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The Bantam microRNA Is Associated with Drosophila Fragile X Mental Retardation Protein and Regulates the Fate of Germline Stem Cells

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Abstract

Fragile X syndrome, a common form of inherited mental retardation, is caused by the loss of fragile X mental retardation protein (FMRP). We have previously demonstrated that dFmr1, the Drosophila ortholog of the fragile X mental retardation 1 gene, plays a role in the proper maintenance of germline stem cells in Drosophila ovary; however, the molecular mechanism behind this remains elusive. In this study, we used an immunoprecipitation assay to reveal that specific microRNAs (miRNAs), particularly the bantam miRNA (bantam), are physically associated with dFmrp in ovary. We show that, like dFmr1, bantam is not only required for repressing primordial germ cell differentiation, it also functions as an extrinsic factor for germline stem cell maintenance. Furthermore, we find that bantam genetically interacts with dFmr1 to regulate the fate of germline stem cells. Collectively, our results support the notion that the FMRP-mediated translation pathway functions through specific miRNAs to control stem cell regulation.


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Introduction

Stem cells, which can self-renew and produce different cell types, are known to be regulated by both extrinsic signals and intrinsic factors [1]. In Drosophila ovary, a very small population of germline stem cells (GSCs) is maintained in a well-defined microenvironment, which provides an attractive system for investigating the regulatory mechanisms that determine the fate of stem cells [2,3]. Studies from multiple laboratories have identified the genes that are essential for GSC fate determination [4,5]. Recently, the microRNA (miRNA) pathway was also found to be required for controlling GSC self-renewal, since mutations in Dicer-1, Ago1, and loquacious, which are involved in miRNA production and function in Drosophila, lead to rapid loss of GSCs [6,7,8,9]. MiRNAs could regulate gene expression through translational repression and mRNA degradation by binding to the 3’ untranslated region (UTR) of their target miRNAs [10]. However, the specific miRNAs required for the regulation of GSC self-renewal and fate specification are yet to be determined.

Fragile X syndrome, the most common cause of inherited mental retardation, is caused by the loss of functional FMRP [11]. FMRP is an RNA-binding protein and is known to bind to specific mRNAs and regulate their translation both in vitro and in vivo [12]. FMRP is largely cytoplasmic, incorporated into large messenger ribonucleoprotein (mRNP) particles [12]. A growing body of work from several groups now suggests that the microRNA pathway is the major molecular mechanism by which FMRP regulates translation. In Drosophila and mammals, FMRP, as well as its autosomal homologs in mammals, FXR1P and FXR2P, is found to be a part of the RNA-induced silencing complex (RISC) [13,14,15,16]. However, what role if any FMRP plays in RNA interference is unclear. In the miRNA pathway, FMRP is associated with miRNAs in both Drosophila and mammals, and the genetic interaction between dFmr1 and the miRNA pathway has been demonstrated in Drosophila [13,17]. Therefore, FMRP is one component of the miRNA pathway involved in miRNA-mediated translational control. Recently, we also showed that dFmr1 is required for both GSC maintenance and repressing differentiation [17]. Furthermore, we demonstrated that in Drosophila ovary, dFmr1 protein (dFmrp) interacts with Argonaute protein 1 (AGO1), a key component of the miRNA pathway. Hence dFmr1 could modulate the fate of GSCs, likely via the miRNA pathway. Nevertheless, whether dFmrp could use specific miRNAs to regulate the fate of GSCs has remained unclear.

