Maternally inherited piRNAs direct transient heterochromatin formation at active transposons during early Drosophila embryogenesis

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Abstract

The PIWI-interacting RNA (piRNA) pathway controls transposon expression in animal germ cells, thereby ensuring genome stability over generations. In Drosophila, piRNAs are intergenerationally inherited through the maternal lineage, and this has demonstrated importance in the specification of piRNA source loci and in silencing of I- and P-elements in the germ cells of daughters. Maternally inherited Piwi protein enters somatic nuclei in early embryos prior to zygotic genome activation and persists therein for roughly half of the time required to complete embryonic development. To investigate the role of the piRNA pathway in the embryonic soma, we created a conditionally unstable Piwi protein. This enabled maternally deposited Piwi to be cleared from newly laid embryos within 30 minutes and well ahead of the activation of zygotic transcription. Examination of RNA and protein profiles over time, and correlation with patterns of H3K9me3 deposition, suggests a role for maternally deposited Piwi in attenuating zygotic transposon expression in somatic cells of the developing embryo. In particular, robust deposition of piRNAs targeting roo, an element whose expression is mainly restricted to embryonic development, results in the deposition of transient heterochromatic marks at active roo insertions. We hypothesize that roo, an extremely successful mobile element, may have adopted a lifestyle of expression in the embryonic soma to evade silencing in germ cells.
Introduction

Transposable elements (TEs) are mobile genomic parasites that can change their genomic position or increase in copy number, and therefore pose a threat to genome integrity. Many TEs have evolved mechanisms that promote their activity specifically in gonads, thereby introducing new insertions that are inherited by future generations (Kim et al., 1994; Leblanc et al., 2000; Wang et al., 2018). Accumulation of insertional mutations in germ cells can lead to decreased population fitness and increased risk of disease (Hancks and Kazazian, 2016; Payer and Burns, 2019). Germ cells, however, harbour protective systems that substantially decrease the likelihood of transposition events.

In animal gonads, the main transposon defence mechanism is the PIWI-interacting RNA (piRNA) pathway (reviewed in (Czech et al., 2018; Ozata et al., 2019)). At its core, this system depends on 23- to 30-nt piRNAs to distinguish transposon-derived RNAs from host-encoded transcripts and to direct their associated PIWI proteins to active TE targets. In Drosophila, PIWI-guided repression involves cytoplasmic post-transcriptional mRNA cleavage by Aubergine (Aub) and Argonaute-3 (Ago3) and nuclear P-element induced wimpy testis (Piwi) that engages nascent transposon transcripts and instructs co-transcriptional gene silencing (coTGS) through heterochromatin formation. coTGS requires additional factors acting downstream of Piwi, including Panoramix (Panx), Nuclear Export Factor 2 (Nxf2), NTF2-related export protein 1 (Nxt1) and Cut-up (Ctp) that together form the PICTS complex (also known as SFiNX) (Batki et al., 2019; Eastwood et al., 2021; Fabry et al., 2019; Murano et al., 2019; Schnabl et al., 2021; Sienski et al., 2015; Yu et al., 2015; Zhao et al., 2019). PICTS interfaces with general chromatin silencing factors including Su(var)205/HP1a, SETDB1/Eggless (Egg) and its cofactor Windei (Wde), Su(var)3-3/Lsd1 and its cofactor coRest, Mi-2, Rpd3, Ovaries absent and Su(var)2-10 (Czech et al., 2013; Handler et al., 2013; Koch et al., 2009; Muerdter et al., 2013; Mugat et al., 2020; Ninova et al., 2020; Osumi et al., 2019; Rangan et al., 2011; Sienski et al., 2015; Yang et al., 2019; Yu et al., 2015). While the detailed mechanisms of transcriptional silencing remain to be established, loci targeted by Piwi are decorated in repressive chromatin marks including trimethylated H3K9 (H3K9me3) (Klenov et al., 2014; Le Thomas et al., 2013; Rozhkov et al., 2013; Sienski et al., 2012; Wang and Elgin, 2011). Loss of Piwi in Drosophila ovaries results in de-repression of TEs and correlates with a severe reduction in H3K9me3 deposition at their corresponding genomic loci.

Piwi and Aub, and to a lesser degree Ago3, have been detected as maternally deposited proteins in Drosophila embryos (Brennecke et al., 2007; Brennecke et al., 2008; Gunawardane et al., 2007; Mani et al., 2014; Megosh et al., 2006; Rouget et al., 2010). Considering that pluripotent progenitor cells give rise to multiple cell lineages, including the germline, maintaining genome integrity during the early stages of embryogenesis is potentially critical. Consistent with their adult roles, maternally inherited PIWI proteins have been observed in the pole plasm of syncytial embryos and in pole cells, the germ cell progenitors, after cellularization (Brennecke et al., 2008; Dufourt et al., 2017; Mani et al., 2014; Megosh et al., 2006).

Though, in adult flies, the piRNA pathway is restricted to the gonad, during the early phases of embryogenesis Piwi is also present in somatic nuclei (Brennecke et al., 2008; Mani et al., 2014; Megosh et al., 2006). This has long been taken as an
indication that the piRNA pathway could play roles also in the developing soma, for example helping to establish its epigenetic landscape (Gu and Elgin, 2013; Seller et al., 2019; Yuan and O’Farrell, 2016). However, probing piRNA pathway function during early embryogenesis has been hampered by a lack of suitable experimental approaches. Disrupting Piwi or other piRNA pathway factors in the female parent either via mutation or RNAi leads to oogenesis defects and often results in sterility or patterning defects that would confound the outcome of analyses (Cox et al., 1998; Czech et al., 2013; Handler et al., 2013; Khurana et al., 2010; Klattenhoff et al., 2007; Klenov et al., 2011; Li et al., 2009a; Malone et al., 2009; Mani et al., 2014; Muerdter et al., 2013; Pane et al., 2007; Park et al., 2019). RNAi-mediated depletion in embryos or generation of homozygous mutant embryos carrying piRNA pathway defects enables analysis of later developmental stages (Akkouche et al., 2017; Gu and Elgin, 2013; Marie et al., 2017), but not time windows where maternally deposited proteins predominate and generally drive development.

Here, we exploit a conditional protein degradation strategy to explore the function of maternally deposited piRNAs during Drosophila embryonic development. We find that Piwi-piRNA complexes present in the embryo are primarily derived from the oocyte, whereas, components of the PICTS complex are both maternally deposited and zygotically expressed. An embryonic burst of transposon expression in somatic cells as the zygotic genome becomes active precedes the transient decoration of normally active elements in repressive chromatin marks. Rapid and efficient degradation of maternally deposited Piwi protein in embryos leads to earlier and increased activity of zygotically expressed TEs in concert with loss of repressive marks during the affected developmental stages. Although loss of transposon control in the embryonic soma does not result in an overt morphological phenotype, our results suggest that the piRNA pathway indeed plays a role in regulating the somatic chromatin structure during early embryogenesis. Through these mechanisms, a wave of expression, primarily of the roo transposon is attenuated, though substantial expression of the transposable element remains.
Results

A transient burst of transposon expression during *Drosophila* embryogenesis

The maternal deposition of Piwi, Aub and Ago3, noted more than a decade ago (Brennecke et al., 2007; Brennecke et al., 2008; Gunawardane et al., 2007; Mani et al., 2014; Megosh et al., 2006; Rouget et al., 2010), has long suggested a possible role for the piRNA pathway during embryogenesis. Prior studies have indicated that maternal instructions transmitted via piRNAs are important for defining piRNA clusters in the subsequent generation and/or provide critical information for gaining control over at least some transposons in daughters (Akkouche et al., 2013; Akkouche et al., 2017; Brennecke et al., 2008; de Vanssay et al., 2012; Hermant et al., 2015; Khurana et al., 2011; Le Thomas et al., 2014a; Le Thomas et al., 2014b). Both of these functions are relevant in gonadal cells. Yet, prior studies highlighted the presence of maternally deposited Piwi protein in the somatic nuclei of developing embryos (Brennecke et al., 2008; Mani et al., 2014; Megosh et al., 2006), leading to suggestions that piRNAs might help set the global epigenetic landscape of the embryonic soma (Gu and Elgin, 2013). To investigate the role of the piRNA pathway during embryogenesis, we first focused on its most well-established role, that of transposon control. Toward this end, we first characterized the expression of transposons throughout *Drosophila* embryogenesis by RNA-seq and quantitative mass spectrometry (Figure 1A).

Transcriptomes of 0-2h after egg laying (AEL) embryos represent the maternally inherited mRNA pool. Maternal transcripts are cleared and the zygotic genome is activated (zygotic genome activation, ZGA) around nuclear cycle 14 (NC14; 2-2.5h AEL), and we generated RNA-seq data spanning one hour intervals of development from this point up to 10h AEL (stage 13). For comparison, we also included two late-stage embryo time points (12-13h and 17-18h AEL), as these were times when our prior data indicated that maternal Piwi was no longer detectable in somatic nuclei (Brennecke et al., 2008). We benchmarked our dataset by comparing the expression of selected embryonic genes in our RNA-seq to reported transcriptomes in Flybase (Graveley et al., 2011). We found highly similar expression patterns of genes that are dynamically regulated during embryogenesis (Figure 1-figure supplement 1A, B). Furthermore, well validated maternal (e.g., *fs(1)N* and *gammaTub37C*) and zygotic (e.g., *Ultrabithorax* (*Ubx*)) and *wingless* (*wg*)) genes demonstrated their expected expression patterns in our datasets (Figure 1-figure supplement 2A).

We detected only very few transposon transcripts in pre-ZGA embryos (0-2h AEL), as might be expected from their effective silencing by the piRNA pathway in ovaries. TE expression steadily increased following ZGA and peaked between 4-6h AEL (Figure 1B), similar to what was noted in prior reports (Batut et al., 2013). At the peak, transposon RNAs correspond to ~1.7% of the total embryonic transcriptome, with levels at the later studied time points decreasing to below 1% of the overall transcriptome. Transposons often show highly dynamic spatio-temporal expression; thus, we analysed the contribution of individual TE families to the embryonic transcriptome. Interestingly, the majority of transposon expression could be attributed to one single transposon family, *roo* (Figure 1C). At its peak at 4-6h AEL, reads derived from the *roo* TE accounted for more than 1% of the entire embryonic transcriptome, corresponding to more than 70% of all TE-derived reads overall.
its expression peak, roo mRNA levels declined strongly before levelling off at around 12h AEL. While less pronounced, other transposons, such as copia and 297, also showed dynamic expression changes during embryogenesis (Figure 1-figure supplement 2F).

