MVH in piRNA processing and gene silencing of retrotransposons

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Keywords: Spermatogenesis; MVH; small RNA; piRNA; retrotransposon; epigenetic regulation

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Supplemental material is available at http://www.genesdev.org.

Received January 6, 2010; revised version accepted March 5, 2010.

There are many evolutionarily conserved proteins that play essential roles in germ cell development. Some proteins—such as VASA, PIWI, and NANOS in Drosophila, and their mammalian homologs—are involved in the processing and binding of RNA, suggesting the importance of RNA processing in germ cell development and differentiation [Saga 2008; Siomi and Kuramochi-Miyagawa 2009]. VASA, a germ cell-specific DEAD-box RNA helicase of Drosophila, is essential for germ cell development and oogenesis through regulation of target mRNAs such as Nanos [Noce et al. 2001; Becalska and Gavis 2009]. Expression of its murine homolog, MVH (mouse vasa homolog), is also restricted to germ cell lineage [Toyooka et al. 2000]. Primordial germ cells [PGCs] develop in MVH-deficient mice, but spermatogenesis is blocked at the first meiotic cell division [Tanaka et al. 2000]. In round spermatids of the adult testis, MVH is localized in the chromatoid body, a unique cloud-like structure of male germ cells that contains mRNA, microRNA [miRNA], and various proteins, including MVH [Kotaja and Sassone-Corsi 2007]. As the chromatoid body would be an intracellular focal domain necessary for RNA processing, MVH is likely to have some pivotal role(s) in RNA processing in male germ cells. However, its molecular role has not been elucidated.

Piwi family genes also show germ cell-specific expression and are essential for germ cell maintenance and spermatogenesis in Drosophila and mammals, respectively [Lin 2007; Peters and Meister 2007; Siomi and Kuramochi-Miyagawa 2009]. Piwi was originally identified as a gene essential for germ stem cell maintenance in Drosophila, and its gene family is found in a wide range of organisms from Arabidopsis to mammals [Cox et al. 1998]. The three mouse Piwi homologs Miwi, Mil, and Miwi2 are all essential for spermatogenesis [Deng and Lin 2002; Kuramochi-Miyagawa et al. 2004; Carmell et al. 2007]. The phenotypes of Miwi and Miwi2 gene targeted mice were essentially the same and showed male sterility due to apoptosis of the germ cells at early pachytene phase [Kuramochi-Miyagawa et al. 2004; Carmell et al. 2007]. In addition, both mouse mutants showed enhanced retrotransposon expression in the male germ cells due to defective de novo DNA methylation of the genes [Kuramochi-Miyagawa et al. 2008]. Piwi proteins are bound to a novel class of germ cell-specific small RNAs called Piwi-interacting RNAs [piRNAs] [Aravin et al. 2006; Girard et al. 2006; Grivna et al. 2006; Lau et al. 2006; Watanabe et al. 2006]. Miwi, which is expressed from PGCs at embryonic day 12.5 [E12.5] to round spermatids, binds with 26- to 27-nucleotide (nt) piRNAs [Kuramochi-Miyagawa et al. 2001; Aravin et al. 2006]. On the other hand, Miwi2, which is expressed in fetal gonocytes from E15.5 until soon after birth, binds to 28- to 29-nt piRNAs [Aravin et al. 2008; Kuramochi-Miyagawa et al. 2008]. Previously, we showed that most piRNAs at the fetal stage were derived from repetitive retrotransposon genes, and that the production of piRNA was markedly impaired in Miwi- and Miwi2-deficient mice [Kuramochi-Miyagawa et al. 2008]. These data suggest that Miwi and Miwi2 are involved in piRNA production in the fetal male gonads, and that the piRNA would play some important role(s) in gene silencing of retrotransposons via DNA methylation.

Many proteins are involved in piRNA production in Drosophila [Malone et al. 2009]. A feed-forward loop to mediate the generation of piRNAs was originally postulated for Drosophila piRNA production [Brennecke et al. 2007;
Gunawardane et al. 2007). This ping-pong amplification cycle is mediated by two Drosophila PIWI family proteins, AUB and AGO3, which bind primarily to antisense primary piRNAs and secondary sense piRNAs, respectively. Based on the observation that MIWI2 binds preferentially to secondary antisense piRNAs compared with MIWI, a similar ping-pong cycle would presumably involve MIWI and MIWI2 in the mouse fetal testes instead of AUB and AGO3 in Drosophila [Aravin et al. 2008]. It is conceivable that the ping-pong cycle cannot proceed by the actions of MIWI and MIWI2 alone, and we attempted to identify other molecules essential for the ping-pong cycle.

