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Small RNAs meet their targets: When methylation defends miRNAs from uridylation

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Small RNAs are incorporated into Argonaute protein-containing complexes to guide the silencing of target RNAs in both animals and plants. The abundance of endogenous small RNAs is precisely controlled at multiple levels including transcription, processing and Argonaute loading. In addition to these processes, 3' end modification of small RNAs, the topic of a research area that has rapidly evolved over the last several years, adds another layer of regulation of their abundance, diversity and function. Here, we review our recent understanding of small RNA 3' end methylation and tailing.

Introduction

Small silencing RNAs (small RNAs) of 20–30 nucleotide (nt) in length are key regulators of gene expression in both animals and plants. Together with their effector protein called Argonaute (AGO), small RNAs repress gene expression at either the transcriptional or the post-transcriptional level and play important roles in various biological processes, such as cell differentiation and transgenerational inheritance. Small RNAs can be classified into microRNAs (miRNAs), small interfering RNAs (siRNAs) and Piwi (a class of AGO protein)-interacting RNAs (piRNAs) based on their origin, biogenesis and/or the type of AGO they are associated with, although the boundaries of this classification become blurred and sometimes even difficult to discern.^{1,2} Dicer-like proteins, a class of RNaseIII-type

enzymes, release miRNAs and siRNAs from their precursors as duplexes with 2 nt 3' overhangs at each end.³ By contrast, piRNAs are processed via a Dicer-independent pathway.⁴ More details regarding the biogenesis and function of small RNAs have been reviewed elsewhere.^{3,4} After biogenesis, plant small RNAs as well as some animal small RNAs are 2'-*O*-methylated at their 3' end by HUA ENHANCER1 (HEN1) in plants or its homologues in animals. 2'-*O*-methylation is critical for small RNA stability. Besides methylation, small RNAs are also subject to 3' untemplated nucleotide addition (tailing), which in turn affects their function and stability. In this point of view, we summarize our recent understanding on small RNA methylation and tailing and discuss their biological relevance.

Methylation of small RNAs

Plant miRNAs and siRNAs contain a 2'-*O*-methyl group at their 3' end. This modification is catalyzed by HEN1.^{5,6} HEN1 contains 2 double-stranded RNA binding domains (dsRBD1 and dsRBD2), a La-motif containing domain (LCD) and a methyltransferase domain (MTase).⁷ HEN1 specifically recognizes 21–24 base-pair (bp) dsRNAs with 2nt overhangs and deposits a methyl group to the 2' OH position of the 3' end in each strand *in vitro*.⁸ Structural analysis reveals that HEN1 recognizes dsRNAs via its dsRBD1 and dsRBD2 domains. The substrate length of HEN1 is determined by the distance between the LCD domain and the MTase domain.⁷ Loss of 2'-*O*-methylation in the

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hen1 mutants not only leads to the reduced small RNA abundance but also results in 3' end untemplated uridine addition (uridylation) and trimming of small RNAs.⁹⁻¹¹ These results demonstrate that methylation stabilizes small RNAs by preventing uridylation and degradation. A single uridine addition at miR171 in the *hen1* mutant results in the production of secondary siRNAs from target RNAs that undergo miR171-mediated cleavage.¹¹ This result indicates that methylation may also block miRNA-dependent secondary siRNA production, and, therefore, protects plants from undesired regulation by small RNAs.