Here we show that dFmrp is associated with specific miRNAs, including the bantam miRNA, in Drosophila ovary. Like dFmr1, the bantam miRNA is not only required for repressing primordial germ cells (PGCs), but also functions as an extrinsic factor for GSC maintenance. Furthermore, we show that bantam genetically
**Author Summary**

Fragile X syndrome, the most common cause of inherited mental retardation, results from the loss of functional Fragile X mental retardation protein (FMRP). FMRP is an RNA-binding protein and is known to bind to specific mRNAs and regulate their translation both in vitro and in vivo. Previous studies have suggested that FMRP is one component of the miRNA pathway involved in miRNA-mediated translational control. In the past we found that, as a translational regulator, dFmrp can modulate the proliferation and fate specification of stem cells in Drosophila, likely via the miRNA pathway. However, whether dFmrp could use specific miRNAs to regulate the fate of GSCs has remained unclear. Here, we show that dFmrp is associated with specific microRNAs (miRNAs), including the bantam miRNA, in Drosophila ovary. Furthermore, we show that bantam genetically interacts with dFmrp to regulate the fate of GSCs. Our findings imply that FMRP could utilize specific miRNAs to regulate the translation of its mRNA targets.

interacts with dFmr1 to regulate the fate of GSCs. These results support the notion that FMRP-mediated translational control functions through specific miRNAs to control stem cell behavior.

**Results**

Identification of Specific miRNAs Associated with dFmrp in Drosophila Ovary

Given that dFmr1 plays an important role in the fate determination of germ cells, and that dFmrp physically associates with dAGO1, a key component in the miRNA pathway, we proposed that dFmr1 regulates the fate of GSCs through the miRNA pathway in Drosophila ovary [17]. To test this, we first determined whether dFmrp is associated with the endogenous miRNAs. We used the previously developed anti-dFmrp antibody to perform immunoprecipitation from the lysates of both wild-type (WT, w1118) and dfmr1 mutant ovaries (Figure 1A) [18]. RNAs from both the immunoprecipitated (IP) complex and input were isolated for RNA TaqMan assays. We used TaqMan assays available from ABI that could detect a total of 72 known individual miRNAs. To identify the miRNAs specifically associated with dFmrp, we determined the level of each miRNA in both IP and input RNAs from WT and dfmr1 mutant flies. We identified the miRNAs that were consistently enriched by more than two-fold in WT-IP over both WT-Input and dfmr1-IP in two independent experiments, since such miRNAs are likely to be the ones specifically associated with dFmrp based on previous IP experiments (Figure 1B) [19]. Among the 72 miRNAs examined here, only selective miRNAs were enriched in WT-IP compared with both WT-Input and dfmr1-IP, suggesting that dFmrp is indeed associated with specific miRNAs, but not with endogenous miRNAs in general (Figure 1B). Since the bantam miRNA is among those miRNAs specifically associated with dFmrp in ovary, we further confirmed their specific association by quantitative RT-PCR and Northern blot (Figure 1C and 1D). The rest of this study focuses on the role of the bantam miRNA and its potential interaction with dFmr1 in regulating GSCs.

Generation of New Alleles for bantam

To determine whether dFmr1 regulates germline development through the bantam miRNA, we first explored whether the bantam miRNA plays similar roles to dFmr1 in repressing primordial germ cells (PGCs) and GSC differentiation during the larval and adult stages. Since the hypomorphic allele of bantam (ban), banF{P}[22], was fertile and exhibited no apparent defects in germ cells, including PGCs and GSCs, we performed a mutagenesis through imprecise mobilization of the P-element from banF{P}[22] to generate stronger alleles for the ban gene. From 50 imprecise excision lines, we isolated one sterile line, ban20, and one lethal line, ban12. Both of these alleles failed to complement the deficiency line of Df(3L)Fmceu-E12, which deletes the chromosomal segment that covers the ban locus. The breakpoints of these two mutant lines were determined by PCR and DNA sequencing (Figure 2). Furthermore, we detected no mature bantam miRNA by miRNA TaqMan assay in these mutant lines (data not shown).

