MILI, a PIWI-interacting RNA-binding Protein, Is Required for Germ Line Stem Cell Self-renewal and Appears to Positively Regulate Translation*§

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The Argonaute/PIWI protein family consists of Argonaute and PIWI subfamilies. Argonautes function in RNA interference and micro-RNA pathways; whereas PIWIs bind to PIWI-interacting RNAs and regulate germ line development, stem cell maintenance, epigenetic regulation, and transposition. However, the role of PIWIs in mammalian stem cells has not been demonstrated, and molecular mechanisms mediated by PIWIs remain elusive. Here we show that MILI, a murine PIWI protein, is expressed in the cytoplasm of testicular germ line stem cells, spermatogonia, and early spermatocytes, where it is enriched in chromatoid bodies. MILI is essential for the self-renewing division and differentiation of germ line stem cells but does not affect initial establishment of the germ line stem cell population at 7 days postpartum. Furthermore, MILI forms a stable RNA-independent complex with eIF3a and associates with the eIF4E- and eIF4G-containing m7G cap-binding complex. In isolated 7 days postpartum seminiferous tubules containing mostly germ line stem cells, the mili mutation has no effect on the cellular mRNA level yet significantly reduces the rate of protein synthesis. These observations indicate that MILI may positively regulate translation and that such regulation is required for germ line stem cell self-renewal.

Stem cells are defined by their dual ability to self-renew and to produce numerous differentiated daughter cells. Among adult tissue stem cells, germ line stem cells in mammalian testes serve as the source of spermatogenesis, a robust stem cell-driven process that generates millions of sperm hourly (1). In the testis, germ line stem cells can be identified as a subpopulation of spermatogonia that reside in the basal layer of the seminiferous epithelium. These stem cells divide in a self-renewing fashion to produce differentiating spermatogonia, which then further divide and differentiate into numerous spermatocytes. Spermatocytes undergo meiosis to produce haploid round spermatids, which then undertake drastic morphological changes via a process called spermiogenesis to become motile sperm. Germ line stem cells are not only of paramount importance in reproduction but also are effective models for studying molecular mechanisms that define the fate and behavior of stem cells in general.

Germ line stem cell division and differentiation are regulated by both intercellular signaling and cell-autonomous mechanisms (1). The requirement of intercellular signaling is exemplified by mutations in c-Kit receptors or its ligand, stem cell factor, which causes spermatogonial arrest prior to the differentiating (A1) spermatogonia stage (2, 3). Intercellular signaling also involves dose-sensitive regulation by Glial cell line-derived neurotrophic factor, since mice heterozygous for its gene (Gdnf) display aberrant germ line stem cell division (4). Perhaps the best illustrated cell-autonomous requirements for germ line stem cell maintenance are Pten and Sox3 genes, both of which encodes transcriptional factors (5–7).

The cell-autonomous mechanism for mammalian germ line stem cell self-renewal probably involves translational regulation, as inferred from studies of germ line stem cells in lower model systems and spermatogenesis in mammals. In Drosophila, translational repression mediated by Pumilio and Nanos is crucial for maintaining the undifferentiated states of germ line stem cells and their precursors, primordial germ cells (8–10). In Caenorhabditis elegans, ATX-2, a homolog of mammalian ATAXIN 2, binds to cytoplasmic poly(A)-binding protein and is required for translational regulation of germ line stem cell proliferation (11). In mice, translational regulation has been shown to be crucial for spermiogenesis (12), although its role in germ line stem cells is not yet determined. However, studies of a mutation that affects germ line stem cell maintenance, Jsd (juvenile spermatogonial depletions), hint at a role for translation in mammalian germ line stem cell self-renewal. Jsd mutant mice can only undergo one round of spermatogenesis, followed by early (A1) spermatogonia arrest (13). Molecular analysis indicates that the Jsd mutation affects mItp14b, a testis-spe-
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specific retroposed copy of the ubiquitously expressed gene mUtp14a (14). This gene is homologous to the yeast UTP14 gene, which is an essential component of a large ribonucleoprotein complex containing the U3 small nuclear RNA (15). Deletion of UTP proteins in yeast inhibits 18 S rRNA production, indicating that they are part of the active pre-rRNA-processing complex. Similarly, mice lacking Dazl (deleted in azoospermia-like) contain no differentiated spermatogonia (16), and Dazl has been implicated in translational regulation of meiotic genes but not germ line stem cells (17, 18).

Translational regulation of germ line stem cells may involve the PIWI/ARGONAUTE (AGO) protein family, in addition to their known role in transposition repression and epigenetic silencing (19, 20). The PIWI/AGO family was first discovered to be required for stem cell maintenance in diverse organisms, such as Drosophila melanogaster, C. elegans, and Arabidopsis (21). This stem cell function has also been demonstrated in planaria (22). Recently, the role of piwi genes in C. elegans spermatogenesis has been further defined (23, 24). The PIWI/AGO protein family consists of AGO and PIWI subfamilies. AGO proteins bind to ~21-nucleotide small interfering RNAs and micro-RNAs (miRNAs). They function in RNA interference and miRNA pathways as essential components of the RNA-induced silencing complex (RISC) and RNA-induced silencing of transcription (RITS) complex to negatively regulate gene expression at the posttranscriptional and transcriptional level, respectively. Recently, mammalian AGO2 has been shown to up-regulate translation via the AU-rich elements (ARE) in mRNA 3′-untranslated regions (25). In contrast, PIWI proteins bind to 24–31-nucleotide PIWI-interacting RNAs (piRNAs) and regulate germ line development, stem cell maintenance, epigenetic regulation, and transposition (26–34). Two PIWI proteins in Drosophila, AUBERGINE and PIWI itself, have been genetically implicated in translational regulation (35–37). In addition, MIWI, a mouse homolog of PIWI, binds to piRNAs as well as to the mRNA cap and polysomes, with the polysome association correlating to the translational activity during spermatogenesis (26, 38).

