In *Drosophila melanogaster* the COM Locus Directs the Somatic Silencing of Two Retrotransposons through both Piwi-Dependent and -Independent Pathways

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**Background.** In the Drosophila germ line, repeat-associated small interfering RNAs (rasiRNAs) ensure genomic stability by silencing endogenous transposable elements. This RNA silencing involves small RNAs of 26-30 nucleotides that are mainly produced from the antisense strand and function through the Piwi protein. Piwi belongs to the subclass of the Argonaute family of RNA interference effector proteins, which are expressed in the germline and in surrounding somatic tissues of the reproductive apparatus. In addition to this germ-line expression, Piwi has also been implicated in diverse functions in somatic cells. **Principal Findings.** Here, we show that two LTR retrotransposons from *Drosophila melanogaster*, *ZAM* and *Idefix*, are silenced by an RNA silencing pathway that has characteristics of the rasiRNA pathway and that specifically recognizes and destroys the sense-strand RNAs of the retrotransposons. This silencing depends on Piwi in the follicle cells surrounding the oocyte. Interestingly, this silencing is active in all the somatic tissues examined from embryos to adult flies. In these somatic cells, while the silencing still involves the strict recognition of sense-strand transcripts, it displays the marked difference of being independent of the Piwi protein. Finally, we present evidence that in all the tissues examined, the repression is controlled by the heterochromatin COM locus. **Conclusion.** Our data shed further light on the silencing mechanism that acts to target *Drosophila* LTR retrotransposons in somatic cells throughout fly development. They demonstrate that different RNA silencing pathways are involved in ovarian versus other somatic tissues, since Piwi is necessary for silencing in the former tissues but is dispensable in the latter. They further demonstrate that these pathways are controlled by the heterochromatin COM locus which ensures the overall protection of *Drosophila* against the detrimental effects of random retrotransposon mobilization.

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**INTRODUCTION**

Genome sequencing projects have revealed that eukaryotic genomes contain large numbers of repetitive sequences and mobile genetic elements. Retrotransposons, which are mobile genetic elements that move through an RNA intermediate in a process termed retrotransposition, are highly abundant and comprise nearly half of the human and a third of the *Drosophila* genome. It is thus essential that eukaryotic cells retain tight control over these potential invaders in order to protect their genomes from the mutational threat that they pose. This control is especially critical in the germline, where retroelement activity can create a mutational burden that is then transmitted to subsequent generations. As numerous metazoan elements display expression that is restricted to the reproductive apparatus, including the germline and in surrounding somatic tissues, genomes have evolved specific mechanisms to protect these tissues by further restricting the expression of these elements.

It has become clear in the last few years that RNA interference (RNAi) plays a major role in ensuring this type of protection [1–3]. Three main RNA silencing pathways act, at the post-transcriptional level and involving three distinct populations of small RNAs—siRNAs, miRNAs and piRNAs—have been reported to date. siRNAs (small interfering RNAs) are derived from processed double-stranded RNAs (dsRNAs) into siRNAs of 20–24 nucleotides (nt) in length. These siRNAs are loaded onto an RNA-induced silencing complex (RISC) as single-stranded siRNA molecules which then bind and cleave the target RNA [4–6]. miRNAs (microRNAs) of 22 nts in length are endonucleaseolytically processed from endogenous non-coding transcripts. After their production, they bind the miRISC to mediate RNA silencing. miRNAs are developmentally regulated and play an important role in gene silencing throughout development [7,8]. Whereas siRNAs and miRNAs are derived from both the sense and antisense strands of their double-stranded precursors, piRNAs (for PIWI-interacting RNAs) are mainly derived from antisense strands and are produced from discrete genomic loci [9,10]. piRNAs are from 26–30 nts in length and have been reported in germine cells of *drosophila*, mice, rats and humans [11,12]. In mice and human, they are required for male fertility. In *Drosophila*, a subset of...
piRNAs, called rasiRNA for repeat associated small interfering RNAs, has been directly implicated in the protection of the fly germ line against selfish genetic elements such as retrotransposons and repetitive sequences [13–15].

