

Evidence for maternally transmitted small interfering RNA in the repression of transposition in *Drosophila virilis*

Justin P. Blumenstiel*[†] and Daniel L. Hartl[†]

Department of Organismic and Evolutionary Biology, Harvard University, Cambridge, MA 02138

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Hybrid dysgenesis in *Drosophila* is a syndrome of gonadal atrophy, sterility, and male recombination, and it occurs in the progeny of crosses between males that harbor certain transposable elements (TEs) and females that lack them. Known examples of hybrid dysgenesis in *Drosophila melanogaster* result from mobilization of individual families of TEs, such as the *P* element, the *I* element, or *hobo*. An example of hybrid dysgenesis in *Drosophila virilis* is unique in that multiple, unrelated families of TEs become mobilized, but a TE designated *Penelope* appears to play a major role. In all known examples of hybrid dysgenesis, the paternal germ line transmits the TEs in an active state, whereas the female germ line maintains repression of the TEs. The mechanism of maternal maintenance of repression is not known. Recent evidence suggests that the molecular machinery of RNA interference may function as an important host defense against TEs. This protection is mediated by the action of endogenous small interfering RNAs (siRNAs) composed of dsRNA molecules of 21–25 nt that can target complementary transcripts for destruction. In this paper, we demonstrate that endogenous siRNA derived from the *Penelope* element is maternally loaded in embryos through the female germ line in *D. virilis*. We also present evidence that the maternal inheritance of these endogenous siRNAs may contribute to maternal repression of *Penelope*.

hybrid dysgenesis | maternal effect | RNA interference | transposable element

In sexual organisms, genetic parasites, such as transposable elements (TEs), can proliferate in populations despite being deleterious to the host (1). Harmful effects include mutations in genes (2, 3), chromosomal abnormalities arising from ectopic recombination (4, 5), and costs associated with the production of transcripts that divert resources from the host (6). A prime example of the proliferative ability of TEs is the finding that half of the human genome is composed of these elements (7). Hybrid dysgenesis, a phenomenon in which mobilized TEs can induce sterility, represents an extreme example of the harm TEs can inflict as a result of their proliferation. In *Drosophila melanogaster*, when males carrying *P*, *I*, or *hobo* elements are mated with females lacking them, these elements become mobilized, resulting in sterility of the offspring (8–12). However, in the reciprocal cross between females carrying these elements and males lacking them, TE repression is maintained in the progeny. Thus, the salient pattern of TE-induced hybrid dysgenesis in *Drosophila* is that the maternal germ line maintains repression, whereas the male germ line does not. The mechanism by which maternal TE repression is epigenetically inherited is not well understood. However, it is clear that a genetic phenomenon known as the “homology effect” plays an important role. For *P* and *I* elements, TE repression can be maintained in a manner dependent on maternal copies of the particular TE, and maternal repression can be transmitted, even if the maternal copies are defective and unable to produce functional transcripts (13–15). Furthermore, TE repression can be epigenetically inherited maternally even if copies of the TE are not transmitted to the next generation (16).

Recent studies suggest a potential role for RNA interference (RNAi) in the maintenance of maternal repression of TEs. The RNAi machinery processes dsRNA into smaller 21- to 25-nt double-stranded fragments that mediate the destruction of homologous RNA molecules (17, 18). Mutations in genes necessary for RNAi often lead to increased TE activity, and it is now clear that the RNAi machinery functions as a defense mechanism to reduce the harmful effects of TEs and viruses in a wide variety of organisms, including nematodes (19, 20), fruit flies (21, 22), plants (23, 24), and *Chlamydomonas* (25). Endogenous silencing of TEs by the RNAi machinery likely depends on the action of 21- to 25-nt small interfering RNAs (siRNAs) derived from the same repeat sequences liable to silencing. These species have been designated repeat-associated siRNAs (26).

Given that repeat-associated siRNAs likely play a role in silencing endogenous TEs and that maternal repression can be mediated by TE copies unable to produce functional transcripts, maternal transmission of siRNA molecules may contribute to the maternal transmission of TE repression. Here we examine the hybrid dysgenesis syndrome in *Drosophila virilis* to determine patterns of TE-derived siRNA expression and test the hypothesis that they are maternally inherited.

