Coupled RNA Processing and Transcription of Intergenic Primary MicroRNAs

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The first step in microRNA (miRNA) biogenesis occurs in the nucleus and is mediated by the Microprocessor complex containing the RNase III-like enzyme Drosha and its cofactor DGCR8. Here we show that the 5′→3′ exonuclease Xrn2 associates with independently transcribed miRNAs and, in combination with Drosha processing, attenuates transcription in downstream regions. We suggest that, after Drosha cleavage, a torpedo-like mechanism acts on nascent long precursor miRNAs, whereby Xrn2 exonuclease degrades the RNA polymerase II-associated transcripts inducing its release from the template. While involved in primary transcript termination, this attenuation effect does not restrict clustered miRNA expression, which, in the majority of cases, is separated by short spacers. We also show that transcripts originating from a miRNA promoter are retained on the chromatin template and are more efficiently processed than those produced from mRNA or snRNA Pol II-dependent promoters. These data imply that coupling between transcription and processing promotes efficient expression of independently transcribed miRNAs.

It is well documented that efficient and regulated mRNA biogenesis is ensured by coupling processing to transcription within the framework of an “mRNA factory” comprising the elongating RNA polymerase II (Pol II) and the associated processing factors. This machinery relies on a complex network of protein interactions leading to the release from chromatin of properly modified and “marked” mRNAs (19, 24, 27). Notably, this complex apparatus also provides a quality control mechanism that prevents “incorrect” molecules from progressing along the maturation pathway (1). It has been previously shown in yeast (Saccharomyces cerevisiae) that snRNA biogenesis also relies on a specific “factory” in which transcription and termination events are intimately coupled to processing and release of correctly assembled snRNP particles (2, 22). In eukaryotes, Pol II is also responsible for the transcription of a different class of transcripts, the microRNAs (miRNAs). The tiny single-stranded miRNA molecules are transcribed as long precursors, pri-miRNAs. More than half of the human miRNA genes are carried within introns of both coding and noncoding genes, while others are transcribed as independent monocistronic or polycistronic units (16). The Microprocessor complex, containing the RNase III-like enzyme Drosha and its cofactor DGCR8, converts the nascent pri-miRNAs into small nucleotide pair 196-F and 196-R). In

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MATERIALS AND METHODS

Plasmid construction. All constructs were generated in pCDNA3.1 (+ Invitrogen) without the T7 promoter. Derived plasmids were generated by replacing the cytomegalovirus (CMV) promoter with miR-23a-1, PGK, and U1 promoters amplified from genomic DNA (pr23a-F and pr23a-R, prPGK-F and prPGK-R, and prU1-F and prU1-R oligonucleotide pairs, respectively). The BGH pA site was deleted by inverse PCR. miR, PGK, U1, and CMV constructs were generated by inserting the miR-223 containing region (amplified with the 223WT-F and 223-200R oligonucleotide pair) in all derived plasmids. Constructs t-WT0.2, t-WT1.5, and t-WT2.0 were obtained by fusing miR-223 (oligonucleotide pair 223F and 223R) and miR-507 (oligonucleotide pair 506-F and 507-R). The miR-507–miR-506 cluster was cloned in pcDNA-miR-23P (oligonucleotide pair 507–506-F and 507–506-R), and 507-X constructs were generated by inverse PCR (oligonucleotide pair 223-506-F and 223-2000-R) in t-WT2.0. A region downstream of miR-223 coding sequence (oligonucleotide pair 223-2000-F and 223-2000-R) was inserted upstream of miR-196a-1. t-mut and t-X were obtained by inverse PCR (oligonucleotide pair pr23a-F and 223SHORFT-F and oligonucleotide pair pr23a-F and 223Δ-R). The miR-507–miR-506 cluster was cloned in pcDNA-miR-23P (oligonucleotide pair 507–506-F and 507–506-R), and 507-X constructs were generated by inverse PCR (oligonucleotide pair pr23a-R and 2000-R), and 507-X constructs were extended by inserting a common stuffer region (oligonucleotide pair pr23a-R and 223-506-F and 223-506-R). 507-WTX constructs and t-3.0 constructs were extended by inserting a common stuffer region (oligonucleotide pair 223-506-F and 223-2000-R), and 507-X constructs were extended by inserting a common stuffer region (oligonucleotide pair 223-506-F and 223-2000-R). Oligonucleotide sequences are listed in Table S1 in the supplemental material.

Cell culture, RNA extraction, and analysis. HeLa cells were grown and trans-
Probes. Denstometric analysis was performed using Typhoon Imager (GE Healthcare) and O planctiqut software.

miRNA quantification was performed by TaqMan miRNA detection (Applied Biosystems). RNU6b was used as a calibrator. The relative quantity of each miRNA was calculated using SDS system software version 1.4 (Applied Biosystems).