HEN1 is an evolutionarily conserved protein in eukaryotes and bacteria.¹² In bacteria and animals, HEN1 lacks dsRNA-binding domains and methylates single-stranded RNAs.^{13,14-21} In bacteria, Hen1 is involved in an RNA repair process.^{22,23} In animals, HEN1 methylates some small RNAs in a germ cell-specific manner.¹⁴⁻²¹ Intriguingly, among 297 animals homozygous for zebrafish *hen1*, only one was identified as female, suggesting a critical role of HEN1 in sex determination.¹⁸ In mice, HEN1 is specifically expressed in testes and methylates piRNAs.^{15,17} In addition, the zebrafish HEN1 is expressed both in ovary and testis, and is required for piRNA stability.¹⁸ Besides piRNAs, AGO2-associated siRNAs in *Drosophila* and ERGO-1 (a PIWI clade AGO protein)-associated 26G RNAs in *C. elegans* are also subject to 2'-*O*-methylation by DmHEN1/Pimet (piRNA methyltransferase) and HENN-1 (HEN1 of Nematodes 1), respectively, in a germline-specific manner. In all cases, loss-of-function mutations in HEN1 cause uridylation and 3'-to-5' exonucleolytic trimming of small RNAs,^{16,18-21} reminiscent of the observations in *Arabidopsis*.⁹⁻¹¹ Animal HEN1 homologs appear to act on AGO-bound small RNAs in vivo and fly HEN1 interacts with PIWI,^{14,16} suggesting that the physical interaction between HEN1 and AGOs may determine its substrate specificity in animals.

Uridylation of small RNAs

In contrast to piRNAs, animal miRNAs are not methylated. Comprehensive

small RNA deep sequencing analyses reveal that uridylation or adenylation (usually mono- or di-nucleotide additions) at the 3' end of miRNAs is widespread and conserved across a variety of animal species.²⁴⁻²⁶ Uridylation of small RNAs is catalyzed by the terminal uridylyltransferases (TUTase) (Fig. 1). While many TUTases show overlapping activities on some miRNAs, some of them act on miRNAs in a sequence-specific manner.^{24,26,27} Uridylation often serves as an RNA decay signal whereas adenylation appears to increase RNA stability.²⁸ Besides stability, 3' modification can also regulate small RNA activities. For example, uridylation of miR-26a prevents the miRNA from repressing its mRNA target, without affecting its abundance.²⁵ PAPD4/GLD-2 mediated 3' end adenylation has little effect on miRNA abundance but reduces their AGO loading efficiency.²⁴ Interestingly, miRNAs derived from the 3' arms of their pre-miRNAs are more frequently uridylated, suggesting that at least some of the uridylation occurs to the 3' end of pre-miRNAs before they are channeled into Dicer processing.²⁴ Indeed, mono-uridylation of group II pre-miRNAs (with a 1nt 3' overhang instead of the canonical 2nt 3' overhang after Droscha cleavage) is crucial for their further maturation.²⁹ In *C. elegans*, 22G secondary siRNAs are not methylated and are extensively oligo-uridylated by CDE-1, a germline-specific ncPAP.³⁰ CDE-1-mediated uridylation of CSR-1 bound 22G siRNAs prevents their over-accumulation through active degradation, which is important for both mitotic and meiotic chromosome segregation.³⁰

Lack of HEN1 function causes extensive tailing and trimming of small RNAs in higher plants.^{9,11} An early study demonstrates that untemplated nucleotides added to the 3' end of small RNAs in *hen1* are predominantly uridines,⁹ which is subsequently shown to be the same in all other tested plant species.⁹⁻¹¹ We recently showed that HESO1 acts in small RNAs uridylation.^{31,32} HESO1 is a TUTase (Fig. 1) and possesses robust RNA poly(U) polymerase activities in vitro.^{31,32} The activity of HESO1 is completely blocked by 2'-*O*-methylation.^{31,32} This may be due to that HESO1

