Dynamic Regulation of ARGONAUTE4 within Multiple Nuclear Bodies in Arabidopsis thaliana

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DNA methylation directed by 24-nucleotide small RNAs involves the small RNA-binding protein ARGONAUTE4 (AGO4), and it was previously shown that AGO4 localizes to nucleolus-adjacent Cajal bodies, sites of snRNP complex maturation. Here we demonstrate that AGO4 also localizes to a second class of nuclear bodies, called AB-bodies, which are found immediately adjacent to condensed 45S ribosomal DNA (rDNA) sequences. AB-bodies also contain other proteins involved in RNA-directed DNA methylation including NRPD1b (a subunit of the RNA polymerase IV complex, RNA PolIV), NRPD2 (a second subunit of this complex), and the DNA methyltransferase DRM2. These two classes of AGO4 bodies are structurally independent—disruption of one class does not affect the other—suggesting a dynamic regulation of AGO4 within two distinct nuclear compartments in Arabidopsis. Abolishing Cajal body formation in a coilin mutant reduced overall AGO4 protein levels, and coilin dicer-like3 double mutants showed a small decrease in DNA methylation beyond that seen in dicer-like3 single mutants, suggesting that Cajal bodies are required for a fully functioning DNA methylation system in Arabidopsis.


Introduction

RNA-directed DNA methylation (RdDM) is a phenomenon discovered in plants in which small RNAs cause DNA methylation and transcriptional gene silencing at complementary sequences in the genome [1–3]. Frequent targets of RdDM include regions which contain direct and/or inverted repeats such as those present in transposable elements, some intergenomic regions, and a small percentage of promoters at endogenous genes [4]. Numerous studies have linked components of the RNA interference (RNAi) pathway to RdDM [5]. In RNAi, double stranded RNA substrates are cleaved by the endonuclease Dicer to form small interfering RNAs (siRNAs) which are then bound by an Argonaute protein [6]. Using the siRNA as a guide, Argonautes can direct gene silencing through transcriptional or post-transcriptional mechanisms [7,8].

In Arabidopsis, genes involved in the production of siRNAs associated with DNA methylation include NRDP1a, RNA-DEPENDENT RNA POLYMERASE2 (RDR2), and DICER-LIKE3 (DCL3) [9–12]. Downstream genes which carry out the transcriptional silencing function directed by the siRNAs include ARGONAUTE4 (AGO4) and NRDP1b [13–15]. NRDP1a and NRDP1b encode isoforms of the largest subunit of the plant-specific RNA polymerase IV complex (RNA PolIV), but depending on which subunit is present in the complex, RNA PolIV can function upstream in siRNA biosynthesis (NRDP1a) or downstream in causing DNA methylation (NRDP1b) [9,10,14,15]. The final step of the RdDM pathway is performed by DOMAINS REARRANGED METHYLTRANSFERASE2 (DRM2), a de novo DNA methyltransferase that is required for the establishment of DNA methylation and for the maintenance of DNA methylation in asymmetric CHH (where H=A, T, or C) and CHG sequence contexts [12,16]. Mutations in these RdDM genes result in decreased transcriptional silencing of certain repetitive loci.

Past studies investigating the subcellular distribution of nuclear proteins have uncovered a spatial partitioning of functions within the nucleus [17]. Distinct nuclear compartments or bodies with functions such as splicing, RNA metabolism, or complex formation have been characterized [18]. One such nuclear compartment is the nucleolus-adjacent Cajal body, at which small nuclear RNAs (snRNAs) are modified and assembled into small nuclear ribonucleoprotein (snRNP) complexes that are involved in splicing in the nucleoplasm [19–21]. Localization studies in Arabidopsis revealed that several RdDM components also formed nuclear bodies which can be found either adjacent to or within the nucleolus [21,22]. Proteins which formed such discrete nuclear bodies include RDR2, DCL3, AGO4, and NRDP1b. More recently, Dicing bodies (D-bodies), which contain microRNA processing factors DCL1 and HYL1, were also discovered in plant nuclei [23,24]. These D-bodies are present in the nucleoplasm but are not associated exclusively with the

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Author Summary

Argonautes are components of the RNA interference (RNAi) pathway that bind small interfering RNAs (siRNAs) of 21–24 nucleotide length. In the flowering plant Arabidopsis thaliana, ARGONAUTE4 (AGO4) is involved in gene silencing at the transcriptional level in a process called RNA-directed DNA methylation (RdDM), during which siRNAs cause transcriptional gene repression at complementary sequences. Previously, we found that AGO4 localized to nucleolus-adjacent Cajal bodies, sites of snRNP complex assembly. In this study, we show the existence of a second class of AGO4 nuclear foci, which we call the “AB-bodies,” that is distinct from the Cajal body and is immediately adjacent to the condensed 45S ribosomal DNA (rDNA) loci. In addition to AGO4, AB-bodies also contained NRPD1b and NRPD2 (subunits of the plant-specific RNA polymerase IV complex) and the DNA methyltransferase DRM2. The two different classes of AGO4 nuclear foci are structurally distinct, since the loss of one class did not affect the other. Losing Cajal bodies resulted in the enhancement of the DNA methylation defects seen in the RNA silencing mutant dicer-like3, suggesting Cajal bodies are required for a fully functioning RdDM pathway leading to gene silencing.