MILI is a murine PIWI protein that we previously reported to be essential for the early prophase of meiosis (39). Most recently, MILI has also been shown to bind to piRNAs (27) and been implicated in transposon control and DNA methylation (28, 40). Here, we show that MILI is a cytoplasmic protein specifically expressed in the germ line during early spermatogenesis. Deleting the mili gene severely affects the self-renewing division and differentiation of germ line stem cells, in addition to its meiotic defects. Furthermore, we provide evidence that MILI positively regulates translation in prepubertal testes that contain only stem cells in the germ line, in contrast to the negative role of AGO proteins and their associated small interfering RNAs and miRNAs. These results suggest a function of MILI in promoting germ line stem cell division and differentiation via translational regulation.

EXPERIMENTAL PROCEDURES

Antibody Generation—The MILI antibody for immunoprecipitation was generated by Anaspec Inc. (San Jose, CA). Amino acid residues 108–122 of MILI were synthesized, purified, and conjugated to keyhole limpet hemocyanin for immunization in rabbits. The resulting antisera were affinity-purified using MILI peptide conjugated to activated immunoaffinity support (Affigel 15; Bio-Rad). The purified antibody was bound to protein A-Sepharose beads (Sigma) for immunoprecipitation. A different antibody, DU-PG15, generated in a guinea pig against a 483-amino acid peptide (52.8 kDa) from bp 1634 to 2117 of the Mili open reading frame, was used for immunostaining and Western blotting.

Immunoprecipitation and Protein Identification—Testicular lysates were prepared as follows. Mice were asphyxiated with CO₂, and testes were dissected. The testes were then decapsulated and homogenized in 1 ml of lysis buffer (100 mM KOAc, 0.1% Triton X-100, 50 mM HEPES, pH 7.4, 2 mM MgOAc, 10% glycerol, 1 mM dithiothreitol, 20 units/ml RNaseOut (Invitrogen), 1× complete mini EDTA-free protease inhibitor mixture (Roche Applied Science)) in a Dounce homogenizer for 20 strokes. The lysate was adjusted with lysis buffer to 10 mg/ml. Antibody-bound protein A-Sepharose (washed with lysis buffer minus RNaseOUT and protease inhibitor) was added to the adjusted lysate. The mixture was tumbled end over end for 2 h at 4 °C. The beads were then washed, and protein samples were eluted by boiling in 2× SDS sample buffer for 5 min. For the blocking experiment, MILI peptide was added to the beads at a final concentration of 100 μg/ml and incubated for 1 h at 4 °C prior to the addition of lysates. RNase A (Sigma) was added to a final concentration of 10 μg/ml prior to a 2-h, 4 °C incubation. The RNase A-treated sample and other samples were tumbled end over end for an additional 10 min at room temperature to ensure RNase activity. Co-immunoprecipitated MILI-interacting proteins were identified by mass spectrometry at the Proteomics Center at Duke University Medical Center.

RNA in situ Hybridization and Immunofluorescence Microscopy—RNA in situ hybridization and immunofluorescence microscopy was performed as described by Deng and Lin (41). Affinity-purified polyclonal guinea pig MILI antibody (GP15 against amino acid residues 483–644) was used at 1:15 dilution. Plzf mouse monoclonal antibody from Calbiochem was used at a 1:40 dilution. Human anti-GE-1 serum IC6 from Ref. 42 was used at a 1:200 dilution to recognize GE-1 in cells.

The m7GTP Cap Binding Assay—Testicular extract was prepared as described above and incubated for 2 h at 4 °C with either 60 μl of protein A-Sepharose beads (Sigma) or m7GTP Sepharose beads (Amersham Biosciences) in lysis buffer. The beads were then washed, and protein samples were eluted by boiling in 2× SDS sample buffer for 5 min. For the competition experiment, m7GTP was added to the testicular extract to a final concentration of 200 μg/ml and rotated end over end at 4 °C for 1 h prior to incubation with m7GTP-Sepharose.

RNA Isolation and Labeling—RNAs were isolated from the bound fraction of immunoprecipitation using TRizol reagent (Invitrogen) according to the manufacturer’s instructions. For small RNA labeling, one-half of the RNA co-immunoprecipi-
tated with MILI from 6 mg of testicular extract was dephosphorylated using calf intestinal phosphatase (New England Biolabs) at 10 units/30-μl reaction in 1× calf intestinal phosphatase reaction buffer at 37 °C for 2 h. The dephosphorylated RNA was then precipitated overnight at −20 °C by ethanol and NaAc, supplemented by Glyco Blue glycogen (Invitrogen). End-labeling of small RNAs was performed using polynucleotide kinase (New England Biolabs), at 10 units/30-μl reaction, using 2 μl of γ-ATP (6000 Ci/mmol). The reaction was performed in 1× T4PNK buffer at 37 °C for 2 h and terminated by adding 1 μl of 0.5 M EDTA. The sample was then put through a Sephadex G-50 RNA purification column (Roche Applied Science) to remove the unincorporated radioactive nucleotides. Formamide sample buffer was added to one-half of the labeled RNA sample, which was resolved on 15% urea polyacrylamide gel.

