

A *C. elegans* Piwi, PRG-1, Regulates 21U-RNAs during Spermatogenesis

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Summary

Background: Epigenetic regulation by diverse classes of small RNAs is mediated by the highly conserved Argonaute/Piwi family of proteins. Although Argonautes are broadly expressed, the Piwi subfamily primarily functions in the germ line. Piwi proteins are associated with germline-specific ribonucleoprotein (RNP) granules in *Drosophila*, zebrafish, and mouse. Depending on the species and on the specific family member, Piwis play important roles in spermatogenesis and/or in maintaining germ cell and stem cell totipotency. Piwis bind to a newly discovered class of small RNAs, called piRNAs. *C. elegans* contains a large set of Argonaute/Piwi-related proteins, including two closely related to *piwi* called *prg-1* and *prg-2*. The function of *prg-1* and *prg-2* and whether piRNAs exist in *C. elegans* is unknown.

Results: Here, we demonstrate that the Piwi-like protein PRG-1 is localized to P granules in germ cells entering spermatogenesis and is required for successful spermatogenesis. Loss of *prg-1* causes a marked reduction in expression of a subset of mRNAs expressed during spermatogenesis, and *prg-1* mutant sperm exhibit extensive defects in activation and fertilization. Moreover, *prg-1* activity is required for the presence of the small RNAs called 21U-RNAs.

Conclusions: Our data suggest that PRG-1 promotes expression, processing, or stability of 21U-RNAs, which, in turn or in concert with PRG-1, promote proper expression of spermatogenesis transcripts.

Introduction

Piwi proteins bind to and are required for the accumulation of small RNAs known as piRNAs (reviewed in [1]). piRNAs are derived from a few long precursor transcripts expressed from heterochromatic or noncoding regions of the genome. The biogenesis and function of piRNAs remains mysterious, in part because piRNA sequence and Piwi function differ between species. For instance, in *Drosophila*, most piRNA sequences correspond to transposons, and the Piwi proteins act primarily to suppress transposon activity in the germ line [2, 3]. In mice the three Piwi-related proteins Miwi, Mili, and Miwi2 primarily function in spermatogenesis and only Miwi2 and Mili have an apparent role in transposon silencing [4–7]. Mouse piRNAs fall into two groups, one expressed premeiotically that contains transposon-associated sequences and one expressed meiotically that consists of unique sequences [8–10]. This second group of piRNAs does not have sequence

complementarity to coding or 3'UTR sequences, and their function in spermatogenesis remains unclear.

C. elegans contains a large set of Argonaute/Piwi-related proteins, including two genes closely related to *piwi* called *prg-1* and *prg-2* (*prg* = *piwi*-related gene [11]). *prg-1* mutants display a temperature-sensitive defect in fertility that is currently uncharacterized [12]. Although piRNAs have not been identified in *C. elegans*, a class of small RNAs called 21U-RNAs was recently discovered [13]. 21U-RNAs bear several similarities to piRNAs: they have a uridine at the 5' position; their sequences are diverse, nonconserved, and do not obviously match mRNAs or transposons; and the loci encoding 21U-RNAs are tightly clustered in the genome [13]. However, 21U-RNAs are shorter than piRNAs (21 nt rather than 26–31 nt) and appear to be individually transcribed rather than derived from a long precursor transcript [13], so their relationship to piRNAs or other small RNA species in other organisms is uncertain. In this report we investigate the function of PRG-1 and PRG-2 in the germ line of *C. elegans* and define both a set of mRNAs and a class of small RNAs that require PRG-1 for their expression.

Results

Disruption of *prg-1* Causes Temperature-Sensitive Sterility

The key functional motifs of Argonaute/Piwi proteins include an N-terminal PAZ domain that binds the 3' end of a small RNA and a C-terminal PIWI domain that has RNase H catalytic activity (Figure 1A). The PIWI domain also coordinates binding of the 5' phosphate of the small RNA, along with a basic pocket domain called the MID domain [14]. To study the function of the *C. elegans* *piwi*-related genes *prg-1* and *prg-2*, we obtained deletion mutants [11]. *prg-1(tm872)* deletes 640 bp encoding most of the MID domain and part of the PIWI domain, and *prg-2(tm1094)* removes 1065 bp, which includes the entire PAZ domain. Both *prg-1(tm872)* and *prg-2(tm1094)*, therefore, should be null or loss-of-function alleles.

A previous report demonstrated that treating animals with RNAi to reduce *prg-1* and *prg-2* levels resulted in decreased germline stem cell proliferation [11]. We therefore first examined *prg-1(tm872)*, *prg-2(tm1094)*, and *prg-1(tm872); prg-2(tm1094)* mutants for defects in proliferation by monitoring mitotic germ cell number with phosphoS10 histone H3 staining but did not see any statistically significant differences from wild-type. Moreover, both DIC microscopy and DAPI staining of adult *prg-1*, *prg-2*, and *prg-1; prg-2* double mutant gonads revealed essentially normal germ cell number and development (Figures 1B and 1C). However, as previously reported, *prg-1(tm872)* mutants exhibit temperature-sensitive sterility: they have reduced brood size at 20°C and are essentially sterile at 25°C (less than five progeny per animal [12]) (Figure 1D). *prg-2(tm1094)* single mutants do not display any obvious defects compared to wild-type and do not enhance or otherwise affect *prg-1(tm872)* sterility at 25°C.