Results

Two Distinct Classes of AGO4 Nuclear Bodies in Arabidopsis Nuclei

Utilizing a plant line containing Myc-tagged genomic AGO4 that is under the control of its endogenous promoter, we previously found that AGO4 shows very good colocalization with Cajal body markers, with colocalization seen in roughly 95% of the nuclei [21]. Here we undertook an examination of a large population of nuclei in order to study the small fraction of nuclei in which AGO4 and Cajal body markers did not colocalize. Within about 200 nuclei isolated from Myc-AGO4 seedlings, we confirmed that a majority of nuclei contained AGO4 colocalizing with U2B99, a marker for the Cajal body (Figure 1A) [21,25,26]. However, we also observed a small population (4%) of nuclei that contained AGO4 foci that did not colocalize with U2B99, and an even smaller population (1%) of nuclei which displayed both types of AGO4 localization (one colocalized with U2B99 and one not colocalized with U2B99). We confirmed this result by examining AGO4 localization relative to SmD3 and SmB, which are two proteins also enriched at the Cajal body [21,27]. Non-colocalization between AGO4 and SmD3 or SmB was observed in a small population of nuclei (Figure S1; Table S1), again suggesting that AGO4 forms nuclear foci distinct from the Cajal body.

An examination of AGO4 relative to NRPD1b localization within nuclei was also performed to determine if there are AGO4 foci that are separate from NRPD1b. As observed previously (see Table S1 in [21]), NRPD1b colocalized with AGO4 in a small percentage of cells, while the remainder of the nuclei contained only AGO4 foci and no NRPD1b (Figure 1B). Also as previously observed, when analyzing those nuclei in which clear NRPD1b foci were present, those NRPD1b foci always colocalized with AGO4. However, we observed a small population of nuclei (2% in the experiment shown in Figure 1B) in which, in addition to the NRPD1b and AGO4 containing bodies, a second focus of AGO4 staining was found that did not colocalize with NRPD1b.

These results suggest the possibility of multiple different AGO4 bodies, and suggest that NRPD1b might not localize to Cajal bodies as was previously assumed [21]. To test this hypothesis, we examined U2B99 localization relative to NRPD1b in nuclei and found that these two proteins were indeed localized to separate foci adjacent to the nucleolus (Figure 1C). Utilizing three fluorescent colors to look at Myc-AGO4, GFP-U2B99, and NRPD1b simultaneously, we confirmed that multiple AGO4 foci within the nucleus either colocalized with NRPD1b or U2B99, but we never observed all three proteins colocalizing together (Figure 1D). Further characterization of the two different AGO4 bodies by fluorescence measurements using confocal microscopy showed that AGO4 immunostaining at the NRPD1b body was generally more intense and punctate, while AGO4 immunostaining at the Cajal body was usually less intense and more diffuse (Figure S2). We also tested whether any of the AGO4 foci corresponded to Dicing bodies which contain HYL1 and DCL1 [23,24]. By co-immunofluorescence analysis, GFP-HYL1 did not colocalize with AGO4 or NRPD1b, suggesting none of the different AGO4 bodies are Dicing bodies (Figure S3A and S3B; Table S2). GFP-HYL1 also did not colocalize with U2B99, consistent with published results which showed Dicing bodies are not Cajal bodies (Figure S3B; Table S2) [23,24]. These results suggest there are two classes of AGO4 nuclear bodies: a first class that colocalizes with the Cajal body in a majority of nuclei, and a second class that is with NRPD1b in a smaller population of cells (called “AB-bodies” for AGO4/NRPD1b bodies).
Figure 1. AGO4 Localizes to Two Distinct Nuclear Bodies

(A) Fluorescent microscopy analysis of Myc-AGO4 and Cajal body localization in nuclei isolated from Myc-AGO4 seedlings. A monoclonal antibody to endogenous U2B^99 was used to detect the Cajal body. Three different AGO4 localization patterns relative to U2B^99 are shown. Nuclei containing observable AGO4 nuclear foci were examined.

(B) Localization of Myc-AGO4 and NRPD1b in Myc-AGO4 nuclei. A polyclonal antibody was used to detect endogenous NRPD1b. Three representative nuclei are shown. Nuclei containing observable AGO4 nuclear foci were examined.

(C) Localization of U2B^99 relative to NRPD1b within wild type Ler nuclei.

(D) Three color fluorescent microscopy analysis of Myc-AGO4, U2B^99-GFP, and NRPD1b localization within the same nucleus. One representative nucleus containing two different AGO4 bodies is shown: colocalization with NRPD1b (top of the nucleus) versus colocalization with the Cajal body (bottom of the nucleus). All AGO4 foci colocalized with either NRPD1b or Cajal body (n = 156).

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Cajal Body Integrity Affects AGO4 Localization and Is Required for a Fully Functioning RdDM System

The presence of the two classes of AGO4 bodies led us to ask whether the Cajal body and AB-body might be dependent on one another for stability. To investigate this, Myc-AGO4 and NRPD1b localization were examined in a coillin mutant that was unable to form visible Cajal bodies (Figure S4). Mutations in COILIN were shown to completely disrupt Cajal body formation, and interestingly, this disruption did not cause any obvious morphological defect in plant development [28]. We found that in the coillin mutant background, only a small fraction of nuclei (14%) contained AGO4 foci, and these AGO4 foci all colocalized with NRPD1b (Figure 2A and 2B; Table S3). This indicates that AGO4 is no longer able to form nuclear foci outside of the AB-bodies in the absence of the Cajal body. This finding that only AB-bodies were observed in the coillin mutant suggests that the formation of the AB-body is not affected by Cajal body integrity or by the ability of AGO4 to localize to the Cajal body. Interestingly, in a plant line that overexpressed COILIN-mRFP and contained enlarged Cajal bodies [28], AGO4 enrichment at the Cajal body also expanded to the same size as the enlarged Cajal body, while no major effect on the AB-body was observed (Figure 2C).

In addition to the effects on AGO4 localization, we also observed less intense staining of Myc-AGO4 both within nuclear foci and throughout the nucleoplasm in the coillin mutant (Figure 2A and 2B). This decrease in AGO4 detection is reminiscent of the effects of rdr2, del3, and nrd1a on AGO4, where a mutation in any of these genes decreased overall protein levels of AGO4 [21]. To confirm that the observed decrease in AGO4 staining intensity in coillin nuclei is also due to lower AGO4 protein levels, a semi-quantitative western blot analysis was used to examine Myc-AGO4 levels in coillin or a wild type sibling. Consistent with the immunofluorescence analysis, the overall protein level of Myc-AGO4 was slightly decreased in coillin when compared to wild type (Figure 2D). This finding suggests that the Cajal body is required for maintaining full levels of the AGO4 protein.