MVH is expressed in the male germ cells from E10.5 to around spermatid [Toyooka et al. 2000], which covers the period of de novo DNA methylation of retrotransposons. In addition, we reported previously that the defective spermatogenesis and impairment of gene expression in MILI-deficient mice were similar to those of MVH-deficient mice [Kuramochi-Miyagawa et al. 2004]. We also found that both MILI and MIWI bound to MVH. Therefore, we postulated that MVH may play some role(s) in piRNA production and subsequent DNA methylation of retrotransposons. Here, we showed that MVH plays essential roles in de novo DNA methylation of retrotransposons, presumably due to the defective piRNA production, and that MVH is an essential factor in the piRNA processing pathway.

Results and Discussion

Expression and DNA methylation of intracisternal A particle (IAP) and Line-1 retrotransposons in MVH-deficient testes

First, we analyzed the expression of IAP and Line-1 retrotransposons, both of which were derepressed in the MILI- and MIWI2-deficient testes. As shown in Figure 1A, expression levels of the 5.4-kb transcript IΔ-type IAP and the Line-1 transcripts were elevated in the testes of MVH-deficient mice. As heavy methylation of the 5′ long terminal repeat (LTR) of IAP and the promoter region of Line-1 was the major cause of transcriptional silencing of these genes, the methylation status of the regulatory regions of these retrotransposons was examined. Methylation-sensitive Southern blotting of Line-1 [Fig. 1B] and bisulfite sequencing analyses of IAP and Line-1 [Fig. 1C] in male germ cells clearly showed that DNA methylation was significantly impaired in MVH-deficient male germ cells at the postnatal stage.

As reported previously, de novo DNA methylation of retrotransposons was reduced in MILI- and MIWI2-deficient male germ cells [Kuramochi-Miyagawa et al. 2008]. We carried out bisulfite sequencing of the regulatory regions of the Line-1 and IAP retrotransposons in MVH-deficient fetal germ cells at E16.5, when de novo DNA methylation occurred [Fig. 1D]. Compared with the control male germ cells, DNA methylation of the MVH-deficient male germ cells was severely impaired. Two paternal imprinted genes, H19 and Dlk-Gtl2, were methylated normally in the MVH-deficient male germ cells, suggesting that de novo DNA methyltransferases function normally even without Mvh [Supplemental Fig. 1]. These observations indicated that MVH plays essential roles in de novo DNA methylation and subsequent silencing of the retrotransposons.

Expression and binding of piRNAs in MVH-deficient fetal testes

The abnormalities of spermatogenesis and the regulation of retrotransposon gene expression in MVH-deficient mice were quite similar to those of MILI- and MIWI2-deficient mice. As piRNA expression in MILI- and MIWI2-deficient fetal male germ cells was significantly reduced [Aravin et al. 2008; Kuramochi-Miyagawa et al. 2008], we analyzed the expression of piRNA in fetal male germ cells of MVH-deficient mice by microarray analysis [Fig. 2A]. Although the expression of miRNA was not altered, the expression of piRNA of repetitive sequences was significantly impaired under MVH-deficient conditions. General reduction of piRNA expression in MILI-deficient male germ cells was significantly more severe than that of MIWI2-deficient cells [Kuramochi-Miyagawa et al. 2008]. Gross comparison of the piRNA microarray analyses also showed that expression of piRNA in MVH-deficient fetal male germ cells was intermediate between those of MILI- and MIWI2-deficient cells. Reduced expression of arbitrary chosen piRNAs was confirmed by Northern blot analysis [Supplemental Fig. 2].

We examined whether the piRNA was bound to MVH. Extracts of E16.5 wild-type fetal male germ cells were
immunoprecipitated with the appropriate antibodies, and the bound small RNAs were radiolabeled and subjected to gel electrophoresis. As shown in Figure 2B, 26- to 27-nt-long and 28- to 29-nt-long piRNAs were bound to MILI and MIWI2, respectively. However, the band of piRNA was undetectable in the complex immunoprecipitated with the anti-MVH antibody.

piRNA expression was reduced in the MVH-deficient testes, but small amounts of piRNA remained. We analyzed the binding of residual piRNAs to MILI and MIWI2 in E16.5 MVH-deficient testes. As shown in Figure 2C, MILI-bound piRNA was detected in the MVH-deficient testes, but MIWI2-bound piRNA was not. These observations were not due to the disappearance of MIWI2 caused by defective MVH expression, because MIWI2 was expressed in MVH-defective mice (Supplemental Fig. 3). Taken together, these results indicate that, although piRNA did not bind to MVH, piRNA expression was significantly decreased in the MVH-deficient fetal testes. In addition, MILI-bound piRNA was present in the MVH-deficient fetal testes, whereas MIWI2-bound piRNA was not.