The Sterility of *prg-1(tm872)* Mutants Arises from a Defect in Spermatogenesis

The germ line of adult *C. elegans* hermaphrodites contains both sperm and oocytes. Sperm are made only during the

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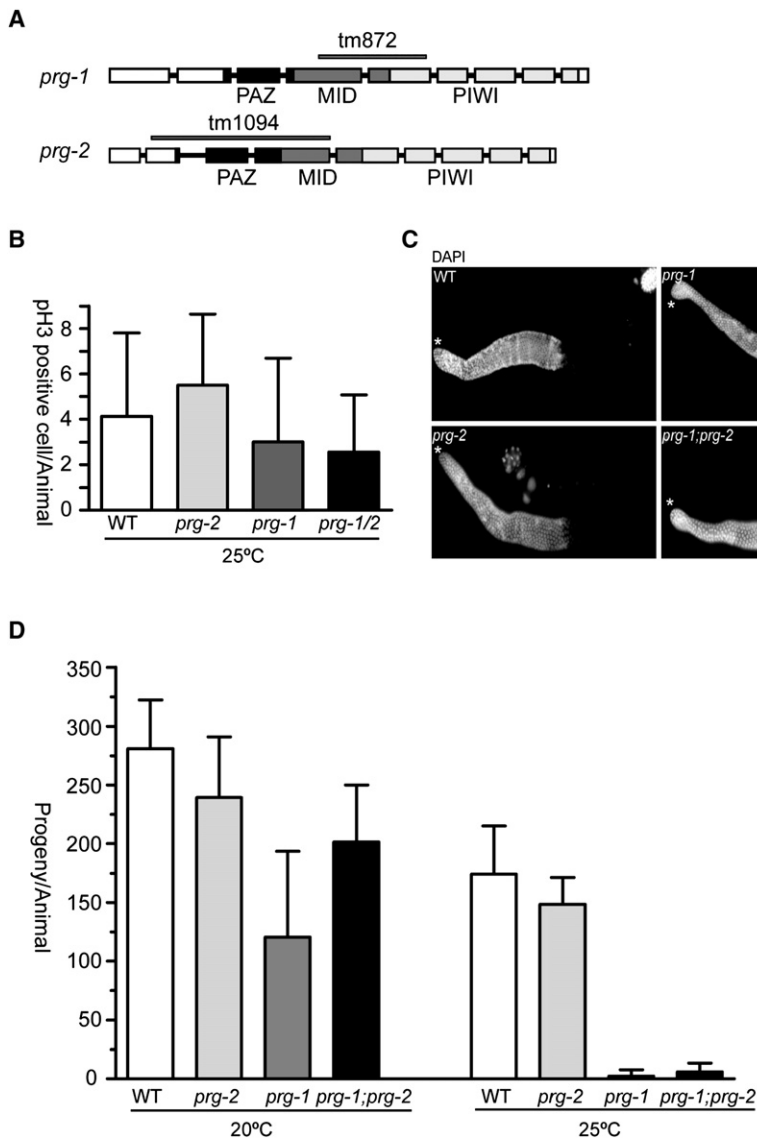


Figure 1. Disruption of *prg-1* Causes Reduction in Brood Size and Temperature-Sensitive Sterility (A) Schematic of *prg-1* and *prg-2* genomic structure. Box, exon; connecting line, intron. Functional domains (PAZ, MID, and PIWI) of PRG proteins are indicated below. The solid lines above indicate the deletions in *prg-1(tm872)* and *prg-2(tm1094)*.

(B) The average number of germ cells per gonad positive for phosphorylated Ser10 of histone H3 at 25°C is graphed. The x axis indicates genotype; the y axis represents mean number of pH3⁺ cells. n > 10 gonads. Error bars represent standard deviation.

(C) Dissected gonad arms from different genotypes stained with DAPI. Asterisk marks distal end of gonad.

(D) *prg-1* mutants have a temperature-sensitive sterile phenotype. Brood size of wild-type or mutant worms was determined at 20°C or 25°C. The x axis indicates corresponding genotypes; the y axis is the mean value of total number of progeny per animal. Greater than eight animals per trial, average of three trials. Error bars represent the standard deviation.

fourth stage (L4) of larval development, and oocytes are subsequently produced throughout early adulthood. In males sperm production persists into adulthood. Even though hermaphrodites are self-fertile, male sperm are preferentially used over self-sperm for oocyte fertilization if mating occurs.