\(\text{ZAM}\) and \(\text{Idefix}\) are two LTR retrotransposons that are generally silent in the genome of \(\text{Drosophila melanogaster}\) [16]. In the vast majority of lines, these elements do not start their replication cycle and no mobilisation is observed. These lines are denoted S (stable) lines. However, in certain lines, called unstable (U) lines, this control has been perturbed and both elements are highly expressed in the ovaries. As a result of this expression, multiple copies of \(\text{ZAM}\) or \(\text{Idefix}\) become integrated into the germ line and are transmitted through successive generations [17,18]. Characterisation of the U line has shown that an active transposon-silencing process that targets \(\text{ZAM}\) and \(\text{Idefix}\) has been mutated in this line. The mutated locus has been identified and shown to be a heterochromatic locus called COM, which is located at position 20A2-3 on the X-chromosome [16]. Analysis of the replication cycles of \(\text{ZAM}\) and \(\text{Idefix}\) in U and S lines has indicated that \(\text{ZAM}\) transcripts are only present in the posterior follicular cells of the ovaries of U lines, and not in the ovaries of S lines. Similarly, \(\text{Idefix}\) transcripts are detected in the germarium of U line ovaries, but not in S lines. Overall, our previous work indicated that both elements are subjected to two types of control in \(\text{Drosophila}\): first, a regulatory pathway silences the elements and prevents them from initiating their replication cycles. Second, if this silencing is lost, another control mechanism that relies on specific cis-regulatory sequences present in the elements themselves restricts their expression to specific somatic cells of the ovaries [19].

In this study, we have carried out an in-depth analysis of the intrinsic regulatory properties of \(\text{ZAM}\) and \(\text{Idefix}\), examining the pathways that promote their silencing in the S lines. We show that the control of \(\text{ZAM}\) and \(\text{Idefix}\) is mediated by a homology-dependent trans-silencing pathway that displays characteristics of the rasiRNA pathway. At the same time, it displays three unique features: i) in addition to the reproductive apparatus, the silencing is exerted in most if not all of the somatic tissues of flies; ii) Piwi is required in cells of the reproductive apparatus, but it does not play a role in other tissues; iii) The function of COM is ubiquitous.

**RESULTS**

The U5 region of the \(\text{ZAM}\) LTR is sufficient for the differential regulation between S and U lines

We have previously reported that the transgene denoted pZ499, which contains the full-length LTR of \(\text{ZAM}\) and the first 26 bp of its 3′UTR (499 bp) fused to a \(\text{lac}\)-\(\text{Z}\) reporter gene, responds to the two types of control over \(\text{ZAM}\) expression: i) repression, which depends on the fly genotype (U or S); and ii) tissue-specific activation, which drives expression in a very specific group of cells located at the posterior pole of the follicle [19]. The LTR is composed of a U3 region spanning nucleotides 1 to 325, a central R region from nucleotides 326 to 347, and a U5 region from nucleotides 348 to 473. The transcription initiation site defines the boundary between the U3 and R regions, and the polyadenylation site corresponds to the boundary between the R and U5 regions. To investigate the specificity of \(\text{ZAM}\) transcription in the different lines and to localise the sequences involved in its regulation, we analysed the expression of two additional transgenes placed in an S or U genetic background. These transgenes, pZ310 and pZ475, contain \(\text{ZAM}\) fragments extending from nucleotides 1 to 310 or 1 to 475, respectively, fused to the \(\text{lac}\)-\(\text{Z}\) reporter gene. We found that pZ475 responds to both the strain- and tissue-specific controls that have been previously described for the full-length LTR [19]: its expression is restricted to the follicle cells of the U line and is absent in the S line (Fig. 1). By contrast, pZ310 which is expressed from a minimal heat shock promoter responds to the tissue-specific control that restricts its expression to the posterior follicle cells, but is insensitive to the line-specific control, since it is expressed in both the U and S genetic backgrounds (Fig. 1).

Therefore, we concluded based on this analysis of the pZ310, pZ475, and pZ499 lines that the line-specific expression of \(\text{ZAM}\) is
controlled by sequences present in the R and/or U5 regions of its LTR.

**ZAM and Idefix are regulated by a homology-dependent gene-silencing mechanism**

Transgenes pZ475 and pZ499 both initiate transcription from the \(ZAM\) promoter at nucleotide 326. When these transgenes are transcribed, 149 and 173 nucleotides, respectively, of \(ZAM\) are present at the 5’ end of the transcripts (Fig. 1, table). By contrast, the \(ZAM\) promoter is absent in pZ310, and no \(ZAM\) sequence is transcribed. To determine whether the \(ZAM\) promoter or the presence of a \(ZAM\) fragment within a chimeric transcript is responsible for the differential expression of \(ZAM\) in the S and U lines, we designed constructs in which a GFP reporter gene, driven by the UASt promoter, is fused downstream of its coding sequence and upstream of the polyadenylation site to diverse fragments of \(ZAM\) (Fig. 2 A). Transgenic lines were established and tested for GFP expression in the ovaries. A 720 bp fragment from within the third ORF of \(ZAM\) was tested. This fragment corresponds to the region encoding the Env protein, spanning nucleotides 6385 to 7105 of the \(ZAM\) sequence. This \(ZAM\) fragment was inserted in an orientation such that transcription of the transgene would give rise to mRNA corresponding to the sense-strand fragment of \(ZAM\). Furthermore, the \(5\prime\) UTR fragment was flanked by FRT elements, which are targets for \(flp\) recombinase.