Figure 1: A transient burst of transposon expression during Drosophila embryogenesis. (A) Schematic of Drosophila embryogenesis indicating Bownes stages and collected time points. (B) Bar graphs showing contribution of transposon derived reads to the transcriptome of control w^{1118} embryos at the indicated time points in percent. Error bars show standard deviation (n=2). (C) Line graphs showing the RNA expression (in rpm) for the 30 most expressed transposons during the indicated time points of embryogenesis. Error bars show standard deviation (n=2). (D) Confocal fluorescent microscopy images of control w^{1118} embryos showing nuclei stained with DAPI and roo transcripts by RNA-FISH at the indicated embryonic stages (also see Figure 1-figure supplement 2B). Scale bar=100µm.
Figure 1-figure supplement 1: Correlation of embryo collection time points between data sets. (A) Heatmaps showing expression changes in RNA-seq (in log2 reads per million; n=2) of control w1118 embryos for the indicated time points and selected genes with dynamic expression during embryogenesis. (B) As in (A) but showing reanalysed data sets published by Graveley and colleagues (Graveley et al., 2011). Grey squares=not detected.
Figure 1-figure supplement 2: Transposon protein and the known cohort of piRNA coTGS factors are present during embryogenesis. (A) Heatmaps showing expression changes in RNA-seq (left; in log2 reads per million; n=2) and protein abundance in TMT-MS (right; log2 signal intensity; n=3) of control w1118 embryos for indicated time points and selected maternal/zygotic genes, chromatin factors and the indicated piRNA pathway components. Grey squares=not detected. (B) Confocal fluorescent microscopy images of embryos showing immunofluorescence staining for Piwi protein, roo transcripts by RNA-FISH and nuclei with DAPI at the indicated embryonic stages. Scale bar=100µm. (C) Line graphs showing fold-changes in protein abundance (relative to 0-2h AEL) measured by quantitative mass spectrometry for indicated time points of control w1118 embryos. Shown are genes with increasing protein abundance (over 25% with p<0.01) between 0-2h and 5-7h AEL. (D) As in (C) but showing genes with protein abundance decreasing (over 25% with p<0.01). (E) Schematic indicating the LTR-flanked 272 kDa ORF of roo and its coding potential. Gag=Group-specific antigen-like protein, pol=reverse transcriptase, env=envelope protein, Pep=two peptidases, zf=zinc finger associated domain. Peptides detected by TMT-MS are indicated in yellow, regions not detected are shown in grey. (F) Heatmap showing protein signal intensity (arbitrary units) of the indicated transposons and time points in control w1118 embryos (n=3).
The roo expression peak at 4-6h AEL could be due to transcription from germ cell precursors, which become transcriptionally active around 3.5h AEL (stage 8) (Van Doren et al., 1998; Zalokar, 1976). However, the sheer abundance of roo and other transposon transcripts argued strongly that they must emanate at least in part from somatic nuclei, as these vastly outnumber the germ cell precursors. To test directly the origin of roo transcripts during embryogenesis, we performed RNA fluorescence in situ hybridisation (RNA-FISH). In agreement with our RNA-seq data, roo transcripts were detected as early as stage 6 (in gastrulating embryos ~3h AEL) and localised predominantly to yolk cell nuclei (Figure 1D, Figure 1-figure supplement 2B). Stage 11 embryos (~5h AEL) showed strong roo RNA signal in somatic cells of the mesoderm, similar to earlier reports (Bronner et al., 1995; Ding and Lipshitz, 1994). In contrast, roo transcript levels were undetectable by FISH in late-stage embryos (>10h AEL). These data indicate a transient somatic burst of roo expression during early Drosophila development.

TEs rely on proteins encoded in their open reading frames (ORFs) for mobilization. roo is an LTR retrotransposon and, as has been proposed for gypsy in ovarian follicle cells (Kim et al., 1994; Leblanc et al., 2000; Song et al., 1997), could potentially be packaged into virion-like particles, possibly enabling infection of germ cell precursors as a propagation mechanism. To determine whether roo-encoded proteins are expressed in embryos, we mined quantitative proteomic data from three developmental intervals (Figure 1A). The first, 0-2h AEL, represents the time before ZGA when the proteome is derived from maternal protein deposition and zygotic translation of maternal mRNAs. The second, 5-7h AEL, represents an interval where zygotic roo expression had become robust, and the third, 10-12h AEL, is a time at which roo RNA levels had substantially declined.

In transcriptionally silent embryos (0-2h AEL) we detected over 6,400 unique proteins. Compared to 0-2h embryos, the signal intensity of ~17% or 1,114 proteins significantly increased (p<0.01) by over 25% in 5-7h AEL embryos (Figure 1-figure supplement 2C). We also detected 490 (or ~8% of) proteins that significantly decreased (p<0.01) by over 25% in 5-7h AEL embryos (Figure 1-figure supplement 2D). The majority of proteins (4,652 or 72%), however, did not change by more than 25% between 0-2h and 5-7h AEL embryos. As with transcriptome analyses, known maternally deposited and zygotically expressed proteins showed their expected patterns of presence in the datasets.

Compared to the early time point (0-2h AEL), 5-7h AEL embryos showed significant accumulation of roo peptides (p<0.01) corresponding to its expression peak. roo encodes a single ORF (with a predicted protein weight of 272 kDa), which contains a Group-specific antigen-like protein (gag), a reverse transcriptase (RT/pol), an envelope protein (env), two peptidases-like domains (Pep), and a zinc finger (Figure 1-figure supplement 2E). We detected peptides corresponding to the gag, pol and env proteins (Figure 1-figure supplement 2E, bottom), indicating potential competence for retrotransposition. We additionally detected proteins derived from other transposons including copia and 297. Of note, roo ORFs remained detectable at 10-12h AEL (Figure 1-figure supplement 2F), possibly suggesting substantial stability, as this was a time at which roo mRNA levels had diminished.
The known cohort of piRNA coTGS factors is present during embryogenesis

The decline in transposon expression from 4-6h to 10-12h intervals of embryogenesis could potentially involve the piRNA pathway. However, piRNA-guided post-transcriptional or co-transcriptional silencing also requires a growing list of additional proteins (reviewed in (Czech et al., 2018; Ozata et al., 2019)). We therefore probed the expression of known piRNA pathway components during various stages of embryogenesis in our transcriptomic and proteomic datasets.

With the exception of Piwi, genes involved in coTGS were both maternally deposited and zygotically expressed during the first ~10h of embryogenesis (Figure 1-figure supplement 2A). Components of the the PICTS complex, comprising Panx, Nxf2, Nxt1 and Ctp, showed abundant protein expression in the 5-7h and 10-12h AEL time intervals. piRNA-mediated coTGS also depends on several general chromatin modifiers, including Egg and its co-factor Wde (Osumi et al., 2019; Rangan et al., 2011; Sienski et al., 2015; Yu et al., 2015). Both of these proteins are required for heterochromatin formation in the embryo, and Egg in particular has previously been implicated in embryonic repeat silencing (Seller et al., 2019). Similar to piRNA-specific coTGS factors, proteins involved in general chromatin silencing were both maternally deposited and zygotically expressed and detected at all studied time points, as expected based on their ubiquitous functions (Figure 1-figure supplement 2A). Of note, Piwi mRNA appears to be primarily maternally deposited, with zygotic transcript levels remaining low throughout embryogenesis (Figure 1-figure supplement 2A, Figure 2A, B).

In contrast, we noted little or no maternal deposition and low zygotic expression of key components of the piRNA precursor expression and export machinery and of critical piRNA biogenesis factors (Figure 1-figure supplement 2A). Considered together, our expression analyses are consistent with the potential of maternally-instructed Piwi protein acting through coTGS during Drosophila embryogenesis.

Components of the piRNA-guided coTGS machinery are enriched in somatic and pole cell nuclei during embryogenesis

To assess the potential role of the piRNA pathway in regulating the transposon burst during Drosophila embryogenesis, we examined the spatial and temporal expression of coTGS proteins in the developing embryo using light-sheet live fluorescence microscopy. For this purpose, we used two previously published lines carrying a modified BAC expressing either GFP-Piwi or GFP-Panx from its endogenous regulatory region (Handler et al., 2013; Sienski et al., 2015) and a GFP-Nxf2 knock-in line that we generated by CRISPR/Cas9 (Fabry et al., 2019). As Ctp and Nxt1 have many additional functions, we did not examine their localization in this study. We also crossed in a transgene carrying H2Av-RFP to enable tracking of nuclei. Pre-blastoderm stage embryos (0.5h AEL) were continuously imaged for >10h of embryogenesis. As previously reported (Brennecke et al., 2008; Mani et al., 2014; Megosh et al., 2006) and consistent with its maternal deposition, we detected GFP-Piwi during the pre-blastoderm stage (NC1-9, ~0-30min AEL) localized to the posterior pole where it formed a crescent-like structure (Video S1, Figure 2C).
Figure 2: piRNA coTGS factors are maternally inherited and localize to somatic cells of the *Drosophila* embryo. (A) Bar graphs showing Piwi RNA expression (in rpm) at the indicated time points in control w^{1118} embryos. Error bars show standard deviation (n=2). (B) Bar graphs showing Piwi protein signal intensity (arbitrary units) at the indicated time points in control w^{1118} embryos. Error bars show standard deviation (n=3). (C) Stand-still images from Movie S1 obtained by lightsheet fluorescent live microscopy of embryos derived from parents expressing GFP-Piwi (green) and H2Av-RFP (red) for the indicated time points. Scale bar=50µm. (D) As in (C) but showing the transition from NC13 to NC14. Scale bar=50µm. (E) Confocal fluorescent microscopy images of embryos derived from females expressing GFP-AID-Piwi crossed to control w^{1118} males probing for GFP and DAPI. Shown are embryos at the blastoderm stage (stage 5) and late-stage embryos (>12h AEL). Scale bar=100µm. Zoom of the indicated regions showing developing germ cells. Scale bar=10µm. (F) As in (E) but showing embryos derived from control w^{1118} females crossed to GFP-AID-Piwi males. (G) Bar graph showing small RNA-seq from Piwi immunoprecipitation of 0-8hour control w^{1118} embryos (green, n=1) or adult ovaries (orange, n=1). Shown are antisense piRNAs of the 30 most abundant TE families in embryos as percentage of reads mapping to indicated transposons relative to all TE-mapping antisense piRNAs.
Figure 2-figure supplement 1: Maternally inherited coTGS factors localise to nuclei of pole and somatic cells during embryogenesis. (A) Stand-still images from Movie S2 obtained by light-sheet fluorescent live microscopy of embryos derived from parents expressing GFP-Panx (green) and H2Av-RFP (red) for the indicated time points. Scale bar=50µm. (B) As in (A) but showing stand-still images from Movie S3 for GFP-Nxf2 (green) and H2Av-RFP (red) for the indicated time points. Scale bar=50µm. (C) Quantification of co-localisation between GFP-Piwi and H2Av-RFP. Plot profile shows relative intensity of GFP-Piwi (green) and H2Av-RFP (red) protein for the indicated section (white line) of stage 4 (2h AEL) embryos. (D) As in (C) but showing embryos expressing GFP-Panx and H2Av-RFP. (E) As in (C) but showing embryos expressing GFP-Nxf2 and H2Av-RFP. (F) As in (A) but showing the transition of GFP-Panx from NC13 to NC14. Scale bar=50µm. (G) As in (B) but showing the transition of GFP-Nxf2 from NC13 to NC14. Scale bar=50µm. (H) Bar graphs showing Panx RNA expression (in rpm) at the indicated time points in control w^{1118} embryos. Error bars show standard deviation (n=2). (I) Bar graphs showing Panx protein signal intensity (arbitrary units) at the indicated time points in control w^{1118} embryos. Error bars show standard deviation (n=3). (J) As in (H) but showing the Nxf2 RNA expression. (K)
As in (I) but showing Nxf2 protein expression. (L) Confocal fluorescent microscopy images of developing germ cells of late-stage embryos derived from females expressing GFP-AID-Piwi crossed to control w^{1118} males probing for GFP and DAPI. Scale bar=10µm. (M) As in (L) but showing developing germ cells of late-stage embryos derived from control w^{1118} females crossed to GFP-AID-Piwi males. (N) Bar graph showing size distribution of Piwi-associated small RNAs (in reads per million, rpm) from 0-8h w^{1118} embryos (n=1). Blue=sense, red=antisense. (O) As in (N) but showing Piwi-associated small RNAs from adult w^{1118} ovaries (n=1).
As embryogenesis progressed and somatic nuclei migrated to the surface (NC 9-14, ~1.5-3h AEL), Piwi localised to somatic nuclei and to the pole plasm surrounding the nuclei of germline progenitor cells, as we and others reported earlier based on immunofluorescence staining of fixed embryos (Brennecke et al., 2008; Mani et al., 2014; Megosh et al., 2006). In agreement with an earlier report (Mani et al., 2014), our dynamic data revealed that nuclear Piwi signal strongly decreased during mitotic cycles (Video S1, Figure 2D). We continued to detect Piwi expression in somatic nuclei throughout the first 10h of embryogenesis; however, signal intensity decreased over time. This observation was consistent with transcriptomic and proteomic measurements taken over a comparable time course (Figure 2A, B).

