Conversion of pre-RISC to holo-RISC by Ago2 during assembly of RNAi complexes

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ABSTRACT

In the *Drosophila* RNA interference (RNAi) pathway, small interfering RNAs (siRNAs) direct Argonaute2 (Ago2), an endonuclease, within the RNA-induced silencing complex (RISC) to cleave complementary mRNA targets. In vitro studies have shown that, for each siRNA duplex, RISC retains only one strand, the guide, and releases the other, the passenger, to form a holo-RISC complex. Here, we have isolated a new Ago2 mutant allele and provide, for the first time, in vivo evidence that endogenous Ago2 slicer activity is important to mount an RNAi response in *Drosophila*. We demonstrate in vivo that efficient removal of the passenger strand from RISC requires the cleavage activity of Ago2. We have also identified a new intermediate complex in the RISC assembly pathway, pre-RISC, in which Ago2 is stably bound to double-stranded siRNA.

Keywords: RNAi; RISC assembly; Ago2; siRNA; *Drosophila*

INTRODUCTION

RNA interference (RNAi) is a post-transcriptional gene silencing process during which endogenous messenger RNA is destroyed upon introduction of the corresponding double-stranded RNA (dsRNA) (Fire et al. 1998). RNAi and other forms of RNA silencing have been observed in a wide variety of organisms such as animals, plants, fungi, and protozoa (Meister and Tuschl 2004).

The initiation of RNAi occurs with the processing of dsRNA into siRNAs (small interfering RNAs), by an RNase III enzyme called Dicer (Meister and Tuschl 2004). In *Drosophila*, Dicer-2 (Dcr-2) generates these duplex siRNAs. An siRNA then binds to a heterodimer of R2D2 and Dcr-2 proteins to form a R2D2/Dcr-2 initiator complex (RDI) (Liu et al. 2003; Lee et al. 2004; Pham et al. 2004; Tomari et al. 2004a). It has been shown that RDI orients the siRNA duplex so that the 5' end with lower melting temperature preferentially interacts with Dcr-2 (Tomari et al. 2004b). The strand with the lower melting temperature becomes the single-stranded guide RNA of RISC (RNA-induced silencing complex). RISC uses this single-stranded guide RNA to find

complementary mRNA and endonucleolytically cleave target mRNA. Previous studies have identified Ago2 as the catalytic subunit of RISC cleavage activity (Liu et al. 2004; Song et al. 2004). The Piwi domain of Ago2 is a RNase H-like endonuclease that cleaves a single phosphodiester bond in the target RNA backbone (Liu et al. 2004; Song et al. 2004).

One of the essential steps of RISC formation is dissociation of the paired siRNA strands, and selective retention of one strand (guide) and loss of the other (passenger) from RISC. It has long been thought that double-stranded siRNAs are unwound by an ATP-dependent RNA helicase, activated or recruited with formation of RDI. Consistent with the idea, assembly of RISC in a *Drosophila* embryo lysate is reduced upon depletion of ATP (Nykanen et al. 2001). Likely candidates have been identified using both genetic and biochemical studies, yet their biochemical roles have not been demonstrated.

Recent biochemical studies have presented a model in which the endonucleolytic activity of Ago2 is responsible for passenger strand dissociation in vitro by means of passenger strand cleavage (Matranga et al. 2005; Rand et al. 2005; Leuschner et al. 2006). In this paper, we used a genetic screen to isolate a new Ago2 allele and demonstrate in vivo that efficient removal of the passenger strand from RISC requires the cleavage activity of Ago2. With this mutant allele, we have also identified a new intermediate complex in the RISC assembly pathway in which Ago2 is stably bound to siRNA before unwinding.

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RESULTS

Genetic identification of *Ago2*^{V966M} mutant allele

We previously performed an EMS-induced mutagenesis screen to identify new and essential components of the RNAi pathway in *Drosophila* (Lee et al. 2004). Using eye-specific mitotic recombination, we screened animals that had homozygous mutant compound eyes, while all other tissues, including the germ-line, were heterozygous (Stowers and Schwarz 1999; Newsome et al. 2000). Animals with constitutive RNAi against the *white* gene were examined for variants that had de-repressed *white* gene activity in the eye (Fig. 1A,B).