The Bantam miRNA Is Required for Repressing PGC Differentiation

Our previous study demonstrated that dFmr1 plays a role in repressing PGC differentiation [17]. To determine whether ban

The Bantam miRNA Is Required for GSC Maintenance

A typical Drosophila ovary is composed of 16-20 distinct units known as ovarioles. Each ovariole consists of an anterior functional unit, called a germinarium, and a linear string of differentiated egg chambers posterior to the germinarium. At the tip of the germinarium, GSCs normally divide asymmetrically to ensure that one daughter cell remains attached to the niche cells for self-renewal, while the other is displaced from the niche, becoming a cystoblast (CB) that initiates differentiation and sustains oogenesis (Figure 3A) [20]. We have previously shown that dFmr1 is involved in the regulation of GSC fate [17]. To compare the biological effects of ban with dFmr1 in GSCs, we examined whether ban is also required for GSC maintenance by quantifying the number of GSCs in ban mutant germa...
plays a similar role in PGC development, we examined the behavior of PGCs in ban mutants. PGCs are the precursors of germ cells during the three larval stages; the number of PGCs simply grows from 12 cells to more than 100 cells, but without further differentiation before the pupa stage. Using anti-Vasa and anti-Hts antibodies as previously described, we stained the third instar gonads of both WT and ban mutants to visualize germ cells and fusomes, respectively [21]. As shown in Figure 4A, in WT gonads from late third instar larvae, ~80% PGCs carried a single spherical fusome, and the other PGCs were dividing with two spherical fusomes associated between two PGC cells. By contrast, in the ban mutant, most gonads (84%, n = 25) from the third instar larva contained differentiated PGC clusters that were marked by branched fusomes (Figure 4B and 4C), reminiscent of other allelic combinations of ban (ban12/Df(3L)emc-E12) (75%, n = 12). Together, these results suggest that, like dfmr1, ban plays a similar role in repressing PGC differentiation.

The Bantam miRNA Plays a Non-Autonomous Role in GSC Maintenance

The loss of germline stem cells in ban mutant ovaries indicates that ban is required by either GSCs or somatic cells. To analyze
whether ban functions as a cell-autonomous factor in maintaining GSC fate, we used a FLP-FRT-mediated mitotic recombination technique to generate marked mutant GSCs [17,22], then calculated the life span of the marked mutant GSCs by quantifying their loss rate. The marked mutant GSCs were identified by a lack of GFP fluorescence in the nuclei and by their positions directly attaching to the base cells of the terminal filament or cap cells. The ban loss-of-function alleles (ban20 and ban21) were used to generate marked mutant GSC clones for an analysis of ban function in GSCs. The rates of GFP-marked GSCs were measured at two days, seven days, 12 days, and 15 days after heat-shock treatment (AHT). As shown in Figure 5E, compared with FRT control GSC clones, the marked clone rates of both ban20 and ban21 were not reduced significantly during the testing period, indicating that ban is dispensable for GSCs. Together with the phenotypic analyses of ban, our data indicate that, as with dfmr1, ban functions as an extrinsic factor for GSC maintenance (Figure 5).

**Bantam Genetically Interacts with dfmr1**

Given that ban plays a similar role to dfmr1 in the regulation of GSC fate and that Ago1 genetically enhances the phenotype of dfmr1 in the maintenance of GSCs [17,22], we next investigated whether ban genetically interacts with dfmr1. Interestingly, we found that compared with the control flies, double heterozygotes of dfmr1 and ban displayed significantly reduced fertility in female flies (Table 1), which suggests that, besides the physical association, ban also genetically interacts with dfmr1 in Drosophila oogenesis. To test whether the partial loss of both the ban miRNA and dfmr1 could affect the normal maintenance of GSCs, we further quantified the number of GSCs in the single or double heterozygotes of ban and dfmr1 mutants at days two and 12 after eclosion. As shown in Figure 6, the double heterozygotes displayed a significant increase in the number of germaria with one or no GSCs, suggesting that a reduction of both ban and dfmr1 leads to greater loss of GSCs. In both experiments, we found that the balancer chromosomes (TM6) did not cause any phenotype while they are in trans to either ban or dfmr1 null allele (data not shown). These data suggest that dfmr1 could potentially regulate the fate of GSCs through the bantam miRNA.