Transgenic lines were established and denoted pGFP-Zenv. The expression of the UASt transgenes was induced by crossing with flies containing a ubiquitous somatic actin-Gal4 driver in the S genetic background. Data obtained are presented in Fig. 2 A, B and C.

We examined the expression of pGFP-Zenv in the ovaries of three independent transgenic lines. The genotype of these lines was [S/S; actin-Gal4/CyO; pGFP-Zenv/pGFP-Zenv]. No fluorescence was detected in either of the lines (Fig. 2A). A number of these flies were subjected to \(flp\) recombinase action; to do this, they were crossed with flies expressing the \(flp\) recombinase under the control of a heat-shock promoter. Embryos laid by these females were then exposed to two heat-shocks at 38°C for 1 hr [20]. In these conditions, the efficiency of recombination between the two FRTs and the consequent deletion of the intervening \(env\) fragment is near 100%. The resulting flies were denoted pGFP-Zenv (Fig. 2B). Analysis of the expression of the pGFP-Zenv and pGFP-ZUtransgenes, activated by the actin-Gal4 driver in the ovaries, showed that whereas GFP is never expressed in the follicle cells when the pGFP-Zenv transgene is present in an S genetic background (Fig. 2A), fluorescence was never present in the same S background when the \(ZAM\) fragment was flipped out (pGFP-ZUtransgene Fig. 2B).

In a second series of tests, we compared GFP expression from the pGFP-Zenv construct in the S and U genetic backgrounds. We found that GFP is only expressed in U line flies, as shown in Fig 2C for a follicle with genotype [U/U; actin-Gal4/CyO; pGFP-Zenv/pGFP-Zenv]. It should be noted that because of the non-functionality of the UASt promoter in germinal cells, the absence of a signal in the nurse cells or the oocyte is not informative in these experiments.

When GFP expression was compared between the different transgenic lines using the same settings on the confocal microscope, we found that the GFP signals were always lower in the transgenic U lines bearing the pGFP-ZUtransgene than they were with the flipped-out transgenes. This suggests that full expression of GFP might not be recovered in U lines, which may only partially release \(ZAM\) silencing. Alternatively, the pGFP-Zenv transcripts might be less stable than the shorter pGFP-ZUtransgene transcripts. Although this latter hypothesis cannot be excluded, results obtained when analyzing expression of these transgenes through northern blot experiments favour the former one (see below).

In additional assays, 1 kb of the 5’ UTR of \(ZAM\) was placed downstream of the GFP gene, in the 5’ to 3’ orientation with respect to \(ZAM\) transcription. Similar tests as those performed with the pGFP-Zenv transgenic lines were performed. The regulation of the GFP gene in the so-called pGFP-ZU transgenic lines gave the same results as for pGFP-Zenv (data not shown).

We conclude that the silencing of the pGFP-Zenv and pGFP-ZU transgenes is under the control of a transposon silencing pathway that targets endogenous \(ZAM\) retrotransposons and that is absent in the U line.

As Idefix is also repressed in S lines and is active in U lines, we used P-element transformation to introduce additional constructs containing UAS repeats located upstream of the GFP reporter gene and different fragments of Idefix into the genome of S line flies. We tested two different fragments. One corresponded to a non-coding region from the 5’ UTR of the gene, and the second to a coding region taken from its gag gene. Similarly to the pGFPZU and pGFPZenv transgenes, the Idefix fragments were inserted in an orientation such that transcription of the transgenes would give rise to mRNA corresponding to the sense-strand fragment of Idefix.

These constructs were respectively denoted pGFP-IdU1 and pGFP-Idgag. Expression of the GFP reporter gene under the control of the actin-Gal4 driver was assayed in the ovaries of flies that were homozygous for both the X chromosome of the stable S line and the pGFP-Id transgene. Results obtained when the gag gene from nucleotides 1003 to 1422 was placed downstream of GFP are presented in Figure 2 D, E, and F. Similar results were obtained with the pGFP-IdU1 lines (data not shown). No fluorescence was detected in the ovaries of this S line (Fig. 2D). This absence of fluorescence depends upon the S status of the line, because fluorescence was clearly observed when the stable X chromosome was replaced by an X chromosome from a U line (Fig. 2F). A \(flp\) recombinase assay was conducted on the transgenic S/S line, as depicted for pGFP-Zenv. When the gag fragment of Idefix was flipped out with the \(flp\) recombinase, giving rise to flies denoted pGFP-Idgag, fluorescence was clearly recovered in the ovaries of S line flies (Fig 2E). These assays were performed on four independent transgenic lines for each construct, giving the same results.

