genes native to heterogeneous regions and TEs increased with age in fly heads and fat bodies. A dietary restriction regimen, known to extend life span, repressed the age-related increased expression of genes located in heterochromatin, as well as TEs. We also observed a corresponding age-associated increase in TE transposition in fly fat body cells that was delayed by dietary restriction. Furthermore, we found that manipulating genes known to affect heterochromatin structure, including overexpression of Sir2, Su(var)3–9, and Dicer-2, as well as decreased expression of Adar, mitigated age-related increases in expression of TEs. Increasing expression of either Su(var)3–9 or Dicer-2 also led to an increase in life span. Mutation of Dicer-2 led to an increase in DNA double-strand breaks. Treatment with the reverse transcriptase inhibitor 3TC resulted in decreased TE transposition as well as increased life span in TE-sensitized Dicer-2 mutants. Together, these data support the retrotransposon theory of aging, which hypothesizes that epigenetically silenced TEs become deletoriously activated as cellular defense and surveillance mechanisms break down with age. Furthermore, interventions that maintain repressive heterochromatin and preserve TE silencing may prove key to preventing damage caused by TE activation and extending healthy life span.

TEs are mobile genetic elements that make up a large percentage of eukaryotic genomes and that are particularly enriched in regions of heterochromatin (reviewed in ref. 9). Because TEs function by either excising or copying themselves from the genome and integrating into new genomic locations, there is a high potential for deleterious effects on cellular function. Over-expressing TE RNAs is sufficient to cause cytotoxicity, and inhibiting expression of these TEs causes reversal of both senescence and cytotoxic phenotypes (10, 11). TE activation has been observed with age in mice (8, 12), as well as in *C. elegans* (13), yeast (14), *Drosophila* (15, 16), and senescent human tissue culture cells (7).

To combat the genomic threat of activation of TEs and loss of heterochromatic gene silencing, somatic cells have evolved mechanisms to suppress TE transcription, primarily through establishment and maintenance of repressive heterochromatin at TE sites (17). In *Drosophila*, the small interfering RNA (siRNA) pathway, through the proteins Dicer-2 and Argonaute 2 (AGO2), recruits the histone H3K9 methyltransferase Su(var)3–9 to catalyze formation of repressive heterochromatin at TE sites (18–20). Mutation of Dicer-2, the double-stranded RNAse responsible for generating siRNAs, results in TE reactivation and heterochromatin disorganization (19, 20). The RNA-editing enzyme ADAR (adenosine deaminase acting on RNA) is thought to modify the effectiveness of siRNA-directed heterochromatinization, and therefore the efficacy of TE silencing, by editing the dsRNA substrates used for siRNA biogenesis, thereby preventing proper processing by Dicer-2 (21). Sir2/Sirt1, the founding member of the sirtuin family of proteins, is a protein deacetylase important in maintaining gene

Significance

Most eukaryotic genomes contain abundant transposable elements (TEs), mobile DNA elements that can replicate and move within the genome. Because of the deleterious nature of active TEs, cells have mechanisms to suppress and prevent TE activation, including formation of repressive heterochromatin. In this report, we show that many TEs become activated with age in *Drosophila*, and this activation is prevented by dietary restriction, an intervention known to extend life span. We also show TE activation is blocked by genetic manipulations that stabilize heterochromatin and increase life span. This study provides evidence that a breakdown in TE silencing and repression may be a contributing factor to aging, and preventing TE activation may be a significant method of ameliorating the diseases of aging.


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silencing in heterochromatin regions of yeast and flies through deacetylation of specific histone lysines required for establishment and spreading of heterochromatin (3, 6, 22). Increasing Sir2 expression has been shown to repress the normal age-related loss of gene silencing in heterochromatin regions in both yeast and flies (3, 6), as well as the loss of gene silencing seen in mammalian embryonic stem cells undergoing genotoxic stress (23). In addition to their roles in heterochromatin formation and maintenance, these pathways have also been linked to aging. Decreasing activity of the siRNA pathway through Ago2 and Dicer-2 mutants leads to de-repression of TE activity and shortened life span (15, 24). Sir2 has a central role in aging, with increased gene dosage shown to extend life span in yeast (2) and in nine of 10 studies in nematodes and flies (25–28). Hypomorphs of Adar, a negative regulator of the siRNA pathway, also exhibit an extension of life span (21). These pathways share an ability to affect heterochromatin formation and life span.