Hybrid dysgenesis in *D. virilis* occurs in the progeny of a mating between two strains, known as strains 9 and 160. The hybrid dysgenesis is unique in that at least five unrelated families of TEs become mobilized (27, 28). When females of strain 9 are mated with males of strain 160, a large proportion of male and female progeny are sterile, but in the reciprocal cross the progeny are normal (29). The element *Penelope*, a member of a recently described class of retroelements that possess an intron (30), appears to play a primary role (31). It is abundant in strain 160, but aside from several ancient and highly degraded copies (ref. 32 and our unpublished data), is entirely absent from strain 9. Furthermore, when *Penelope* is injected into a strain lacking the element, mobilization of unrelated TEs has been reported (31). Aside from *Penelope*, at least four other unrelated elements are mobilized in the dysgenic cross: *Paris*, *Helena*, *Telemac*, and *Ulysses*. *Paris* is a DNA element in the *mariner/Tc1* family, and *Helena* is a long interspersed nuclear element-like non-LTR retroelement. Each of these families is more abundant in strain 160 and may play some causal role in comobilization. *Telemac*, a BEL-like LTR-retrotransposon, and *Ulysses*, a gypsy-like LTR retroelement, are evenly distributed between the strains and, thus, are unlikely to play a role (27, 28). The mechanisms of maternal repression and comobilization are not known. Here we

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Abbreviations: RNAi, RNA interference; siRNA, small interfering RNA; TE, transposable element.

*Present address: Stowers Institute for Medical Research, 1000 East 50th Street, Kansas City, MO 64110.

[†]To whom correspondence may be addressed: jpb@stowers-institute.org or dhartl@oeb.harvard.edu.

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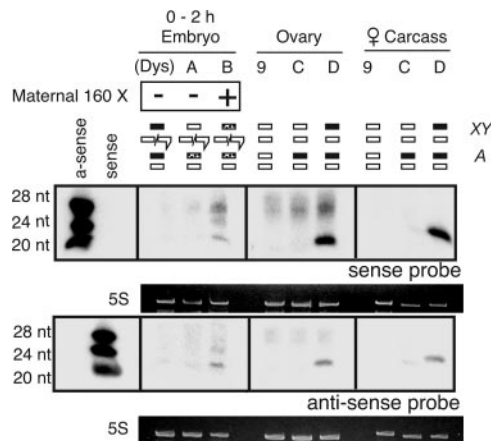


Fig. 3. The X chromosome is the source of *Penelope* siRNA within the female germ line. In 0- to 2-h embryos, *Penelope* siRNA is found only if mothers possess the 160 X chromosome. In females, the X chromosome drives the expression of *Penelope* siRNA in the germ line and soma. Females that lack this chromosome do not demonstrate *Penelope* siRNA in either tissue. Five micrograms of enriched RNA was used per lane.

The Strain 160 X Chromosome Is Necessary for Maternal Transmission of *Penelope* siRNA. Because the primary source of *Penelope* siRNA in whole adults is the 160 X chromosome, we determined whether this expression is limited to either the germ line or the soma. Fig. 3 (see Fig. 6, which is published as supporting information on the PNAS web site, for crossing schemes) shows that *Penelope* siRNA expression in 0- to 2-h embryos depends on the presence of the 160 X chromosome in the mother. Dysgenic embryos (Fig. 3, lane Dys) or embryos whose mother lacks the 160 X chromosome (Fig. 3, lane A) do not exhibit *Penelope* siRNA, but embryos whose mother does carry the 160 X chromosome (Fig. 3, lane B) do show this siRNA. Furthermore, adult females from strain 9 or those lacking the 160 X chromosome (Fig. 3, lane C) do not express the *Penelope* siRNA in ovaries or soma, whereas adult females possessing the 160 X chromosome (Fig. 3, lane D) exhibit this siRNA in ovaries and soma.