Thus, it appears that both \(ZAM\) and Idefix are controlled by a common silencing mechanism that has several definable properties. First, it is active in the follicle cells of S lines. Second, it does not function through a specific sequence present within both elements, but can rather target regions all along their lengths, from their 5’ to the 3’ ends. Finally, this silencing mechanism is disrupted in the U lines.

**Sense-strand transcripts of \(ZAM\) and Idefix are specifically targeted by the silencing machinery**

A next set of experiments was performed with similar sensor GFP transgenes, but in which the fragments of \(ZAM\) and Idefix were inserted in the opposite orientation. Specifically, the 720 bp fragment within the third ORF of \(ZAM\) and the 456 bp fragment corresponding to the 5’ UTR of Idefix were tested in these experiments. These transgenes were denoted pGFP-ZenvAS and pGFP-IdUAS, respectively (Fig. 3). When transcribed, the resulting transgenes gave rise to transcripts which were antisense with respect to the endogenous \(ZAM\) or Idefix genomic RNAs. The ability of GFP to be expressed in the different lines was then assayed by introducing the actin-Gal4 transcription driver by
crossing. While no fluorescence was observed when the expression of the (sense-strand) pGFP-Zenv or pGFP-IdU transgenes was assayed, strong GFP fluorescence was detected in ovarian somatic tissues of the three independent (antisense) transgenic lines established with either pGFP-ZenvAS or pGFP-IdUAS (Fig 3 A and B). The intensity of the fluorescence was very similar to that observed with pGFP transgenes containing no ZAM or Idefix sequences, indicating that no silencing was exerted on these sensor transgenes.

Therefore, these transgenes are not sensitive to the repression exerted by the S genetic background. Further, this result suggests that the silencing mechanism that targets ZAM and Idefix is only directed against mRNAs containing sequences homologous to their sense-strand transcripts.

Figure 2. Transgenes with a GFP reporter gene fused to a ZAM sequence act as sensors of the repression. The genomic structures of the transgenes pGFP-Zenv and pGFP-Idgag used in this study are presented at the tops of both panels: The grey boxes correspond to the UAS promoter, the dotted boxes to the GFP gene, and the white box to the env fragment of ZAM or the gag fragment of Idefix. Triangles indicate the FRT sites. Focal plane of the follicles dissected from a line in which the pGFP-Zenv transgene is driven by the ubiquitous Actin-Gal4 driver. Expression of the pGFP-Zenv transgen in an S genetic background before (A) or after (B) flip-recombinase action, or in a U genetic background before the flip treatment (C). GFP expression in the ovarioles of a transgenic line bearing the pGFP-Idgag transgene driven by the ubiquitous Actin-Gal4 driver. Expression of the pGFP-Idgag transgene in an S genetic background before (D) or after (E) flip-recombinase action, or in a U genetic background before the flip treatment (F). No GFP is detected in ovaries of the S lines. Its expression is recovered after the flip treatment or when the COM locus is mutated, as in the U genetic background.

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Silencing of ZAM and Idefix occurs at the RNA level and involves small RNAs with characteristics of rasiRNAs

The trans-silencing phenomenon described above was analysed through GFP expression of the transgenes. To determine whether this silencing acted at the translational level or at the RNA level, we looked for GFP RNA encoded by the transgenic lines. Total RNA was extracted from ovaries of transgenic lines in either the S or U genetic backgrounds, and northern blots were performed. The nylon filters were probed first with a riboprobe corresponding to sense-strand and anti-sense strand RNAs of ZAM or Idefix. A typical experiment is presented in Figure 4. GFP transcripts revealed by a riboprobe with this more sensitive assay, we could only barely detect small rasiRNAs [13,15,21]. Therefore, we searched for putative small RNAs that are homologous to ZAM and Idefix. These RNAs are detected in S lines and, at a much lower level, in the U line. Small RNAs homologous to the antisense strand of the 5'UTR of ZAM are presented in C. 20 to 30 nt long antisense strand RNAs (−) homologous to the 5'UTR or the gag gene of Idefix are detected. Sense strands (+) are absent or present in very small amounts. A typical experiment is presented in D. Signs (+) and (−) indicate respectively sense-strand and anti-sense strand RNAs of ZAM or Idefix revealed by the riboprobes. These results indicate that the S line control does not act at the translational level but rather at the RNA level. As observed in the analysis of the sensor transgenes, the silencing is strictly directed against transcripts with sense-strand RNAs of ZAM or Idefix, and not against their antisense strands. Further, the silencing is released, although only partially, in the U genetic background.