**Discussion**

Fragile X syndrome, the most common cause of inherited mental retardation, is due to the loss of functional FMRP [11]. As an RNA-binding protein, FMRP is known to regulate the translation of specific mRNAs [23]. Recent work has suggested that FMRP could regulate translation via the miRNA pathway [24,25]; however, the experimental evidence to elucidate the specific role, if any, FMRP plays in small RNA-mediated post-transcriptional gene regulation is still lacking, and whether specific miRNAs are used in FMRP-mediated translational control remains to be determined. In this study, we found that dfmrp is associated with specific miRNAs, including the bantam miRNA, in Drosophila ovary. Interestingly, like dfmr1, ban also functions as an extrinsic factor in the maintenance of GSCs. Even more importantly, though, we show that bantam genetically interacts with dfmr1 to regulate the fate of GSCs. These data together suggest that dfmrp can use specific miRNA(s) to regulate the behavior of stem cells. These findings also implicate that FMRP could be associated with selective miRNAs, and utilize specific miRNA(s) to regulate the translation of specific mRNA targets.

Studies of Drosophila germline stem cells have yielded heuristic examples to help us understand the molecular regulatory mechanism of stem cells. Previous investigations have demonstrated that the self-renewal of GSCs requires both extrinsic and intrinsic mechanisms. Gene products, such as Piwi, Yb, Dpp, and Gbb, are produced from niche cells and function as extrinsic factors for GSC maintenance. On the other hand, intrinsic factors, including Nanos (Nos), Pumilio (Pum), the Dpp-dependent receptors, and transcription factors, are also important for GSC self-renewal [2,4]. In Drosophila, dfmr1 is known to be required for cyst formation and oocyte specification, potentially via the regulation of Orb mRNA translation [26]. Our recent study demonstrated that dfmr1 is involved in modulating the fate of GSCs [17]. The phenotypic assay of dfmr1 mutants revealed that the loss of dfmr1 function leads to a defect in the maintenance of GSCs; however, clonal analyses of dfmr1 via FLP/FRT-mediated recombination demonstrated that dfmr1 is dispensable in GSC fate regulation, suggesting dfmr1 functions as an extrinsic factor. In this paper, we found that the bantam miRNA is associated with dfmrp
Figure 3. The bantam miRNA is required for GSC maintenance. A. A schematic diagram of a *Drosophila* gerarium with different cell types labeled by different colors: GSCs, cystoblast (CB) and cysts, spectrosomes (SS), terminal filaments (TF), cap cells (CPC), inner gerarium sheath cells (IGC), follicle cells (FC), SSC (somatic stem cells) and fusomes. B–G: Ovaries from wild-type (B), *ban* mutant flies at different ages (C–F), and *ban* mutants carrying transgene *P*{banP-ban} (G) were stained with anti-Vasa (Green) and anti-Hts (Red) antibodies. H. Quantitative analyses of the number of GSCs in *ban* mutant; the x-axis shows the day of examination post-eclosion, while the y-axis shows the average number of GSCs per germarium in *ban* mutants and WT. P<0.001 when *ban* mutant was compared with WT at different time points. Arrows indicate GSCs. doi:10.1371/journal.pgen.1000444.g003
in Drosophila ovary. To determine the role of the bantam miRNA, we generated null alleles of \textit{ban} and examined the role of the bantam miRNA in the maintenance and fate specification of GSCs. Interestingly, we found that \textit{ban} plays a similar role to \textit{dFmr1} in the regulation of GSC fate. More importantly, the double heterozygotes of \textit{ban} and \textit{dFmr1} mutants displayed a greatly reduced number of GSCs per germarium and lower fertility in general, suggesting that \textit{ban} strongly interacts with \textit{dFmr1} genetically in regulating the maintenance and fate specification of GSCs. Furthermore, the transgene \textit{P\{banP-ban\}} could not rescue the GSC phenotype associated with the loss of \textit{dFmr1} (data not shown). The role of \textit{ban} in GSC maintenance and its genetic interaction with \textit{dFmr1} appear specific since another miRNA that is associated with dFmrp, miR-1, has no effect on GSCs (Chen and Jin, unpublished data). Although it is possible that \textit{ban} could potentially function upstream of \textit{dFmr1}, however, given the strong genetic interaction that we observed, it is more likely that \textit{ban} functions in concert with \textit{dFmr1} to repress GSC differentiation by