Next we tested whether proper Cajal body formation was required for AGO4 dependent RNA-directed DNA methylation. Thus, we examined DNA methylation and siRNA production in the coillin mutant and COILIN-mRFP overexpression line. No obvious change in DNA methylation was observed at the 5S rDNA loci or at the euchromatic locus MEA-ISR in DNA methylation analysis, in coillin SALK mutant lines that we have isolated, or in the previously described Cajal body mutant lines (ncb-1, ncb-2, and ncb-4) (Figure S5A and S5B) [28]. Similarly, there was no effect on 45S and 5S siRNAs when the Cajal body was absent or expanded (Figure S5C). This lack of a general effect on gene silencing pathways is consistent with coillin mutants also lacking observable developmental defects even though Cajal bodies are a major site of snRNA modification and snRNP assembly [28].

Because the effect of coillin on AGO4 levels was only slight, we reasoned that we may only see an effect of losing the Cajal body on DNA methylation in a sensitized background in which the DNA methylation pathway has already been partially affected. Thus, we crossed a coillin mutant (ncb-1) to del3. The del3 mutant was shown previously to have weaker effects than rdr2, nrd1a, or ago4 on DNA methylation [11,12], which in part could be due to DCL redundancy [21,29–31]. At MEA-ISR, the ncb-1 del3 double mutant contained further decreased DNA methylation levels beyond that seen in del3 alone (Figure 2E). This enhancement of a weak del3 phenotype was slight but consistently reproducible using independent biological replicates (Figure 2E). Similar to MEA-ISR, we also observed a slight decrease in DNA methylation levels at the 5S rDNA loci in the ncb-1 del3 double mutant relative to del3 alone (Figure S5D). Hence, Cajal body integrity appears to have some role in RdDM efficiency, possibly through an effect on AGO4 protein levels.

Stability of the AB-Body in RdDM Mutants

We investigated the factors that are important for the stability of the AB-body by examining the genetic requirements for its formation. In the upstream RdDM mutants del3, rdr2, nrd1a, and del3 rdr2, NRPD1b nuclear bodies were still seen in a small percentage of nuclei similar to the percentage seen in wild type, despite slightly decreased fluorescence intensity (Figure 3; Table S4). Thus, NRPD1b does not require NRPD1a, RDR2, or DCL3 for localization to nuclear bodies.

We also examined AGO4 localization to the AB-body in the upstream mutants. As seen previously, AGO4 nuclear levels were severely decreased in a portion of del3 nuclei and in all of the rdr2, del3 rdr2, and nrd1a nuclei (Figure 3; Table S4) [21]. In the case of del3, only about one fourth of total nuclei from del3 retained normal AGO4 levels, while the remaining majority of nuclei contained reduced AGO4 signal (Figure 3; Table S4). The del3 mutant is known to have a weaker effect than rdr2, nrd1a, or ago4 on DNA methylation [11,12]. This lack of a stronger effect may be a result of the loss of AGO4 protein in only a fraction of del3 nuclei, in addition to redundancies in DCL function [29–31]. The remaining del3 cells that retain normal AGO4 levels would presumably allow some activity of RdDM to occur, giving an intermediate phenotype that falls between wild type and stronger mutants such as rdr2, nrd1a, and ago4. Thus, some of the DNA methylation defects seen in del3 may be an indirect effect of having less AGO4 protein, in addition to the more direct effect of losing siRNAs that are normally produced by DCL3 [11,29].

Within the population of del3 nuclei that contained decreased AGO4 levels, no AGO4 or NRPD1b foci were observed (Figure 3; Table S4). In contrast, del3 nuclei which had normal AGO4 levels contained both AGO4/Cajal bodies and AB-bodies (Figure 3; Table S4). It remains unclear what cell types within the del3 mutant contain normal AGO4 levels, and why AGO4 protein should be stabilized in these specific cells. One explanation would be that another DCL is expressed in these cells, and in the absence of DCL3, replaces the function of DCL3 in stabilizing AGO4 protein and localization [29–31]. Alternatively, the del3 mutation might result in a stochastic loss of AGO4 protein within certain nuclei that may or may not be dependent on cell type (discussed in [21]).

In the rdr2, del3 rdr2, and nrd1a mutants, despite the low overall fluorescence levels, we observed a faint AGO4 body that colocalized with NRPD1b in all nuclei that contained NRPD1b bodies (Figure 3; Table S4). This is in contrast to del3, where AB-bodies were found in del3 nuclei containing normal AGO4 levels but not in del3 nuclei that lost the majority of AGO4 protein. This would suggest that the population of
nuclei that contain AB-bodies in the \( \text{rdr2, dcl3, rdr2, or nrpd1a} \) mutant backgrounds are among those that also show normal AGO4 levels in a \( \text{dcl3} \) mutant (see Discussion). In summary, these results suggest that localization of both NRPD1b and AGO4 to AB-bodies still occurs in the \( \text{nrpd1a, rdr2, or dcl3} \) mutants.

To examine the effects of downstream mutations in the formation of the two classes of AGO4 bodies, we investigated whether AGO4/Cajal bodies or AB-bodies were present in \( \text{nrpd1b, ago4, and drm2} \) mutant nuclei. In \( \text{nrpd1b} \), a majority of cells still contained AGO4 foci (Table S4), consistent with previous results [21]. However, all of the remaining AGO4 foci present in \( \text{nrpd1b} \) colocalized with U2B\(^99\), indicating that AGO4 is not able to form nuclear foci outside of the Cajal

![Figure 3](https://example.com/figure3.png)

**Figure 3.** Presence of the AB-Body in Upstream RdDM Mutants

Immunofluorescence analysis of Myc-AGO4 and NRPD1b in wild type, \( \text{dcl3, rdr2, dcl3, rdr2, or nrpd1a} \) nuclei. White arrows indicate the presence of a faintly staining Myc-AGO4 body that colocalized with NRPD1b. Two populations of nuclei are shown for each mutant: nuclei with and nuclei without AB-bodies.