Similar to Piwi, both Nxf2 and Panx were detected in somatic and pole cell nuclei from the syncytial blastoderm stage (Video S2, S3, Figure 2-figure supplement 1A-E). In contrast to Piwi, Panx and Nxf2 showed strong co-localisation with H2Av-RFP during mitotic cycles (Video S2, S3, Figure 2-figure supplement 1F, G), suggesting that while Piwi is predominantly excluded, Nxf2 and Panx are retained in the nucleoplasm during mitosis. Consistent with our RNA-seq and TMT-MS data (Figure 2-figure supplement 1H-K), as embryogenesis progressed, Panx and Nxf2 remain detectable for several hours (>10h AEL), closely matching the protein expression of Piwi.

Piwi carries epigenetic information in the form of piRNAs (Brennecke et al., 2008; Le Thomas et al., 2014b). However, it is unclear if Piwi-piRNA complexes are assembled during oogenesis prior to maternal deposition into the embryo, or whether zygotic piRNA biogenesis and Piwi loading also occurs. We therefore analysed the expression of GFP-tagged Piwi from reciprocal crosses with control w^{1118} flies by immunofluorescence staining in early and late-stage embryos. Embryos derived from females expressing GFP-Piwi showed strong maternal deposition of Piwi during early embryogenesis (Figure 2E, Figure 2-figure supplement 1L), with GFP fluorescence in later stage (>12h AEL) embryos restricted to the germline cells. Consistent with maternal deposition of Piwi, embryos derived from the reciprocal cross showed no GFP signal in the early embryos (Figure 2F, Figure 2-figure supplement 1M). Instead, we only observed GFP-Piwi signal in the developing gonads of late-stage embryos, likely as a result of zygotic expression. Strikingly, Piwi of zygotic origin localised exclusively to the cytoplasm of the germ cell progenitors and was not detected in nuclei, suggesting that zygotically transcribed Piwi is likely not relevant for coTGS until later in development.

The decline in embryonic transposon expression is correlated with hallmarks of piRNA-dependent co-transcriptional silencing.

Piwi proteins are guided by their piRNA cofactors to recognize and co-transcriptionally silence active transposons in the Drosophila ovary (Le Thomas et al., 2013; Post et al., 2014; Sienski et al., 2015; Sienski et al., 2012; Yu et al., 2015). If this pathway was relevant in the embryonic soma, maternally deposited Piwi would require instructions to recognize embryonically expressed elements. To examine this possibility, we immunoprecipitated Piwi from 0-8h control w^{1118} embryos as well as from adult ovaries and sequenced the associated small RNAs. Piwi in both tissues existed in complex with 23- to 28-nt piRNAs and showed nearly indistinguishable size profiles that were biased for antisense reads (Figure 2-figure supplement 1N, O).
Closer inspection by aligning the reads to transposon consensus sequences revealed similar piRNA levels for the majority of TEs, however, we detected some notable differences (Figure 2G). Piwi in ovaries showed higher levels of antisense piRNAs targeting the TEs mdg1 and 412, in agreement with the majority of these small RNAs originating from the soma-specific flam piRNA cluster (Brennecke et al., 2007; Malone et al., 2009; Zanni et al., 2013). Piwi in embryos showed high levels of antisense piRNAs targeting roo (~16% or all TE-targeting reads) and 297 (~9%), consistent with an ability of maternally deposited Piwi to potentially recognize these TEs when expressed in the embryo.

In Drosophila ovaries, coTGS depends on Piwi-mediated recruitment of the PICTS/SFINX complex and correlates with the deposition of H3K9me3 marks at TE insertions and surrounding genomic regions (Batki et al., 2019; Eastwood et al., 2021; Fabry et al., 2019; Murano et al., 2019; Schnabl et al., 2021; Sienski et al., 2015; Yu et al., 2015; Zhao et al., 2019). Due to the poor conservation of the genomic locations of transposon insertions between different Drosophila strains, we used whole genome sequencing to de novo identify the TE insertion sites present in our control w^{1118} flies (see methods). This data enabled us to identify over 600 euchromatic transposon insertions that are absent from the dm6 reference genome, and these were used for our chromatin analyses, as most annotated insertions in the dm6 genome assembly were absent from our strain.

In order to determine the fate of transposon loci throughout embryogenesis, we performed H3K9me3 chromatin immunoprecipitation followed by sequencing (ChIP-seq) on control w^{1118} embryos at two-hour intervals covering the period when transposon expression is dynamic (0-10h AEL) and a later time point (16-18h AEL) well after maternal Piwi protein was no longer detectable in somatic nuclei (Figure 1A, 2C). We included adult ovaries, which show piRNA-guided coTGS, as well as adult heads, a somatic tissue without active piRNA pathway, to compare the changes of this repressive chromatin mark across different stages and tissues of Drosophila development.

Early embryos (0-2h AEL) showed low levels of H3K9me3 signal at 117 euchromatic roo insertions (Figure 3A). However, as development progressed, H3K9me3 levels increased with a peak at 6-10h AEL (Figure 3A, Figure 3-figure supplement 1A). Thus, deposition of repressive chromatin marks correlated with the RNA expression of roo, yet the maximum of H3K9me3 accumulation lagged behind the RNA expression peak by approximately two hours. These data are consistent with a requirement for nascent transcription for efficient conversion of a TE insertion into heterochromatin, as previously reported in yeast (Buhler et al., 2006; Shimada et al., 2016) and for the recognition of transposon loci by the piRNA pathway (Le Thomas et al., 2013; Post et al., 2014; Rozhkov et al., 2013; Sienski et al., 2015; Sienski et al., 2012; Yu et al., 2015). Of note, the deposition of repressive marks trailed the direction of transcription and showed higher signal enrichments in the regions downstream of the transposon insertions, as previously observed for piRNA-dependent silencing in cell culture systems (Fabry et al., 2019; Sienski et al., 2015; Sienski et al., 2012).
Figure 3: Transposon insertions targeted by piRNAs in embryos show epigenetic changes characteristic of co-transcriptional gene silencing. (A) Heatmaps (top) and metaplots (bottom) showing H3K9me3 ChIP-seq signal (in rpm) for the indicated control w1118 embryonic stages and adult tissues at 177 euchromatic roo insertions (n=2). Signal is shown within 10-kb from insertion site and sorted from 5’ to 3’. (B) UCSC genome browser screenshot showing H3K9me3 ChIP-seq signal for the indicated genes on chromosome 2R carrying w1118-specific roo and Doc insertions.
Figure 3-figure supplement 1: Transposons targeted by maternally inherited piRNAs show hallmarks of coTGS. (A) Bar graphs showing H3K9me3 signal intensity for the indicated time points at roo insertions. Error bars show standard deviation (n=2). (B) Heatmaps (top) and metaplots (bottom) showing H3K9me3 ChIP-seq signal (in rpm) for the indicated control w^{118} embryonic stages and adult tissues at 20 euchromatic 297 insertions (n=2). Signal is shown within 10-kb from insertion site and sorted from 5' to 3'. (C) Metaplots showing H3K9me3 ChIP-seq signal (in rpm) for the indicated control w^{118} embryonic stages and adult tissues at 18 euchromatic mdg1 (top) and at 22 euchromatic 412 insertions. Signal is shown within 10-kb from insertion site and sorted from 5' to 3' (n=2). (D) Bar graphs showing H3K9me3 signal intensity for the indicated time points at constitutive heterochromatin. Error bars show standard deviation (n=2). (E) As in (D) but showing chromosome 4 regions.
Interestingly, H3K9me3 signal at euchromatic roo insertions of 16-18h AEL embryos, which lacked maternal Piwi in somatic nuclei and no longer express roo, showed diminished intensities compared to earlier time intervals. Similarly, heads and ovaries, both tissues from adult flies, showed no enrichment of H3K9me3 at euchromatic roo insertions, despite the presence of a functional piRNA pathway in ovaries. Considered together, these data suggest that maternal piRNAs program a response to a burst of roo expression during embryonic development but that the deposition of H3K9me3 marks, likely directed via coTGS, no longer occurs at developmental time points and in tissues where roo is not expressed. This is consistent both with the known requirement for active transcription for targeting by Piwi and with the observed need for continuous engagement of the PICTS/SF1NX complex to maintain H3K9me3 marks on transposon loci (Batki et al., 2019; Eastwood et al., 2021; Fabry et al., 2019; Le Thomas et al., 2013; Murano et al., 2019; Post et al., 2014; Rozhkov et al., 2013; Schnabel et al., 2021; Sienski et al., 2015; Sienski et al., 2012; Yu et al., 2015; Zhao et al., 2019).