requires 2' OH for its substrate recognition or catalysis. Alternatively, 2'-*O*-methylation may block substrate recognition or catalysis although HESO1 does not need 2' OH for its activity. Loss-of-function mutations in *HESO1* lead to a general reduction of U-tail length in *hen1*, accompanied by an increase of normal-sized, 3' trimmed, and/or short-tailed small RNAs.^{31,32} In contrast, over-expression of *HESO1* in the *hen1* background accelerates miRNA turnover and causes more severe developmental defects.³¹ These results demonstrate that uridylation triggers small RNA destabilization. As in *Arabidopsis*, the *Chlamydomonas* TUTase MUT68 has also been implicated in both siRNA and miRNA uridylation.³³ The increased extent of 3' trimming in the *hen1 heso1* mutants suggests that uridylation may antagonize the action of 3'-to-5' trimming, of which the biological meaning is not yet clear.^{31,32} As 3' truncated miRNA species are tailed by HESO1,³² it is likely that one function of uridylation is to remove 3' truncated miRNAs. In *Drosophila*, the 3'-to-5' exonuclease Nibbler trims a proportion of AGO1 associated miRNAs.^{34,35} There are 2 putative homologous proteins in *Arabidopsis* and it will be interesting to determine whether any of them is involved in 3'-to-5' trimming of unmethylated small RNAs.

Uridylation of 5' cleavage products (5' CPs) generated by small RNA-mediated cleavage

The endonucleolytic cleavage of target RNAs is a mechanism used by miRNAs and siRNAs to silence gene expression. Slicing of target RNAs often happens at a position opposite to the middle of miRNAs and siRNAs and results in a 3' cleavage product (3' CP) and a 5' CP.³⁶ In *C. elegans*, 3' CPs and 5' CPs generated by siRNA-mediated cleavage are further removed by the 5'-to-3' exonuclease XRN1 and the exosome, which degrade RNAs from 5'-to-3' and 3'-to-5, respectively.³⁷ In *Arabidopsis*, the XRN1-LIKE exonuclease XRN4 is responsible for the degradation of 3' CPs generated by miRNA-mediated target cleavage.³⁸ Intriguingly, the 3' ends of 5' CPs are often subject to oligo-uridylation or adenylation modifications in several

evolutionarily distant plant species.³⁹ The Chlamydomonas MUT68 adds adenosines to the 3' end of 5' CP of *MAA7* (the endogenous target of siRNAs generated by a transgenic inverted-repeat sequence) and promotes its degradation from 3'-to-5', likely involving the action of the exosome.⁴⁰ In a recent study, we show that HESO1 is responsible for the uridylation of 5' CPs in Arabidopsis.⁴¹ A reduction in uridylation in *heso1-2* increases the accumulation of 5' CP, demonstrating that uridylation triggers degradation of 5' CP.⁴¹ The proportion of 3' truncated 5' CPs is increased by *heso1* relative to wild-type plants (WT), suggesting that uridylation may cause 5' CP degradation through a mechanism different from 3'-to-5' trimming.⁴¹ This resembles the effect of *heso1* on miRNAs in *hen1*.^{31,32} In humans, uridylation can trigger decapping of histone mRNAs followed by 5'-to-3' degradation.⁴² This may be not the case for 5' CPs, as the status of 5'-to-3' truncation of 5' CPs is not altered by *heso1*. 5' CPs can also be degraded by 5'-to-3' and 3'-to-5' trimming since both 5' and 3' truncated 5' CPs exist.⁴¹ In *xrn4*, the amount of 5' CPs is increased compared with that in WT, revealing that XRN4 is an enzyme responsible for the degradation of 5' CPs although it is not clear whether it acts before or after uridylation.⁴¹ The enzymes that trim 5' CPs from 3'-to-5' remain to be identified.

How does uridylation trigger the degradation of small RNAs and 5' CPs?

Both small RNAs and 5' CPs at least in transient associate with the AGO proteins.