Characterization of small RNAs in MVH-deficient fetal testes by deep sequencing

To characterize the piRNAs of the MVH-deficient E16.5 male germ cells in more detail, we sequenced a large number of small RNAs from these cells. In comparison with the controls, the amount of piRNA was reduced in the MVH-deficient fetal male germ cells [Fig. 3A; Supplemental Table 1]. Especially, the number of 28- to 29-nt piRNAs, which were MIWI2-bound piRNAs, was markedly reduced. These observations were consistent with those of microarray (Fig. 2A) and binding (Fig. 2C) analyses. The sequences of small RNAs of control and MVH-deficient E16.5 fetal male germ cells were classified according to the corresponding genes (Supplemental Table 1), and their relative abundances compared with the numbers of miRNAs were calculated (Fig. 3B). The relative amount of piRNAs in the mutant male germ cells was about one-fifth that of the control. piRNAs corresponding to repetitive sequences were markedly reduced in the MVH-deficient fetal male germ cells.
are believed to be produced by amplification through the ping-pong cycle. The absence of MIWI2-bound piRNAs in the MVH-deficient male germ cells indicated clearly that the ping-pong cycle with MILI and MIWI2 did not progress in the mutant mice. Meanwhile, the first nucleotide of the repeat piRNAs was uridine in >79% of cases (Fig. 3C), indicating that the primary processing pathway functions even in the absence of MVH.

The first step of the ping-pong cycle involves the slicer activity of MILI, which cleaves the 10th nucleotide of the complimentary transcript annealed to the primary piRNA. This cleavage gives rise to adenine at the 10th nucleotide from the 5’ end, which corresponds to the first uridine of the primary piRNA (Brennecke et al. 2007; Gunawardane et al. 2007). As shown in Figure 3D, the percentage of piRNAs from MVH-deficient germ cells with adenine at the 10th nucleotide was almost the same as that of control cells. Therefore, although the ping-pong cycle with MILI and MIWI2 did not proceed, it is possible that the primary piRNA production and the initial part of the cycle—namely, the cleavage of the mRNA corresponding to the primary piRNA—took place without MVH. Meanwhile, we cannot exclude the possibility that MILI alone can proceed the ping-pong cycle in a very inefficient manner. As no MIWI2-bound piRNA was observed in the MVH-deficient cells, MVH appears indispensable between the first step of the secondary piRNA production and the loading of piRNA to MIWI2.

A significant proportion of piRNAs are derived from clusters scattered on the genome. The relative amounts of piRNAs derived from large piRNA clusters are shown in Figure 3E. Compared with the reduction of total piRNAs (~20% of the control, as shown in Fig. 3A; Supplemental Table 1), the relative abundance of cluster-derived piRNAs was decreased more markedly (~2.5% of the control, as shown in Fig. 3E; Supplemental Table 2). RNA transcripts from the clusters were postulated to play significant roles in the ping-pong cycle. Thus, the marked reduction of cluster-derived piRNAs is consistent with the defective ping-pong cycle with MILI and MIWI2.

**Alterations in the subcellular localizations of MVH, MILI, and MIWI2 in deficient mice**

Nuage or germ cell granules are cytoplasmic ribonucleoprotein (RNP) aggregates with amorphous ultrastructural granules localized specifically in the male germ cells (Eddy 1975; Chuma et al. 2009). It has been reported that germ granules have a close relationship with piRNA production and subsequent retrotransposon silencing in the germline. Two groups, including our group, have reported recently that MILI and MIWI2 are components of two distinct subcellular compartments in fetal post-spermatagonia/gonocytes in which de novo DNA methylation is established (Aravin et al. 2009; Shoji et al. 2009). MILI is localized to the intermitochondrial cement, a form of germ granule, and MIWI2 is a component of processing bodies [P-bodies; an mRNP assembly], which are proposed to be involved in RNA degradation/translational control, including miRNA- and siRNA-mediated pathways (Aravin et al. 2009; Shoji et al. 2009). Aravin et al. (2009) proposed naming the former and the latter pi-body and p-body, respectively. In addition, two Tudor domain-containing proteins, TDRD1 and TDRD9, both of which are important for piRNA production and gene silencing, belong to the intermitochondrial cement and processing bodies, respectively (Aravin et al. 2009; Shoji et al. 2009). We next analyzed the subcellular localizations of MVH, MILI, MIWI2, TDRD1, and TDRD9 in the mutant gonocytes (Fig. 4A).