To investigate whether this mechanism is specific to roo or more general, we examined the transposon 297, which is also expressed during embryogenesis (Figure 1-figure supplement 1F) and showed high targeting potential by maternally inherited piRNAs (Figure 2G). Genomic loci in close proximity to euchromatic 297 insertions (n=20) showed dynamic deposition of H3K9me3 similar to roo (Figure 3-figure supplement 1B). However, while H3K9me3 levels at roo insertions peaked between 6-10h AEL, 297 insertions showed the maximum H3K9me3 signal intensity between 2-8h AEL, suggesting that these loci are targeted by coTGS earlier than roo insertions. In contrast, H3K9me3 occupancy at transposons such as mdg1 and 412 that were expressed during embryogenesis but lacked substantial maternal deposition of piRNAs, retained low H3K9me3 levels throughout embryogenesis, though they showed a strong enrichment in ovaries (Figure 3-figure supplement 1C).

To determine whether the deposition of repressive chromatin marks at euchromatic 297 and roo insertions was specific, rather than reflecting a general trend of H3K9me3 accumulation genome-wide, we analysed genomic regions not targeted by maternally inherited piRNAs. H3K9me3 signal at constitutive heterochromatin remained stable throughout the sampled time points (Figure 3-figure supplement 1D), while H3K9me3 levels on chromosome 4 increased steadily throughout development (Figure 3-figure supplement 1E). Of note, while ovaries showed no coTGS signature at roo insertions, other transposons, such as Doc, showed a clear accumulation of H3K9me3 marks that was absent in embryos during all assayed time points (Figure 3B). Considered together, these results are consistent with piRNA-guided chromatin modification of a subset of transposons that show activity during Drosophila embryonic development.

An auxin-inducible degron enables rapid depletion of Piwi in ovaries and early embryos

Though embryonically repressed transposons bore hallmarks of piRNA-guided heterochromatin formation, the reliance of the pathway on maternally deposited Piwi-piRNA complexes prevented a demonstration that silencing depended on the pathway through conventional genetics. Ovaries that lack key piRNA pathway silencing factors show substantial expression changes and produce morphologically altered eggs that...
largely fail to develop normally (Cox et al., 1998; Czech et al., 2013; Handler et al., 2013; Khurana et al., 2010; Klattenhoff et al., 2007; Klenov et al., 2011; Li et al., 2009a; Malone et al., 2009; Mani et al., 2014; Muerdter et al., 2013; Pane et al., 2007; Park et al., 2019).

To investigate the effect of Piwi depletion on *Drosophila* embryogenesis without affecting oogenesis, we used the auxin-inducible degron (AID) system (Nishimura et al., 2009). This protein degradation system is comprised of an AID-tag, fused to the protein of interest, and the plant-derived F-box protein transport inhibitor response 1 (TIR1). AID and TIR1 associate with each other in an auxin-dependent manner, with binding of TIR1 to the AID-tagged target leading to the recruitment of the cellular ubiquitination machinery and target protein degradation via the proteasome (Figure 4A). This conditional degradation system has proven effective in several model organisms including *Drosophila* where it was recently shown to enable degradation of the germ cell-specific protein Vasa (Bence et al., 2017).

We used CRISPR/Cas9 to insert an amino-terminal GFP-AID tag at the *Drosophila piwi* locus and crossed these flies to a line expressing the *Oryza sativa*-derived TIR1 (OsTIR1) protein under the control of the ubiquitin promoter. As a proof of concept, we tested the auxin-induced degradation of Piwi in adult ovaries of flies homozygous for both GFP-AID-Piwi and OsTIR1. Feeding flies for 24h with 5mM auxin-containing yeast paste was sufficient to induce complete degradation of Piwi in ovaries (Figure 4B, C, Figure 4-figure supplement 1A), and this depletion resulted in the de-repression of transposons (Figure 4D). Notably stronger changes were observed following longer treatments, possibly implying a lag between loss of piRNA pathway function and that of repressive chromatin marks. Following a 1-day treatment, embryos laid by Piwi-depleted females developed without defects and showed similar hatching rates as their control treated siblings (Figure 4E, F). Longer auxin treatments resulted in more frequent deformation of embryos that was accompanied by reduced hatching rates (Figure 4E, F), likely due to patterning defects as a result of Piwi depletion from follicle cells.

*Drosophila* embryos develop within a relatively impermeable chorion, and treatment of embryos directly with auxin showed little impact. However, in dechorionated embryos we observed a near complete degradation of Piwi protein following 30min auxin treatment of embryos collected for 0-30min AEL (Figure 5A, B). To investigate the dynamics of auxin-mediated Piwi depletion in embryos, we used light-sheet fluorescence live microscopy. Early blastoderm embryos treated with 5mM auxin showed rapid degradation of GFP-AID-Piwi signal, which was undetectable after 25min of treatment (Figure 5C, Video S4). Of note, the removal of maternal Piwi in this time window did not significantly affect the embryo hatching rate (Figure 4-figure supplement 1B).
Figure 4: Degradation of Piwi protein in ovaries resembles mutant phenotypes. (A) Cartoon illustrating the Piwi protein degradation strategy using the auxin-inducible AID-TIR1 system. (B) Confocal fluorescent microscopy images showing ovary egg chambers of GFP-AID-Piwi; OsTIR1 flies fed with yeast paste containing 5mM auxin for the indicated time (also see Figure 4-figure supplement 1A). Blue=DAPI. Green=GFP-AID-Piwi. (C) Western blot of ovaries from females treated with 5mM auxin-containing yeast paste for the indicated time period or control females probing for Piwi and Tubulin as a loading control. (D) Bar graphs showing rps49-normalised steady-state RNA levels of the indicated TEs and control genes in ovaries of GFP-AID-Piwi; OsTIR1 flies fed with yeast paste containing 5mM auxin for the indicated time. Error bars show standard deviation (n=3). Asterisk denotes significant changes compared to control (p<0.05, unpaired t-test). (E) Bar graphs showing the percentage of embryo deformation phenotypes laid by GFP-AID-Piwi; OsTIR1 females fed with yeast paste containing 5mM auxin for the indicated time. (F) As in (E) but showing the hatching rate in percent.
Figure 4-figure supplement 1: Piwi degradation in ovaries resembles knockdown and mutant phenotypes. (A) Confocal fluorescent microscopy images showing ovary egg chambers of GFP-AID-Piwi; OsTIR1 flies fed with yeast paste containing 5mM auxin for the indicated time. Blue=DAPI, Green=GFP-AID-Piwi. (B) Bar graphs showing the percentage of hatched embryos laid by auxin- or control-treated females after 24h (left) and of embryos treated for 2.5h with 5mM auxin following dechorionation (right). Statistical significance was determined by unpaired (two sample) t-test.
Maternally deposited Piwi directs heterochromatin formation at active transposon insertions during early embryogenesis

We next investigated the impact of degrading maternal Piwi from early-stage embryos on transposons. Embryos derived from flies expressing GFP-AID-Piwi and OsTIR1 were collected across a 30min period and treated for an additional 2.5h with or without 5mM auxin before RNA extraction and library generation (Figure 5A). These embryos corresponded to 2.5-3h AEL, the point at which we began to observe zygotic roo transcripts (Figure 1C). The majority of transposons showed no significant expression change upon Piwi depletion, however, roo and 297 were significantly de-repressed (p<0.05) by more than 2-fold (Figure 5D), suggesting that Piwi impacts their expression during embryogenesis. Previous studies suggested that auxin in small concentrations has a negligible impact on Drosophila development (Bence et al., 2017; Trost et al., 2016) but to control for effects of auxin itself on TE regulation, we also evaluated transposon expression in auxin-treated GFP-AID-Piwi embryos that lack OsTIR1. Without OsTIR1, 2.5-3h embryos treated with 5mM auxin showed no significant changes in transposon expression, compared to control siblings treated with PBS (Figure 5-figure supplement 1A).