The characteristics of this silencing were reminiscent of silencing involving rasiRNAs [13,15,21]. Therefore, we searched for putative small RNAs that are homologous to ZAM and Idefix. As northern blot experiments failed to identify any such small RNAs, we employed the more sensitive RNase protection assay. Even with this more sensitive assay, we could only barely detect small RNAs, indicating that they are very low in number. However, in
view of the consistency of the results—obtained in at least three independent experiments performed with each sense and antisense probe from the 5′UTR, gag, pol, and env genes from both elements—certain general observations concerning the small RNA populations detected can be made as illustrated fig 4 C and D. First, short RNAs of 20 to 30 nt and homologous to ZAM and Idefix were detected, but no clear populations of any specific length can be defined at this stage of our detection assays (Fig. 4C and D). Second, these small RNAs were found in S lines as well as, although at lower levels, in U lines (Fig. 4C). Third, the abundance of short antisense RNAs was always higher than the almost undetectable short sense RNAs, suggesting that most of the detected small RNAs are antisense to ZAM and Idefix and single-stranded (Fig. 4D). Finally, small antisense RNAs were detected regardless of the probe used. Typical results obtained with the 5′UTR of ZAM (Fig. 4C) and with the 5′UTR or the gag gene of Idefix (Fig 4D) are presented.

These results show that the silencing machinery acting against ZAM and Idefix is associated with the presence of a small population of 20–30 nt RNAs, most of them being complementary to ZAM and Idefix mRNA.

The silencing mechanism targeting ZAM and Idefix involves the PIWI Argonaute protein in the reproductive apparatus

The Piwi protein has been shown to be involved in the rasiRNA pathway to maintain transposon silencing in the germline [1]. To test whether Piwi is also necessary for the repression of ZAM and Idefix, we first investigated the effect of the piwi3 mutation on the expression of endogenous ZAM and Idefix elements in the ovaries of S-line flies (Fig. 5). Because the morphology of homozygous piwi3 ovaries is severely affected in adult flies, it was impossible to investigate ZAM and Idefix expression in adult ovaries. We thus performed experiments in the gonads of third instar larvae. By in situ RNA analysis using strand-specific riboprobes for ZAM and Idefix, we found that both of the elements are expressed in female gonads of third instar larvae from the U line, as shown in a homozygous [U/U; piwi+/+] genetic background (Figure 5, middle). No staining corresponding to ZAM or Idefix RNA was ever detected in the gonads of larvae having the corresponding genotypes in the S line [S/S; piwi+/+] (not shown) or [S/S; piwi+/+] (Figure 5, left). By contrast, clear expression of ZAM and Idefix was observed in the homozygous genetic background [S/S; piwi+/+], displaying a pattern of expression similar to that detected in the U line (Fig. 5, right).

These findings provide evidence that Piwi is a component of the pathway that silences ZAM and Idefix in the ovarian somatic tissue.

The silencing mechanism controlling ZAM and Idefix is active in somatic tissues throughout fly development

We next investigated whether the silencing mechanism involved in the repression of ZAM and Idefix is strictly restricted to follicular cells, where proper ZAM and Idefix enhancers are active, or if it is more widely present and active in other tissues. To address this question, the expression of the pGFP-ZAM and pGFP-Idefix transgenes was examined throughout fly development, in embryos, larvae, and adult flies. Two Gal4 drivers were used in these experiments: the ubiquitous actin-Gal4 driver, as described above, and the 24B-Gal4 driver, which is specifically expressed in mesodermal cells [22].

In the S/S genetic background, no fluorescence was detected with any of the transgenes (pGFP-ZU, pGFP-Zenv, pGFP-IdU or pGFP-IdGag), regardless of the driver used (actin-Gal4 or 24B-Gal4). It should be noted that, if the microscope settings are optimized, a very faint level of fluorescence can be detected at each stage of development. This transgene silencing was observed in all the cells examined and throughout fly development, including in embryos, larvae, and adult flies. As an example, results obtained with pGFP-IdU driven by 24B-Gal4 are presented Fig. 6, column A. In contrast, when the X-chromosome in S-transgenic lines was replaced by one from a U line, clear fluorescence resulting from the expression of the GFP reporter gene driven by 24B-Gal4 was detected in embryos, larvae, and adult flies (Fig. 6 column B).

To determine whether the silencing mechanism that is active in these somatic tissues specifically targets sense-strand RNAs of ZAM and Idefix, similar experiments were conducted using the pGFP-ZenvAS and pGFP-IdUAS transgenes described in Fig. 3. We found that no silencing occurred on these transgenes, with both giving rise to clear ubiquitous GFP expression in all the examined stages. Results obtained with pGFP-IdUAS are shown in Fig. 6, line C.