**Microarray Analysis of the Levels of Total and MILI-associated mRNAs in Wild Type and MILI−/− Mutant**—Total RNA from groups of 9-dpp milii+/− or milii−/− testes was isolated using TRIzol reagent (Invitrogen) without any nucleic acid carrier. The high molecular weight RNAs were then further enriched against the low molecular weight RNAs (<200 nucleotides) by selective precipitation in 12.5% polyethylene glycol 8000 and 1.25 M NaCl. The samples were treated with RQ1 RNase-free DNase (Promega) to remove any contaminating DNA and cleaned with RNeasy columns (Qiagen) according to the manufacturer’s instructions. The quantity and quality of the RNA samples were evaluated by both spectrophotometry and gel electrophoresis. For the microarray analysis, 10 μg of each sample was reverse transcribed with oligo(dT)20 primers and SuperscriptII reverse transcriptase (Invitrogen). cDNA samples were labeled with Cy5-conjugated nucleotides during the reverse transcription and directly hybridized to a mouse AROS version 3.0 oligonucleotide array (Operon) representing gene transcripts in the mouse. Cy5-labeled oligonucleotides complementary to the probes on the array were externally included in the sample during hybridization as a reference set for initial normalization of the raw data. Likewise, MILI-associated mRNAs were isolated using the TRIzol reagent from the anti-MILI immunoprecipitates from adult (2–4 months old) wild type testicular extract. After one round of amplification with the MessageAmp II RNA amplification kit (Ambion Inc.), the cDNAs were prepared, labeled, and subjected to the microarray chips as above. For the negative control, anti-MILI was blocked with the peptide antigen before applying for immunoprecipitation. The negative control immunoprecipitates were processed and analyzed side by side with the experimental sample in the same way. The same amount and batch of antibodies and extract were used for the negative control as for the experimental sample. The enriched mRNAs in the experimental sample relative to the negative control were dubbed MILI-associated mRNAs, and their expression levels were evaluated in the milii+/− and milii−/− microarray samples. The represented results were derived from five independent replicas for total RNAs from milii+/− and milii−/− and four replicas for MILI-associated mRNAs (reverse transcription, amplification, and hybridization were conducted by the Duke Institute of Genome Sciences and Policy).

**Bioinformatic Analysis of Microarray Data**—Data on relative abundances of mRNAs within total testicular lysates, MILI immunoprecipitation samples, and mock immunoprecipitation samples were generated as follows. Microarray signals from replicates of total lysates were mean-normalized and averaged, respectively, to generate standard gene expression profiles of mouse testicular tissue. The ratio of the signal of each gene from each MILI immunoprecipitation sample to the standard gene expression profile was then calculated. Based on this ratio, a percentile rank of each gene relative to all genes in the genome was calculated. The expression level of each gene was assigned a percentile rank based on the number of other genes with lower expression levels. Additionally, each sample was normalized to wild type samples, and the ratio of the raw signal from mutant to wild type was calculated and then normalized to an average of 100 controls. A percentile rank of each gene relative to all genes in the wild type samples was calculated.
each immunoprecipitated replicate was calculated. The percentile ranks in the three replicates of MILI-immunoprecipitated samples were averaged. Student’s t test was utilized to determine if the average percentile ranks of enrichment of individual genes were significantly higher (p value) than the mean enrichment of all genes in the immunoprecipitation samples.

To determine the MILI-associated genes, the following criteria were used: 1) average percentile ranks of enrichment were greater than the mean enrichment of all genes in MILI immunoprecipitation with \( p < 0.01 \); 2) average signal in MILI immunoprecipitation replicates was significantly greater than the background signal (including \( 2 \times \text{S.D.} \)), with background signal and S.D. calculated based on signals from empty spots on each microarray; 3) criterion 1 was not satisfied for the same gene in the corresponding control immunoprecipitation. Using the above criteria, 802 mRNAs were designated as MILI-associated mRNAs. We further compared the expression profiles of MILI-associated mRNAs as well as nonassociated mRNAs in wild type versus \( \text{mili}^{-/-} \) testis. All cDNA microarray signals were median-normalized. The relative abundances of individual mRNAs from replicates were averaged.

RESULTS

**MILI Is Required for Germ Line Stem Cell Self-renewal**—We previously reported that \( \text{mili} \) is required for meiosis (39), which echoes post-germ line stem cell function of \( \text{piwi} \) and \( \text{pumilio} \) in the Drosophila ovary (8, 46). Although the \( \text{mili} \) mutant displays a terminal arrest phenotype at the pachytene stage of meiosis, we notice that only 2 ± 2.5% of the mutant spermatogonia actually reach this stage of differentiation (analysis detailed below), indicating an earlier spermatogenic defect prior to meiosis.

To characterize the early spermatogenic defect, we examined the mutant testes at critical stages of early spermatogenesis. In mouse testes, the precursors of germ line stem cells, called gonocytes, start proliferation at 3 dpp to establish the population of germ line stem cells (also known as primitive Type A spermatogonia) at 6 dpp. These stem cells then undergo self-renewing divisions to produce differentiated Type A spermatogonia (47). To determine whether this defect occurs prior to stem cell formation at 6 dpp, we examined the number and proliferation rate of gonocytes at 5 dpp by immunofluorescence microscopy, using the gonocyte/spermatogonium-specific EE2 antibody (48) to label gonocytes and anti-phosphohistone 3 antibody to label mitotic cells. The \( \text{mili}^{-/-} \) and \( \text{mili}^{+/-} \) tubules were similar in size (51.0 ± 4.0
versus 48.1 ± 5.1 μm in diameter) and contained an equivalent number of gonocytes that proliferated at a similar rate (Fig. 1, A and M). This indicates that the mili mutation does not affect gonocyte proliferation. At 8 dpp, the number of spermatogonia in mili−/− and mili+/+ tubules remained similar (Fig. 1, C and D), indicating that the germ line stem cell population had been established normally. However, at 8 dpp, the mili−/− germ line stem cells already showed a reduced mitotic rate (2.1 ± 0.40% versus 3.6 ± 0.4%; p < 0.006; Fig. 1M), indicating that spermatogonia division was compromised in the mili mutant.