Our screen of the third chromosome identified a mutation that partially de-repressed *white* gene activity (Fig. 1C,D). Animals that were homozygous for the mutation also exhibited *white* de-repression, and they were both viable and fertile. Complementation analysis with known viable RNAi mutants was performed. The mutation failed to complement a null *Ago2* mutation, *Ago2*⁴¹⁴ (Fig. 1E,F; Okamura et al. 2004). In addition, a single copy of an *Ago2* genomic transgene (Okamura et al. 2004) rescued the

RNAi silencing defect of the mutant chromosome (Fig. 1G,H). Sequencing of the *Ago2* gene from the mutant revealed a base substitution that results in a valine to methionine amino acid substitution in the Piwi domain (Fig. 2B). Based on all of these data, we have identified the mutation as an allele of *Ago2* and have named it *Ago2*^{V966M}. The *white* (RNAi) phenotype of *Ago2*^{V966M} was not as strong as the *Ago2* null phenotype, suggesting that it is not a null mutation. Consistent with this notion, Western blot analysis with anti-Ago2 antibody revealed that there was a normal level of Ago2 protein present in the mutant flies (Fig. 1I).

The Piwi domain of an archaeobacterial Ago (Pf-Ago) protein has been shown to resemble the catalytic domain of RNase H, an RNase that cleaves the RNA strand of RNA/DNA hybrid duplexes (Song et al. 2004). There are two sequence motifs, a GxDV and an RDG motif, within the Piwi domain that are highly conserved in eukaryotic Ago proteins (Fig. 2A). The two aspartate residues of these motifs are structurally equivalent to two aspartate residues that coordinate a metal ion in RNase H (Yang and Steitz 1995). Critically, the metal ion is at the catalytic core of the

RNase H enzyme. A third coordinating carboxylate varies in its position within the active site of RNase H. Studies of Pf-Ago suggested that a glutamate in close proximity to the two aspartates was the third coordinating residue (Parker et al. 2004). However, a recent structural study has determined a nearby histidine residue to be the third residue of Pf-Ago (Rivas et al. 2005). Consistent with these structural predictions based on Pf-Ago, mutation of the histidine or either aspartate residue in human Ago2 abolishes the RNA cleavage activity of RISC (Rivas et al. 2005).

The mutation of *Ago2*^{V966M} changes the GxDV motif into GxDM. This valine residue is highly conserved among all Ago proteins. Modeling of a GxDM variant of the Pf-Ago structure (Song et al. 2004) with PyMol revealed a steric change in the integrity of the structure at the catalytic site (data not shown). We suspect that the V-to-M mutation may alter the metal coordinating properties required for normal catalysis. Recent study of the crystal structure of bacterial Rnase H suggests a two metal ion mechanism (Nowotny et al. 2005). Two metal ions found in the active site of RNase H are important for activating the nucleophile and stabilizing the transition state. In addition,

FIGURE 1. Phenotype of the *Ago2*^{V966M} MutantCompound eyes from *white*⁺ adult flies (A–H). (A) An *Ago2*⁺ adult. (B) An *Ago2*⁺ adult with one copy of *GMR-wIR*. The *white* gene is partially silenced, resulting in a pale orange eye color. (C) An adult with one copy of *GMR-wIR* and heterozygous for *Ago2*^{V966M}. (D) An adult with two copies of *GMR-wIR* and homozygous for *Ago2*^{V966M}. (E) An adult with two copies of *GMR-wIR* and homozygous for *Ago2*⁴¹⁴. (F) An adult with two copies of *GMR-wIR* and *trans*-heterozygous for *Ago2*^{V966M} and *Ago2*⁴¹⁴. (G) An *Ago2*⁴¹⁴ adult homozygote with two copies of *GMR-wIR* and a single copy of the *Ago2* rescue transgene. (H) An *Ago2*^{V966M} adult homozygote with two copies of *GMR-wIR* and a single copy of the *Ago2* rescue transgene. (I) Lysates prepared from wild-type, *Ago2*^{V966M}, *Ago2*⁴¹⁴, and *Ago2*^{V966M}/*Ago2*⁴¹⁴ embryos were separated by SDS-PAGE, blotted onto nitrocellulose membrane, and subjected to Western analysis using antibodies against TSN, VIG, dFXR, and Ago2.