It thus appears that the repression machinery targeting ZAM and Idefix is not restricted to ovarian follicle cells of S lines but is instead active in a broad range of cells (if not all) throughout fly development. This machinery is able to discriminate between sense- and anti-sense strand transcripts of ZAM and Idefix, and is under the control of the COM locus.

The silencing mechanism active in somatic cells does not involve the PIWI Argonaute protein

Since the control exerted by COM is active in the somatic cells outside of the reproductive apparatus, the question arose then to know whether Piwi is also a component of this somatic silencing pathway. Indeed, in addition to its function in the reproductive apparatus of flies, diverse functions have been attributed to Piwi protein in somatic tissues at different stages of fly development [23–26]. Thus, GFP expression of the sensor transgenes pGFP-ZAM or pGFP-Id, driven by 24B-Gal4, was analyzed in larvae, pupae, and adult S flies mutated or not for the piwi gene. piwi+/+, piwi+/+, and transheterozygous piwi+/+ mutations were tested in these experiments. The results obtained indicated that the silencing of the sensor transgenes in the somatic tissues of the fly outside of the...
ovaries does not depend on the presence of the Piwi protein. Indeed, in contrast to what was observed in the gonads, GFP expression was not recovered in the somatic tissues of S lines when the Piwi gene was mutated (see Figure 7 and data not shown). These findings provide evidence that Piwi is not required for the silencing pathway present in somatic tissues outside of the reproductive apparatus.

DISCUSSION

The silencing machinery targeting ZAM and Idefix involves the rasiRNA pathway

The rasiRNA pathway has been implicated in the silencing of several repeated genetic elements in the drosophila genome, such as roo, Ste, and the LTR retrotransposon gypsy [13,27]. In this study, we have shown that the rasiRNA pathway, which is based on the strict recognition of sense-strand RNAs, is likely to be involved in the silencing of two additional LTR retrotransposons from Drosophila melanogaster, ZAM and Idefix. rasiRNAs have been reported to arise mainly from the antisense strand of retrotransposons or repetitive sequences [15]. Consistent with this, a profound strand bias for the silencing of the + strand of both ZAM and Idefix was observed. Further, rasiRNAs have been reported to consist of single-stranded RNAs of 25-30 bases in length, and in our experiments we detected 20-30 nucleotide long small RNAs corresponding to antisense strands of ZAM and Idefix. Most of these RNAs also appear to be single stranded, since most of them were only detected using a probe complementary to antisense strand ZAM and Idefix RNAs. The 30-nucleotide antisense strand RNAs might be effectors of a rasiRNA pathway that silences ZAM and Idefix. At the same time, the detection of RNAs of 20 bases in length is consistent with the possibility that an additional pathway could contribute to the establishment of complete silencing.

The silencing of ZAM and Idefix is not only ensured in the ovaries as already described for other transposable elements, but our data further provide evidence that it is also ensured in somatic tissues of the whole fly from embryos to adults.

In Drosophila cells, a surveillance machinery is thus capable of specifically detecting genomic mRNA from both of these retrotransposons and interpreting their synthesis as an ongoing invasion that has to be countered because it would be ultimately harmful.

Transposable elements are generally viewed as genomic forces that are able to contribute to genomic diversity [28]. With this in mind, it is interesting that the expression of genes which have been subjected to integration of ZAM or Idefix in the antisense orientation will not be affected by the silencing machinery. The strand bias can thus preserve this source of genetic innovation brought by these nucleic invaders.
Figure 7. **ZAM and Idefix are regulated by a PIWI-independent pathway outside of the reproductive apparatus.** The pGFP-ZU sensor transgene driven by 248-Gal4 is not expressed in larvae, pupae, or adult stages in a [S/S; piwi+/+] genetic background (right panel). Only a very faint level of fluorescence, corresponding to the background, is detected. A clear GFP expression is observed in these stages of development in a [U/U; piwi+/+] line (middle panel). In piwi mutant backgrounds, in homozygous [S/S; piwi3/3] lines, the silencing of the sensor transgene is not released. A very faint fluorescence level similar to that observed in homozygous [S/S; piwi+/+] lines is observed (left panel).