**Figure 4. The bantam miRNA is required for repressing primordial germ cell (PGC) differentiation.** Gonads from the third-instar larvae of \textit{w\textsuperscript{1118}} (A) and \textit{ban\textsuperscript{12}} mutants (B, C, and C inset) were stained with anti-Vasa and anti-Hts antibodies. Anti-Hts (Red) was used to outline gonads and morphology of fusomes, while anti-Vasa (green) was used to visualize all germ cells. PGCs carrying a single round fusome are indicated by arrows, while differentiated germ cells are indicated by arrowheads. Scale bar represents 10 \(\mu\)m.

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repressing the translation of common target(s) in the dFmr1-dependent pathway.

The role of the miRNA pathway in GSC fate specification has already been explored. Previous work including our own has demonstrated that Dicer1, Loqs, and Ago1, key components of the miRNA pathway, play intrinsic and essential roles in the maintenance of GSCs [6,7,8,9]. In this study, we chose to focus on a specific miRNA, the bantam miRNA. We found that, besides the intrinsic role of the miRNA pathway in general, specific miRNAs, such as the bantam miRNA, could also function as a niche factor to regulate GSCs. Given our findings, the next important step becomes identifying the mRNA target(s) of the bantam miRNA that could regulate GSCs.

In summary, here we show that, as an extrinsic factor regulating GSCs, dFmrp is selectively associated with specific miRNAs, including the bantam miRNA, in Drosophila ovary. The strong genetic interaction between ban and dFmr1 in the regulation of GSCs suggests that dFmrp could use the bantam miRNA to

**Table 1. Relative fertility of different mutant flies.**

<table>
<thead>
<tr>
<th>Genotype</th>
<th>Percentage of fertility</th>
</tr>
</thead>
<tbody>
<tr>
<td>w¹¹¹⁸</td>
<td>98.9% (87/88)</td>
</tr>
<tr>
<td>ban¹²/TM6</td>
<td>97.5% (78/80)</td>
</tr>
<tr>
<td>dfmr1¹⁴¹⁰/TM6</td>
<td>98% (49/50)</td>
</tr>
<tr>
<td>dfmr1¹¹¹³/TM6</td>
<td>97.8% (45/46)</td>
</tr>
<tr>
<td>ban¹²/dfmr1²⁵₁²</td>
<td>7.7% (5/65)*</td>
</tr>
<tr>
<td>ban¹²/dfmr1¹¹¹³</td>
<td>4.6% (3/65)*</td>
</tr>
</tbody>
</table>

Individual female was crossed with wildtype (w¹¹¹⁸) male flies and the number of fertile eggs was quantified. *P<0.01 when the trans-heterozygotes were compared with either WT or single heterozygotes. doi:10.1371/journal.pgen.1000444.t001

Figure 5. The bantam miRNA plays a non-autonomous role in GSC maintenance. GSC clones were induced by heat-shock treatment in adult female flies. Ovaries from FRT control flies (A and B) and FRT, ban flies (C and D) were dissected at day 2 and day 12 following heat-shock treatment; GSC clones were identified by the lack of GFP expression. Scale bar represents 10 μm. (E) Relative percentages of negatively GFP-marked GSC clones in FRT control and two FRT; ban null alleles at days 2, 7, 12, and 15 AHST are shown. P>0.5. doi:10.1371/journal.pgen.1000444.g005
regulate the translation of specific mRNAs in ovary, which in turn modulates the behavior of GSCs. Identifying those mRNAs that are co-regulated by dFmrp and the bantam miRNA in Drosophila ovary will provide an important system for further study of the role(s) that dFmrp plays in miRNA-mediated translational control.