Confirmation That Repression of Dysgenesis in Males Results from a Maternal Effect. Because dysgenic and nondysgenic females are genetically identical and differ only in whether strain 9 or strain 160 was the mother, the repression of hybrid dysgenesis in females is due to a maternal effect. However, it is a formal possibility that the mechanism of TE mobilization in the germ line of males is different from that in females. Nondysgenic sons possess the X chromosome from strain 160, whereas dysgenic sons lack this chromosome. It could be that maintenance of repression in the male germ line is zygotic and due to the presence of the X chromosome from strain 160. Specifically, because the X chromosome drives *Penelope* siRNA expression in adults, it may also drive repressive siRNA expression in the early male germ line. If this possibility were the case, then mobilization of TEs in dysgenic males may result from absence of the “protective” X chromosome from strain 160 rather than absence of maternally transmitted siRNA.

To determine whether TE mobilization in the male germ line is due to the absence of maternal repression, we compared the level of dysgenesis in genetically identical males, all possessing the 160 X chromosome but differing in the genotype of the mother (Fig. 4; see Fig. 7, which is published as supporting information on the PNAS web site, for crosses). Nondysgenic males from the 160 female \times 9 male cross and male progeny from the XX/9 female \times 160 male cross are genetically identical.

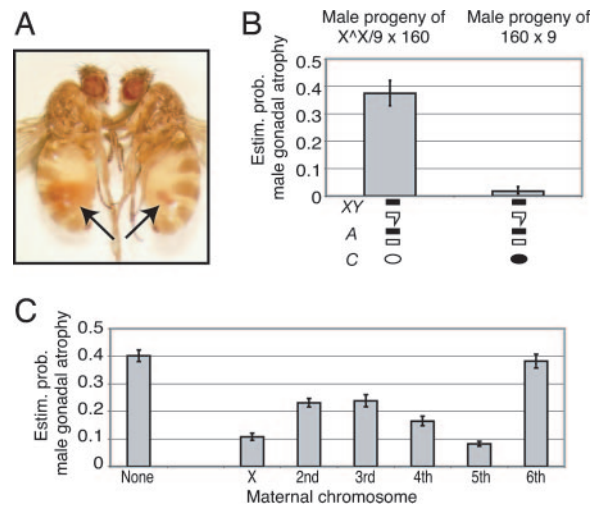


Fig. 4. The contribution of maternal 160 chromosomes to the repression of gonadal atrophy. (A) Wild-type testis (left) and atrophied testis (right). (B) Genetically identical progeny of crosses with or without an attached-X chromosome (XX) provide evidence that hybrid dysgenesis in males results from lack of maternally transmitted repression. Shown is the estimated probability of gonadal atrophy in genetically identical males that differ only in respect to the cytoplasm of the mother. Males that possess the X chromosome from strain 160 but whose mothers lack chromosomes from strain 160 show high levels of gonadal atrophy ($n = 211$). Males whose mothers possess 160 chromosomes do not ($n = 228$). (C) Maternal effects of individual strain 160 chromosomes on suppression of gonadal dysgenesis; the lower the bar, the greater the level of repression. The largest effects are due to the X chromosome and chromosome V.

In particular, both types of males possess an X chromosome from strain 160. However, maternal inheritance differs between these males, because the mothers in the nondysgenic cross 160 female \times 9 male are from strain 160, whereas mothers of the XX/9 female \times 160 male cross possess an attached-X chromosome in a strain 9 background. For both types of males, the degree of sterility was determined by counting atrophied testes as shown in Fig. 4A, and the probability of gonadal atrophy was determined. Nondysgenic males showed nearly complete repression of testicular atrophy, whereas the probability of atrophy for each of the testes of identical males whose mothers lack chromosomes from strain 160 is nearly 40% (Fig. 4B). This result is similar to currently observed levels of gonadal atrophy in the dysgenic cross (see below). Thus, hybrid dysgenesis in males correlates with the absence of a maternal factor rather than the lack of the X chromosome from strain 160 in the zygote.