We additionally examined changes in the repressive chromatin mark H3K9me3 to determine whether these were deposited in a piRNA-dependent fashion, at euchromatic roo and 297 transposon insertions. We again collected embryos for 30min and treated with 5mM auxin (or PBS as a negative control) for 6h which yielded embryos 6-6.5h AEL (Figure 5A) and corresponds to the peak in H3K9me3 signal at roo insertions in control w^1118 embryos (Figure 3A, Figure 3-figure supplement 1A). Piwi depletion severely impacted H3K9me3 signal over the transposon consensus sequence of roo and 297, but not that of other TEs (Figure 5E). Additionally, H3K9me3 levels at individual transposon genomic loci (see methods for identification of TE insertions in our fly stock) showed similar patterns. H3K9me3 signal in genome-wide 5-kb bins predominantly changed when roo or 297 insertions were nearby (Figure 5-figure supplement 1B). We identified 154 bins with significantly reduced (p<0.05) H3K9me3 occupancy, while only two bins showed an increase. Of the bins with lower H3K9me3 signal, 122 and 10 were within 5-kb of roo or 297 insertions, respectively, thereby illustrating the impact of Piwi on chromatin states at genomic loci specifically targeted by maternal piRNAs. Furthermore, individual euchromatic TE insertions of both roo and 297 showed a strong decrease of H3K9me3 levels in Piwi-depleted embryos (Figure 5F, G, Figure 5-figure supplement 1C-E), while H3K9me3 levels at constitutive heterochromatin and on chromosome 4 were not affected (Figure 5H, I). Of note, while roo and 297 TE levels were elevated upon auxin treatment in 2.5-3h AEL embryos, transposon expression returned to baseline levels comparable to untreated embryos in 6-7h and 7-8h AEL time intervals (Figure 5-figure supplement 1C-E). Taken together, these data strongly indicate a relationship between the deposition of repressive H3K9me3 chromatin marks at transposon insertions and maternally deposited Piwi-piRNA complexes (Figure 5J).
Figure 5: Degradation of maternally deposited Piwi in embryos leads to transposon deregulation. (A) Schematic of embryo auxin treatments and sample collection for RNA-seq and ChIP-seq experiments. (B) Western blot showing abundance of GFP-AID-Piwi fusion protein in embryos treated with 5mM auxin for 2h. OsTIR1 and tubulin expression are shown as loading control. (C) Stand-still images from Movie S4 obtained by light-sheet fluorescent live microscopy of embryos derived from parents expressing GFP-AID-Piwi and OsTIR1 treated with 5mM auxin for the indicated time intervals. Scale bar=50µm. (D) MA plot showing base mean expression (log10 scale) of transposon RNAs relative to their fold-change (log2 scale) in GFP-AID-Piwi; OsTIR1 embryos treated with 5mM auxin versus control (n=3). Grey=genes. Green=TEs not changed (p>0.05). Red=TEs significantly changed (p<0.05) and fold-change>2. (E) MA plot showing base mean signal intensity (log10 scale) of TEs relative to the H3K9me3 ChIP-seq signal enrichment (log2 scale) in GFP-AID-Piwi; OsTIR1 embryos treated with 5mM auxin versus control (n=3). Grey=TEs not significantly changed (p>0.05). Purple=TEs significantly changed (p<0.05). Blue=TEs significantly changed (p<0.05) and fold-change<-2. (F) Heatmaps (top) and metaplots (bottom) showing H3K9me3 ChIP-seq signal (in rpm) for control embryos and 5mM auxin treated embryos at 205 euchromatic roo insertions (n=3). Signal is shown within 10-kb from insertion site and sorted from 5’ to 3’. (G) Bar graphs showing H3K9me3 signal intensity (in rpm) for the indicated treatments at roo loci. Error bars show standard deviation (n=3). Statistics were calculated with unpaired (two-sample) t-test. (H) As in (G) but showing constitutive heterochromatin. (I) As in (G) but showing chromosome 4 regions. (J) Model of piRNA-guided chromatin modification at active transposons in somatic cells of the developing Drosophila embryo.
Figure 5-figure supplement 1: Piwi depletion in embryos leads to epigenetic and expression changes at TE insertions targeted by maternally inherited piRNAs. (A) MA plot showing base mean expression (log_{10} scale) of genes and transposons relative to their fold-change (log_2 scale) in GFP-AID-Piwi embryos (without OsTIR1) treated with 5mM auxin versus control (n=3).
Grey=genes. Green=TEs not changed (p>0.05). **(B)** MA plot showing base mean signal intensity (log scale) of TEs relative to the H3K9me3 ChIP-seq signal within 5-kb genomic bins of GFP-AID-Piwi; OsTIR1 embryos treated with 5mM auxin versus control (n=3). Grey=unchanged (p>0.05). Green=significantly changed (p<0.05). Blue=significantly changed (p<0.05) within 5-kb of a roo insertion. Red=significantly changed (p<0.05) within 5-kb of a 297 insertion. **(C)** Heatmaps (top) and metaplots (bottom) showing H3K9me3 ChIP-seq signal (in rpm) for control embryos and 5mM auxin treated embryos at 63 euchromatic 297 insertions (n=3). Signal is shown within 10-kb from insertion site and sorted from 5’ to 3’. **(D)** UCSC genome browser screenshot showing RNA-seq and H3K9me3 ChIP-seq signal of control and auxin-treated embryos at the indicated roo insertion on chromosome 3L. **(E)** As in (D) but showing a genomic locus with two 297 insertion sites. **(F)** Bar graphs showing rp49-normalised steady-state RNA levels of the indicated TEs and control gene in GFP-AID-Piwi; OsTIR1 embryos at the indicated time intervals (AEL) treated with 5mM auxin compared to PBS treated control embryos. Error bars show standard deviation (n=2). **(G)** Scatter plot showing RNA expression levels (in reads per million, log2 scale) of transposons from ovaries of nxf2 heterozygote and homozygote mutants (n=3). Transposons whose abundance change more than two-fold compared to heterozygotes are highlighted in red, roo is shown in orange. Data re-analysed from (Fabry et al., 2019). **(H)** As in (G) but showing H3K9me3 ChIP-seq levels. Transposons with more than two-fold reduced levels in the mutant compared to heterozygotes are shown in blue.
Discussion

Here, we have examined the role of the Piwi-directed coTGS arm of the piRNA pathway during early embryogenesis in Drosophila. By far, the majority of our insight into the function of piRNAs has derived from studies in germ cells or in the support cells of reproductive tissues. The intriguing observation that piRNAs and their Piwi-family binding partners are maternally deposited has led to speculation regarding potential roles for piRNAs in inter- and transgenerational epigenetic inheritance. Indeed, maternal piRNAs are critical in the suppression of hybrid dysgenesis induced by paternal transmission of I- or P-elements in matings with females that lack these transposons (Brennecke et al., 2008; Khurana et al., 2011). Epigenetic modifications induced by piRNAs appear to aid in piRNA cluster definition in the germline (Akouche et al., 2017). Additionally, maternally deposited Aub-piRNA complexes have been implicated in embryonic gene regulation (Barckmann et al., 2015; Dufourt et al., 2017; Rouget et al., 2010). Last, maternally inherited piRNAs control transposon expression in interspecies hybrids between D. melanogaster and D. simulans (Kelleher et al., 2012) and regulate the TE tirant in the somatic compartment of the female gonad in D. simulans (Akouche et al., 2013). Yet, to date, the lack of mechanisms to rapidly deplete maternally deposited PIWI proteins specifically from early embryos has hampered our ability to broadly assess their zygotic roles. By fusing a chemically inducible degron to Piwi, we were able to deplete Piwi-piRNA complexes from dechorionated embryos within less than 30min of treatment and well before the nuclear accumulation of Piwi that is observed following activation of zygotic transcription.

Though nuclear localisation of Piwi correlates with the appearance of its potential targets, nascent transcripts of transposons, it is unclear what triggers movement of Piwi into the somatic nuclei. Notably, nuclear translocation of Piwi lags behind in germ cell precursor nuclei, and this correlates with the observation that these nuclei activate transcription of their genomes later in embryogenesis than somatic nuclei do (Van Doren et al., 1998; Zalokar, 1976). Our dynamic imaging of Piwi localization also revealed that it shuttles out of nuclei during mitosis, as previously observed (Mani et al., 2014). Since other factors of the pathway, namely components of the PICTS/SF\(i\)NX complex, remain nuclear, it is likely that Piwi is actively excluded. Several studies have previously shown that nuclear localization of Piwi is conditional upon its binding to a piRNA partners (Klenov et al., 2011; Saito et al., 2009; Yashiro et al., 2018), but we have no indication that Piwi is unloaded and reloaded during mitotic cycles. Rather, we hypothesize that another mechanism regulates the activity of the Piwi nuclear localization signal, though what purpose this might serve and whether it also occurs in germline and follicle cells or is restricted to embryogenesis remains unclear.

It has been suggested that the evolution of the abbreviated piRNA pathway in ovarian follicle cells arose as a consequence of the lifestyle adopted by gypsy family elements. These retrotransposons show their highest expression levels in the support cells surrounding the developing germline. gypsy family elements encode an envelope protein and have been shown to assemble into virus-like particles (Kim et al., 1994; Leblanc et al., 2000; Song et al., 1997). This has led to the hypothesis that their ancestral propagation strategy combined evasion of TE repression mechanisms present in germ cells with an ability to create particles that could infect the germline,
where the element could insert into the genome of the developing oocyte following reverse transcription (Kim et al., 1994; Leblanc et al., 2000; Song et al., 1997).

While this remains speculative, it does provoke questions of whether a similar strategy is adopted by roo in the embryo. roo is a quite successful element, as indicated by it being the element with the highest copy number of individual insertions in our sequenced strains (9.4% of all identified TE insertions in the w^{1118} strain and 9.9% in our degron line). How this is achieved remains mysterious, since roo expression is extremely low in the ovary. Moreover, roo does not appear to be a target of the ovarian piRNA pathway, since its gonadal expression is not increased nor does its HP1a enrichment and H3K9me3 levels change in piRNA pathway mutant animals (Figure 5-figure supplement 1G, H) (Senti et al., 2015; Wang and Elgin, 2011). In the embryo, the expression of roo is restricted to somatic cells, especially cell lineages giving rise to the adult mesoderm. Previous studies have suggested that roo expression is activated by twist (twi) and snail (sna), which are highly expressed in the embryonic mesoderm (Bronner et al., 1995), and this is consistent with the spatial expression pattern that we also observe. roo expresses the full repertoire of proteins needed to form virus-like particles, and its high expression levels (exceeding 1% of the transcriptome at its peak) might enable a strategy of propagation by infection in trans, even if rates of transmission to the germ cell precursors are relatively low.

Our data strongly suggests that only maternally deposited piRNAs engage Piwi in the soma of the developing embryo. Since roo is not regulated by the piRNA pathway in the ovary, evolutionary pressures must have driven the development of a set of maternal instructions that are inherited to dampen the burst of roo expression in the developing embryo. Indeed, 16% of maternally deposited piRNAs target roo. Though there are differences between the populations of piRNAs observed in ovaries as compared to embryos, this is mostly driven by the presence of follicle cell piRNAs in in samples taken from the gonad. In our small RNA analyses, we collapse all stages of oogenesis. Thus, it is not clear whether the composition of piRNA populations shifts as the ovarirole matures and whether any such shifts enrich for embryonically expressed elements late in oocyte maturation. Irrespective, a set of instructions from maternal piRNA clusters clearly builds a transgenerational ability to recognize roo and other embryonically expressed elements.

Consistent with its recognition by Piwi-piRNA complexes and recruitment of the PICTS/SF1NX complex, H3K9me3 marks build at presumably active, euchromatic roo insertions as embryos progress toward stage 13 (10h AEL). The peak of H3K9me3 abundance lags about 2h behind the peak of transcription. Since we have little other information on the dynamics of piRNA-mediated silencing, it is not clear whether this is an expected observation or whether there may be mechanisms that antagonize the ability of the piRNA pathway to immediately recognize and direct heterochromatinization of expressed roo insertions. Of note, we see a shorter interval between the embryonic peak of 297 expression and its peak of H3K9me3 accumulation. Imposition of a repressive chromatin state on roo is transient during somatic development. By 13h AEL, H3K9me3 peaks over roo insertions have disappeared, but roo expression has not returned. The lack of H3K9me3 also correlates with the absence of critical piRNA pathway proteins in the soma. Overall, this suggests that both the expression of these TEs and the host response via small RNAs is transient.
Though zygotic depletion of maternal Piwi during early embryogenesis does produce a statistically significant change in roo expression (roughly 2-fold), this transposon remains highly expressed reaching up to 1% of the entire transcriptome in control animals, despite being targeted by the piRNA pathway. This provokes the question of whether targeting of roo by the piRNA pathway is biologically relevant. In favour of this hypothesis are several observations. Roo is expressed in ovaries at very low levels, yet the hallmarks of piRNA-dependent silencing, specifically H3K9me3, are absent from euchromatic roo insertions. This strongly indicates that roo is not controlled by the piRNA pathway in this tissue. Nonetheless, ovaries produce abundant roo piRNAs, and these are overwhelmingly in the antisense orientation. Additionally, the only uni-strand cluster expressed in germ cells, cluster 20A, has collected roo insertions in the antisense orientation. These piRNAs are abundantly maternally transmitted (16% of all piRNAs in embryos) and persist throughout the time during early embryogenesis when high level roo expression is proposed to be driven by mesodermal transcription factors. An argument against biological significance is the lack of a clearly observable phenotype in flies following embryonic depletion of maternal Piwi. However, technical limitations enable us to only measure impacts within a single generation. It is entirely possible that the fitness cost of roo occupying 2% of the embryonic transcriptome might be substantial over time or in conditions flies might experience in the wild compared to the controlled rearing conditions in the lab.