FIGURE 2. Structural characteristics of Ago2^{V966M}. (A) Protein sequence alignment of Argonaute Piwi domains of *Drosophila* (Dm), Human (Hs), *Pyrococcus furiosus* (Pf). Aligned sequences are color-coded: 100% conserved residues (black), 75% conserved residues (light gray), and well-conserved hydrophobic or hydrophilic residues (dark gray). The two conserved motifs, GxDV and RDG, are indicated with a bar below the sequence. Three conserved catalytic residues within the Piwi domain are marked with circles at the top of the alignment. The mutation in Ago2^{V966M} is marked with an arrow above the alignment. (B) Schematic representation of *Drosophila* Ago2 protein with known domains. The mutation of Ago2^{V966M} is labeled.

efficient catalysis takes place only when the two metal ions are separated by an ideal distance and arranged in appropriate geometric positions (Nowotny et al. 2005). The valine residue affected in Ago2^{V966M} may disrupt the coordination of the metal ion required for RNA cleavage activity. Alternatively, the mutation may disrupt the positioning of target substrate within the catalytic site.

Ago2^{V966M} is defective in holo-RISC formation

To determine how RNAi is defective in the Ago2^{V966M} mutant, we performed an in vitro target cleavage assay. Embryo lysates, either from Ago2^{V966M} homozygotes or Ago2^{V966M}/Ago2⁴¹⁴ heterozygotes, were greatly impaired in their ability to cleave target mRNAs (Fig. 3A). This suggested that holo-RISC is not functional and/or the mutant is unable to form a structurally intact holo-RISC.

To test if Ago2^{V966M} is able to form holo-RISC, we analyzed RISC formation in Ago2^{V966M} mutant embryo lysate by native gel electrophoresis of siRNA–protein complexes. Ago2^{V966M} lysate formed a complex closely corresponding to holo-RISC (Fig. 3B). However, the complex was of slower electrophoretic mobility when compared with wild-type holo-RISC (Fig. 3B). Lysate made from Ago2^{V966M}/Ago2⁴¹⁴ embryos showed reduced abundance of the slower complex, indicating that its formation is dependent upon Ago2 (Fig. 3B).

Ago2^{V966M} forms a pre-RISC complex containing duplex siRNA

We examined RISC formation using a different siRNA, which is highly asymmetric for siRNA strand retention into holo-RISC. This siRNA preferentially retains one strand (top) over the other strand (bottom) due to a G:U wobble at the 5' end of the guide strand (Fig. 3C). Upon incubation in wild-type lysate, a siRNA containing radiolabeled guide strand formed two discrete low-mobility complexes (Fig. 3C). The slower complex (S) was less abundant than the faster complex (F). Ago2^{V966M} mutant lysate also formed the two low-mobility complexes, but there were higher levels of the S complex relative to the F complex. When a siRNA containing radiolabeled passenger strand was incubated in wild-type or Ago2^{V966M} lysate, only the S complex was observed, with more of this complex observed in the mutant reaction (Fig. 3C). The inability to observe the F complex with labeled passenger strand indicates that the F complex selectively associates with guide strand RNA. In contrast, the S complex associates with either strand equivalently. This would suggest that the F complex contains unwound single-stranded siRNA whereas the S complex contains duplex siRNA.

If our hypothesis is correct about the RNA composition of the two RISC complexes, then we would predict that target mRNA (complementary to the guide siRNA strand) should be able to associate with the F complex but not the S complex. We performed a RISC formation assay with

FIGURE 3. Biochemical characterization of Ago2^{V966M}. (A) In vitro target RNA cleavage assay. Lysates from wild-type, Ago2^{V966M}, Ago2⁴¹⁴, and Ago2^{V966M}/Ago2⁴¹⁴ embryos were incubated with 5'-labeled *Pp-luc* target mRNA and unlabeled cognate siRNA duplex, as indicated. The 5' product from RISC activity is indicated and comigrates on the gel with a cleavage product from siRNA-independent nuclease activity. (B) RISC formation assay. Lysates from wild-type and Ago2 mutant embryos were incubated with labeled *Pp-luc* siRNA, and complexes were resolved by native gel electrophoresis. RDI and RLC intermediate complexes are indicated. The complex (arrow) closely corresponding to holo-RISC in Ago2^{V966M} mutant was of slower electrophoretic mobility when compared with wild-type holo-RISC. Ago2^{V966M}/Ago2⁴¹⁴ embryo lysate showed reduced abundance of the slower complex. (C) RISC formation with a highly asymmetric siRNA. Either the guide (top) or the passenger (bottom) strand was 5'-radiolabeled. Both wild-type and Ago2^{V966M} lysates form two discrete low-mobility complexes when incubated with radiolabeled guide strand. In wild-type lysate, the slower complex (S) was less abundant than the faster complex (F). In Ago2^{V966M} lysate, the relative amount of S complex increased while F complex decreased. When incubated with radiolabeled passenger strand, both wild-type and Ago2^{V966M} lysates formed only the S complex due to asymmetric loading of unlabeled guide strand into holo-RISC. Note that more of the S complex was formed in Ago2^{V966M} lysate. (D) Radiolabeled guide siRNA strand was incubated with either unlabeled cognate or noncognate 2'-O-methyl target mRNA in wild-type lysate. Addition of cognate target mRNA resulted in the depletion of the F complex, whereas the S complex was not affected. Addition of noncognate mRNA had no effect on either complex.