To determine whether the sterility of *prg-1* mutants arises from a defect in oogenesis or in spermatogenesis, we tested whether providing wild-type sperm to *prg-1* hermaphrodites by mating would rescue the brood size defects at 20°C and 25°C. After mating to wild-type males, *prg-1(tm872)* hermaphrodites exhibited substantially improved fertility at both temperatures; in particular, at 20°C the brood size increased to the same extent as seen in mated wild-type hermaphrodites (Figure 2A). This observation suggests that *prg-1(tm872)* mutant germ cells do not have major impairments in proliferation or oogenesis and that the sterility at 25°C arises from a defect in spermatogenesis.

prg-1 mutant males also exhibit a temperature-sensitive defect in spermatogenesis. *prg-1(tm872)* males raised at 20°C are capable of producing cross progeny when mated to a self-fertile hermaphrodite, but not at 25°C (Figure 2B). At

25°C, *prg-1* males could generate only ~2 cross progeny in 10% of matings, whereas wild-type males generated ~100 cross progeny in 83% of matings. *prg-1* males are physically capable of mating, as we could visualize sperm from the *prg-1* male in a *fem-1* mutant hermaphrodite, which lacks self-sperm. Although mating occurred, *prg-1* males could not rescue the infertility of *fem-1* mutants or produce cross progeny after mating to self-fertile *unc-32* hermaphrodites at 25°C. During spermatogenesis, undifferentiated germ cells undergo meiotic divisions with incomplete cytokinesis. Most cellular contents are dumped into a shared residual body from which the haploid spermatids eventually bud. These spermatids then undergo further morphological changes to become activated, ameboid spermatozoa that are competent for oocyte fertilization. Microscopic examination of spermatogenesis revealed that *prg-1* mutant hermaphrodites and males raised at 25°C possessed differentiated spermatocytes but relatively few mature spermatids (Figure 2C). Because some spermatids were present, we tested whether they were deficient in activation by treatment with pronase, which stimulates pseudopod formation [15]. *prg-1* mutant sperm almost always failed to produce normal pseudopodia (17% in *prg-1* compared to 84% in wild-type; Figures 2D and 2E). Taken together, our data suggest that a defect at very late stages in spermatogenesis is the principal reason for the reduced fertility in *prg-1(tm872)* hermaphrodites and males at both 20°C and 25°C.

PRG-1 Is Localized to P Granules during Spermatogenesis

To determine the distribution and subcellular localization of PRG-1 during germline development, we generated two transgenic lines expressing full-length PRG-1 fused to GFP under the control of the *pie-1* promoter, which is permissive for