Here we report a characterization of the heterochromatic transcriptome in adult Drosophila tissues and show activation of TEs and increased transposition with age. We provide evidence that the aging process is characterized by a failure to properly control and repress expression of heterochromatin-based genes and TEs in the soma. We show that genetic interventions positively modifying heterochromatin structure or negatively modulating TE activity extend metazoan life span. Finally, we show that attenuating TE activity pharmacologically decreases transposition and extends life span in Dicer-2 mutants. This study offers the first comprehensive characterization of the effect of known life span-extending interventions on heterochromatin regions and explores new interventions for controlling age-associated TE reactivation.

**Results**

**Expression of Genes Located Within Heterochromatin Increases with Age and Is Attenuated Under Dietary Restriction.** Because heterochromatin regions change in structure with age, specifically in female heads and fat bodies (5), we reasoned that genes located within these regions may also show changes in silencing or gene expression with age. We used RNA-seq to assay the expression of the ~250 genes natively located within the Drosophila heterochromatin.
compartmentalization (29) from young (10 d) and old (40 d) female fly heads and fat bodies. The vast majority of cells in the head are neurons and glial cells. The fly fat body is the functional homolog of mammalian adipose tissue and liver, and a central organ in the fly immune system. We found a number of genes located in heterochromatic regions increased expression with age in both wild-type heads and fat bodies (Fig. 1 A and B). Gene ontology analysis reveals no significantly enriched pathways among these genes, suggesting the age-related increase in expression of these genes is likely to be a consequence of loss of heterochromatic silencing, rather than a specific compensatory or correlational response tied to aging.

Dietary restriction (DR) is an intervention known to extend life span and delay numerous age-related phenotypes. We previously observed that DR can oppose age-related loss of gene silencing of reporter genes in constitutive heterochromatin (6). On this basis, we examined whether the age-dependent increase in expression of native genes in the constitutive heterochromatin regions were also suppressed under a DR regimen. The vast majority (95% in heads, 89% in fat bodies) of heterochromatin genes that increased expression with age on high-calorie food increased less or decreased in expression with age on DR food [high calorie (HC) vs. DR] in both heads (Fig. 1 A) and fat bodies (Fig. 1 B). Although we observed a similar fraction of euchromatin genes that increased expression with age, this pattern of suppression by DR was not maintained in the most highly up-regulated euchromatin genes (Fig. S1 A and B), suggesting DR has a selective effect on heterochromatin genes. Similar to previous studies using reporter genes in both yeast (2) and flies (6), these data suggest that gene expression in heterochromatic regions becomes dysregulated with age, and that a known life span-extending intervention, DR, can prevent this.

### TE Expression Increases with Age and Is Attenuated Under DR.

One of the most notable attributes of heterochromatin is the abundant enrichment of TEs within these regions. The *Drosophila* genome is rich in retroviral-like long terminal repeat (LTR) and long interspersed nuclear elements (LINEs) (30), and activation of both LINEs and LTRs has been observed in the fly nervous system and fat body during aging (15, 16). Because heterochromatin integrity declines with age in multiple model systems (1), we hypothesized that a loss of heterochromatin in regions highly enriched with TEs would show a correlated increase in TE expression with age. To test this hypothesis, we examined the expression profile of TEs with age and diet in female fly heads and fat bodies, using whole transcriptome RNA-seq. Similar to our observations for native heterochromatin genes, we also found many TEs that increased their expression with age, in both heads and fat bodies (Fig. 1 C and D). We observed a high degree of overlap in the TEs that increased expression most with age between these two tissues, most of which were retrotransposons (Fig. S2 A). In both tissue types, as observed with the heterochromatin genes, DR attenuated the age-related increase in expression observed in HC food (Fig. 1 C and D, right “DR” columns). TEs that had the highest increase in expression with age on HC food tended to increase less or even decrease in expression with age under DR conditions (57% lower in DR relative to HC in both heads and fat bodies; Dataset S1). These data suggest that, similar to reporter genes in heterochromatin (2, 6) and native heterochromatin genes (Fig. 1 A and B), TE silencing is lost with age, and DR opposes this.