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body in this mutant (Figure 4A and 4B; Table S4). Because only AGO4/Cajal bodies remain in the \textit{nrpd1b} mutant, this suggests that the formation of the AB-body is dependent on NRPD1b while the localization of AGO4 to the Cajal body is not. In the \textit{ago4} mutant however, NRPD1b bodies were still observable, although the overall fluorescence appeared to be slightly weaker than wild type (Figure 4C; Table S5). In the \textit{drm2} mutant, we observed the presence of NRPD1b and both classes of AGO4 bodies, indicating that both AB-bodies and AGO4/Cajal bodies are unaffected in this mutant (Figure 4A and 4B; Table S4). These results suggest NRPD1b is upstream of both AGO4 and DRM2 in AB-body formation, and NRPD1b localization to nuclear bodies is mostly independent of AGO4.

**Localization of NRPD2 to AB-Bodies**

\textit{NRPD2} encodes the second largest subunit of RNA PolIV complexes containing NRPD1a or NRPD1b [9,14,15]. NRPD2 was shown to interact with NRPD1a and NRPD1b in vivo, and was observed to be colocalized with NRPD1a and NRPD1b in small speckles outside of the nucleolus [22]. To determine if NRPD2 is also present at the AB-body, the localization of NRPD2 within nuclei was examined using an antibody which detects endogenous NRPD2 [22]. By immunofluorescence analysis, we observed NRPD2 localized to a nucleolus-adjacent body that colocalized with AGO4 in a small population of nuclei (Figure 5A and S6). However, unlike AGO4, these NRPD2 nuclear bodies did not colocalize with

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**Figure 4. Effects of Downstream RdDM Mutations on the AB-Body**
(A,B) Myc-AGO4 nuclear localization in wild type, \textit{nrpd1b}, or \textit{drm2} mutant nuclei is shown relative to the immunostaining of (A) NRPD1b or (B) U2B\textsuperscript{99}. (C) Immunolocalization of NRPD1b and U2B\textsuperscript{99} in wild type (Ler) or an \textit{ago4} mutant. The white arrow indicates a less intense NRPD1b body that still remained in \textit{ago4}.

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indicating NRPD2 is not enriched at the Cajal body (Figure 5B; Table S6). Since NRPD2 is at an AGO4 focus that is not a Cajal body, this strongly suggests that NRPD2 is also targeted to the AB-body. Although we could not directly test colocalization of NRPD2 with NRPD1b since antibodies to NRPD2 and NRPD1b are both rabbit polyclonal, our observed NRPD2 body is most likely the AB-body given the finding that NRPD2 interacts with NRPD1b in vivo [22], and NRPD2 also colocalized with a fourth protein present at the AB-body, DRM2 (see below).

To determine whether NRPD2 displays the same genetic requirements as AGO4 and NRPD1b for localization to the AB-body, the ability of NRPD2 to form nuclear bodies was examined in nrpd1b, rdr2, ago4, and nrpd1a mutants. Similar to NRPD1b, NRPD2 nuclear bodies were still visible at wild type fluorescence levels in nrpd1a (Figure 5B; Table S6) and rdr2 (not shown), and still present with slightly decreased intensity levels in ago4 (Figure 5B; Table S6). However, in the nrpd1b mutant, no obvious nuclear bodies were seen for NRPD2 (Figure 5B; Table S6). We performed the reciprocal experi-
ment and found no obvious NRPD1b nuclear body in the nrpd2 mutant (Figure 5C; Table S5). This lack of NRPD1b or NRPD2 bodies in nrpd2 or nrpd1b, respectively, is likely in part due to the fact that NRPD1b and NRPD2 depend upon each other for protein stability [14]. However, it is also possible that the absence of NRPD1b or NRPD2 bodies is due to a defect in targeting to the AB-body in the nrpd2 or nrpd1b mutant background, respectively. These results suggest NRPD1b and NRPD2 show similar requirements with regards to localization to AB-bodies.

**AB-Bodies Are Always Adjacent to Condensed 45S rDNA Loci**

To further characterize the subnuclear localization of AB-bodies, DNA fluorescence in situ hybridization (FISH) in combination with protein immunofluorescence analysis was performed to visualize the location of major repetitive genomic regions (45S rDNA, 5S rDNA, and centromeric repeats) relative to the AB-body. As seen previously, a significant population of Myc-AGO4 nuclei showed AGO4 foci separate from the condensed 45S rDNA loci (or nucleolar organizer regions, NORs) (Figure 6A, bottom panel; Table S7) [21,32], since most of the nuclei show AGO4 at Cajal bodies. However, we also observed a population of AGO4 foci that were immediately adjacent to or slightly overlapping with the NORs (Figure 6A, top panel and Figure S7; Table S7). One AGO4 focus per NOR was frequently observed, although two or three AGO4 foci surrounding one NOR were also seen (Figure S8A).

To confirm that the AGO4 foci associated with the condensed 45S rDNA loci were AB-bodies and not Cajal bodies we performed DNA FISH together with NRPD1b immunolocalization. In all nuclei containing NRPD1b bodies, NRPD1b was always found adjacent to or overlapping with the NORs, indicating that the NOR-adjacent AGO4 foci are AB-bodies (Figure 6A; Table S7). Consistent with this result, we did not find the Cajal body, either marked by GFP-SmD3 or GFP-SmB fluorescence, to be closely associated with the NORs (Figure S8B; Table S7). In contrast to the NORs, AGO4 or NRPD1b foci were not found to be closely associated with the 5S rDNA loci or centromeric repeats (CEN) (Figure 6B, 6C, and S7; Table S7). These results suggest that the AB-body is specifically situated adjacent to the NORs.