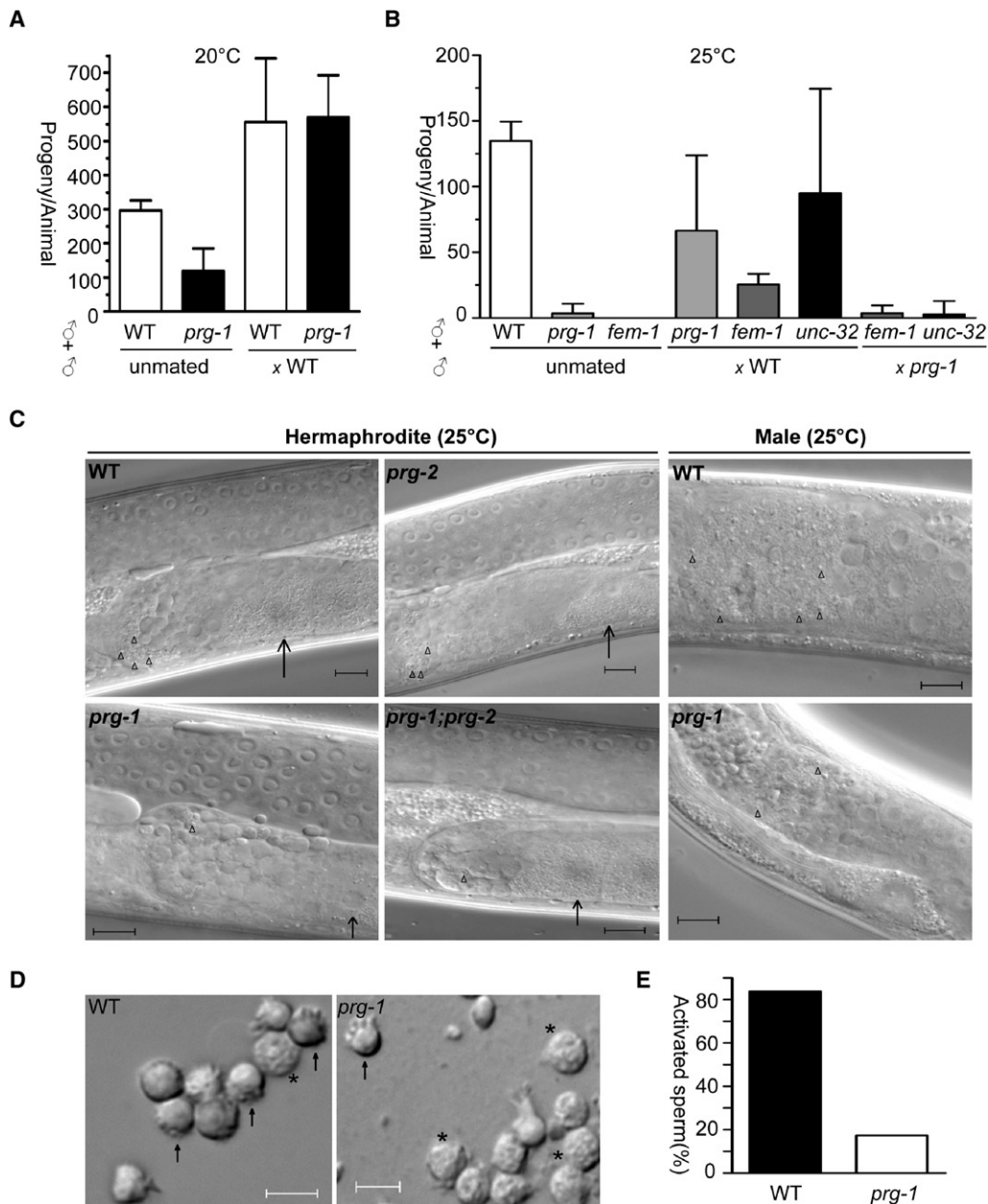


Figure 2. *prg-1(tm872)* Mutants Exhibit Defects in Spermatogenesis

(A) Wild-type or *prg-1(tm872)* L4 hermaphrodites either were left unmated or were mated to wild-type males at 20°C, and the number of progeny was counted. x axis = genotypes, y axis = mean progeny per animal. $n \geq 16$. Error bars represent the standard deviation.

(B) Wild-type or *prg-1(tm872)* males were mated with *prg-1*, *fem-1*, or *unc-32* mutant hermaphrodites at 25°C, and the number of progeny was counted (cross progeny only for *unc-32*). x axis = genotypes, y axis = mean progeny per animal. Greater than eight animals per trial, two or three trials averaged. Error bars represent the standard deviation.

(C) Young adult hermaphrodites or males were examined by microscopy under DIC. Arrows indicate the first oocyte produced after completing spermatogenesis in hermaphrodites. Triangles indicate spermatids. Scale bar: 10 μ m.

(D) Image of pronase-treated spermatids in wild-type and *prg-1* mutants. Scale bar: 5 μ m. Arrow, activated spermatids; asterisk, unactivated spermatids.

(E) Quantification of spermatid activation, $n > 137$ spermatids.

germline expression [16]. These lines showed identical spatial and temporal expression patterns. GFP was detectable in the cytoplasm but was greatly enriched in perinuclear granular structures in pachytene germ cells in males and in L4 hermaphrodites during spermatogenesis (Figure 3A). PRG-1 was preferentially expressed during spermatogenesis because expression decreases over two-fold in hermaphrodite

adults after the switch from spermatogenesis to oogenesis and the granular localization completely disappeared (Figures 3B and 3C). This transgenic expression pattern likely reflects endogenous expression because the *pie-1*-driven PRG-1::GFP transgene partially rescues the *prg-1* mutant sterile phenotype at 25°C from two to 40 progeny ($n = 11$). Moreover, we have generated transgenic strains expressing PRG-1::GFP