We then investigated the effects of the mili mutation on spermatogonial division and differentiation at representative stages of postnatal development. At 10 dpp, mili−/− spermatogonia underwent active divisions and started to produce spermatocytes (Fig. 1E and Fig. S1A), yet mili−/− spermatogonia became further reduced in division rate (Fig. 1F and Fig. S1B). By 35 dpp, a complete complement of spermatogenic cells, resulting in drastically enlarged tubules (Fig. 1G and Fig. S1C). In contrast, mili−/− spermatogonia were mostly arrested in mitosis and differentiation and only occasionally produced early spermatocytes (Fig. 1H and Fig. S1D). By 98 dpp, wild type testes contained 32.9 ± 12.2 spermatocytes/tubule, whereas the mili mutant testes only contained 0.7 ± 1.5 spermatocyte/tubule, 47-fold fewer than the wild type situation (Fig. 1N). No spermatid was ever observed (Fig. 1N). The identities of these spermatocytes were confirmed by staining with the spermatocyte-specific antibody BC7 (Fig. S2, E and F). Furthermore, as we previously reported (39), these spermatocytes are arrested at zygotene and pachytene stages, as judged by the characteristic morphology of their chromosomes (Fig. S1D). The defects in spermatogonial division and differentiation persisted through 52 dpp (Fig. S1, E and F), 63 dpp (Fig. S2, C–F), and 98 dpp (Fig. 1, I and J, quantified in Fig. 1, N and O), without any loss of Sertoli cells or interstitial cells (Fig. 1N and Fig. S1, C–F). Along this time line, the arrested mutant spermatogonia increasingly underwent apoptosis (Fig. 1, K and L), so that most tubules were depleted of spermatogonia by 180 dpp, resulting in a Sertoli-only phenotype (Figs. S1, G and H and S2, G–J). These defects clearly indicate that mili mutant spermatogonial stem cells are blocked in both proliferation and differentiation. Thus, MILI is essential for the self-renewing division of these stem cells.

MILI is Expressed in Gonocytes, Spermatogonia, and Early Spermatocytes—To examine whether MILI is directly involved in germ line stem cell functions, we examined the expression of mili during germ line development. We first conducted Northern analysis of mili expression in adult tissues. mili RNA was detectable as a 3.7-kb mRNA in the testis but not in other tissues (Fig. 2A), consistent with its spermatogenesis-specific phenotype. To further investigate the expression of mili during spermatogenesis and germ line development, we examined the expression of mili mRNA in embryonic gonads and adult testes. mili was expressed in primordial germ cells starting at 12.5 days post coitum in sexually dimorphic male and female gonads (Figs. 2, B and C). At 13.5 days post coitum, when primordial germ cells in the male and female gonad further differentiate into gonocytes and oocytes, respectively (49), mili expression became stronger in both male and female gonads (Fig. 2, D and E). In the testis, mili was expressed in all cells in the testicular cord, with strong expression in centrally located gonocytes and weak expression in the basally located Sertoli cells in the cord (Fig. 2D). During subsequent fetal development, mili expression persisted at this high level in the male gonad (Fig. 2F) but was abolished in the female gonad (Fig. 2G). This reduction of mili expression in the female gonad correlates with the entry of female primordial germ cells into meiosis at 13.5 days post coitum. In adult mice, mili was not expressed in the ovary but was strongly expressed in the testis at the basal and sub-basal layers of the seminiferous epithelium, where spermatogonia and primary spermatocytes reside (Fig. 2, H and I). This expression pattern suggests that mili is probably involved in early spermatogenesis, from spermatogonial stem cell division to early meiosis.

To examine the expression of MILI protein, we generated an anti-MILI antibody that specifically recognizes MILI but not its homologs, as evident by Western blot analysis (Fig. S3; also see “Experimental Procedures”). Immunofluorescence staining of adult mouse testicular sections using this antibody revealed that MILI was specifically expressed in the cytoplasm of spermatogonia and pachytene stage spermatocytes but not in more differentiated germ cells (Fig. 3, A and B), similar to the mili mRNA expression pattern (Fig. 2, H and I). MILI expression was not detectable in Sertoli cells, which can be readily identified either by their expression of the Ttx antigen (Fig. 3C) or by their distinct chromatin morphology as visualized by the DNA dye DAPI (Fig. 3, C and E). In the germ line cytoplasm, MILI is enriched in spherical structures, most of which contain the Ge-1 protein, a marker for the chromatoid body and nuage, perinuclear structures related to the P-body of somatic cells, and was proposed to serve as an RNA storage and processing center (Fig. 3D) (38, 50), yet a small number of MILI-enriched spheres do not have Ge-1, and vice versa. The chromatoid

![Figure 1](image-url)
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FIGURE 2. The expression of mili mRNA during mouse germ line development. A, a Northern blot containing total RNAs prepared from various adult tissues as indicated and probed for the mili mRNA. B–I, in situ mili RNA hybridization of embryonic and adult testes and ovaries at specific stages as indicated. Red, mili mRNA; green in B and G, laminin, which outlines testicular cord (t. cord) in B, G, and F; green in H and I, DNA (stained with DAPI).

body/nuage localization of MILI resembles that of MIWI (38), suggesting that MILI might be involved in mRNA storage and processing.