The 160 X Chromosome Is Sufficient But Not Necessary for Suppression of Dysgenesis. Because the X chromosome from strain 160 drives the expression of *Penelope* siRNA in whole adults, we determined whether this chromosome is sufficient by itself and when placed into a strain 9 genetic background to maternally suppress hybrid dysgenesis. Chromosomes from strain 160 were made heterozygous in the genetic background of strain 9, either alone or in combination with other chromosomes from strain 160, by means of two generations of crossing with strain 9 (see Fig. 8, which is published as supporting information on the PNAS web site, for the crossing schemes and sample sizes). Females of each genotype were then singly mated with males of strain 160 to determine the degree to which each female genotype could maternally transmit factors suppressing dysgenesis. The mother’s genotype was assessed post hoc by examining the progeny for the presence of recessive markers on all five autosomes.

Fig. 4C shows that a maternal X chromosome from strain 160 contributes substantially to suppression of sterility in the male

offspring. The estimated probability of gonadal atrophy is reduced from 40% to 11% when a maternal X chromosome alone is provided. The second, third, and fourth chromosomes exhibit a weaker maternal effect on sterility, although the fifth chromosome has a similar effect on suppression as the X chromosome (estimated 8% gonadal atrophy). Interestingly, the tiny sixth chromosome, which is homologous to the fourth chromosome of *D. melanogaster*, has a negligible effect on the suppression of dysgenesis. This chromosome in strain 160 lacks copies of *Penelope*, whereas all other chromosomes from strain 160 possess copies of *Penelope* (28).

The data in Fig. 4C also indicate that, although the X chromosome and autosomes 2–5 all contribute to maternal repression, no individual chromosome is sufficient to maximally repress hybrid dysgenesis. Rather, the chromosomes act cumulatively (Fig. 8), and, when the mother's genotype includes all six chromosomes from strain 160, complete repression is observed (proportion of atrophied testes, 0 of 248 males). Indeed, strong repression is observed when the mother's genotype includes chromosomes 2–6 from strain 160 but the X chromosome from strain 9 (proportion of atrophied testes, 1% of 368 males). These data imply that the X chromosome from strain 160, as a source of *Penelope* siRNA, has a strong repressing effect, but that other chromosomes from strain 160, which also carry *Penelope*, also are capable of mediating repression.

Discussion

We have demonstrated that small (21–25 nt) sense and antisense RNA species homologous to *Penelope* can be identified in strain 160, and, in females, these species are found both within the germ line and soma. We estimate the size of the small RNA molecules to be ≈ 23 nt, which is similar to the size found for most TE-derived siRNAs found in *D. melanogaster* (26). These species of endogenous RNA derive from a locus (or loci) on the X chromosome and are transmitted to offspring through the female, but not male, germ line. Additionally, the X chromosome provides substantial repression of dysgenesis in the offspring when present in the mother. Thus, *Penelope* siRNA complexes may function as an important maternal factor that mediates repression in nondysgenic embryos.

The X-linked locus (or loci) that functions as the source of siRNA molecules in the female germ line is not known, but it presumably generates double-stranded *Penelope* species that are processed by the RNAi machinery. We have shown definitively that this presumptive *Penelope* siRNA is maternally, but not paternally, transmitted. We have no direct evidence that these *Penelope* siRNAs account for the strong suppression of dysgenesis observed for the X chromosome, although the circumstantial evidence supports this hypothesis.

The mechanism or mechanisms by which the other chromosomes mediate their maternal effect is not known. All chromosomes that possess *Penelope* elements contribute cumulatively to maternal repression (Figs. 4 and 8). There are a number of possibilities. The autosomes may contribute siRNA molecules at a level below the limit of detection. By way of precedent, RNAi silencing has been observed in *Caenorhabditis elegans* in the absence of detectable levels of siRNA (34). Alternatively, sequence divergence between autosomal copies of *Penelope* in strain 160 may preclude their detection by our probe, or they may originate primarily from regions of *Penelope* not included in the probe. Independently of any siRNA effects, maternal repression of *Penelope* may be mediated by means that are reminiscent of the "poison subunit" effect on *P* element transposition caused by certain internally deleted elements designated *KP* (35, 36). Finally, the hybrid dysgenesis system in *D. virilis* is unique in that multiple TEs become mobilized, and, although *Penelope* appears to be the primary cause of the comobilization, the other TEs may also contribute (28). The failure of the 160 X chromosome to

completely suppress dysgenesis and the cumulative effects of the autosomes may simply reflect the complex nature of TE repression and mobilization.