Perhaps more importantly, our study demonstrates that recognition of a locus by the piRNA pathway does not necessarily impose the creation of a mitotically heritable epigenetic state. This is consistent with observations made by many groups in follicle cells wherein heterochromatin-mediated silencing of somatic transposons requires the continuous presence of the piRNA machinery (Batki et al., 2019; Clark et al., 2017; Donertas et al., 2013; Fabry et al., 2019; Muerdter et al., 2013; Murano et al., 2019; Ohtani et al., 2013; Saito et al., 2009; Sienski et al., 2015; Sienski et al., 2012; Zhao et al., 2019). These data are at odds with prior observations and speculation that the maintenance of silenced epigenetic states at transposon insertions can be primed by Piwi but maintained in a Piwi-independent mechanism throughout adult life (Gu and Elgin, 2013). It must be considered that the prior study examined a different set of genomic loci in different Drosophila strains and depleted Piwi to a different degree using alternative methods; however, we do not observe changes in H3K9me3 on the transposable elements they examined upon zygotic depletion of maternal Piwi.

A recent detailed and elegant study examined the patterns of H3K9me3 accumulation during early embryogenesis in D. miranda (Wei et al., 2021). Though overall, deposition of H3K9me3 did not correlate with the abundance of maternally deposited piRNAs, a set of the earliest heterochromatin nucleating elements were associated with abundant piRNAs. These targeted elements had high copy numbers and showed evidence of recent transposition activity, suggesting that they were under evolutionary pressure for robust silencing both in the ovary and the soma. It should be noted that precise nucleation sites did not necessarily overlap with abundant piRNAs, suggesting that multiple silencing mechanisms might collaborate to repress these transposon families.

Considered as a whole, our data are consistent with a role for maternally deposited piRNAs in the recognition of transposon families which have focused their expression...
and activity during early embryogenesis. However, our data does not support a model wherein the piRNA pathway nucleates heritable patterns of heterochromatin formation that broadly pattern the epigenetic landscape of the adult Drosophila soma, and this is perhaps consistent with our failure to observe consequential developmental abnormalities upon negation of embryonic Piwi function.
### Material and Methods

#### Key Resources Table

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**Fly stocks and handling**

All flies were kept at 25°C on standard cornmeal or propionic food. Flies expressing GFP-Nxf2 from the endogenous locus were generated by CRISPR/Cas9 (Fabry et al., 2019). Transgenic flies carrying a BAC transgene expressing GFP-Panx and GFP-Piwi were generated by the Brennecke lab (Handler et al., 2013; Sienski et al., 2015) and obtained from the Vienna Drosophila Resource Center. Control w<sup>118</sup> flies were a
gift from the University of Cambridge Department of Genetics Fly Facility and flies
expressing His2Av-RFP were a gift from the St Johnston lab. Flies between 3 to 14
days after hatching were used for experiments.

Generation of fly strains
GFP-AID-Piwi flies were generated by CRISPR/Cas9 genome engineering. Homology
arms of 1-kb flanking the targeting site were cloned into pUC19 by Gibson Assembly
and co-injected with pCFD3 (addgene # 49410) containing a single guide RNA (Port
et al., 2014) into embryos expressing vas-Cas9 (Bloomington Drosophila Stock Center
# 51323). Flies expressing OsTIR1 under the Drosophila melanogaster Ubiquitin-63E
promoter were generated by phiC31 integrase-mediated transgenesis by injection of
plasmids containing expression cassettes for proteins into embryos of genotype "y w
P[y[+t7.7]=nos-phiC31\int.NLS\]X #12;; P[y[+t7.7]=CaryP]attP2" resulting in transgene
integration on chromosome 3. Microinjection and fly stock generation was carried out
by the University of Cambridge Department of Genetics Fly Facility. Transgenic and
knock-in flies were identified by genotyping PCRs and confirmed via sanger
sequencing.

Western blot
Protein concentration was measured using a Direct Detect Infrared Spectrometer
(Merck). 20µg of proteins were separated on a NuPAGE 4–12% Bis-Tris gel (Thermo
Fisher Scientific). Proteins were transferred for 2h at 100V, 400mA, 100W on an
Immun-Blot Low Fluorescent PVDF Membrane (BioRad) and blocked for 1h in 1x Licor
TBS Blocking Buffer (Licor). Primary antibodies were incubated over night at 4°C. Licor
secondary antibodies were incubated for 45min at room temperature (RT) and images
acquired with an Odyssey CLx scanner (Licor).

Drosophila ovary immunofluorescence
Fly ovaries were dissected in ice-cold PBS and fixed in 4% PFA diluted in PBS for
15min at room temperature while rotating. Following 3 rinses and three 10min washing
steps in PBS-Tr (0.3% Triton X-100 in PBS), ovaries were blocked for 2h at RT while
rotating in PBS-Tr +1% BSA. Primary antibody incubation was carried out in blocking
buffer overnight at 4°C while rotating, followed by three washing steps for 10min each
in PBS-Tr. All following steps were performed in the dark. Secondary antibodies were
dilated in blocking buffer and incubated overnight at 4°C while rotating. Ovaries were
washed 4 times for 10min in PBS-Tr and stained with 0.5µg/ml DAPI (Thermo Fisher
Scientific) for 10min. Following two additional washing steps for 5min in PBS, ovaries
were mounted in ProLong Diamond Antifade Mountant (Thermo Fisher Scientific) and
imaged on a Leica SP8 confocal microscope using a 40x Oil objective.

Drosophila embryo immunofluorescence
Embryos were collected and dechorionated in 50% bleach for 1min. Embryos were
transferred into 1ml fixing solution (600µl 4% PFA in PBS, 400µl n-heptane) and fixed
for 20min at RT while rotating. The lower aqueous phase was removed and 600µl
methanol added. The tube was vortexed vigorously for 1min to remove vitelline
membranes. Embryos were allowed to sink to the bottom of the tube and all liquid was
removed, followed by two washes with methanol for 1min each. Embryos were stored
at -20°C at least overnight or until further processing. In order to rehydrate embryos,
three washes each 5min with PBST (0.1% Tween20 in PBS) were performed and
embryos blocked for 1h at RT in PBST + 5% BSA. Primary antibodies were incubated
overnight at 4°C while rotating in blocking buffer followed by 3 washes for 15min each
with PBST. All following steps were performed in the dark. Secondary antibodies were
diluted in blocking buffer and incubated at RT for 2h. Embryos were rinsed 3 times
and washed 2 times for 15min. Nuclei were stained with 0.5µg/ml DAPI (Thermo Fisher Scientific) for 10min. Following two additional washing steps for 5min in PBS, embryos were mounted in ProLong Diamond Antifade Mountant (Thermo Fisher Scientific) and imaged on a Leica SP8 confocal microscope using a 40x Oil objective.

Combined RNA-FISH and IF in embryos
Embryos were collected, dechorionated and processed as described above until secondary antibody incubation. For all steps containing BSA addition, RNAsin Plus RNase inhibitors were added (1:1,000, Promega). Following secondary antibody incubation, cells were washed 3 times for 15min in PBST at RT while rotating. Embryos were fixed in 4% PFA in PBST solution for 25min and rinsed 3 times with PBST for 5min each. Embryos were pre-hybridised in 100µl hybridisation buffer (50% formamide, 5x SSC, 9mM citric acid pH 6.0, 0.1% Tween20, 50µg/ml heparin, 1x Denhardt’s solution (Sigma Aldrich), 10% dextran sulphate) for 2h at 65°C. Probes were hybridised in hybridisation buffer supplemented with 2nM of each FISH probe at 45°C overnight. Following washing twice with probe wash buffer (50% formamide, 5x SSC, 9mM citric acid pH 6.0, 0.1% Tween20, 50 µg/ml heparin) for 30min and twice for 30min at 45°C, embryos were incubated in amplification buffer (5x SSC, 0.1% Tween20, 10% dextran sulphate) for 10min at RT. Hairpins were prepared as described above and embryos incubated in fresh amplification buffer with 120nM of each probe at RT overnight in the dark. Embryos were washed twice with 5x SSC for 5min. Nuclei were stained with 0.5µg/ml DAPI diluted in 2x SSC for 15min. Following washing twice with 2x SSC for 10min, embryos were mounted in ProLong Diamond Antifade Mountant (Thermo Fisher Scientific) and imaged on a Leica SP8 confocal microscope using a 40x Oil objective.

Light Sheet Fluorescent Microscopy (LSFM) of Drosophila embryos
Embryos were collected and dechorionated as described above. 1ml of 1% low melting point (LMP) agarose was prepared and embryos transferred into capillaries (catalogue number 100003476381, Brand) using a fitting plunger. Embryos were attempted to be positioned vertically in the capillary by twisting until agarose solidified. Capillaries were stored in PBS at RT until imaging. LSFM was performed on a Zeiss Lightsheet Z.1 (Carl Zeiss, Germany) at 25°C with a 20x/1.0 Plan-Apochromat water-immersion objective lens. Embryos were lowered carefully out of the capillary into the imaging chamber filled with PBS and positioned directly between the light sheet illumination objectives (10x/0.2, left and right). Z-stack images for GFP and RFP (excitation at 488 and 561 nm, respectively) were acquired every 2min for >10h with the lowest possible laser intensity (2.5% for GFP and 10% for RFP). Generated data was analysed in Zeiss ZEN Imaging Software and Fiji (ImageJ).