Since germ line stem cells are a subset of spermatogonia, the expression of MILI in all spermatogonia indicates that it is expressed in germ line stem cells. This was confirmed by the colocalization of MILI with PItf, a germ line stem cell marker (Fig. 3, E–H) (5, 6). To examine whether MILI is expressed in germ line stem cells as soon as they are established, we conducted Western blot analysis on protein extracts from 4-, 6-, 8-, and 10-dpp testes. Since MILI was expressed at all these time points (Fig. S4A), yet MILI expression was germ line-specific (Fig. 3, B and C), this suggests that MILI is expressed in gonocytes (4 dpp), newly formed germ line stem cells (6 dpp), differentiating spermatogonia (8 dpp), and possibly leptotene spermatocytes (10 dpp). This result is also consistent with the detection of mili mRNA in embryonic gonocytes, adult germ line stem cells, spermatogonia, and primary spermatocytes but not in somatic cells (Figs. 2 and 3). The germ line-specific expression of MILI implicates that the function of MILI in germ line stem cell division and meiosis is cell-autonomous.

MILI Associates with Polysomes in an RNA-dependent Manner—To understand how MILI regulates spermatogonial division and differentiation, we explored the biochemical role of MILI. Recent works have shown that two Argonaute subfamily members, human eIF2C2 (also known as Ago2) and trypanosome TbAGO, associate with polysomes and, in the case of Ago2, stimulate translation (25, 51, 52). Moreover, we have shown that MIWI, a close homolog of MILI, is a cytoplasmic protein associated with polysomes in a manner correlated to active translation (26, 38). Since MILI is also a cytoplasmic protein, these reports raised the possibility that MILI may be involved in translational regulation. To explore this possibility, we first performed sucrose gradient fractionation of adult testicular lysates to examine if MILI co-fractionates with monosomes and/or polysomes (see “Experimental Procedures”). Testicular lysates displayed prominent polysome profiles, reflecting active translation (Fig. 4A). As expected, MILI co-sedimented not only with RNP fractions (fractions 1 and 2; Fig. 4A), which contain chromotoid body components, as we have previously shown (38), but also with polysome fractions (fractions 10–15; Fig. 4A), similar to MIWI (38).

To determine whether the co-sedimentation of MILI with polysome fractions is due to unrelated co-migration or reflects true association between MILI and polysomes, we added EDTA to chelate Mg$^{2+}$ required for polysome assembly. EDTA eliminated large polysome fractions (fractions 11–15; Fig. 4A). Correspondingly, MILI disappeared in these fractions (Fig. 4A). Instead, MILI was mostly concentrated in the monosome and subunit fractions (fractions 3–9); very little was in the RNP fractions (fractions 1 and 2), where chromotoid body components are most abundantly segregated under this condition (38). To further eliminate the possibility that MILI co-migrates with polysomes in other EDTA-sensitive complexes, we treated the lysates with micrococcal nuclease, which digested mRNAs and reduced some polysomes to monosomes (Fig. 4B). Correspondingly, MILI in the polysome fractions was also reduced. These analyses indicate that the co-sedimentation of MILI with polysome reflects its association with polysomes.

MILI Interacts with eIF3a and eIF4G—To further investigate the role of MILI association with polysomes, we used co-immunoprecipitation to search for specific components of the translational machinery that interact with MILI. We first developed a new rabbit antibody that specifically recognizes and precipitates endogenous full-length MILI from mouse testicular extracts (Fig. 5A; also see “Experimental Procedures”). This immunoprecipitation was not seen for mili mutant lysate (Fig. 6B) and was completely blocked by adding the antigen-peptide (Fig. 5A). Proteins co-immunoprecipitated with MILI were separated by SDS-polyacrylamide gels and silver-stained (Fig. 5B). Bands that were present only in the co-immunoprecipitant without the blocking peptide were excised from the gel and analyzed by matrix-assisted laser desorption-ionization-time of
flight mass spectrometry (see “Experimental Procedures”). We identified two major MILI-associated proteins as eukaryotic translational initiation factors eIF4GII and eIF3a (Fig. 5B). The eIF4GII protein migrated at the expected size, whereas eIF3a migrated at 122 kDa, smaller than its reported molecular mass of 170 kDa (53).

To verify the eIF3a-MILI interaction, we used an anti-eIF3a antibody to probe a Western blot of the MILI co-immunoprecipitant (Fig. 5C). The antibody detected an abundant presence of eIF3a in the co-immunoprecipitant at the expected size of 170 kDa (Fig. 5C). The addition of the blocking peptide drastically reduced eIF3a in the co-immunoprecipitant to an almost undetectable level (Fig. 5C). The eIF3a-MILI interaction was further confirmed by reciprocal co-immunoprecipitation of the testicular extract using the anti-eIF3A antibody. MILI was specifically co-immunoprecipitated by the eIF3a antibody but not by actin or T7 antibodies, as indicated by Western blot analysis (Fig. 5D). Thus, eIF3A interacts with MILI in the same protein complex. The smaller size of eIF3a in the mass spectrometry sample is presumably due to the breakdown of the 170-kDa protein into a 122-kDa fragment during the sample preparation for electrophoresis and mass spectrometry. The MILI-eIF4GII interaction was further verified by co-immunoprecipitation in the m7GTP cap-binding complex, as described below.

To examine whether the association between MILI and eIF3a is mediated by RNA, we treated the testicular extract with RNase A prior to co-immunoprecipitation by either MILI antibody or eIF3a antibody. RNase A treatment caused complete RNA degradation, as assessed by visualization of rRNA by ethidium bromide on agarose gel (data not shown). However, the association between MILI and eIF3a was still maintained (Fig. 5, C and D). These results indicate that MILI and eIF3A form a complex in vivo in an RNA-independent manner.