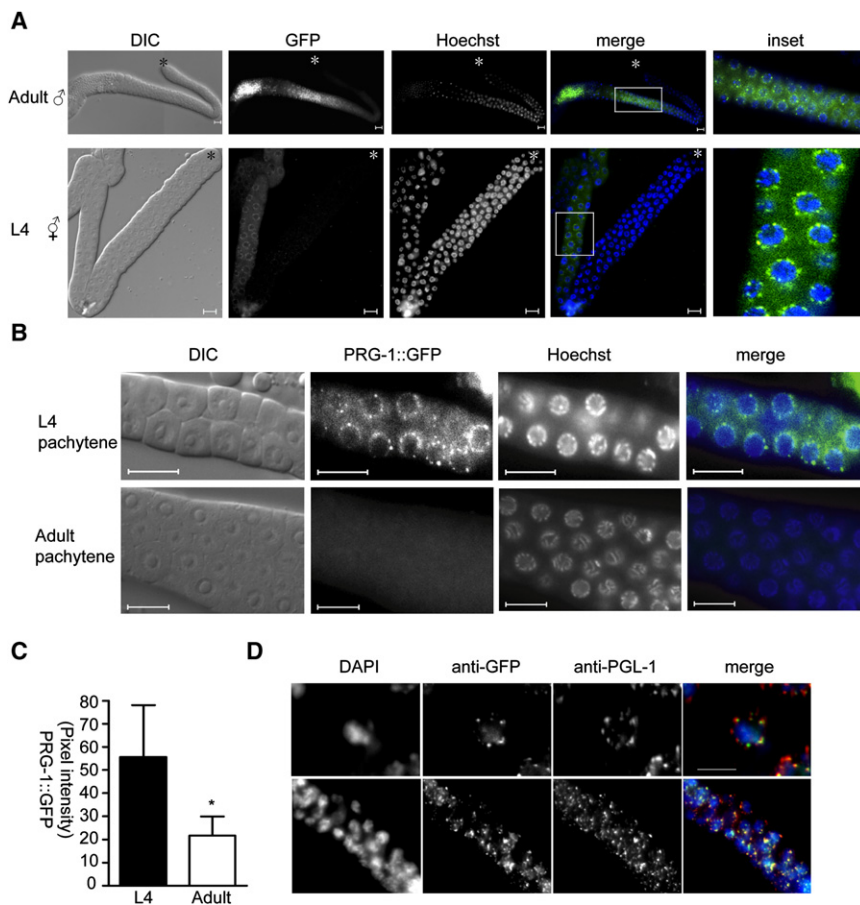


Figure 3. PRG-1 Localizes to P Granules during Spermatogenesis

(A) Live animals expressing the PRG-1::GFP transgene were exposed to Hoechst dye and then gonads directly dissected and mounted for viewing. DIC and fluorescent images were collected. Scale bar: 10 μ m.

(B) The pachytene region of L4 and adult gonads from PRG-1::GFP transgenic animals are shown. Green, PRG-1::GFP; blue, Hoechst; scale bar: 10 μ m.

(C) PRG-1::GFP intensity was measured from equivalent exposures of pachytene germ cells from ten animals each at L4 and adult stages. Error bar represents the standard deviation. Asterisk, Student's t test, $p < 0.01$.

(D) L4 hermaphrodite gonads were stained with anti-GFP (green) and anti-PGL-1 (red). Upper panels: 1000 \times magnification; scale bar, 5 μ m; lower panels: 400 \times magnification.

under the control of its own promoter that exhibited identical spatial, temporal, and subcellular localization to *pie-1*-driven PRG-1::GFP (data not shown).

C. elegans germ cells possess RNP granules called P granules. Similar to germ granules in other species, P granules likely regulate mRNA stability and/or translation, perhaps to correctly localize germline determinants or suppress somatic fates. To determine whether the perinuclear granular structures containing PRG-1::GFP are P granules, we costained the dissected gonads of PRG-1::GFP animals with antibodies to GFP and the constitutive P granule component PGL-1 [17]. We found extensive colocalization between PGL-1 and PRG-1::GFP (Figure 3D), indicating that PRG-1 was indeed localized to P granules. This pattern was consistent with the localization of Piwi family proteins to germ granules in *Drosophila*, zebrafish, and mammals [2, 18–20].

prg-1 Mutant Males Exhibit Reduced Levels of Spermatogenesis-Expressed Transcripts

To further explore the function of PRG-1 during spermatogenesis, we analyzed *prg-1*-dependent gene expression. We performed microarray analysis, comparing dissected gonads from wild-type and *prg-1* mutant males grown at 25 $^{\circ}$ C, in triplicate on biologically independent preparations. Examination of differentially expressed genes identified several telling trends (Table S1). First, 529 transcripts are downregulated ($\geq 1.5\times$, Z test, $p < 0.05$) in *prg-1* mutants, whereas only 48 are upregulated, suggesting that PRG-1 is primarily involved in promoting transcript accumulation. Of the downregulated

set, 63% are preferentially expressed during spermatogenesis, based on previous microarray experiments [21], compared to 10% of the upregulated set. Moreover, *prg-1*-regulated genes displayed a very strong preference for residence on chromosome IV as well as a striking exclusion from chromosome X, consistent with the known distribution of genes expressed during spermatogenesis [21, 22] (Figure 4). Consistent with the *prg-1* mutant phenotype, the downregulated genes included several

that are known to act relatively late during spermatogenesis. For instance, *spe-15* acts during budding of spermatocytes from the residual body, and *spe-11*, *spe-41*, and *spe-42* all act during spermatid activation [23].