Since the 45S rDNA repeats are present on both chromosome 2 (NOR2) and chromosome 4 (NOR4) [33], it is possible that AGO4/NRPD1b may be preferentially targeted to one NOR versus another. We examine this possibility by performing dual DNA FISH analysis looking at both condensed 45S and 5S rDNA loci simultaneously in addition to AGO4 or NRPD1b immunofluorescence. Since the 5S rDNA loci are present on chromosome 4 but not on chromosome 2 [34,35], NOR4 can be detected by the close proximity of the 45S signal to 5S signal, while NOR2 will only display 45S signal and no adjacent 5S signal [33]. Using this method, we found AB-bodies that were adjacent to both NOR2 (45S only) and NOR4 (45S with 5S) (Figure 6D; Table S8). This suggests that the AB-body is present at both NORs.

**The AB-Body Associates with DRM2**

The finding that both AGO4 and NRPD1b are associated with the NORs suggests that the AB-body might be a site of active transcriptional silencing at the 45S rDNA repeats. The 45S rDNA loci consist of both silent and active copies of the 45S rDNA gene [32]. It is possible that some of these active rDNA copies may undergo de novo silencing by the RdDM machinery in the form of the AB-body, especially since 45S rDNA silencing in *Arabidopsis* is known to involve DNA methylation at promoter sequences [36]. If this is the case, the de novo DNA methyltransferase DRM2 might also colocalize with the AB-body. To test this, we utilized a stable plant line expressing Myc-tagged genomic DRM2 which is under the control of its endogenous promoter [21]. We examined the localization of DRM2 in nuclei which contained AB-bodies marked with NRPD1b or NRPD2. As previously observed, a majority of nuclei contained DRM-Myc localized diffusely throughout the nucleoplasm with less intense signal at chromocenters (Table S9) [21]. In a small population of nuclei where NRPD1b or NRPD2 bodies were present, DRM2 was also observed as a nucleolus-adjacent body that colocalized with NRPD1b or NRPD2 (Figure 7A and 7B; Table S9). In contrast, no DRM2 body was seen colocalized with U2B′, indicating DRM2 is not enriched at the Cajal body (Figure S9) [21]. Interestingly, the *ago4* mutation caused a drastic reduction in the number of nuclei containing DRM2 bodies, suggesting that AGO4 has a role in targeting DRM2 to the AB-body (Figure 7A; Table S9). We also attempted to determine whether DRM2 localization is affected in *nrpd1b*. However, due to the speckle-like nucleoplasmic staining of DRM2 that is only slightly fainter than the DRM2 signal at the AB-body, and without a second marker (such as NRPD1b or NRPD2) to confirm localization, it is difficult to conclusively determine whether DRM2 is still able to form a nuclear body that colocalizes with the AB-body in *nrpd1b*. However it is unlikely that DRM2 would be targeted to the AB-body in the absence of NRPD1b, since AGO4 localization to the AB-body requires NRPD1b, and DRM2 localization to the AB-body requires AGO4. In sum, the presence of DRM2 at the AB-body and its dependence on AGO4 are consistent with the idea that the AB-bodies might represent active sites of gene silencing.

**Discussion**

Our findings demonstrate the existence of two independent populations of AGO4 nuclear bodies that are structurally distinct. This is in contrast to the model that we proposed previously in which we assumed that all AGO4 bodies were Cajal bodies, including AGO4 bodies that colocalized with NRPD1b [21]. Examination of a large population of nuclei combined with direct testing of the localization of NRPD1b relative to the Cajal body demonstrated that our initiation interpretation was incorrect, and that the regulation of AGO4 distribution within the nucleus is more complex than first conceived. Our new data show that AGO4 localizes independently to the Cajal body or to the 45S rDNA-adjacent AB-body, and that AB-bodies are distinct from AGO4/Cajal bodies.

The majority of AGO4 foci within the nucleus represents the Cajal body, a nucleolus-adjacent compartment in which the maturation of snRNP complexes occurs. Our analysis with a *collin* mutant showed that AGO4 levels were affected by the absence of Cajal bodies. This suggests that the Cajal body may play a role in stabilizing AGO4 protein, just as RdDM components RDR2, DCL3, and NRPD1a are also involved in AGO4 stability. Although no obvious DNA methylation defects were seen in the *collin* mutants alone, the *ncb-1*
Figure 6. Localization of AB-Bodies to the Immediate Proximity of the 45S rDNA Loci
(A) DNA FISH analysis of the condensed 45S rDNA loci (NORs) relative to Myc-AGO4 (left) or NRPD1b (right) localization in Myc-AGO4 or Ler nuclei, respectively.
(B,C) Localization of AGO4 or NRPD1b is shown relative to (B) 5S rDNA loci or (C) centromeric repeats (CEN).
(D) Dual probe DNA FISH analysis showing 45S and 5S rDNA loci relative to Myc-AGO4 (top panel) or NRPD1b (bottom panel). The 45S probe was labeled with biotin and the 5S probe was labeled with digoxigenin (DIG). NOR4 is observed as a 45S rDNA signal with an adjacent 5S rDNA signal, while NOR2 shows only 45S rDNA signal with no adjacent 5S rDNA signal.
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mutation enhanced the DNA methylation phenotype of del3 at MEA-ISR and the 5S rDNA loci. The ncb-1 del3 double mutant had less DNA methylation than del3 alone but did not decrease methylation to the levels seen in ago4 or the triple DNA methyltransferase mutant drm1 drm2 cmt3. One explanation for the lack of a stronger enhancement by ncb-1 is that the function of the Cajal body may still partially remain in the coilin mutant. It is possible that although visible Cajal body formation is disrupted, Cajal body function may still exist in smaller, less concentrated foci through the nucleus in coilin mutants. This is a likely possibility since Cajal body mutants display no obvious developmental or morphological defects, despite the fact that Cajal bodies are major centers for assembly of splicing complexes that are essential for cellular processes [28]. A similar situation exists in mammals where mice unable to form wild type Cajal bodies as a result of a deletion in coilin are also largely aphenotypic [37]. In turn, AGO4 function at the Cajal body also may be dispersed throughout the nucleus in the absence of a visible concentrated Cajal body. Despite the weak phenotype, our data showing the effects of coilin on AGO4 stability and on enhancing DNA methylation defects in the del3 mutant suggest that the Cajal body plays some role in the efficiency of RdDM.