In Xrn2 and nuclear fractionation experiments, reverse transcription (RT) was performed in the presence of random examers. cDNA was amplified by quantitative real-time RT-PCR (qRT-PCR). Absolute quantification of each specific fragment was performed using Power SYBR green (Applied Biosystems). The oligonucleotide pairs were C-F and C-R, 5'-let7 and 3'-let7, neo-F and neo-R, 3'-ACT-F and 3'-ACT-R, actEX3-F and actEX3-R, p23-tr-F or CMV-tr-F and miR23a1WT-R, and GFP-F and GFP-R (green fluorescent protein) (see Table S1 in the supplemental material). All reactions were performed in triplicate.

ChIP assay. Chromatin immunoprecipitation (ChIP) and qRT-PCR analyses on endogenous loci were carried out as previously described (11). Drosha (Abcam), Pol II (N-20; Santa Cruz) or Xrn2 antibodies (14) were utilized. PCR amplifications were performed with oligonucleotide pairs miR-23a1-F and miR-223 WT-R. The occupancy of the immunoprecipitated factor was estimated by normalizing for amplification efficiency and subtraction of background (5).

RESULTS

The majority of clustered miRNA have short spacers. Previous bioinformatics analysis classified as clustered those miRNAs spaced less than 50 kb apart, with 89% of these clusters spaced less than 5 kb apart (20). Using the mirbase database (13), we filtered the data for intergenic clusters and analyzed their distance distribution up to 10 kb apart (distances between Drosha cleavage sites). We found that the different units follow specific distributions in which ~45% of the miRNAs are <0.5 kb apart, ~37% are between 0.5 and 1.5 kb apart, and ~36% are between 1.2 and 2.0 kb apart (Fig. 1). In a minority of cases, miRNAs are spaced >2.0 kb apart. However, since no expression studies have been performed, it cannot be assumed that these widely spaced miRNAs represent polycistronic units.

These findings indicate a preferential distribution of clustered miRNAs to closely spaced units, suggesting the existence of a selection constraint that restricts the distance between miRNAs of the same transcriptional unit.

Drosha cleavage affects downstream transcription. To examine whether the expression of clustered miRNAs is affected by the spacer length, we generated minigene constructs containing dicistronic transcriptional units using two miRNAs, miR-223 and miR-196a-1, that are not expressed in HeLa cells (10, 34). These were separated by increasing distances with their transcription driven by the constitutive miR-23a WT-27a~24-2 gene promoter (18) (Fig. 2A). After transfection into HeLa cells, miR-223 and miR-196a-1 abundance was measured by qRT-PCR using miRNA-specific TaqMan probes (Fig. 2B). While the distance did not affect miR-223 levels, a progressive reduction of miR-196a-1 accumulation was observed in the constructs with longer spacers, with a residual 20% accumulation when the distance was increased to 3.0 kb (Fig. 2B; t-WT constructs). Since it has been shown that Drosha processing occurs during pri-miRNA transcription (23), we tested whether efficient cotranscriptional processing of miR-223 affected the decrease in miR-196a-1 accumulation. Notably, miR-196a-1 levels were less affected when the upstream pri-miR-223 contained a mutation in the Drosha cleavage site (t-mut constructs) which reduced miR-223 levels to 27% (Fig. 2C). Finally, abrogation of upstream cleavage, by deletion of the entire pre-miR-223 sequence (t-WT constructs), maintained high levels of miR-196a-1 in all constructs (Fig. 2B). While for t-WT and t-mut, the expression of miR-196a-1 was compared...
with that of the cotranscribed miR-223, for t-X constructs, this was normalized to the levels of a cotransfected miR-223-expressing plasmid. These data indicate that the accumulation of the downstream miRNA inversely correlates with the distance and integrity of the upstream miRNA Drosha cleavage site.

Primer extension and 3′/H11032 rapid amplification of cDNA ends analyses indicated the occurrence of correct processing at the Drosha cleavage site and the absence of cryptic polyadenylation in the spacer region utilized (see Fig. S1A and B in the supplemental material).

ChIP analysis with Pol II antibodies was performed on t-WT, t-mut, and t-X constructs containing 0.5-, 2.0-, and 3.0-kb spacer sequences. Figure 2D shows that when the upstream hairpin is a canonical substrate for Drosha, Pol II density on the miR-196a-1 region decreases with the distance. A smaller reduction of Pol II loading was observed when the upstream miRNA was a subcanonical Drosha substrate, whereas, no significant changes were observed with the t-X constructs. Therefore, these data indicate that efficient Drosha cleavage is able to induce transcriptional attenuation.