### TE Transposition Increases with Age and Is Delayed by DR.

An age-dependent increase in TE expression and transposition in the *Drosophila* brain associated with decline in neural function has been reported (15). Similar increases in TE expression levels and copy numbers have been observed in aging mouse tissue (8). To assay for age-related transposition in our flies, we used the gypsy-TRAP reporter system (15). This system is designed such that green fluorescent protein (GFP) becomes expressed when an endogenous gypsy retrotransposon jumps into an engineered site containing known hotspot sequences from the *ovo* locus (31). We monitored flies as they aged and visually sorted them into either low, medium, or high categories based on the number of GFP-positive cells observed in the fat body (Fig. 2 A). Longitudinal fly cohorts were assayed for transposition every 7 d on either HC or DR foods (Fig. 2 B). We observed a clear age-associated increase in transposition, and DR delayed this increase relative to HC controls (Fig. 2 B and C). Similar results were seen in two additional genetic backgrounds (Fig. S3 A–D). We examined the life spans of these three different genetic background gypsy-TRAP lines and observed that life span correlated with the timing of increased transposition. Lines exhibiting earlier transposition showed shorter life span, and those with later transposition longer life span (Fig. S3 E). Similarly, the delay in age-related transposition resulting from DR correlated with the ability of DR to extend life span in each line. To ensure the increased GFP expression we observed with age was caused by transposition rather than age-related disruption in the GAL80/GAL4/GFP reporter system, we examined a gypsy-TRAP line with a mutant *ovo* locus not susceptible to transposition and observed no significant increase in GFP expression with age (Fig. S3 F). Rather than a constant linear increase in GFP expression with time, the gypsy-TRAP assay showed a clear age-associated increase in TE mobilization, consistent with the TE theory of aging, which posits an age-related loss of normal cellular surveillance mechanisms and consequent increase in deleterious TE activity. The relationship between the timing of increased gypsy transposition and life span further supports TE mobilization as an age-associated phenomenon.

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**Fig. 2.** DR delays age-related increase in TE transposition. (A) Gypsy-TRAP flies (genotype: *w*1118; *Hf/Hf; r4-GAL4, UAS-GFP/tub-ovo-GAL80), were categorized as low (<20 GFP-positive cells), medium (20–100), or high (>100) based on number of GFP-positive cells observed in fat body (representative images shown). The demarcation between low and medium categories is clear; most flies have either fewer than 10 cells staining, or more than 50. (Scale bar, 200 μm.) (B) Percentage of observed flies falling into each category at indicated points for HC (Top) and DR (Bottom) diets (n = 120 per cohort). (C) Transposition (defined here as medium- or high-category GFP staining) increases with age and is delayed by DR. Percentage of flies falling into either medium or high categories is displayed with time for both HC and DR diets. Log-rank P < 10⁻¹⁰.
Age-Related Increases in TE Expression Are Attenuated by Genetic Interventions That Affect Heterochromatin Structure. Our data demonstrate a correlation between changes in heterochromatin structure with age and expression and transposition of TEs during normal aging. This led us to examine whether genetically manipulating heterochromatin structure and maintenance, either directly or indirectly, could also affect the age-related changes in expression of TEs. To test our hypothesis, we genetically increased the activity of two important systems that maintain repressive heterochromatin in somatic cells: the siRNA pathway and Sir2. Using transgenic fly lines, we increased gene dosage of Sir2, Dicer-2, and Su(var)3-9 and also used an Adar hypomorph, interventions we expect to stabilize heterochromatin. Each of these genes offers a unique and complementary approach to simultaneously manipulating heterochromatin regions and enhancing TE silencing.

We assayed age-dependent loss of silencing of TEs in each of these various transgenic [Sir2, Su(var)3-9, and Dicer-2] or hypomorphic (Adar) fly lines compared with their respective controls. The TEs that showed the greatest increase in expression with age in the control fly lines were overwhelmingly attenuated or decreased with age by these genetic interventions [Fig. 3 A–D; Sir2, 72%; Adar, 84%, Su(var)3-9, 92%; Dicer-2, 68%]. Interestingly, we again observed significant overlap in the TEs that increased with age among the different genetic backgrounds (Fig. S2B). Together, these data suggest that increased expression of Sir2, Dicer-2, or Su(var)3-9, as well as decreased gene dosage of Adar, all prevent the age-related loss of silencing of TEs observed in wild-type and control lines, suggesting a common link between heterochromatin factors and proper maintenance of TE silencing.

TE activation is known to disrupt gene expression and destabilize the genome. Increased DNA damage has been observed in aging fat body nuclei, as assayed by γH2Av immunostaining (16). Our finding of a repression of TE expression with increased Dicer-2 expression led us to investigate whether genomic integrity was similarly compromised in animals with disrupted RNAi/heterochromatin pathways and increased TE activity. Similar to depletion of AGO2 in fat body cells (16), we observed an increase in double-strand breaks in Dicer-2 mutants compared with controls in fat body nuclei (Fig. S44), suggesting deleterious genomic consequences when proper TE silencing is disrupted.