Several aspects of the dysgenic syndrome observed here in *D. virilis* differ from those for the *P* element observed in *D. melanogaster*. First, it appears that repression is maintained when a sufficient number of chromosomes from strain 160 are present in the mother, even if these chromosomes were inherited paternally. Our crossing scheme used to measure the maternal effect of different chromosomes on suppression (Fig. 8) entailed backcrossing hybrid males to strain 9 females. Despite the backcross to strain 9, females that are heterozygous for all chromosomes from strains 9 and 160 still transmit repression to their offspring. This effect is not observed with *P* elements in *D. melanogaster*. In this system, when a mother is mated with a male possessing *P* elements, her ability to transmit repression to her progeny depends, in part, on the genotype of her own mother (8, 16), and the maternal transmission of repression depends on the continued presence of *P* copies in the genome of the mother (37). Over time, when transmitted maternally, the level of repression accumulates. The delay in the accumulation of maternal repression is also observed in the case of the *I* element when repression is mediated by a transgene (13). Here, by contrast, females are capable of transmitting repression as long as they possess a sufficient number of chromosomes from strain 160, even if their mothers were originally from strain 9. Nonetheless, although these results show that a grandmother contribution is not essential to the repression of dysgenesis, it can certainly modulate it (28).

A second aspect that differs between hybrid dysgenesis in *D. melanogaster* and that in *D. virilis* is the comobilization of unrelated TEs. When first observed, it was difficult to imagine how multiple unrelated elements (class I LTR and non-LTR elements as well as class II DNA elements) could be mobilized by similar mechanisms (27). However, it is becoming increasingly clear that the RNAi machinery regulates a diverse array of genetic parasites. For example, in nematodes, a DNA transposon is regulated by the RNAi machinery (20), whereas, in flies, a likely component of the RNAi machinery, *spn-E*, plays a role in repressing LTR and non-LTR elements in ovaries and testes (21). In an analysis of the small RNA profile during *D. melanogaster* development, small species of RNA derived from LTR and non-LTR elements as well as DNA elements were all found (26). The cause of comobilization in *D. virilis* is not known, but it appears that *Penelope* plays a primary role. Many viruses have evolved the ability to antagonize the RNAi machinery (38–40). Furthermore, it has been demonstrated in plants that these suppressors can cause developmental defects by interfering with the proper function of microRNAs important in development (41, 42). It is therefore possible that *Penelope* possesses factors that suppress RNA silencing, and the comobilization of other elements may be a pleiotropic consequence of a compromised RNAi machinery. It is tempting to speculate that the sterility is also a pleiotropic consequence of a compromised RNAi machinery, especially in light of evidence that components of the RNAi machinery, such as *piwi* (43, 44), also play a role in germ-cell maintenance.

Although the *D. virilis* syndrome of hybrid dysgenesis differs in several aspects from the *P* and *I* element system, the most significant similarity is that, in all three cases, the female germ line maintains TEs in a repressed state, whereas the male germ line does not. In *D. melanogaster*, this observation also extends to *lacZ* reporter constructs that only undergo transsilencing in the germ line when inherited maternally (45). From an evolutionary perspective, it may be that, because females invest substantial resources in the production of oocytes, natural selection favors additional investment in maintaining genetic

parasites in a silent state. Interestingly, microarray experimental data (46) imply that, among the core components of the RNAi machinery, seven of nine differentially expressed components are up-regulated in ovaries relative to testes (Fig. 9, which is published as supporting information on the PNAS web site). Furthermore, if silencing is mediated by protein-RNA complexes, males that produce and transmit these complexes may be at a disadvantage to the extent that success in fertilization

requires the production of large numbers of energetically inexpensive sperm (47).

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