To examine whether MILI-eIF3a-eIF4GII interaction occurs in germ line stem cells, we examined the expression of MILI, eIF3a, eIF4GII, and ribosomal protein S6, another key component of the translational initiation complex in 6-dpp testes, which contain only nascent stem cells in the germ line, and 8-dpp testes, which contain only germ line stem cells and early type A spermatogonia in the germ line, by Western blotting analysis. MILI, eIF3a, eIF4GII, and ribosomal protein S6 were all present in the 6- and 8-dpp testicular extracts (Fig. S4A). Moreover, MILI-eIF3a interaction was detected at 8 dpp, the time point we examined (Fig. S4B). These results suggest the likely association between MILI and eIF3a as well as between MILI and eIF4GII in germ line stem cells and early type A spermatogonia.

MILI Is Associated with m7GTP Cap-binding Complex, Poly(A)⁺ mRNA, and piRNA—The interaction between MILI and eIF3a suggests that MILI may regulate translational initiation at an early step, possibly when the eIF3a-containing preinitiation complex binds to the mRNA m7GTP cap prior to mRNA scanning. To explore this possibility, we performed an m7GTP cap binding assay to test if MILI is associated with the cap-binding complex. Testicular extracts were incubated with m7GTP-cap analog conjugated to Sepharose beads, and the bound fraction was analyzed by Western blot. The m7GTP-Sepharose specifically co-purified the cap-binding complex components eIF4E, eIF4GII, and eIF3a, as well as MILI (Fig. 6A). The association of MILI and the eukaryotic initiation factors with m7GTP-conjugated Sepharose is specific, since these proteins did not bind to protein-A-conjugated Sepharose (Fig. 6A); nor did GRP 94, a chaperone protein, bind to the m7GTP-Sepharose (Fig. 6A).

To further test the specificity of MILI association with the cap-binding complex, a m7GTP-cap analog was added to the testicular extract to a final concentration of 200 μg/ml to compete with the interaction between the cap-binding complex and the Sepharose-conjugated m7GTP moiety. This addition drastically reduced the association of eIF4E, eIF3a, and eIF4GII as well as MILI with the m7GTP-Sepharose beads (Fig. 6A, fourth lane), indicating that MILI is associated with cap-binding complex in a specific manner. Since we were unable to observe the interaction between MILI and eIF4e, eIF2a, or eIF4GII by MILI co-immunoprecipitation (data not shown), the association
between MILI and the cap-binding complex is probably through the association between MILI and eIF3a.

The association of MILI with the translational initiation complex predicts that MILI is also associated with mRNAs. To investigate this possible association, we searched for the presence of polyadenylated mRNAs in the MILI-co-immunoprecipitated complexes by reverse transcription, using oligo(dT) primers (see “Experimental Procedures”). Radioactively labeled cDNA products were then fractionated by gel electrophoresis. Many poly(A)+ RNA species ranging from 0.5 to 4 kb in size were co-immunoprecipitated with MILI from the wild type but not mili mutant testicular lysate (Fig. 6B). This suggests that MILI is probably associated with many mRNAs in the testis. Under the same immunoprecipitation conditions, we also detected abundant association of piRNAs with MILI, as previously reported (Fig. 6C). These results point to the enticing possibility that some piRNAs are involved in translational regulation.

*mili* Mutant Testes Display Globally Reduced Translational Activity—To assess the function of MILI in translation, we performed [35S]methionine metabolic labeling to measure total protein synthesis in seminiferous tubules isolated from 7-dpp wild type versus mili mutant testes, based on the method of Kleene (43, 45) (see “Experimental Procedures”). We chose 7 dpp, because both wild type and mili mutant testes contain mostly, if not exclusively, stem cells in the germ line at this stage (Fig. 1 and Figs. S1 and S2). In addition, the mutant testes are phenotypically similar to the wild type testes at this time point, and both contain ~80% germ line stem cells and only 20% Sertoli cells within seminiferous tubules (Fig. 1 and Figs. S1 and S2). Hence, by isolating seminiferous tubules, we eliminated extratubular somatic cells, thus allowing our measurements to reflect largely the status of protein synthesis in germ line stem cells yet under a physiologically normal condition that could not be achieved if we used isolated germ line stem cells (43, 45).

We pulse-labeled the isolated seminiferous tubules with [35S]methionine for just 30 min at 32 °C to allow the labeling of only nascent peptides. Total proteins were then extracted and resolved by SDS-PAGE. Protein synthesis occurs robustly in isolated wild type tubules, yet there is a decrease in the synthesis of many protein bands in the mutant tubules (Fig. 7A). To quantify the decrease in total protein synthesis, we performed scintillation counting on the radioactivity of

**FIGURE 4.** A subpopulation of MILI is associated with polyribosomes. Sucrose density gradient analysis of adult mouse testicular extracts, with and without EDTA and micrococcal nuclease (MNase) treatment. Absorbance at 260 nm indicates the polyribosome profile. The presence of polyribosomes in various fractions was verified by the presence of 28 and 18 S rRNAs in the corresponding fractions. MILI in individual fractions was detected by Western blot analysis. A, effect of EDTA treatment. B, effect of micrococcal nuclease treatment.
trichloroacetic acid (20%)-precipitated protein samples from \textit{mili}^{+/-} or \textit{mili}^{-/-} testes to measure the total levels of protein synthesis. The experiments were repeated twice with independent samples. The variation among the three different experiments of either \textit{mili}^{+/-} or \textit{mili}^{-/-} testes were negligible (Fig. 7C). However, the total protein synthesis in \textit{mili}^{-/-} testes was consistently decreased to 59% of the \textit{mili}^{+/-} control level (Fig. 7C). This decrease is statistically highly significant ($p = 0.004$), indicating that global protein synthesis is reduced in the \textit{mili} mutant.