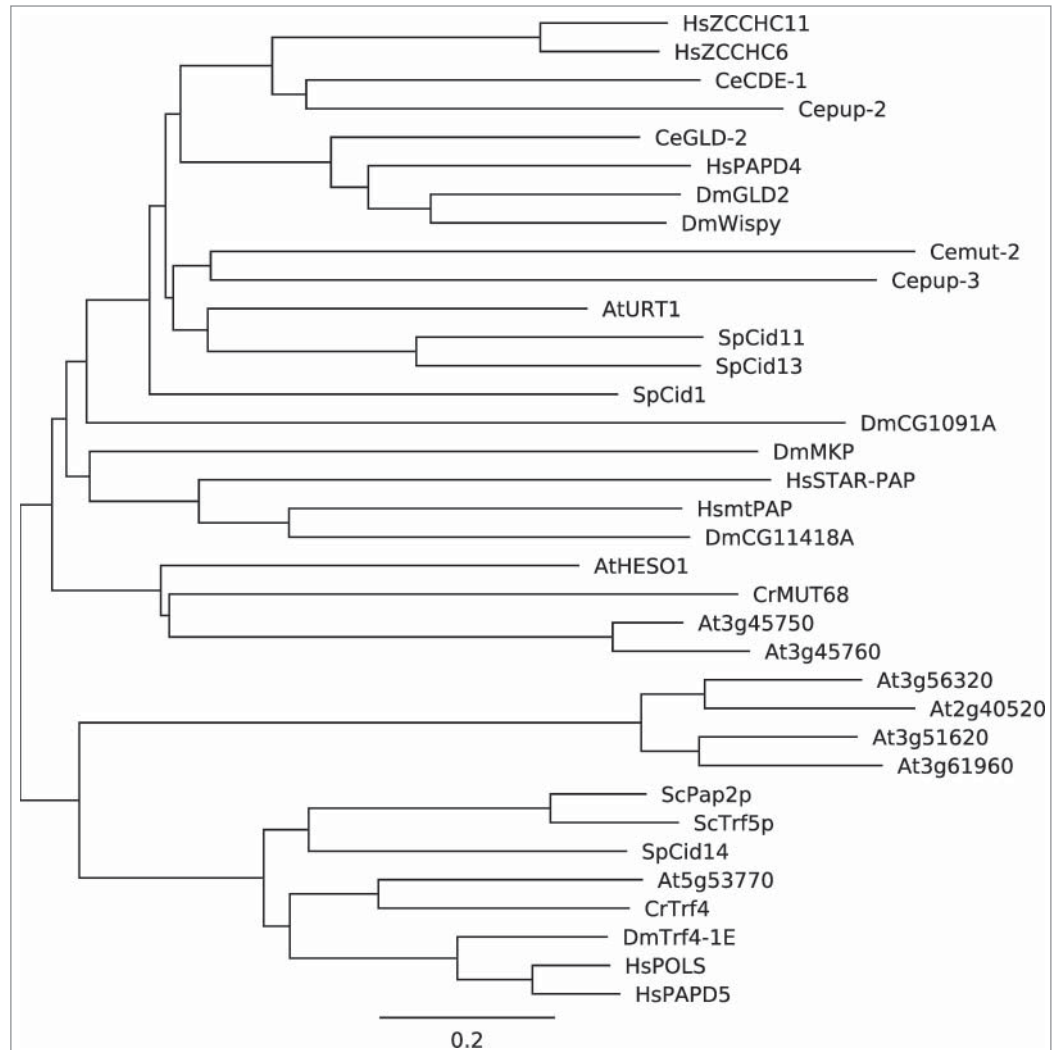


Figure 1. A phylogenetic tree of selected non-canonical poly(A) polymerases. Full-length proteins were downloaded from the Genbank database (<http://www.ncbi.nlm.nih.gov/genbank>). A neighbor joining phylogenetic tree was constructed using MEGA 5.1.52 and displayed using Geneious v7.1.5 (<http://www.geneious.com/>). Hs: Homo sapiens; Ce: Caenorhabditis elegans; Dm: Drosophila melanogaster; At: Arabidopsis thaliana; Sp: Schizosaccharomyces pombe; Cr: Chlamydomonas reinhardtii; Sc: Saccharomyces cerevisiae.