Notably, over 1000 genes known to be expressed during spermatogenesis are not significantly affected by loss of *prg-1*. This set includes an extremely abundant and highly variable set of genes encoding major sperm protein (MSP). Fourteen probes on the array detecting different *msp* transcripts had an average fold regulation of 0.95 ± 0.13 . The general stability of most spermatogenesis transcripts, and most especially the stability of *msp* transcripts, in *prg-1* mutants argues against the likelihood that the gene expression changes are a downstream consequence of disrupted spermatogenesis and suggests a more direct role for the differentially regulated genes in causing the mutant phenotype.

prg-1 Mutants Fail to Express 21U-RNAs

We hypothesized that PRG-1 might function in the production of a small RNA species related to piRNAs based on the role of Piwi-like proteins in other species [1]. A recently described class of small *C. elegans* RNAs called 21U-RNAs resembles piRNAs in two respects: they have a uridine at the 5' position, and the corresponding 21U-RNA genes are tightly clustered in the genome [13]. We noted that the 21U-RNA gene clusters are all located on chromosome IV near regions that encode many sperm-enriched transcripts and that 21U-RNAs were present at slightly higher frequency in a population containing males [13]. These observations suggested to us that 21U-RNA expression might be regulated by PRG-1.

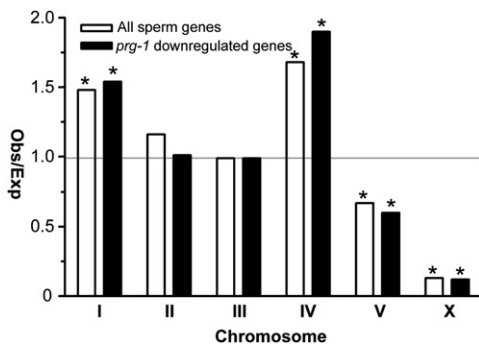


Figure 4. *prg-1*-Regulated Genes Exhibit the Same Chromosome Bias as Do Genes Expressed during Spermatogenesis

A total of 1300 genes known to be expressed during spermatogenesis and the 529 genes downregulated by *prg-1* were mapped to their chromosomal locations, and the numbers were compared to the expected number given the representation of each chromosome on the microarray. Asterisk = $p < 0.01$ (Chi square).

We therefore isolated RNA from wild-type and a set of mutant hermaphrodites with various germline defects. We enriched for RNAs <200 bp and used northern analysis to examine the expression profile of the 21U-RNA *21ur-1* [13]. *21ur-1* RNA can be detected in wild-type hermaphrodites at L3, L4, and adult stages, but not in *prg-1* mutants at those stages (Figure 5A). *21ur-1* was still expressed in *prg-2* mutants, but not in *prg-1*; *prg-2* mutants or in *glp-1* mutants, which lack germ cells (Figure 5B). A *fem-1(lf)* mutant lacking sperm but containing oocytes displayed reduced levels of *21ur-1*, whereas a *fem-3(gf)* mutant that contains sperm and no oocytes had normal levels. The effect of loss of *prg-1* activity is specific, as expression of the miRNA *let-7* was unaffected. Thus, *21ur-1* displays the expected characteristics of a small RNA expressed during spermatogenesis that requires PRG-1 for its expression. Another 21U-RNA, *21ur-1353*, also depended upon *prg-1* for its expression (data not shown), indicating that the expression of 21U-RNAs generally require PRG-1.

Discussion

Based on our data, we propose that PRG-1 functions in a manner analogous to mammalian Miwi. Both PRG-1 and Miwi proteins achieve peak expression during meiosis ([4]; this work). Like *prg-1*, *miwi* mutations cause a relatively late defect at the onset of spermiogenesis (the transformation of round spermatids to mature sperm), whereas the other mammalian Piwi-like proteins, *mili* and *miwi2*, act at an earlier stage, in prophase of meiosis I. Moreover, Miwi-associated piRNAs do not have additional sequence matches in the genome and do not correspond to transposons [10, 24], similar to the 21U-RNAs regulated by PRG-1.

This model of PRG-1 function differs from *Drosophila*, where Piwi, along with related proteins Ago3 and Aubergine, regulate piRNAs to silence transposable elements through an amplification mechanism [2, 25]. 21U-RNAs do not show sequence similarity to transposons and do not have the 10 bp sequence overlap with each other that is characteristic of the Ago3/Aubergine-mediated amplification mechanism [13]. Additionally, our data do not support the notion that PRG-1 and PRG-2 function together to amplify 21U-RNAs because 21U-RNAs

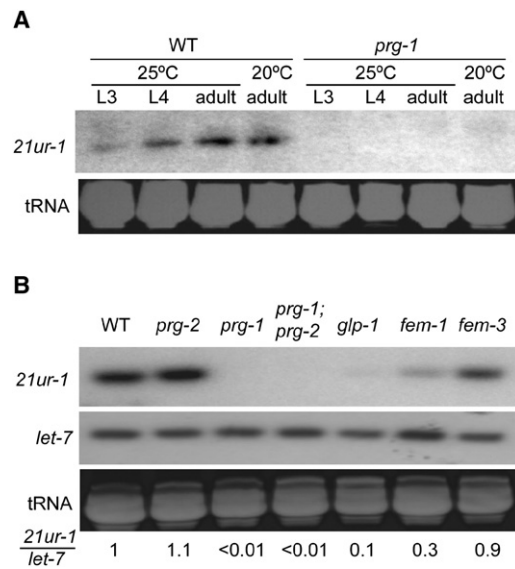


Figure 5. *prg-1* Mutants Fail to Express 21U RNAs

(A) Temporal expression of *21ur-1* in wild-type and *prg-1(tm872)* worms at 20°C and 25°C.