To determine whether the decreased protein synthesis is due to decreased levels of mRNAs in the mutant, we first quantified the total poly(A)$^+$ mRNA in 24-dpp \textit{mili}^{+/-} and \textit{mili}^{-/-} testes by isolating total RNA and reverse transcribing poly(A)$^+$ mRNA into $^{32}$P-labeled cDNA using oligo(dT) primers. The reason for choosing 24 dpp is that it was difficult to collect a sufficient amount of RNAs from the tiny 7-dpp testes, yet the \textit{mili} mutant displays a clear phenotype at 24 dpp, thus providing a more stringent first evaluation. Electrophoretic examination of the resulting cDNAs indicated the success of the reverse transcription and purification procedure (Fig. 7B). Scintillation quantification of the cDNAs, again repeated twice with independent samples, indicated that \textit{mili}^{+/-} or \textit{mili}^{-/-} testes contain very similar levels of mRNAs.

We then further investigated whether the levels of individual mRNAs are changed in the \textit{mili} mutant, especially whether there is an increase in retrotransposon mRNAs that compensates for a decrease in the “regular” transcripts, given the role of \textit{mili} in transposon silencing (28, 40). Our previous Affymetrix GeneChip analysis revealed that most mRNAs, including transposon mRNAs, are not changed (40). Only five IAP retrotransposon genes are up-regulated more than 3-fold. To confirm this result and to further quantify mRNAs for the change of their expression in the \textit{mili} mutant, we compared the expression of 15,647 genes in the 9-dpp \textit{mili}^{+/-} and \textit{mili}^{-/-} mice, using an AROS version 3.0 oligonucleotide array (Operon; see “Experimental Procedures”). Among these genes, only 365 showed a greater than 2-fold decrease in expression in \textit{mili}^{-/-} mice, representing a 0.65% decrease in the total
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mRNA pool (Fig. 7D). Likewise, only 381 genes showed a greater than 2-fold increased expression in mili−/− mutants, representing a 0.3% increase in the total RNA. The net change in the mRNA pool is only a 0.35% decrease. These results further indicate that the levels of most individual mRNAs are not affected in the mili mutant. We then further examined whether MILI-associated mRNAs are somewhat more affected than nonassociated mRNAs by co-immunoprecipitation of MILI and its associated mRNAs from the adult mili+/− or mili−− testicular extract, followed by extraction of the MILI-associated mRNAs and quantification of their expression by an AROS version 3.0 oligonucleotide array (Operon). Again, of the 802 MILI-associated mRNAs that we detected, only 21 genes showed a greater than 2-fold decrease of expression in the mili−/− mutant, and only 23 genes showed a greater than 2-fold increased expression in the mili−/− mutant (Fig. 7D; see “Experimental Procedures”). Thus, by all of these criteria, the reduction of global protein synthesis in mili mutant seminiferous tubules reflects a novel and probably positive role of MILI in translational regulation.

DISCUSSION

MILI Is Essential for Germ Line Stem Cell Self-renewal and Differentiation—Although the piwi/ago gene family was first discovered for its function in stem cell self-renewal (21), such a function has not been demonstrated in any vertebrate system. Zebrafish PIWI (ZIWI) prevents excessive apoptosis in the germ line but not germ line stem cell division (54). The disruption of ago2 in mice leads to embryonic lethality early in development after the implantation stage (55). Among three murine PIWI proteins, MIWI is a key regulator of spermiogenesis (41); whereas MIW12 is predominantly involved in meiosis and transposon control (32). Our previous work identified a function of MILI in meiosis (39). The earlier and essential function of MILI in germ line stem cell self-renewing division and differentiation reported here extends the stem cell function of the PIWI/AGO protein family to vertebrate systems.

This stem cell function of MILI is unique in that it is required for both self-renewal and differentiation, which is different from other known germ line stem cell genes. Plzf mutant mice produce a limited number of sperm but progressively lose their germ line afterward (5, 6), suggesting that Plzf is only required for self-renewal and not for differentiation. On the other hand, c-Kit and stem cell factor are only required for the differentiation of spermatogonia and not for self-renewal (2, 3). Sox3 appears to be a survival factor for germ line stem cells, since Sox3-deficient mice begin to lose germ cells as early as at 10 dpp (7). By 14 dpp, Sox3 null mice are nearly germ lineless. The dual function of MILI in controlling both the self-renewal and differentiation of germ line stem cells is reminiscent of that of PIWI, which affects both the self-renewal of germ line stem cells and sometimes the differentiation of their progeny in Dro sophila (46), although the differentiation function of PIWI has not been extensively characterized.

In mili mutants, the escape of ~2% of spermatogonia from differentiation block (see “Results”) has provided an opportunity to identify its subsequent function during early meiosis (39). The function of MILI in germ line stem cells and prophase spermatocytes matches nicely with its expression pattern during development.

MILI Represents a Novel Mechanism That Appears to Positively Regulate Translation—Recent advancements in noncoding small RNA research have shed light on the crucial role of PIWI/Argonaute proteins and miRNAs in diverse biological...
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processes. Argonaute subfamily proteins have been implicated in small interfering RNA-mediated mRNA degradation, epigenetic silencing, and miRNA-mediated translational repression, all of which negatively regulate gene expression. Likewise, PIWI proteins have been shown to participate in the transcriptional and post-transcriptional transgene silencing as well as epigenetic regulation in various organisms (33, 34, 56–59). Particularly, MILI itself has been implicated in repressing DNA methylation and transposition (28, 40). Although a very small number of MILI molecules might exit in the nucleus for these functions, the cytoplasmic localization of MILI suggests that MILI may indirectly regulate these nuclear events. Our results suggest that MILI positively regulates the translation of many, if not all, genes during early spermatogenesis. Some of the MILI target genes could in turn regulate epigenetic state and repress transposition. Furthermore, our microarray analyses show that the increase of IAP retrotransposon RNAs reflects a minor change in the total levels of retrotransposon mRNAs, which had little impact on the total mRNA level in the mili mutant. In fact, the increased IAP mRNAs in the mili mutant should further contribute to the synthesis of the total pools of proteins that we measured.