guide-labeled siRNA and added an unlabeled 2'-O-methylated target mRNA. Addition of cognate target mRNA resulted in depletion of the F complex, presumably due to a shift in complex mobility, whereas the S complex was unchanged (Fig. 3D). Addition of noncognate mRNA had no effect on either complex. This result indicates that the F complex has the potential to specifically associate with a target mRNA, which is a property of holo-RISC.

To directly visualize the nature of the siRNAs in the two complexes, we performed a two-dimensional gel electrophoresis experiment (Fig. 4A). A RISC formation reaction

was performed with guide-labeled siRNA. After resolving the complexes in the first dimension by native gel electrophoresis, we excised the entire lane and placed it horizontally on a SDS-polyacrylamide gel. Labeled siRNAs from complexes were then subjected to electrophoresis in the second dimension. We predicted that ribonucleoprotein complexes resolved in the first dimension would dissociate in the second dimension due to SDS-dependent denaturation. This would permit siRNAs to electrophorese as free species. However, SDS treatment in the second dimension did not denature siRNA duplexes, which were easily distinguishable from single-stranded siRNA (Fig. 4B). Thus, we were able to

FIGURE 4. *Ago2*^{V966M} RISC complexes contain primarily duplex siRNA. (A) Schematic representation of two-dimensional gel electrophoresis. RISC formation reaction was performed with guide-labeled siRNA duplex. The complexes were resolved in the first dimension by native gel electrophoresis. The entire lane was then excised and placed horizontally on a 15% SDS-polyacrylamide gel. Labeled siRNAs from complexes were then subjected to electrophoresis in the second dimension. (B) Both wild-type and *Ago2*^{V966M} lysates show RDI and RLC complexes that contain duplex siRNA. The low-mobility complexes observed in wild-type lysate primarily contain single-stranded siRNA while those formed in *Ago2*^{V966M} lysate primarily contain duplex siRNA.

visualize the conformation of siRNAs associated with various complexes formed in the RISC reaction.

When wild-type lysate was subjected to two-dimensional analysis, we observed that both RDI and RLC (RISC loading complex) complexes contained duplex siRNA (Fig. 4B), which is consistent with previous observations (Pham et al. 2004). The S and F complexes contained single-stranded siRNA though some duplex siRNA was also detected (Fig. 4B). However, when *Ago2*^{V966M} lysate was analyzed, the low-mobility complexes predominantly contained duplex siRNA with a peak of single-stranded siRNA detected in the F complex (Fig. 4B). Thus, Ago2 stimulates conversion of duplex siRNA into single-stranded siRNA within RISC complexes.

Ago2^{V966M} is defective for unwinding of siRNAs and passenger strand cleavage

Based on all of our observations, it is likely that the F complex corresponds to holo-RISC. Several lines of evidence are offered in support. First, formation of the F complex requires Ago2, a key subunit of holo-RISC. Second, both holo-RISC and the F complex contain single-stranded siRNA, and both preferentially retain the guide strand of a siRNA duplex. Finally, holo-RISC and the F complex are able to specifically bind to target mRNA substrate. Our native gel analysis has also identified

a distinct Ago2-dependent complex, the S complex. The formation of the S complex requires the presence of Ago2 and is more pronounced with deficient catalytic activity of Ago2, which is in contrast to holo-RISC. The S complex appears to contain siRNA in duplex form, and it is unable to bind to target mRNA substrate.

A simple model is that the S complex is a holo-RISC precursor that contains duplex siRNA, and siRNA unwinding accompanies the conversion of S complex to holo-RISC. If this model is correct, then siRNA unwinding should correlate with complex conversion. Therefore, we determined whether *Ago2*^{V966M} mutant lysate shows defective siRNA-unwinding activity comparable to its defective holo-RISC-forming activity. When we examined the rate of single-stranded siRNA production, *Ago2*^{V966M} mutant lysate showed a two-fold reduced rate of siRNA unwinding (Fig. 5A). A null *Ago2* mutant did not produce any single-stranded siRNAs. The strength of the *Ago2*^{V966M} mutant defect is comparable to the relative changes in S and F complex abundance in the mutant (Fig. 3C). Taken together, our data most strongly support a model in which the S complex is a holo-RISC precursor. Thus, we have renamed it pre-RISC.