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The **COM** locus directs silencing through either a Piwi-dependent or -independent silencing pathway in different tissues

To ensure the silencing of endogenous genetic elements in the germ line or in surrounding somatic cell lineages, rasiRNAs function through Piwi, a protein of the *Drosophila* Argonaute family [1,3,13–15]. Consistent with data reporting the tissue-specific expression of Piwi in ovaries [14,29], our data indicate that Piwi is required for the silencing of **ZAM** and **Idefix** in ovarian tissues. However, several data have reported that Piwi is also active outside of the reproductive apparatus. It has been implicated in the regulation of transgenes expressed in tissues where the Adh gene is expressed [24], in the salivary glands [30] or recently in the eyes [26]. Furthermore, Piwi has been found to colocalize with PcG bodies, and it was suggested that it could regulate the nuclear organization of PcG chromatin targets [23]. Despite these somatic functions of Piwi, our data indicate that the somatic silencing of **ZAM** and **Idefix** which is also based on the strict recognition of their sense-strand RNAs, is Piwi-independent.

Our results further demonstrate that in the whole fly, the silencing of **ZAM** and **Idefix** is under the control of the **COM** locus [16,31]. This discrete genomic locus, **COM**, is located at the heterochromatic 20A locus of the *Drosophila* X-chromosome and suppresses the activities of **ZAM** and **Idefix** [16,31]. It displays several specific characteristics. It is located in the same region as the *flamenco* gene which controls the mobilisation of the retrotransposon *Gypsy* [31]. Further, it is mainly composed of defective transposon copies (BDGP release 5). Recently, it has been shown to be a piRNA cluster, and piRNAs homologous to defective copies of **ZAM**, **Idefix**, and *Gypsy* emitted by this locus have been reported [9]. Brennecke et al further reported that a link exists between *flamenco*-derived piRNAs and *gypsy* suppression. Based on their results, they proposed an amplification loop, the “ping-pong” model, to account for piRNA biogenesis. According to this model, sense transcripts from transposons are cleaved by Piwi or Aub RISC with a piRNA guide. The cleaved transcript is not merely degraded but is also used to program Ago3 RISC. This complex, in turn, cleaves the antisense transcripts that originate from master loci such as the 20A locus. Again, the cleaved RNA serves to program Piwi or Aub RISC. Thus sense and antisense transcripts fuel an amplification cycle. This scenario is consistent with most of the characteristics of the silencing mechanism that targets **ZAM** and **Idefix** in the ovaries. However, if a link exists between the **COM**-derived small RNA and the silencing of **ZAM** and **Idefix**, it should implicate another protein than Piwi in the soma. Furthermore, an important piece of data originating in our study remains obscure. Although our genetic analysis points to **COM** as a master regulator of **ZAM** and **Idefix** silencing, some sequences from endogenous **ZAM** are absent from the **COM** locus (release 5 and Hadi Quesneville personal communication). For example, none of them is complementary to the 5′UTR fragment of **ZAM**, whose sequence has been directly implicated in the silencing of the pGFP-ZU transgene. A direct interaction between rasiRNAs derived from the 20A locus and these targeted transgenes seems thus to be excluded. In the ping-pong model, the need for mutual complementarity keeps the production confined to one pair of complementary piRNAs, preventing piRNA generation from spreading along a primary transcript as allowed by RNA-dependent RNA polymerase (RdRP)-mediated amplification. If piRNA emitted from the **COM** locus cannot spread along primary **ZAM** or **Idefix** transcripts, then an additional step in the piRNA mechanism should exist to ultimately direct destruction of transcripts bearing any fragment of **ZAM** or **Idefix**.

More data are then necessary to understand how such piRNAs are generated and what their exact role is in the control. Our present data, however, already implicate the ubiquitous activity of **COM** coupled to different factors in various tissues.

**MATERIALS AND METHODS**

**Drosophila strains**

The S line *w*^1118^ and the U line Rev were from the collection of the Institut National de la Santé et de la Recherche médicale UMR 384. The ubiquitous actin-GAL4 and the mesodermic 24B-Gal4 drivers used are both located on chromosome 3. The 24B-Gal4 driver was a gift from K. Jagla.

All transgenic lines were obtained by injection of indicated transformation vectors into *w*^1118^. All stocks were maintained at 20°C. Expression of transgenes in a genetic context allowing **ZAM**
and Idefix mobilisation was obtained in the progeny of crosses performed with transgenic lines (S genotype) and the U line described in [32]. Flies used for analysis of expression were raised and kept at 25°C.