This suggests that a common mechanism may be used to degrade both uridylylated small RNAs and 5' CPs. Agreeing with this notion, the exosome is responsible for the degradation of both uridylylated small RNAs and adenylated 5' CPs in Chlamydomonas.^{33,40} Thus, it is possible that a nuclease may simultaneously target both uridylylated 5' CPs and small RNAs in higher plants. Recently, Dis3l2, which is a 3'-to-5' RNase II nuclease encoded by a paralog of RRP44, a core component of the exosome, was shown to degrade uridylylated pre-let-7 (precursor of let-7 miRNA).^{43,44} In yeast, the Dis3l2

homolog can trigger rapid degradation of uridylylated RNAs in a tail-length dependent manner.⁴⁵ Arabidopsis SOV (SUPPRESSOR OF VARICOSE) is the closest homologous protein of Dis3l2.⁴⁶ It is worth testing if SOV is the enzyme responsible for the degradation of uridylylated small RNAs and 5' CPs in the near future. HESO1 can add long U-tails (Up to hundreds of nt long) to its substrates in vitro.^{31,32,41} However both miRNAs and 5' CPs only contain short U-tails (less than 15 nt) in vivo.^{32,41} This provides an alternative explanation for the degradation of uridylylated miRNAs and 5' CPs: long

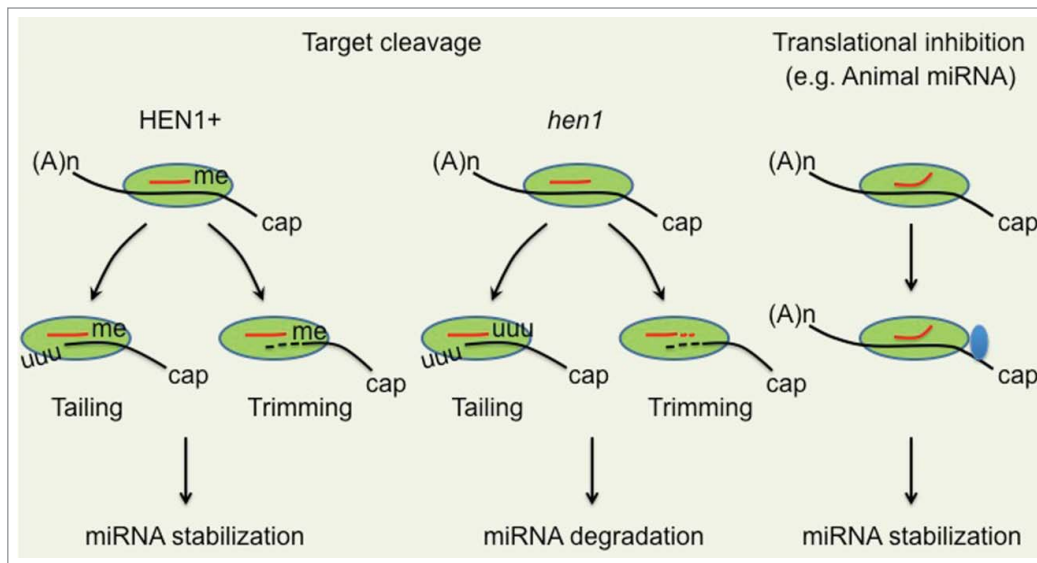


Figure 2. A proposed model for target cleavage mediated small RNA destabilization. In this model, small RNA mediated target cleavage (**A and B**) but not translational inhibition (**C**) may provide a signal in the recruitment both tailing and trimming enzymes to AGO for the modification and further degradation of 5' cleavage products. (**A**) HEN1 mediated 2'-O-methylation protects small RNAs from both tailing and trimming activities. (**B**) In the absence of HEN1, small RNAs become frequently tailed and trimmed. (**C**) In the case of partially complementary target RNA (e.g. animal miRNAs), small RNAs predominantly suppresses gene expression via translational inhibition. In this scenario, tailing and trimming enzymes will be barely recruited to AGO and small RNAs are less tailed and trimmed, regardless of methylated or not. Green oval, AGO protein; Blue oval, translational machinery.