Genetic Interventions That Affect Heterochromatin Structure Also Affect Life Span. In addition to their effects of delaying age-related heterochromatin gene and TE silencing loss, both increased Sir2
expression and decreased Adar expression have been shown to extend life span in flies (21, 25), whereas Dicer-2 and Ago2 mutants are short-lived (15, 24). Given our findings presented earlier, we examined whether increased expression of Su(var)3-9 or Dicer-2 also have a positive effect on life span. We found that two different Su(var)-3-9 transgenic lines, SV2 and SV3, which constitutively increase Su(var)3-9 expression to 160–180% of the wild-type level in the absence of a driver (Fig. S4A), extend life span significantly compared with their genetically matched control (Fig. 4A). Similarly, increasing Dicer-2 expression in adult life in either all tissues (Fig. 4B) or selectively in adult neurons (Fig. 4C), using the conditional inducible GeneSwitch system, was able to extend life span, with flies fed the gene-activating drug RU486 outliving a genetically identical control cohort fed only the diluent control EtOH. Together with previously published data (21, 25), these observed life span extensions suggest a coordinated response integrating maintenance of heterochromatin structure, TE suppression, and longevity.

Pharmaceutical Suppression of Transposable Element Activity Delays Age-Related Transposition and Extends Life Span. To test whether the age-related effects we observed were a result of increased TE activity, we sought to directly modulate transposition pharmacologically. We used the reverse transcriptase inhibitor lamivudine (3TC) to directly block TE transposition. Using the gypsy-TRAP reporter system in the fat body, we observed that flies treated with 3TC had a significant delay in age-related transposition compared with controls (Fig. 5A). This effect was observed whether the drug was administered constantly throughout life (Fig. 5A) or started during midadulthood at 20 d of age (Fig. S4C).

As our evidence suggests that maintenance of TE silencing is important for ensuring longevity, we tested whether attenuating TE activity directly could extend life span. In the Dicer-2 L1116X mutant background, which exhibits high levels of TE activity because of defective RNAi silencing (32), we found that administration of 3TC was able to extend life span significantly compared with controls (Fig. 5B). Together, these data suggest that heterochromatin-altering pathways that extend life span may do so at least in part by suppressing TE activity to ensure healthy longevity, and confirm that suppression of TE activation is likely to be an attractive target mechanism for pharmacological intervention.

Discussion

Aging is characterized by a breakdown in homeostasis throughout multiple cellular and organismal systems. Genomic and epigenomic stability is vital for somatic cells to maintain proper gene expression and silencing. A number of studies have shown changes in chromatin structure during organismal aging in various model systems, especially in heterochromatin (1, 2). Although heterochromatin regions contain some genes, a likely reason for the central importance of maintaining repressive heterochromatin is that heterochromatin in replete with TEs, and unchecked TE activation has particularly damaging ramifications for genomic stability. Indeed, the expression and mobilization of TEs is disruptive enough that cells have evolved RNAi pathways responsible for silencing and preventing TE activation. These cellular defenses against TE activation include siRNA-directed induction and maintenance of repressive heterochromatin to prevent TE expression, and degradation of target transcripts. siRNA pathway mutants exhibit both an activation of TEs and a shortening of life span (15, 24). Here we provide evidence that many heterochromatin genes and TEs do in fact increase expression with age in two different tissues. Increased transposition has been previously reported in the aging fly brain (15). Similarly, we also observe an increase in somatic TE transposition with age in the fat body. DNA double-strand breaks increase with age in the fat body (16), and we show this DNA damage also accumulates at a younger age in Dicer-2 mutant flies with high levels of TEs. The age-related increase in TE transcript levels as well as TE mobilization were suppressed by DR. Together, these data suggest that an increase in TE expression and transposition is an important hallmark of somatic aging, and preventing this increased TE activity is likely to preserve genomic stability and cellular homeostasis.

We demonstrate in a whole organism model that aging is associated with a loss of silencing of both TEs and genes natively located in heterochromatin regions, with a resultant increase in TE mobilization and an associated increase in DNA damage.