Recent in vitro studies have shown that unwinding of siRNA duplex is accompanied by passenger strand cleavage to form holo-RISC (Matranga et al. 2005; Rand et al. 2005; Leuschner et al. 2006). Since *Ago2*^{V966M} shows defective siRNA unwinding activity, it was possible that passenger

FIGURE 5. Ago2^{V966M} is defective in siRNA duplex dissociation. (A) Guide-labeled siRNA duplex was incubated in wild-type, Ago2^{V966M}, or Ago2⁴¹⁴ lysate. Reactions were resolved on a 15% SDS-polyacrylamide gel and radioactive signal was quantified by PhosphorImager. The percentage of single-stranded siRNA generated from siRNA duplex was plotted against reaction time. (B) Passenger-labeled siRNA duplex was incubated in the same lysates as A. Reactions at various time points were removed, deproteinized, and electrophoresed to visualize passenger strand cleavage products. The 9-nt 5' cleavage product accumulated early in the reaction and disappeared. Ago2^{V966M} mutant lysate produced little passenger strand cleavage product. The percentage of the 9-nt product cleaved from labeled passenger strand was plotted against reaction time.

strand cleavage is also inhibited. To test if Ago2^{V966M} is able to convert passenger strand siRNAs into 9-nucleotide (nt) cleavage products, we performed an in vitro passenger strand cleavage assay. Cleavage of 5'-labeled passenger strand by wild-type lysate yielded a 9-nt labeled product (Fig. 5B). However, in the Ago2^{V966M} mutant lysate, the rate of passenger strand cleavage was strongly reduced (Fig. 5B). These data further support a model where siRNAs are initially loaded as duplexes into Ago2-containing pre-RISC. The catalytic activity of Ago2 is then required to cleave the passenger strand, stimulating it to dissociate from the guide strand and form holo-RISC.

DISCUSSION

We have isolated a mutation in *Drosophila* Ago2 that specifically affects its ability to cleave target RNA when the target is associated with holo-RISC. The mutation changes a highly conserved amino acid residue that neighbors one of the nucleophile-coordinating residues within the Piwi domain. Structural modeling predicts that the change disrupts precise positioning of the nucleophile within the catalytic site. Experimentally, this change results

in an eightfold reduction in target cleavage activity by holo-RISC. The experimental observations are consistent with the structural predictions and provide further evidence that Ago2 has RNA slicer activity in vitro. Moreover, it provides, for the first time, in vivo evidence that endogenous Ago2 slicer activity is important to mount an RNAi response in *Drosophila*.

The assembly of RISC in vitro follows a pathway of siRNA-protein complexes from RDI to RLC to holo-RISC (Fig. 6). A previous model proposed RLC as the direct precursor to holo-RISC, which harbors both Ago2 and single-stranded guide siRNA (Pham et al. 2004; Tomari et al. 2004a; Pham and Sontheimer 2005). In the present study we have identified a new Ago2-dependent complex, pre-RISC. The difficulty of visualizing pre-RISC in *Drosophila* embryo extract explains why this complex was not described earlier. One reason for this difficulty requires an understanding of the differences between pre-RISC and holo-RISC. The pre-RISC complex contains duplex siRNA whereas holo-RISC contains just the guide strand of the siRNA, the passenger strand having dissociated from holo-RISC. Thus, a key difference between pre-RISC and holo-RISC is the conformation of the associated siRNA.

FIGURE 6. Model for RISC assembly in *Drosophila* in vitro. The siRNA is bound to R2D2/Dcr2 heterodimer to form a RDI complex. The RDI complex then enters a RISC-loading complex (RLC), associating with other RNAi components. RLC then recruits Ago2 to form a pre-RISC complex that contains duplex siRNA. Endonucleolytic activity of Ago2 within the Piwi domain promotes release of the passenger strand from pre-RISC to form holo-RISC. Holo-RISC can then base-pair to complementary single-stranded mRNA substrates for cleavage.