ChIP-seq for Drosophila embryos
50µl of embryos were collected and dechorionated as described above and transferred in 1ml Crosslinking solution (1% formaldehyde in PBS, 50% n-heptane) and vortexed on high speed for precisely 15min. 90µl 2.5M glycine solution was added to quench excess formaldehyde and incubated for 5min at RT while rotating. Embryos were allowed to sink to the bottom of the tube and all liquid was removed. Embryos were washed three times for 4min with ice-cold buffer A (60mM KCl, 15mM NaCl, 4mM
MgCl2, 15mM HEPES pH 7.6, 0.5% DTT, 1x PI supplemented with 0.1% Triton X-100 (A-Tx buffer). All liquid was removed and embryos flash-frozen and stored at -80°C until further processing. Crosslinked embryos were transferred to a 2ml Dounce homogeniser in 1ml A-TBP (Buffer A + 0.5% Triton X-100). Following an additional washing step with A-TBP, embryos were lysed in 1ml A-TBP using 10 strokes with a tight-fitting pestle. Lysate was centrifuged at 3,200g for 10min at 4°C and supernatant removed. The pellet was resuspended in 1ml Lysis buffer (15mM HEPES, 140mM NaCl, 1mM EDTA, 0.5mM EGTA, 1% Triton, 0.5mM DTT, 10mM Sodium Butyrate, 0.1% Sodium Deoxycholate, 1x PI) and incubated at 4°C for 15min while rotating. Following centrifugation at 3,200g for 10min at 4°C, the pellet was washed twice with Lysis buffer and centrifuged again. All liquid was removed, and the pellet resuspended in 300µl LB3 (10mM Tris-HCl, pH 8, 100mM NaCl, 1mM EDTA, 0.5mM EGTA, 0.1% Na-Deoxycholate, 0.5% N-lauroylsarcosine, 1x PI). Sonication was carried out using the Bioruptor pico (Diagenode) for 6 cycles (30sec on, 30sec off settings). Debris was removed from the chromatin containing supernatant by spinning down at full speed for 10min at 4°C. Prepared chromatin was either frozen down in liquid nitrogen and stored at -80°C or used immediately. 5% of the chromatin fraction was flash-frozen as an input sample. 100µl magnetic Protein A-coupled Dynabeads (Thermo Fisher Scientific) were washed three times in 1ml blocking solution (0.2% BSA in PBS). The blocking solution was removed using a magnetic rack. 5µl of anti-H3K9me3 or anti-H3K4me2 polyclonal antibody was diluted in 250µl blocking solution and incubated with 100µl washed beads by rotating at 4°C for at least 4h up to overnight. The supernatant was removed and beads washed three times in blocking solution as described above. The chromatin solution was added to the beads and incubated at 4°C while rotating overnight. Following 4 washing steps for 2min each using ice-cold Lysis Buffer (15mM HEPES, 140mM NaCl, 1mM EDTA, 0.5mM EGTA, 1% Triton, 0.5mM DTT, 10mM Sodium Butyrate, 0.1% Sodium Deoxycholate, 1x PI, 0.05% SDS), beads were washed two additional times with ice-cold 1x TE buffer. All liquid was removed and beads resuspended in 200µl Elution buffer (50mM Tris-HCl, pH 8; 10mM EDTA; 1% SDS). Input samples were thawed and brought up to 200µl with Elution buffer. Samples were transferred into 200µl Maxymum Recovery PCR tubes (Axygen) and incubated at 65°C for 16-18h for reverse crosslinking. RNA contamination was removed by adding 200µl 1x TE buffer and 8µl of 1 mg/ml RNase A (Ambion) to ChIP and input samples followed by incubation at 37°C for 30min. Proteins were digested using 4µl Proteinase K (800U/ml, NEB) and incubation at 55°C for 2h. Reverse crosslinked DNA was recovered using the MinElute PCR purification Kit (Qiagen) according to the manufacturer’s recommendation and eluted in 15µl nuclease-free water. DNA recovery was verified and quantified using 1µl for Bioanalyzer (Agilent) electrophoresis.

ChIP-seq for Drosophila ovaries and heads

50 Drosophila ovaries were dissected in ice-cold PBS. Heads were dislodged by pouring liquid nitrogen over whole flies in a dish followed by shaking and collecting 50µl broken-off heads in 1.5ml tube. Samples were homogenised in 100µl Buffer A1 (60mM KCl, 15mM NaCl, 4mM MgCl2, 15mM HEPES pH 7.6, 0.5% DTT, 0.5% Triton X-100, 1x PI) using a rotating pestle. The volume was brought up to 1ml with buffer A1 and formaldehyde added to a final concentration of 1.8% for crosslinking. Samples were rotated for exactly 15min at RT and glycine solution added to a final concentration of 225mM. Samples were allowed to rotate for an additional 5min and were centrifuged at 4,000g for 5min at 4°C. The supernatant was removed and the pellet washed twice.
with buffer A1 and once with buffer A2 (140 mM NaCl, 15 mM HEPES pH 7.6, 1 mM EDTA, 0.5 mM EGTA, 0.5 mM DTT, 0.1% sodium deoxycholate, 10 mM sodium butyrate, 1x PI) at 4°C. The pellet was then resuspended in 100µl A2 buffer supplemented with 1% SDS and 0.5% N-lauroylsarcosine and incubated at 4°C for 2h while shaking vigorously. Lysate was sonicated using the Bioruptor pico for 16 cycles (30sec on, 30sec off). The sonicated lysate was spun at full speed for 10min at 4°C and the supernatant transferred to a new tube. The volume was brought up to 1ml with A2 buffer supplemented with 1% SDS and 0.5% N-lauroylsarcosine and incubated at 4°C while rotating and centrifuged at full speed for 10min to pellet debris. Supernatant was transferred to a new tube and protein concentration determined by Direct Detect (Millipore). 1mg of lysate per immunoprecipitation was used for the following steps. 50µl Protein A Dynabeads (Thermo Fisher Scientific) were washed with lysis buffer 3 times for 3min each. Washed beads were resuspended in 400µl lysis buffer and 5µl anti-Piwi (Hannon Lab) or rabbit IgG antibodies (Abcam, ab37415) added. Following overnight incubation at 4°C while rotating, beads were washed 3 times for 5min in 500µl lysis buffer. Antibody-coupled beads were added to lysates and volume brought up to 1ml with lysis buffer. The solution was incubated at 4°C overnight while rotating. Supernatant was removed and saved for quality control western blotting analysis. Beads were washed 6 times for 10min with 1ml wash buffer (10mM HEPES ph 7.3, 150mM NaCl, MgCl2, 10% glycerol, 1% Triton x-100, 1mM EDTA, 0.1mM PMSF, 1x PI, 1:1,000 RNasin (Promega)) using a 2ml Dounce homogeniser. Material was lysed with 5 strokes with a loose pestle and 5 strokes with a tight pestle on ice. Lysate was incubated for 1h at 4°C while rotating and centrifuged at full speed for 10min to pellet debris. Supernatant was transferred to a new tube and protein concentration determined by Direct Detect (Millipore). 1mg of lysate per immunoprecipitation was used for the following steps. 50µl Protein A Dynabeads (Thermo Fisher Scientific) were washed with lysis buffer 3 times for 3min each. Washed beads were resuspended in 400µl lysis buffer and 5µl anti-Piwi (Hannon Lab) or rabbit IgG antibodies (Abcam, ab37415) added. Following overnight incubation at 4°C while rotating, beads were washed 3 times for 5min in 500µl lysis buffer. Antibody-coupled beads were added to lysates and volume brought up to 1ml with lysis buffer. The solution was incubated at 4°C overnight while rotating. Supernatant was removed and saved for quality control western blotting analysis. Beads were washed 6 times for 10min with 1ml wash buffer (10mM HEPES ph 7.3, 150mM NaCl, MgCl2, 10% glycerol, 1% Empigen BB Detergent (Merck), 1x PI). For the first wash, 1µl RNasin was added to the wash buffer and tubes were changed between each wash. 10% of beads were set aside for quality control and 90% resuspended in 1ml Trizol (Thermo Fisher Scientific) and stored at -80°C until further processing.

Whole genome sequencing (WGS)

100 flies were collected in a 1.5ml tube and frozen at -80°C for at least 1h. High molecular weight genomic DNA was isolated using the Blood and Cell Culture DNA Mini kit (Qiagen). Flies were homogenised using a rotating pestle on ice for 1min. 700µl G2 and 50µl Proteinase K (800U/ml, NEB) was added to each tube and incubated at 50°C for 2h with occasional tube inversions. Tubes were spun at 5,000g for 10min at 4°C and supernatant transferred to new tube avoiding debris. A Qiagen Genomic-tip 20/G was equilibrated with 1ml QBT buffer and allowed to empty by gravity flow. The supernatant containing digested proteins and genomic DNA was added to the column and allowed to flow through. The column was washed 3 times
with 1ml QC buffer. Elution was carried out with 1ml QF elution buffer and repeated once. Flow through was transferred to two new tubes (1ml each) and 700µl isopropanol added. Tubes were inverted 10 times and centrifuged at full speed for 15min at 4°C. The pellet was washed with 70% ethanol twice and air-dried for 5min.

25µl RNase-free water was added and DNA resuspended by flicking tube gently several times while incubating at 37°C for 2h. DNA was stored at 4°C. DNA was sheered using a Covaris S220 (Covaris). 3µg of genomic DNA was diluted in RNase-Free water and transferred to an AFA Fiber Crimp-Cap (PN520052, Covaris) microtube. Sonication was carried out with the following settings: Peak incident Power (W) 105, Duty Factor 5%, Cycles per Burst 200, Treatment time 80sec. This resulted in sheared DNA fragments peaking at 500bp. DNA was recovered using the QIAquick PCR Purification Kit (Qiagen).

RNA extraction
RNA for RNA-seq and qRT-PCR experiments was isolated using the RNeasy Mini kit (Qiagen) with on-column DNA digestion (RNase-free DNase Set, Qiagen) according to the manufacturer’s recommendations. RNA for small RNA-seq experiments were isolated using Trizol following the manufacturer’s instructions.

Library preparation
1µg of total RNA was used as input material for RNA-seq library preparation. The NEBNext Poly(A) mRNA magnetic Isolation Module (NEB) was used to isolate poly(A) RNAs. Libraries were generated with the NEBNext Ultra Directional RNA Library Prep kit for Illumina (NEB) according to manufacturer’s instructions. Small RNA libraries were generated as described previously (Jayaprakash et al., 2011). In short, 19- to 28-nt sized small RNAs were purified by PAGE from Piwi-bound RNA isolated from ovaries or embryos. Next, the 3’ adapter (containing four random nucleotides at the 5’ end) was ligated overnight using T4 RNA ligase 2, truncated KQ (NEB). Following recovery of the products by PAGE purification, the 5’ adapter (containing four random nucleotides at the 3’ end) was ligated to the small RNAs using T4 RNA ligase (Abcam) for 1h. Small RNAs containing both adapters were recovered by PAGE purification, reverse transcribed and PCR amplified. WGS libraries were generated using the NEBNext Ultra II DNA Library Prep kit (NEB) according to manufacturer’s recommendation with 1µg input material. 3 PCR amplification cycles were performed. Libraries were quantified using the Library Quantification Kit for Illumina (Kapa Biosystems).