The smaller population of AGO4 foci, the AB-bodies, is present in nuclei at a frequency that varies from experiment to experiment, typically ranging from 4% to 20% of total nuclei. The AB-bodies are present in every major tissue examined, such as flowers, leaves, and whole seedlings (data not shown). This suggests that AB-bodies might be present in cell types that constitute a minority of cells within most major tissues of the plant.

In addition to AGO4 and NRPD1b, AB-bodies also contain RdDM factors NRPD2 and DRM2. The AB-body was always observed to be associated with the NORs by DNA FISH analysis, suggesting that AB-bodies might be sites of on-going 45S rDNA silencing, especially since DRM2 is also enriched at these particular nuclear bodies. Some of the active rDNA repeats may be specifically targeted for de novo silencing by the RdDM machinery as a mechanism to modulate or fine tune total cellular rDNA gene activity. This idea is consistent with earlier findings showing that a subset of the rDNA repeats in Arabidopsis is more highly DNA methylated and less occupied by RNA polymerase I, while another subset is less DNA methylated and more highly associated with RNA polymerase I [36]. Interestingly, FISH analysis showed that the AB-body was most often adjacent to the NORs instead of overlapping them. One explanation for the minimal overlap is that the AB-body might be present at less condensed or active 45S rDNA sequences that require gene silencing, and that these less condensed 45S rDNA sequences are difficult to visualize by DNA FISH. Euchromatic regions were previously shown to loop out of adjacent condensed DNA regions called chromocenters, which contain highly condensed 45S rDNA, 5S rDNA, and/or centromeric sequences [33]. It is thus possible that the AB-body is present at one of these loops, which would then situate the AB-body adjacent to the NORs.

The stability of the AB-body in upstream RdDM mutants...
sets AB-bodies apart from previously characterized nuclear bodies and indicates that AB-bodies are novel nucleolus-associated bodies [22]. Interestingly, while AB-bodies were detectable in nrpd1a, rdr2, and dcl3 rdr2 nuclei that contained very little overall AGO4 protein, we found that in dcl3, nuclei containing decreased overall AGO4 levels did not show any AB-bodies, but dcl3 nuclei with normal AGO4 levels did contain AB-bodies (Figure 3; Table S4). One possible explanation for this observation is that the specific cell types that normally form AB-bodies exist within the population of dcl3 nuclei that still contain normal AGO4 levels, and not within the dcl3 population containing reduced AGO4 levels. While the molecular basis for this correlation is unknown, it suggests a relationship between the particular cell population or cell type that contains AB-bodies and those able to express normal levels of AGO4 despite the dcl3 mutation.

The fact that AB-bodies were still detected in nrpd1a, rdr2, and dcl3 mutants suggests that an alternative pathway exists to target AGO4 and NRPD1b to 45S rDNA or adjacent regions. The finding that NRPD1a and RDR2 are not required for AB-body formation may indicate that the target sequence of the AB-body can produce double stranded RNA and siRNAs in a manner that does not require the activity of NRPD1a-containing RNA PolII complexes or RDR2. In this way these loci would show the same genetic requirements as do a variety of inverted repeat loci in the Arabidopsis genome whose sRNA synthesis does not require NRPD1a or RDR2 [38].

The finding that AGO4 did not localize to AB-bodies in the nrpd1b mutant, but that NRPD1b could still localize to nuclear bodies in an ago4 mutant, suggests a function of NRPD1b in stabilizing AGO4 at the 45S rDNA loci, possibly by NRPD1b interacting with AGO4 directly [21] and/or by generating a nascent RNA strand for AGO4 binding. Our data would further suggest that once stably localized at the target region by NRPD1b, AGO4 would then recruit DRM2 (directly or indirectly), since the localization of DRM2 to the AB-body depends on AGO4. In the future, it will be interesting to determine the precise nature of the sequences targeted by components of the AB-body, and the mechanisms and factors that recruit NRPD1b, NRPD2, AGO4 and DRM2 to 45S rDNA adjacent sequences.

Materials and Methods

Generation of epitope tagged lines. Genomic AGO4 was N-terminally tagged with four copies of c-Myc and expressed under the control of the endogenous AGO4 promoter as described [21]. Genomic DRM2 was C-terminally tagged with nine copies of c-Myc and expressed under its endogenous promoter as described [21]. GFP-SmD3 and GFP-SmB plasmids, and U2B-GFP and COILIN-mRFP plasmid lines were gifts from Peter J. Shaw. Construction of GFP-SmD3 and GFP-SmB plasmids were described previously [39]. GFP-SmD3 and GFP-SmB were transfected into Mco-AGO4 plants using Agrobacterium strain AGL1. Construction and generation of Ramos plasmids were described previously [28]. The U2B-GFP or COILIN-mRFP line was crossed to Mco-AGO4 plants to generate lines containing both epitopes. Construction of the GFP-HYL1 plasmid was described previously [24]. Mco-AGO4 plants were transformed with GFP-HYL1 produce transgenic plants containing both Mco-AGO4 and GFP-HYL1. For mutant analysis, epitope tagged lines were cross to nrpd1a, dcl3 rdr2, ago4–1, nrpd1b, or drm2 and homozygous for the single or double mutations.