Transgenic constructs

pZ499 is the pZ4M construct described in [19]. It contains a ZAM LTR upstream of the LacZ reporter gene. In pZ475 and pZ310, the ZAM LTR is shortened at its 3’ end, resulting in fragments encompassing nucleotides 1 to 475 and 1 to 310, respectively, with respect to the ZAM sequence. pZ310 contains a minimal hsp70 promoter between the ZAM fragment (corresponding to the U3 part) and the LacZ gene. The pUAst-GFP vector was used for sensor transgene experiments. For pGFP-ZU and pGFP-IdU, the ZAM UTR (from nucleotide 475 to 1841) and Idefix UTR (from nucleotide 502 to 1024) were cloned downstream of GFP. pGFP-Zenv includes 720 bp of the ZAM env coding region (6385–7105), and pGFP-Idgag includes 1419 bp homologous to the Idefix gag coding region (1003–1422). In these transgenic constructs, the ZAM or Idefix fragments were cloned to be transcribed in the sense orientation, i.e., in a 5’ to 3’ orientation, with respect to endogenous ZAM and Idefix transcription. In pGFPZenvAS and GFPIdUAS, the ZAM or Idefix fragments were cloned in the opposite orientation.

Histochemical staining for β-galactosidase

Ovaries were dissected in 1× phosphate-buffered saline (PBS), fixed in 0.5% glutaraldehyde in PBS for 5–10 min at 4°C, and rinsed twice in 1× PBS and once in Fe/NaP buffer [0.003 M Na4Fe(CN)6, 0.072 M NaH2PO4, 0.003 M K3Fe(CN)6, 0.003 M K2Fe(CN)6, 0.15 M NaCl, 0.001M MgCl2]. Staining was performed in Fe/NaP buffer with X-Gal (0.2 mg/ml final concentration) at 37°C. All samples were stained simultaneously and for the same length of time (2 hrs). Stained tissues were washed four times in 1× PBS, mounted in 1:1 PBS:glycerol, and examined under an Axioshot microscope (Zeiss) using Nomarski optics.

Fluorescent microscopy

Light and fluorescence microscopy was performed with an Olympus confocal microscope or an Olympus SZX12 binocular and a CCD color view camera. Comparisons between stable and unstable lines were carried out using the same acquisition settings.

Flip-out experiments

The hFLP flies (w1118, hnp-FLP, cka12 Sb/TM6, Ubx e7), kindly provided by Kent Golic, express fLP recombinase under the heat shock promoter hsp70. Virgin hFLP females were crossed with transgenic males for 24 hrs on cornmeal-glucose-yeast media at 20°C. Heat shocks of embryos <24 hrs old were performed as described by Ahmad and Golic [1996].

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Northern blot experiments

Total RNA from adult flies was extracted by Trizol, and 40 μg of total RNA was resolved on 1% denaturing agarose gels and probed with radiolabelled transcribed probes homologous to GFP. An actin 5C probe was used as a loading control. Experiments were repeated three times. GFP signals were quantified with a Biorad S125 phosphorimager.

RNase protection assays

Small RNAs from adult flies were extracted using the Ambion mirVana™ miRNA isolation kit. Aliquots of 5 μg of small RNAs were used in RPA experiments. Radiolabelled RNA probes homologous to the 5’UTR regions of ZAM or Idefix, or to the gag gene of Idefix, were 400 to 500 bases long (ZAM UTR from nucleotide 1027 to 1515, Idefix UTR from nucleotide 567 to 1010, Idefix gag from nucleotide 1028 to 1422). 5×104 cpm of specific activity probe was used. As indicated for the Ambion mirVana™ miRNA detection kit, hybridization was performed overnight at 42°C, and protected fragments were digested for 45 minutes at 37°C by RNase A/RNase T1. After RNase inactivation, protected fragments were precipitated and separated on a 15% acrylamide/polyacrylamide (19:1) gel running in 0.5×TBE. Protected fragments were detected by autoradiography.

In situ hybridization

ZAM and Idefix mRNA expression was detected by in situ hybridization using DIG-labelled RNA probes transcribed from the pBS plasmids containing ZAM (3830 to 8040) or Idefix (4866–7191) fragments, using the kit from Roche. Ovaries from third instar larvae were dissected in phosphate buffer saline (PBS). Dissected ovaries were fixed in 5% formaldehyde for 30 min. Ovaries were rinsed with PB (PBS, 0.1% Tween 20) prior to proteinase K treatment. Hybridization was performed in hybridization solution (50% formamide, 5× SSC, 0.1% Tween 20, 50 μg/ml heparin, 100 μg/ml salmon sperm DNA, and 100 μg/ml yeast tRNA) at 45°C overnight and was followed by washes in a 1:1 mixture of hybridization solution and PB at 45°C for 30 min each, and in PB at room temperature (two washes of 20 min each). The hybridized probe was detected using the DIG nucleic acid detection kit (Roche).

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Author Contributions

Conceived and designed the experiments: CV SD NB CM. Performed the experiments: SD NB CM. Analyzed the data: CV SD NB CM. Contributed reagents/materials/analysis tools: MC. Wrote the paper: CV.
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