Next-generation sequencing
Sequencing was performed by the Genomics Core facility at CRUK CI. RNA-seq, ChIP-seq and small RNA-seq libraries were sequenced on an Illumina HiSeq 4000 according to manufacturer’s recommendations using single-end 50bp runs. WGS libraries were sequenced with paired-end 150bp runs on Illumina HiSeq 4000 or NovaSeq.

Quantitative reverse transcription polymerase chain reaction (qRT-PCR)
Reverse transcription (RT) was performed using the SuperScript IV reverse transcriptase Kit (Thermo Fisher Scientific) with 1µg of total RNA. qRT-PCR was performed on a QuantStudio Real-Time PCR Light Cycler (Thermo Fisher Scientific) in technical triplicates. Expression of targets was quantified using the ddCT method.
(Livak and Schmittgen, 2001). Fold-change was calculated as indicated in the figure legends and normalised to rp49. All primers are listed in Supplementary file 1.

**Protein isolation from whole embryos and quantitative mass spectrometry**

100µl of control w^{1118} embryos for time points 0-2h, 5-7h and 10-12h AEL were collected in three biological replicates on agar plates and dechorionated. Embryos were then lysed in lysis buffer (0.1% SDS, 0.1M triethylammonium bicarbonate (TEAB), 1x Halt Protease and Phosphatase Inhibitor (Thermo Fisher Scientific) using a rotating pestle on ice for 2min or until entirely homogenised. Lysate was heated for 5min at 90°C and probe sonicated for 20sec (20% power with pulse of 1sec). Debris was pelleted by centrifugation at full speed for 10min at 4°C and supernatant transferred to a new tube. Protein concentration was measured using Bradford Assay (Bio Rad). 100µg protein was digested with trypsin overnight at 37°C. TMT chemical isobaric labelling was performed as described previously (Papachristou et al., 2018). Peptide fractions were analysed on a Dionex Ultimate 3000 UHPLC system coupled with the nano-ESI Fusion Lumos mass spectrometer (Thermo Fisher Scientific).

**Treatment of embryos for auxin-induced degradation**

Embryos were collected for 30min and dechorionated. Control embryos were transferred into a fine mesh strainer placed in a plastic dish and submerged in PBS. 1M auxin solution was generated by diluting Indole-3-acetic acid (IAA), a highly permeable small molecule as recently shown for C. elegans embryos (Zhang et al., 2015), in water and stored protected from light at -20°C. Auxin-treated embryos were submerged in PBS with indicated auxin concentrations. Embryos were placed at 25°C for appropriate times and harvested for RNA experiments by transferring into 1ml Trizol followed by RNA extraction. Embryos used for ChIP-seq were processed as described above.

**RNA-seq and ChIP-seq analysis**

Raw fastq files contained 50bp reads. The first and the last two bases of all reads were trimmed using fastx_trimmer ([http://hannonlab.cshl.edu/fastx_toolkit/](http://hannonlab.cshl.edu/fastx_toolkit/)). Reads were first aligned to the consensus sequence for all Drosophila melanogaster transposons using STAR (Dobin et al., 2013) allowing random allocation of multimappers. Unmapped reads were further aligned to Drosophila melanogaster genome release 6 (dm6) keeping uniquely mapping reads. Generated bam files for RNA-seq were further split in reads originating from sense and antisense genomic strands using samtools view options -f 0x10 and -F 0x10 for sense and antisense reads respectively (Li et al., 2009b). Indexes were generated using samtools index function. Coverage files were generated using bamCoverage with normalisation mode --normalizeUsing CPM (Ramirez et al., 2014) and applying a scaling factor (--scaleFactor). Scaling factors for individual files were calculated by dividing the sum of mapped reads contained in the file by the sum of all transposon and dm6 mapping reads of the corresponding library. Reads mapping to protein-coding genes were counted with htseq (Anders et al., 2015) using a feature file downloaded from Ensembl release BDGP6.22. Reads mapping to individual transposons were counted with a custom script using samtools idxtstats function to extract reads mapping to individual sequences of the reference genome/transposon consensus sequence. Count files generated as described above were normalised to reads per million (rpm) and plotted in R using ggplot2. Bar graphs displaying transposon expression profiles and heatmap during embryogenesis were calculated from pooled replicates. Bar graphs and heatmaps were plotted in R using...
ggplot2. Differential expression analysis was performed using DESeq2 (Love et al., 2014). MA plots show base mean RNA expression across conditions and were calculated as previously described by Love and colleagues. ChIP-seq reads were normalised by library size and rpm calculated for concatenated replicates using the deepTools2 bamCoverage function (Ramirez et al., 2016) with bin size 10bp. MA plots displaying H3K9me3 signal intensity fold-changes between auxin-treated and control samples were calculated using DESeq2 for individual replicates (n=3). Metaplots flanking euchromatic transposon insertion sites were calculated using computeMatrix scale-region function from deepTools2 with bin size 10bp. All scripts used for sequencing analysis are available on GitHub (https://github.com/mhf27/hannon_roo_fabry2021).

Small RNA-seq analysis

Reads from small RNA-seq libraries were adapter clipped using fastx clipper with settings -Q33 -I 15 -a AGATCGGAAGAGCACACGTCT. The first and last 4 bases of adapter clipped reads were trimmed using seqtk trimfq (https://github.com/lh3/seqtk). Only high-quality reads with length between 19-31bp were used for further analysis. Small RNAs were aligned as described above and size profiles plotted in R.

Generation of annotation files

Mappability track for dm6 with 50bp resolution was calculated according to a previously published method (Derrien et al., 2012). The de novo transposon insertion calling for the homozygous control w1118 strain and our line carrying both GFP-AID-Piwi and OsTIR1 was performed using the TEMP algorithm (Zhuang et al., 2014). In brief, ~500bp genomic DNA fragments were amplified and sequenced generating 150bp paired-end reads, which were aligned to dm6 using BWA (Li and Durbin, 2009). Reads with only one mate aligned to dm6 were extracted and the unmapped mate uniquely aligned to transposon consensus sequences in order to ensure correct directionality calling. Calculated insertion sites were extracted from generated GTF files if they were supported by reads on both sides (1p1). Transposon insertion files containing coordinates as well as statistical information have been submitted to GEO (GSE160778). Euchromatic regions (chr2R:6460000-25286936, chr2L:1-22160000, chr3L:1-23030000, chr3R:42000000-32079331, chrX:2500000-21500000) were defined by measuring H3K9me3 signal genome-wide in sliding windows of 10kb bins and calculating signal enrichment over input. We identified a total of 632 euchromatic TE insertions in w1118 and 1,738 in our degron strain (GFP-AID-Piwi; OsTIR1). The protein database used to identify peptides from Drosophila genes and transposons was generated by merging an existing database downloaded from FlyBase (dmel-r6.24.fa) with translated ORFs of transposons. ORFs were predicted and translated using prodigal (https://github.com/hyattpd/Prodigal). ORFs with less than 300 amino acids were removed using seqtk -L 300 and the file was converted to fasta format.

Protein domain prediction

Functional analysis of protein sequences was performed using the InterPro web application (https://www.ebi.ac.uk/interpro/). Protein domains and families for open reading frame encoded by roo transcripts were predicted using default settings.

Mass spectrometry raw data processing

Raw data files were processed according to previous reports (Papachristou et al., 2018). Spectral .raw files were analysed with the SequestHT search engine on Thermo
Scientific Proteome Discoverer 2.1 for peptide and protein identification. Data was searched against a modified FlyBase protein database with following parameters:

- Precursor Mass Tolerance 20 ppm, Fragment Mass Tolerance 0.5 Da. Dynamic Modifications were oxidation of methionine residues (+15.995 Da), deamidation of asparagine and glutamine (+0.984 Da) and Static Modifications were TMT6plex at any amino-terminus, lysine (+229.163 Da) and methylthio at cysteine (+45.988 Da). The Reporter Ion Quantifier node included a TMT 6plex (Thermo Scientific Instruments) Quantification Method, for MS3 scan events, HCD activation type, integration window tolerance 20 ppm and integration method Most Confident Centroid. Peptides with an FDR >1% were removed. The downstream workflow included signal to noise (S/N) calculation of TMT intensities. Level of confidence for peptide identifications was estimated using the Percolator node with decoy database search. Strict FDR was set at q-value < 0.01.

Bioinformatics analysis of proteomics data

Processed data files were analysed as described in a previous publication (Papachristou et al., 2018) using qPLEXanalyzer in R with multimapping peptides included in the analysis. Bar graphs showing protein intensities for Piwi and volcano plots with indicated comparisons were plotted in R using ggplot2.

Quantification and statistical analysis

Statistical tests used for individual experiments are indicated in figure legends. Statistical analyses applied to hatching rates, qPCR data sets and ChIP-seq signal intensity were calculated by unpaired (two sample) t Test. Significance of TMT mass spectrometry data was calculated according to (Papachristou et al., 2018). Differential expression of RNA-seq experiments and differential enrichment of ChIP-seq experiments were calculated using DeSeq2 using adjusted p values as described in (Love et al., 2014). The number of biological replicates is indicated in the figure legends.

Data availability

Raw data from high-throughput sequencing experiments are available at GEO under accession number GSE160778. Raw data from proteomics experiments are available on PRIDE with accession number PXD0022409.
Competing interests

The authors declare no competing interests.

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Supplementary files

Supplementary file 1: List of oligonucleotides for qRT-PCR and RNA-FISH.

Figure supplement 1-source data 1: Gene- and TE-mapping reads per million across embryogenesis RNA-seq time points for plots shown in Figure 1B-C, Figure 1-figure supplement 1A,F, Figure 2A and Figure 2-figure supplement 1E,G.

Figure supplement 1-source data 2: Signal intensity of genes and transposon ORFs in TMT-MS time-course experiment for plots shown in Figure 1-figure supplement 1A,C,D,F, Figure 2B and Figure 2-figure supplement 2F,H.

Movie S1: GFP-Piwi live imaging. Light-sheet fluorescent live microscopy of embryos derived from parents expressing GFP-Piwi (green) and H2Av-RFP (red) in developing embryos for indicated time after egg laying (AEL).

Movie S2: GFP-Panx live imaging. Light-sheet fluorescent live microscopy of embryos derived from parents expressing GFP-Panx (green) and H2Av-RFP (red) in developing embryos for indicated time after egg laying (AEL).

Movie S3: GFP-Nxf2 live imaging. Light-sheet fluorescent live microscopy of embryos derived from parents expressing GFP-Nxf2 (green) and H2Av-RFP (red) in developing embryos for indicated time after egg laying (AEL).

Movie S4: GFP-AID-Piwi live imaging. Light-sheet fluorescent live microscopy of embryos derived from parents expressing GFP-AID-Piwi (green) and OsTIR1. Time course shows early embryo treated with 5mM auxin for indicated time points.

Transparent reporting form
References


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