Loss-of-function mutants. The RdDM mutants used in this study are the following: nrpd1a (SALK 143457, ABRC stock center), dcl3–1 and rdr2–1 [11]; ago4–1 [13], drm2-16, nrpd1b-1 [14], and nrpd2a–2 nrpd2b–1 [15]. The DNA methyltransferase triple mutant dmt1 drm2 cmt3 was isolated as described [40]. The collar mutants SALK 148589, SALK 148630, SALK 010395, and SALK 083548 were obtained from the ABRC stock center. Plasmid coloin mutants nb-1, nb-2, and nb-4 were gifts from Peter J. Shaw and were isolated as described previously [28].

Protein immunofluorescence analysis. Isolation of nuclei from bolting seedlings and immunodetection of proteins in plant nuclei were performed as described [21,41]. Primary antibodies used included anti-c-Myc (1:200, Upstate), polyclonal anti-Ago4 (1:200, Upstate), NRPD1b (1:100) [14], NRPD2 (1:200) [22], mAb 4G3 (anti-U2B) (1:100, gift from Gregory Matera), and polyclonal GFP (1:200, Invitrogen). Secondary anti-mouse-Fluorescein (Abcam), anti-rabbit-Rodamine (Jackson ImmunoResearch), anti-mouse-Rodamine (Abcam), and anti-mouse Alexa Fluor 350 (Invitrogen) were used at a dilution of 1:200. DNA was stained using Vectashield mounting medium containing DAPI (Vector Laboratories). Images were captured with the Zeiss AxioImager Z1 microscope with the Hamamatsu Orca-er camera at 100X magnification and analyzed using the Zeiss Axiovision software. Zeiss FL filter sets used in this study: Zeiss 49 (DAPI), Zeiss 38 (EGFP), and Zeiss 43 (Cy 3).

Confocal fluorescence measurements. Nuclei from Myc-AGO4 seedlings were examined for Myc-AGO4 and NRPD1b nuclear bodies using primary antibodies against c-Myc (1:200, Upstate) and NRPD1b (1:100). Secondary antibodies used included anti-mouse-Fluorescein (Abcam) and anti-rabbit-Rodamine (Jackson ImmunoResearch). DNA was stained using Vectashield mounting medium containing DAPI (Vector Laboratories). Confocal fluorescence measurements for AGO4, NRPD1b, and DAPI were obtained using the Zeiss LSM 510 META, 100X/1.40 NA oil immersion objective, 3 PMT, and the 405, 488, and 633 nm lines with monochromatic excitation and were processed as described previously [21]. Image J analysis was performed as described [22,42]. To combine protein immunolocalization with DNA FISH, protein immunofluorescence was first performed as described above, except nuclei slides were post-fixed in 4% paraformaldehyde after primary antibody incubation. After post-fixing, DNA FISH was performed using probes labeled with biotin-dUTP or digoxigenin (DIG)-dUTP [43]. Probes for the 45S rDNA [44], 5S rDNA [45], or the centromeric repeats [46] were generated by PCR. Biotin or DIG labeled probes were detected using FITC-avidin (1:200, Zymed), NeutrAvidin-Alexa Fluor 350 (1:200, Invitrogen), or anti-DIG-Fluorescein (1:200, Roche). Vectashield mounting medium containing DAPI (Vector Laboratories) was used to stain DNA. Images were captured and analyzed as described for protein immunofluorescence except the Apotome system was used during acquisition.

Western blot analysis. Protein extraction from flowers and western blot analysis were performed as described previously [21]. Equal amounts of total protein were loaded for each sample and resolved on a 10% SDS polyacrylamide gel. A monoclonal antibody to c-Myc (Upstate) was used to detect Myc-AGO4. A polyclonal antibody detecting endogenous CRY1 was used as a loading control [47].

Southern blot analysis. Genomic DNA extraction and southern blot analysis examining DNA methylated at ME4-ISR and the 5S rDNA repeats were performed as described [11,48]. Twenty μg of digested genomic DNA was loaded per sample.

Northern blot analysis. Total RNA extraction and northern blot analysis examining small RNAs corresponding to the 45S rDNA, 5S rDNA, and miR159 were performed as described [21,13]. Twenty μg of total RNA was loaded per sample. Probe sequences are as follows: 45S siRNA: 5′-GCTCTGTTGGTGCCAAGAGGGCTAATA-3′; 5S siRNA: 5′-ATGCGAAGTTTGGCCACAGGGGAAAGCGCTAAT-3′; 5′-TAAGGCTCCITCATTCAACAA-3′.

Supporting Information

Figure S1. Immunofluorescence Analysis of SmD3 and SmB Localization

Myc-AGO4 (top panels) or NRPD1b (bottom panels) nuclear localization relative to GFP-SmD3 or GFP-SmB is shown. GFP fluorescence was used to visualize SmD3 or SmB within plant nuclei to mark the Cajal body. Found at doi:10.1371/journal.pgen.0040027.s001 (2.6 MB TIFF).

Figure S2. Intensity Measurements of the Two Classes of AGO4 Nuclear Bodies

(A) Fluorescence measurements of AGO4, NRPD1b, and DAPI at the different AGO4 bodies. One representative nucleus containing AGO4 colocalized with NRPD1b (left) or the Cajal body (right) used for the analysis is shown. The red arrow indicates the location in the nucleus where measurements were taken. Distance is set at zero where the highest AGO4 signal was measured. Measurements were obtained using a confocal microscope.
(B) Average intensity measurements for AGO4, NRPD1b, and DAPI at the two different AGO4 bodies. Fifteen nuclei containing AGO4/NRPD1b or AGO4/Cajal bodies were used for the averaging. Found at doi:10.1371/journal.pgen.0040027.sg002 (1.5 MB TIF).