(B) *21ur-1* RNA expression in mutants with defects in germline development: *glp-1* has almost no germ cells, *fem-1* has oocytes only, and *fem-3* has sperm only. Numbers indicate the relative intensity ratio of *21ur-1* to *let-7* after comparison to wild-type, which was set to one. Ethidium bromide staining of tRNA shows similar loading levels.

are still present in a *prg-2* mutant and the *prg-1* mutant phenotype is not affected by loss of *prg-2*.

Intriguingly, the genomic sites of 21U-RNA loci clusters are likely conserved between *C. elegans* and the related species *C. briggsae* based on the conserved location of a consensus sequence found upstream of *C. elegans* 21U-RNAs, which is also found clustered on the syntenic chromosome in *C. briggsae* [13]. However, in *C. briggsae*, the sequences downstream of this consensus sequence that likely correspond to 21U-RNAs are not conserved [13], suggesting that the sequence of individual 21U-RNAs is not critical for their function. The location of 21U-RNA loci on chromosome IV and the high density of spermatogenesis-enriched genes on the same chromosome initially suggested that 21U-RNAs might affect expression of these genes in *cis*. However, many of the spermatogenesis-enriched genes on IV were unaffected in *prg-1* mutants, including several *msp* genes. Moreover, *prg-1*-regulated genes are located on chromosomes other than IV, and even those on IV are not strongly enriched in the regions from which 21U-RNAs are produced (only about two-fold). These results suggest that PRG-1 and/or 21U-RNAs do not globally or uniformly alter the expression state of the entire chromosome. However, the fact that genes enriched during spermatogenesis are found on the same chromosome as 21U-RNAs suggests that some advantage might exist for the coordinated evolution of a subset of spermatogenesis-enriched genes and 21U-RNA genes.

A temperature-sensitive sterile phenotype, similar to that seen in *prg-1* mutants, is a common phenotype of other genes encoding P-granule-associated proteins, including PGL-1, GLH-1, MEG-1, and DEPS-1 [17, 26–28]. This commonality suggests that the underlying function of P granules is inherently sensitive to temperature. Two reports indicate that P granules might have a role in RNAi-related processes because

mutations in genes encoding the two P granule proteins PGL-1 and DEPS-1 cause defects in germline RNAi [28, 29]. P granule localization of PRG-1, combined with its requirement for 21U-RNA expression, also suggests that at least one function of P granules is small RNA-mediated regulation of gene expression. However, in contrast to RNAi-based mechanisms, PRG-1 appears to function to promote transcript accumulation rather than transcript degradation. Thus, multiple, independent small RNA pathways might converge at P granules.

The localization of PRG-1 to P granules on the outside of the nuclear envelope suggests that it does not directly function in the transcriptional activation of spermatogenesis-expressed genes or 21U-RNAs, but we cannot rule out the possibility that PRG-1 is present in the nucleus below detectable levels. Indeed, *Drosophila* PIWI is found in the nucleus, where it promotes transcriptionally permissive histone modifications as well as the expression of a nuclear piRNA that silences a master regulatory locus [3]. Alternatively, PRG-1 in P granules could play a role in 21U-RNA processing, as the PIWI domain of PRG-1 retains key residues for "slicing" activity [13]. Although 21U-RNAs appear to be individually transcribed [12], whether or where the nascent 21U-RNA transcript undergoes processing to produce a 21 nt final product is unknown. Finally, 21U-RNAs could associate with PRG-1 in an effector complex of unknown function, perhaps at P granules.

To date our experiments have established a link between the *C. elegans* Piwi, PRG-1, and a novel class of 21U-RNAs in the successful completion of spermatogenesis. Future work should address the precise role of 21U-RNAs in mediating PRG-1 function.

Experimental Procedures

Nematode Strains and Maintenance

C. elegans were grown on NGM as described in Brenner [30]. The following strains were used: as wild-type variety Bristol, strain N2; LGI, *prg-1(tm872)*; LGIII, *glp-1(e2141)*, *unc-32(e189)*; LGIV, *prg-2(tm1094)*, *fem-1(hc17)*, *fem-3(q20 gf)*, BS585. Strains were maintained at 20°C, except for the temperature-sensitive mutants *glp-1*, *fem-1*, and *fem-3*, which were maintained at 15°C. Both *prg-1(tm872)* and *prg-2(tm1094)* strains were outcrossed to N2 animals six times prior to analysis.