The overall magnitude of the impact of MILI on translation is 2-fold, which is a value averaged over all cellular mRNAs, including mRNAs that are targets of MILI regulation. It is therefore conceivable that the effect of MILI toward its target mRNAs or a subset of them could be much higher than 2-fold. Although MILI might achieve this broad range positive regulation by repressing the translation of a small number of translational repressors that subsequently inhibit the translation of many other mRNAs in the cell, this scenario, even if it exists, is unlikely to be a major mode of regulation, given that MILI appears to bind directly to many mRNAs (Fig. 6B). The direct positive role of MILI in translational regulation is supported by the strong correlative evidence for a similar role of MIWI, since MIWI also directly complexes with its target mRNAs (41) and becomes increasingly associated with polyribosomes as translational activity increases in spermaticogenic cells (38).

Translational regulation by MILI appears to represent a novel mechanism-positive regulation by a PIWI protein at the initiation stage. In contrast, AGO proteins mediate miRNA-related translational repression that targets the 3′-untranslated region of mRNA. It has been suggested by several groups that this can cause early ribosome exit of repressed mRNAs and/or deadenylation of these mRNAs. It has also been reported that AGO proteins could subject substrate mRNAs to translational inhibition when miRNA-like small RNAs are used as targeting effectors (60) and that miRNA-dependent translational repression could interrupt translation initiation (61, 62). However, these effects on translational initiation are negative. It has recently been reported that AGO2 can up-regulate translation, but that is via the AU-rich element in the 3′-untranslated region rather than the cap (25). The function of MILI is probably different from all of the above mechanisms. It appears to promote translation via the preinitiation complex and the mRNA cap. Particularly, MILI interacts with eIF3a in an RNA-independent manner. eIF3a is the largest subunit of the eIF3 complex. In mammals, eIF3 binds to the 40S ribosomal subunit prior to the recruitment of 5′ m7GTP-capped mRNA. This recruitment is necessary to initiate translation and is accomplished via the interaction between eIF3a and the eIF4F complex, which binds the m7GTP-cap via the eIF4E subunit (63). Our finding that MILI is associated with both the cap-binding complex and poly(A) + RNAs is consistent with this hypothesis. The fact that MILI is also associated with polysomes suggests that the function of MILI is coupled with both the recruitment of the capped mRNA and the formation of polyribosomes on the mRNA.

**MILI Regulates the Translation of Germ Line Stem Cells**—MILI probably regulates translation in germ line stem cells, since MILI interacts with the components of the translational initiation complex in 8-dpp testes that contain mostly germ line stem cells and a small number of early type A spermatogonia (Fig. S4). Furthermore, translational activity is reduced in 7-dpp mili mutant seminiferous tubules, which contain mostly germ line stem cells and a small number of Sertoli cells. In addition to regulating germ line stem cells, MILI may regulate translation in gonocytes, other spermatogonia, and early spermatocytes, as implicated by its expression pattern and phenotype. These roles of MILI highlight the importance of translational regulation in germ line stem cell maintenance, mitosis, and meiosis. For example, the translational repressors NOS and PUM are essential for maintaining the fate of germ line stem cells in *Drosophila*, presumably by repressing the translation of differentiation genes (8–10). Consistent with this notion, the translational control of cyclin B is essential for the progression of cells through mitosis and meiosis (64). In addition, the requirement of eIF3 and eIF5 in meiosis has been demonstrated in *C. elegans* (65). The identification of MILI-target mRNAs should reveal aspects of cellular processes in spermatogenesis that are regulated by MILI.

**Does MILI Regulate Translation via piRNAs?**—Although MILI is associated with the translational initiation factors, mRNAs, and piRNAs, without definitive data indicating one-to-one correspondence at the single-molecule level, it is not clear whether MILI-mediated translational regulation involves piRNAs. It has been reported that ZIW1-associated piRNAs are localized in the cytoplasm (54). Moreover, MIWI-associated piRNAs are present in the polysome fraction, and MIWI is also associated with the cap (38). Extensive and systematic effort is required to demonstrate whether some piRNAs are involved in translational regulation mediated by PIWI family proteins.

**MILI in Chromatoid Bodies; a Function in RNA Sequestration?**—The presence of MILI in both the nuage/chromatoid body and the free cytosol is consistent with its two-phase distribution between RNP and ribosomal fractions in sucrose gradients (Fig. 4). Our previous study of MIWI has demonstrated that RNP fractions contain chromatoid body components, yet ribosomal fractions contain cytoplasmic MIWI (38). Chromatoid bodies resemble P-/GW-bodies in somatic cells that are implicated in small RNA-mediated mRNA degradation (60, 66–68). However, MIWI is required for the stability of its target mRNAs as well as for the integrity of the chromatoid body (41, 69). Thus, the stabilization
function of MIWI might be related to the chromatoid body. If so, this germ line organelle would have a function opposite to its somatic counterpart; it would sequester and protect mRNAs rather than degrade them. The observed MILI/Ge-1 co-localization in spermatocytes suggests that MILI is also a component of the P-/-GW-bodies. In addition, the presence of a small number of MILI-enriched spheres that share the size and morphology of chromatoid bodies but lack Ge-1 suggests the existence of at least two distinct types of P-body-like structures in the germ line. The role of chromatoid body-associated MILI in mRNA stability control was not tested here, since this study focuses on the effect of MILI on germ line stem cells in 6–8-dpp testes, which do not contain chromatoid bodies.

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