Materials and Methods

Drosophila Genetics

All flies were maintained under standard culture conditions. dfmr113, dfmr1130n113 and dfmr1130n113 were null dfmr1 alleles as described previously [18,27]. banEP3622 is a P-element insertion line obtained from the Szeged stock center. The deficiency line Df(3L)emc-E12, containing the deletion of the chromosomal segment covering the ban locus came from Bloomington Stock center. Two ban alleles, ban12 and ban20, were generated by imprecise excision from banEP3622. ban12 carried an ~6-kb deletion including the ban core sequence. ban20 is a hypomorphic allele that carried a ~21-kb deletion (~24.2 kb to ~2.7 kb) upstream of the ban gene. The transgene P(banP-ban) with 7.2-kb genomic rescue fragment, in which about 500 bp ban primary transcript was under the control of the 6.7-kb ban promoter (3.5-kb upstream and 3.2-kb downstream), was used to rescue the ban mutant phenotype. The following primers were used to map the breakpoints of ban12 and ban20 alleles.

Forward primers:
EP-6K: gtagcttgca gtgggcttac atg  
EP-7.5K: cggagtactg tcctccgggc tgg  
ban-24.5K: tcattgaccaaatcccaacgcaag  
ban-4K: attccagaaattcttgcg

Reverse primers:

Figure 6. Bantam genetically interacts with dfmr1 in modulating GSCs. (A) The number of GSCs per germarium was measured from dfmr1 heterozygotes, ban heterozygotes, and dfmr1, ban double heterozygotes at day 2 and day 12 post-eclosion, respectively. P<0.01 when the percentage of type II and III germaria of the trans-heterozygotes were compared with those of either WT or single heterozygotes. (B) Ovaries from the genotypes above were stained with anti-Vasa (Green) and anti-Hts (Red); germaria carrying two GSCs, one GSC, and no GSCs were referred to as type I, II, III-a and III-b, respectively.

doi:10.1371/journal.pgen.1000444.g006
EP: ccaaccttagttcattcag
ban-hoop-2K: agtttagctegagctag
ban-hoop-4.2K: ccgctagctgaagacagtcccagacat
ban-hoop-3K: gtgttagtaaactgac

Immunohistochemistry and Microscopy
Ovaries were prepared for reaction with antibodies as described previously [17]. The anti-Vasa antibody (Santa Cruz) was used at a 1:200 dilution, and monoclonal anti-His antibody was used at a 1:500 dilution. Secondary antibodies were goat anti-mouse Alexa 568, goat anti-rabbit Alexa 488, and goat anti-rat Cy3 (Molecular Probes), all at 1:200. All samples were examined by Zeiss Microscope, and images were captured using the Zeiss Two Photon Confocal LSM510 META system. Images were further processed with Adobe Photoshop 6.0.

Phenotypic Assay for Quantification of GSC Maintenance in Mutant Adult Ovaries
Ovaries isolated from dfmr1 and homozygous mutant flies of different ages were incubated with anti-His antibody and anti-Vasa antibody to identify terminal filament cells, fusomes, and germ cells. We scored as GSCs any Vasa-positive germ cells at the anterior position that appeared close to cap cells or to the basal cells of terminal filaments and also carried spherical fusomes at the anterior position or extending fusomes.

Germline Clonal Analyses
FLP/FRT-mediated recombination was used to generate dfmr1 mutant GSC and PGC clones. To generate GSC clones, 3-day-old females, hs-flp/FRT79D, ubi-gfp/FRT79D, ban, underwent heat-shock treatment at 37°C for 60 min twice daily at 12-h intervals (hs-flp/FRT79D, ubi-gfp/FRT79D as control). After 3–6 days of heat-shock induction monitored by GSC clone efficiency of control, ovaries were dissected for quantification of GSC clones at day 2, day 7, day 12 and day 15 of the post-clonal induction. The % of GSC clones measured at day 2 was also calculated as the initial rate (100 of relative %). GSC clones were identified by a lack of GFP fluorescence in the nucleus and by the carrying of an anterior-positioned dot fusome (spectrosome).