Brood Size and Fertility Test

To measure hermaphrodite brood size, eight to ten self-fertilizing hermaphrodites were plated and permitted to lay eggs at 20°C for 2 hr before removal. Eggs were either kept at 20°C or shifted to 25°C. The brood counts were performed by placing L4 hermaphrodites singly on plates with daily transfer to a fresh plate until egg production ceased (about 3 days); the number of progeny on each plate was counted after all worms reached the L4 stage. To measure male fertility L4 males were placed with *fem-1(hc17)* or *unc-32(q20 gf)* hermaphrodites at 3:2 ratios and the hermaphrodites were cloned to new plates after 16–24 hr. The number of cross progeny was counted after reaching adulthood. Student's t test was used to assess statistical significance.

Sperm Activation Assay

Sperm activation assays were performed as described previously [15]. Briefly, spermatids were dissected from wild-type and *prg-1(tm872)* males separated from hermaphrodites prior to sexual maturity, grown at 25°C, and analyzed in parallel. Dissections were performed in sperm medium (50 mM HEPES [pH 7.8], 50 mM NaCl, 25 mM KCl, 5 mM CaCl₂, and 1 mM MgSO₄) supplemented with 10 mg/ml PVP and 0.2mg/ml of pronase. The percentage of activated spermatids (n > 100) was determined after 20 min.

Construction of PRG-1::GFP Transgene

prg-1 genomic sequence from ATG to 215 bp downstream of the stop codon, which includes the 3'UTR, was amplified by PCR from genomic DNA with primers 5'-GGGGACAACCTTTGTACAAAAAGTTGGCTTGGCATCTGGAAGTGGTCGCGG-3' and 5'-GGGGACAACCTTTGTACAAGAAAGTTGAT

GTCCAACCTCCTTACGAAAC-3', which contain attB1 and attB2 sequences. The resulting PCR product was first cloned into pDONR201 vector (Invitrogen, Carlsbad, CA) with Gateway technology and was sequence verified before recombining into pID3.02B [31]. This construct was transformed into worms by microparticle bombardment as described previously [32]. Expression of GFP was examined by DIC and fluorescence microscopy.

Immunofluorescence

Immunostaining of extruded gonads was carried out as done in [33]. Briefly, gonads were dissected in M9 buffer, placed on slides, and immersed in liquid nitrogen before cracking off the coverslip. The slide was placed in ice-cold methanol for 20 min, followed by immersion in ice-cold acetone for 10 min. We used the following primary antibodies: goat anti-GFP (Rockland, Gilbertsville, PA) diluted 1:500 and mouse anti-PGL-1 (OIC4D [17];) diluted 1:30. Samples were examined by epifluorescence microscopy and images collected using Zeiss Axiovision.

Small RNA Northern Blot Analysis

Total RNA was isolated from L4 stage of wild-type, *prg-1*, *prg-2*, *prg-1;prg-2*, *glp-1*, *fem-1*, and *fem-3* mutant worms. This RNA was enriched for species <200 bp with the mirVana miRNA Isolation Kit (Ambion, Austin, TX) according to the manufacturer's instructions. All RNA samples were separated by electrophoresis by using 15% polyacrylamide (19:1) gels cast in 7M urea and buffered with 20 mM MOPS/NaOH (pH7). After gel electrophoresis, RNAs were transferred to Hybond-N (Amersham, Piscataway, NJ) membrane and crosslinked to membrane by carbodiimide as described previously [34]. Northern hybridization and washes were carried out as done in [35]. Probes used to detect RNA levels of *21ur-1* (5'-GCACGGTAAACGTA CGTACCA-3'), *21ur-1353* (5'-TACCATCATGTGATGTTCTGA-3'), and *let-7* (5'-ACTATACAACCTACTACCTCA-3') were made with the StarFire Oligonucleotide Labeling System (IDT, Coralville, IA).

Gonad Dissection and Microarray Analysis

Wild-type and *prg-1* mutant males were collected by bleaching mated gravid hermaphrodites to collect eggs, which were then hatched in S basal solution in the absence of food. Starved L1 larvae were cultured with food (bacterial strain OP50) at 25°C for 48 hr, then L4 males were picked to fresh plates and cultured at 25°C for another 16–24 hr before gonad dissection. Total RNA was isolated from 75 dissected male gonads as described previously [36] and amplified with MessageAmpII aRNA kit (Ambion). Amplified mRNA (2 µg) was labeled with ULS-Cy3 or ULS-Cy5 dye (Kreatech, Amsterdam) and hybridized to a *C. elegans* whole-genome DNA array from the Washington University Genome Science Center (St. Louis, MO). Three independent wild-type and *prg-1* mutant samples were collected and hybridized. The fold differences from the three experiments were averaged, and a Z test [$Z = (\text{observed} - \text{expected}) / \text{SE}$] was performed. Genes with up- or downregulation greater than 1.5-fold, $p < 0.05$, in any given mutant were selected for inclusion. Chromosome bias was assessed by using the Chi square test.

Supplemental Data

One table is available with this article online at <http://www.current-biology.com/cgi/content/full/18/12/861/DC1>.

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