Two simple models could explain the relationship between pre-RISC and holo-RISC. Pre-RISC could be an abortive reaction product in the holo-RISC assembly reaction, formed independently of holo-RISC itself. Alternatively, pre-RISC could be an intermediate complex in the assembly reaction, and act as a holo-RISC precursor. Pulse-chase experiments to discern between these models are not possible given our inability to specifically pulse-label pre-RISC. However, the *Ago2*^{V966M} mutant provides evidence that pre-RISC is a holo-RISC precursor. Mutant extract shows a threefold increase in pre-RISC abundance and twofold decrease in holo-RISC abundance. Furthermore, siRNA strand unwinding is impaired twofold. The most parsimonious explanation for these results is that pre-RISC converts into holo-RISC by siRNA strand dissociation, and the Ago2 mutant is partially impaired for this conversion, resulting in a build-up of the precursor. Our results suggest a revised model in the RISC assembly pathway (Fig. 6).

Recent in vitro studies have shown that unwinding of siRNA duplex is accompanied by cleavage of the ejected

siRNA strand during holo-RISC formation (Matranga et al. 2005; Rand et al. 2005). In the *Ago2*^{V966M} mutant, the rate of siRNA strand cleavage and unwinding is reduced. These data support a model where siRNAs are initially loaded as duplexes into Ago2-containing pre-RISC. The catalytic activity of Ago2 is then required to cleave one strand of the duplex, stimulating it to dissociate from the uncleaved strand and ultimately dissociate from RISC. The end result is a RISC retaining one siRNA strand, fully competent to down-regulate transcript targets.

If Ago2-dependent catalysis is obligatory for complex conversion, then we might have expected a more profound assembly block in the *Ago2*^{V966M} mutant than the effect actually observed. One explanation is that conversion of pre-RISC to holo-RISC is promoted by several factors including Ago2, with each factor augmenting strand dissociation. For example, it is conceivable that Ago2-dependent siRNA cleavage leads to some dissociation by passive mechanisms such as temperature whereas other factors such as helicases generate unwinding by other means. Alternatively, strand dissociation might be wholly directed by helicase factors, and strand-cleavage by Ago2 prevents stable reassociation of the strands. The identity of such a putative helicase remains unknown. The RNA helicase Armitage is required for RISC assembly (Tomari et al. 2004a) but seems to act in the RLC to pre-RISC step of the pathway before unwinding has occurred (data not shown).

MATERIALS AND METHODS

Fly genetics

The *Ago2*^{V966M} mutant was isolated from a genetic screen as described (Lee et al. 2004). The *GMR-wIR* transgene has been described (Lee and Carthew 2003). Mosaic analysis of the adult eye was performed as described (Lee et al. 2004). *Ago2*⁴¹⁴ null and *Ago2*⁺ rescue flies were gifts from Mikiko Siomi (Okamura et al. 2004). Complementation analysis was performed by crossing *Ago2*^{V966M} mutant flies to *Ago2*⁴¹⁴ mutant flies in *GMR-wIR* background.

Biochemical methods

Embryo lysate preparation and target RNA cleavage assay were performed as described (Pham et al. 2004). RISC assembly was assayed by native gel electrophoresis as described (Pham and Sontheimer, 2005). Passenger cleavage reactions were performed as described (Matranga et al. 2005).

Target binding assay

We used a cognate *Pp-luc* 2'-O-methyl RNA analog with the sequence 5'-AUCAGUACGGCGGAAUACUUCGAdAdA-3' and a noncognate 2'-O-methyl RNA with the sequence 5'-AUCAAG-GAAAGUCAUAUGUUAUGGdAdA-3'. RISC assembly reactions

were performed with ^{32}P -labeled *Pp-luc* siRNA duplex for 15 min in 25°C. After addition of 1 μM cognate or noncognate 2'-O-methyl RNA, reactions were further incubated at 25°C for additional 45 min and then analyzed on a 4% (40:1) native polyacrylamide gel as described (Pham and Sontheimer 2005).

Two-dimensional gel electrophoresis

RISC was assembled by adding ^{32}P -labeled siRNAs to embryo lysates and resolved by native electrophoresis as described (Pham and Sontheimer 2005) in the first dimension. The entire lane was excised and placed horizontally over 15% SDS-polyacrylamide gel. This was then subjected to electrophoresis in the second dimension at 10 watts at 4°C.

Unwinding assay

^{32}P -labeled siRNA duplexes were added to the RISC assembly assay reactions as described above and incubated at 25°C. Reactions were quenched with 1 μL of a heparin mix (Pham et al. 2004) and loaded onto 15% SDS-polyacrylamide gel and then subjected to electrophoresis at 10 watts at 4°C.

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