Figure S3. Localization of HYL1 Relative to AGO4, NRPD1b, and U2B̂. (A) Immunofluorescence analysis of GFP-HYL1 and Myc-AGO4 nuclear localization. GFP fluorescence was used to visualize GFP-HYL1 within the nucleus to mark the Dicing body. An antibody to Myc was used to detect Myc-AGO4. (B) Localization analysis of GFP-HYL1 relative to NRPD1b (top panel) or U2B̂ (bottom panel) within plant nuclei. An antibody to NRPD1b or U2B̂ was used to detect endogenous NRPD1b or U2B̂ protein, respectively. Found at doi:10.1371/journal.pgen.0040027.sg003 (3.0 MB TIF).

Figure S4. U2B̂ Immunostaining in coilin SALK Mutants Nuclei isolated from four SALK lines containing a T-DNA insertion in coilin were examined for Cajal body formation by U2B̂ immunostaining. No Cajal body was observed in SALK 148589, 148630, and 010395 mutant lines. SALK 083448 is a hypomorphic allele in which a small percentage of nuclei contained a Cajal body. The remaining 2% of nuclei from wild type (Col) did not contain an observable Cajal body by U2B̂ immunostaining. Found at doi:10.1371/journal.pgen.0040027.sg004 (2.8 MB TIF).

Figure S5. DNA Methylation and siRNA Levels in the coilin Mutants and Overexressor (A) Southern blot analysis examining DNA methylation at the 5S rDNA repeats. Genomic DNA was digested with methyl-sensitive enzymes HaeIII (GG mCC) or HpaII (C mCGG). The triple DNA methyltransferase mutant dml1 dml2 cmt3 is a control for the loss of DNA methylation. (B) Southern blot analysis examining DNA methylation at MEA-ISR. Genomic DNA was digested with MspI. The methylated DNA band is labeled “M”, and the unmethylated band is labeled “U”. (C) Northern blot analysis examining levels of siRNAs corresponding to the 45S and 5S rDNA repeats. The microRNA miR159 was used as a loading control. (D) Southern blot analysis examining 5S rDNA methylation levels in the ncb-1 del3 double mutant. Genomic DNA was digested with HaeIII. Two independent biological replicates of del3 and ncb-1 del3 were examined. Found at doi:10.1371/journal.pgen.0040027.sg005 (2.7 MB TIF).

Figure S6. NRPD2 Nuclear Bodies in Arabidopsis Nuclei Different NRPD2 localization patterns relative to AGO4 within Myc-AGO4 nuclei are shown. The white arrow indicates an AGO4 focus that did not colocalize with NRPD2. The yellow arrow shows an NRPD2 nuclear body that did not colocalize with AGO4. Nuclei containing observable AGO4 or NRPD2 foci were examined. Found at doi:10.1371/journal.pgen.0040027.sg006 (2.4 MB TIF).

Figure S7. DNasel Controls for the DNA FISH Experiments Sample slides were treated with DNasel (7.5 U, Roche) for 30 min at room temperature after primary antibody incubation and postfixing for DNA FISH. DNasel treated samples did not show hybridization with the 45S rDNA, 5S rDNA, or GCN probe. Found at doi:10.1371/journal.pgen.0040027.sg007 (2.1 MB TIF).

Figure S8. Localization of AGO4 or Cajal Body Relative to the NORS (A) 45S rDNA FISH combined with Myc-AGO4 immunofluorescence analysis. Nuclei containing NORS surrounded by two AGO4 foci (top panel) or three AGO4 foci (bottom panel) were observed. The remaining 80% of nuclei contained one AGO4 focus per NOR. Only nuclei containing NORS with adjacent AGO4 foci were examined. (B) Localization of GFP-SmD3 or GFP-SmB relative to the condensed 45S rDNA loci. A polyclonal antibody to GFP was used to detect GFP-SmD3 or GFP-SmB within nuclei. Found at doi:10.1371/journal.pgen.0040027.sg008 (1.6 MB TIF).

Figure S9. DRM2 and U2B̂ Localization DRM2 localization relative to the Cajal body was examined in DRM2-Myc nuclei. Nuclei containing an observable Cajal body marked by U2B̂ immunostaining were examined. Found at doi:10.1371/journal.pgen.0040027.sg009 (880 KB TIF).

Table S1. Nuclei Counts for SmD3 or SmB Localization Relative to AGO4 Found at doi:10.1371/journal.pgen.0040027.st001 (71 KB PDF).

Table S2. Nuclei Counts for HYL1 Localization Found at doi:10.1371/journal.pgen.0040027.st002 (68 KB PDF).

Table S3. Nuclei Counts for the coilin Mutant Found at doi:10.1371/journal.pgen.0040027.st003 (93 KB PDF).

Table S4. Nuclei Counts for NRPD1b Localization in ago1 and nrdp2 Mutants Found at doi:10.1371/journal.pgen.0040027.st004 (84 KB PDF).

Table S5. Nuclei Counts for NRPD2 Localization Found at doi:10.1371/journal.pgen.0040027.st005 (79 KB PDF).


Table S7. Nuclear Foci Counts for 45S, 5S, and CEN FISH Experiments Found at doi:10.1371/journal.pgen.0040027.st007 (70 KB PDF).

Table S8. Nuclei Foci Counts for NOR2 and NOR4 Found at doi:10.1371/journal.pgen.0040027.st008 (68 KB PDF).


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Author contributions. CFL, IRH, and SEJ conceived and designed the experiments. CFL and IRH performed the experiments and analyzed the data. IRH generated the DRM2-Myc line and performed DRM2-Myc immunofluorescence studies. LS generated the GFP-HYL1 plasmid. TL provided the NRPD1b antibody. LS, NF, and TL contributed reagents/materials/analysis tools. CFL and SEJ wrote the paper.

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References