Immunoprecipitation of dfmrp from Fly Ovary
About 150 adult WT and dfmr1 mutant fly ovaries were dissected in PBS and homogenized in 1 ml of ice-cold lysis buffer (10 mM Tris, pH 7.4, 150 mM NaCl, 30 mM EDTA, 0.5% Triton X-100) with 2× complete protease inhibitors. All further manipulations of the ovary lysates were performed at 4°C or on ice. Nuclei and debris were pelleted at 3,000 xg for 10 min; the supernatant was collected and raised to 300 mM NaCl and clarified at 14,000 xg for 30 min. The resulting supernatant was preclarified for 1 h with 100 µl recombinant protein G agarose (Invitrogen) (washed with lysis buffer first). An aliquot of preclarified input was saved for RNA extraction (200 µl) and protein analysis (100 µl). We incubated 15 µg of Monoclonal Anti-dFMR1 Clone 6A15 (Sigma) with recombinant protein G agarose at 4°C for 2 h and washed 3 times with lysis buffer. RNase Inhibitors (Invitrogen) were added to the remaining lysates. The preclarified lysates were immunoprecipitated with antibody-coated recombinant protein G agarose at 4°C overnight. After a third wash with the lysis buffer, 10% of the immunoprecipitate was saved for protein analysis. The remainder was washed one more time, and the immunoprecipitate was resuspended into Trizol (Invitrogen) for RNA isolation or kept at −80°C until further processing. Western blot analysis was performed using anti-dFmrp antibody 5A11 (Developmental Studies Hybridoma Bank at the University of Iowa) and detected with horseradish peroxidase-coupled secondary antibody (Sigma) using the chemiluminescent (GE) method.

miRNA TaqMan Assay
Total RNA was isolated from the WT and dfmr1 mutant ovari IP as well as Input using Trizol. TaqMan MicroRNA Assays detecting 72 known individual Drosophila miRNAs were obtained from ABI (ABI). cDNA was prepared with a High-Capacity cDNA Archive Kit (ABI); 10× reaction buffer 0.5 µl; dNTP mix (100 mM) 0.05 µl; Recombinant RNase inhibitor 0.07 µl; MLV reverse transcriptase 0.33 µl; RT TaqMan assays primer (5×) 1 µl; RNA sample 1 µl; and nuclelease-free water to 5 µl. This was performed at 16°C for 30 min and at 42°C for 30 min, terminated at 85°C for 5 min and 4°C forever. cDNA was prepared for the endogenous control Dm_Rpl32.1_Q by QuantiTect Primer Assay (Qiagen) according to the manufacturer’s instructions. Real-time PCR was conducted with 2× SYBR Green Master Mix (ABI) as amplification of RPL32. For Real-time PCR of TaqMan MicroRNA Assays, we used TaqMan MicroRNA Assay Primer (20×) 0.5 µL; cDNA (undiluted) 1.33 µl; TaqMan 2×Universal PCR Master Mix, 5 µl; nuclelease-free water 3.17 µl. Real-time PCR reactions performed in triplicate with MicroAmp optical 96-well plates using 7500 Fast Real-Time PCR System (ABI) with the following conditions; an initial step of 10 min at 95°C, followed by 40 cycles of 15 s at 95°C, 1 min at 60°C.

Northern Blot of miRNAs
RNAs were separated on 15% TBE urea gel, transferred, and UV-cross-linked to nylon membrane (Osmonics, Inc.). 32P-labeled probes were prepared with the Ambion mirVana miRNA Probe Construction Kit. Membranes were prehybridized at 65°C for 1 h and hybridized for 12–16 h at room temperature (RT). Membranes were then washed 3 times at RT and 2 times at 42°C. Membranes were exposed and scanned with a Typhoon 9200 PhosphorImager (Amersham Biosciences).

Statistical Method
We performed post-hoc t-tests (two-samples assuming equal variances) to determine significance, and indicated P values.

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Author Contributions
Conceived and designed the experiments: PJ DC. Performed the experiments: YY SX LX JW SW. Analyzed the data: YY SX LX JW SW. Contributed reagents/materials/analysis tools: PJ DC. Wrote the paper: PJ DC.

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