Immunostaining for MVH, MILI, and TDRD1 in wild-type E16.5 male germ cells showed staining with many cytoplasmic granules, which corresponded to the intermitochondrial cement (Fig. 4A). On the other hand, immunostaining with anti-MIWI2 and anti-TDRD9 antibodies showed a few larger granules, presumably corresponding to the processing bodies, and MIWI2 was detected in the nucleus (Aravin et al. 2009). In MVH-deficient mice, nuclear localization and large granular staining of MIWI2 was not detectable, but that of TDRD9 remained, which suggested that the processing bodies would exist, but MIWI2 was mislocalized. As the subcellular localization of MIWI2 has been suggested to be dependent on piRNA expression (Aravin et al. 2008), it is possible that the mislocalization of MIWI2 is due to the decreased piRNA levels.

Granular staining of both MILI and TDRD1 was abolished in the MVH-deficient germ cells. Examination by electron microscopy indicated that this alteration was not due to the mislocalization of the proteins, but was
attributable to the loss of intermitochondrial cement [Fig. 4B]. In the MILI-deficient germ cells, intermitochondrial cement was not detected and the numbers of the MVH-stained granules were reduced (Supplemental Fig. 4), suggesting the functional loss of the intermitochondrial cement. The disappearance of intermitochondrial cement was not caused solely by the loss of piRNAs, as the intermitochondrial cement was essentially normal in MIWI2-deficient male germ cells in which piRNA production was significantly reduced. Therefore, it is possible that the loss of intermitochondrial cement was due to the loss of MVH per se or by some molecular sequence downstream from MVH other than piRNA production. Taken together, these observations indicated that MVH is essential for the construction, and presumably also the function, of the intermitochondrial cement. It is unclear whether the molecular function of MVH as an RNA helicase on RNA per se, the role in the formation of intermitochondrial cement, or both is essential for normal piRNA production. We are carrying out the experiments to clarify this point by producing the transgenic mouse bearing the mutant MVH without helicase activity [Sengoku et al. 2006]. However, MVH and nuage are essential for the ping-pong amplification in which MVH is involved.

Essential proteins in ping-pong amplification cycle and intermitochondrial cement formation

As shown in Figure 4B, the intermitochondrial cement was missing in the MVH-deficient fetal male germ cells. Intermitochondrial cement was also absent in the TDRD1-deficient fetal male germ cells [Chuma et al. 2006]. Recently, several groups have reported that TDRD1 binds physically to MILI, and that the protein is crucial for the production of piRNA [Chen et al. 2009, Kirino et al. 2009, Kojima et al. 2009, Reuter et al. 2009, Vagin et al. 2009]. TDRD1-deficient fetal male germ cells showed decreases in levels of MIWI2-bound piRNAs, antisense piRNAs, and 10th A piRNAs, suggesting impairment of the ping-pong cycle [Vagin et al. 2009]. Taken together, the similarities in impairment of piRNA production in TDRD1-deficient and MVH-deficient fetal male germ cells support the suggestion that appropriate intermitochondrial cement formation is essential for the progression of ping-pong cycle amplification.

The ping-pong amplification cycle cannot proceed with only MILI and MIWI2. TDRD1 and TDRD9 also play essential roles, presumably in the intermitochondrial cement and processing bodies, respectively [Aravin et al. 2009, Shoji et al. 2009]. These two RNP complexes often show neighboring, suggesting that they cooperate and have interdependent functions in piRNA biogenesis. Therefore, it is possible that piRNAs are transferred from one domain to the other during the cycle. The findings of the present study indicating the existence of MILI-bound primary piRNA and the involvement of MVH in the construction and/or function of intermitochondrial cement suggested that MVH is essential for transfer of piRNA from the intermitochondrial cement to the processing bodies.

Materials and methods

RNA extraction, Northern blotting, and microarray analysis

Total RNA was prepared from 2-wk and 2-mo testes and E16.5–E17.5 embryonic testes using Isogen (Nippon Gene). Northern blotting analysis was performed as described in a previous study [Kuramochi-Miyagawa et al. 2008]. The probes of IAP and Line-1 were as follows: the 3′ noncoding region of IAP (nucleotides 6486–6793 of GenBank accession nos. M17551), and the 5′ noncoding region of Line-1 (nucleotides 2621–2625 of D84391).
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