U-tail may disassociate miRNAs and 5' CPs from the AGO1 complex, resulting in their rapid degradation such that no long-tailed products can be detected.

HEN1 protects small RNAs from AGO-associated tailing and trimming activities that normally acts on 5' CPs

The dual activities of HESO1 and MUT68 on both small RNAs and 5' CPs and the fact that both small RNAs and 5' CPs associate with AGO proteins suggest that the terminal uridylyl transferase may recognize their substrates in the AGO complex (Fig. 2). Several evidences demonstrate that HESO1 indeed uridylylates AGO1-associated miRNAs. First, HESO1 associates with AGO1, as revealed by colocalization and co-immunoprecipitation assays.⁴¹ Second, HESO1 is able to uridylylate AGO1-bound miR165/166 in vitro.⁴¹ Third, in *hen1*, AGO1-bound miRNAs are tailed.¹¹ Fourth, suppression of AGO1 function in *hen1* diminishes miRNA uridylation.^{11,41} We envision that HESO1 also recognizes siRNAs in vivo through its physical

interaction with respective AGO proteins. Further analyses show that the PAP domain rather than the unconserved C-terminal domain of HESO1 interacts with the PAZ and PIWI domain of AGO1.⁴¹ These interactions may facilitate HESO1 to recognize its substrates, as PAZ binds the 3' end of small RNAs, which can be released by base pairing between small RNAs and their targets, and PIWI cleaves the targets. In addition, the interaction of terminal uridylyl transferases with AGO may be common among different species since PAP (Fig. 1), PAZ and PIWI are evolutionarily conserved.⁴⁷

Both small RNAs and 5' CPs are also subject to 3'-to-5' trimming activities. By analogy, it is possible that the enzymes catalyzing 3'-to-5' trimming may also associate with AGO proteins. In fact, Nibbler has been shown to interact with AGO in *Drosophila*.³⁴ The tailing and trimming activities are likely recruited to AGO to eliminate the 5' CPs. The presence of these AGO1-associated activities confers the necessity of small RNA methylation by HEN1 to ensure their stability

in the AGO complex in plants (Fig. 2). In flies, AGO2-associated siRNAs are protected by methylation.¹⁶ In contrast, AGO1-associated miRNAs are not methylated¹⁶ and display very limited tailing and trimming. This may be due to the fact that animal miRNAs are less complementary to their targets.⁴⁸ Consistent with this, introduction of artificial target RNAs with high complementarity to some endogenous miRNAs triggers their tailing and trimming.^{49,50} Moreover, siRNAs, which often have highly complementary targets, become tailed and trimmed when misloaded into AGO1.^{49,50} A possible explanation for these observations is that less complementarity between miRNAs and targets inhibits target cleavage, which may signal tailing and trimming. Alternatively, the extensive complementarity between small RNA and its target may help to release its 3' end from PAZ protection.⁵¹

Conclusions and Perspectives

In addition to transcription and processing, 3' end modification also contributes to the accumulation of small RNAs. We are just beginning to understand the underlying biochemical pathway of this process. While it is clear that methylation plays a crucial role in stabilizing small RNAs through antagonizing uridylation and trimming activities, future studies will be required to determine whether the degree of complementarity to targets or target cleavage serves as a signal for the recruitment of tailing and trimming enzymes. There are more challenges in this field. An immediate one is the identification of all the enzymes involved in the small RNAs modification and catabolic pathway, with those responsible for trimming and degradation in particular. Other

outstanding questions include how these enzymes are recruited, and more importantly, whether small RNA stability control can be regulated in response to any developmental or environmental signals. We believe that addressing these questions will greatly advance our understanding of the regulation of small RNAs.

Disclosure of Potential Conflicts of Interest

No potential conflicts of interest were disclosed.

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