Fig. 4. Increasing expression of Su(var)3-9 or Dicer-2 increases life span. (A) SV2 and SV3 lines [overexpressing Su(var)3-9] are long-lived compared with controls. Survivorship of adult female flies is shown (n = 250). WT mean life span = 40.6 d, median = 44 d; SV2 mean = 58.5 d, median = 60 d, 36% median extension, log-rank P < 10−10; SV3 mean = 58.3 d, median = 60 d, 36% median extension, P < 10−10. (B) Whole-adult animal Dicer-2 overexpressing flies are long-lived compared with controls. Survivorship of tubulin-GeneSwitch > UAS-Dicer-2 adult females (n = 250) is shown, either with EtOH (uninduced) or 200 μM RU486 (induced expression). EtOH mean = 52.3 d, median = 54 d; RU486 mean = 63.5 d, median = 66 d, 22% median extension, P < 10−10. (C) Flies expressing Dicer-2 in adult neurons are long-lived compared with controls. Survivorship of elav-GeneSwitch > UAS-Dicer-2 adult females is shown, for EtOH (uninduced) or 500 μM RU486 (induced) (n = 250). EtOH mean = 53.8 d, median = 56 d; RU486 mean = 61.7 d, median = 66 d, 18% median extension, P < 10−10. All life span experiments were repeated with similar results; representative experiments shown.
is highly likely that the pathways and processes of heterochromatin maintenance, RNAi, and gene and TE silencing represent potential homeostatic failure points that lead to the phenotypes of aging. Here we have also presented genetic evidence that these pathways are linked to aging, as manipulating the expression of several genes involved in maintaining heterochromatin structure, TE silencing, and RNAi efficacy directly suppresses age-related TE expression as well as extends life span. Furthermore, we show that DR, an environmental intervention that extends life span robustly and delays most age-related phenotypes, also opposes the loss of silencing we observe during normal aging and delays age-dependent transposition, further lending support to the idea that these processes are associated with and may be relevant to aging. We also demonstrate that inhibition of retrotransposon activity is strongly correlated with extension of life span. Our data support the retrotransposon theory of aging model, where proper maintenance of silent heterochromatin and prevention of TE activation are key to guaranteeing genomic stability and homeostasis during aging and ensuring healthy longevity. Furthermore, our demonstration that dietary (DR), genetic, and pharmacological interventions that reduce the age-related increases in TE activity can also extend life span suggests new and novel pathways for the development of interventions designed to extend healthy life span.

Materials and Methods

Gypsy-TRAP Assay. Flies from the gypsy-TRAP line were aged for 5 and initially examined for the number of GFP-positive cells, then examined every 7 d until age 49 d, using a fluorescent dissecting microscope. Flies were separated into three categories based on visual inspection, according to how many fat body cells were GFP positive: low (<20 cells), medium (20–100 cells), and high (>100 cells). The same cohort of flies was examined longitudinally throughout each of the experiments.

RNA-seq. Total RNA from the indicated genotypes was collected from ~100 fly heads or ~50 dissected fat bodies, using the mirVana RNA prep kit (Ambion). Then, 100 ng total RNA was used as input for stranded RNA-seq library construction, using the Ovation Universal RNA-seq kit (NuGEN) with Drosophila RNA depletion module. Libraries were sequenced on an Illumina HiSeq. 2500 in 1 x 50 bp mode. All samples were prepared using at least three individually isolated biological replicates for each condition.

Bioinformatics. Standard RNA-seq analysis was performed using TopHat 2.0 for sequence alignment, followed by gene-based counting using the easyRNA-seq R package from Bioconductor. Finally, trimmed mean of M-values (TMM) normalization and fold change calculation was performed using edgeR, with a generalized linear model (GLM) correcting for batch effects, as described in the edgeR vignette. To account for repetitive sequencing reads and perform TE quantification, RepEnrich (33) was used to generate a count table of reads for each TE, and edgeR was used to normalize, calculate fold changes, and perform differential expression testing. The RepBase annotation was used as input for RepEnrich.

Heat maps were generated by ranking the TEs or genes in order of log2 fold change of expression with age (old/young) for the reference (control/HC) condition and then plotting log2 fold change with age for all conditions. To show consistent trends in our data and include genes/TEs that behaved consistently but did not meet the threshold of statistical significance, we displayed genes or TEs where log2 fold change was greater than 0.322 (1.25 × the mean fold change value of 0.263 (1.25 × standard deviation of expression values). Plots were generated using the heatmap.2 function in the gplots2 R package.

See SI Materials and Methods for additional details on fly stocks and methods.

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