We then predicted that the distance between miRNAs in endogenous clusters might have been selected during evolution to ensure that each miRNA is in the suitable genomic context for proper expression. We therefore asked whether we could alter the expression of naturally clustered miRNAs by changing their relative distance.

FIG. 2. Drosha cleavage induces transcriptional attenuation. (A) Schematic representation of dicistronic constructs containing pre-miR-223 and pre-miR-196a-1 under the control of the miR-23a promoter (black arrow) Spacer lengths are indicated in the dashed insert. t-WT contains wild-type pri-miR-223, t-mut has a suboptimal Drosha cleavage site, and t-X lacks the miR-223 hairpin. (B) Histograms show the level of miR196a-1 and miR-223 accumulation measured by qRT-PCR analysis on total RNA. For t-X constructs, miR196a-1 values were compared with the levels of miR-223 expressed from a cotransfected plasmid. 2−ΔΔCt, threshold cycle. (C) The upper panel shows human pre-miR-223 and 5′−3′ flanking sequences. The boxed sequence has been deleted in the t-mut constructs. For the lower panel, Northern blot analysis was performed with 10 μg of total RNA from HeLa cells untreated (lane −) or transfected with miR-WT or miR-mut plasmids. U2 snRNA was used as the loading control. Quantitation of the signals is shown below each lane. (D) Pol II ChIP of t-WT, t-mut, and t-X constructs containing 0.5-, 2.0-, and 3.0-kb spacer sequences. The y axis shows ChIP values as ratios of “b” versus “a” regions. Oligonucleotide pairs (“a” and “b”) are positioned as indicated by arrows in panel A. Error bars show standard errors of the mean based on three independent experiments.
The 0.3-kb spacer region of the miR-507–miR-506 cluster was increased to 1.7 kb. In agreement with previous data, the accumulation of the downstream miRNA was reduced by increasing the distance (Fig. 3; 507-WT). Moreover, this effect was almost abolished when the upstream miRNA was deleted (Fig. 3, 507-X).

Taken together, these results suggest that spacers longer than 1.5 kb are detrimental for the synthesis of 3′-located miRNAs, if canonical Drosha sites are present in the upstream miRNAs.

**Xrn2 is involved in Drosha-mediated transcriptional attenuation.** By analogy with the torpedo model (6, 31), a possible candidate for the decrease in Pol II density on downstream regions as a consequence of Drosha cleavage is the 5′→3′ exonuclease Xrn2 (32). ChIP analysis, shown in Fig. 4A, indicated that Xrn2 exonuclease is present at the sites of endogenous miR-23a-1 and let-7a-1 loci. In this experiment, the termination region of the β-actin gene (3′-ACT) was utilized as a positive control (14), while the β-actin exon 3 (ACT Ex3) and a tRNA gene provided background values.

To test whether Xrn2 recruitment depends on the availability of entry sites generated by Drosha cleavage, ChIP analysis with Xrn2 antibodies was performed on the t-WT, t-mut, and t-X-transfected cells (Fig. 4B). The amount of Xrn2 was measured across the miR-196a-1 coding region ("b") and compared to signal obtained on the neomycin ("neo") gene present on the same plasmid. Xrn2 recruitment was detectable on the miR-196a-1 region following efficient miR-223 Drosha cleavage (Fig. 4C; t-WT); whereas, Xrn2 binding to the downstream miRNA was reduced to 30% when processing of the upstream miRNA was inefficient (t-mut). Furthermore, Xrn2 interaction was completely undetectable on the t-X chromatin, where no upstream cleavage occurs.

HeLa cells transfected with t-WT were subjected to Xrn2-specific small interfering RNA treatment (see Fig. S2A in the supplemental material). The effect of Xrn2 knockdown on the relative levels of mature miR-223 and miR-196a-1 in differently-spaced tandem constructs was determined (Fig. 4D and see Fig. S2B in the supplemental material). A recovery of the downstream miRNA was observed in all cases: in particular, the 3.0-kb construct, which displayed the strongest attenuation effect (Fig. 2B), showed a fourfold increase in miR-196a-1 accumulation (Fig. 4D) and almost a twofold increase in stabilization of 3′-cutoff molecules (Fig. 4E). A similar stabilizing effect was observed for the endogenous let-7a-1~let-7f-1 intermediate cleavage product and for the 3′-ACT-positive control (Fig. 4F). These data are indicative of a direct Xrn2 involvement in miRNA biogenesis; in particular in the degradation of 3′-cutoff products generated following Drosha cleavage.

**A miRNA promoter favors efficient cotranscriptional Drosha recruitment.** Since miRNA genes known to be transcribed by Pol II, we tested whether a transcriptional attenuation effect could also occur in the presence of a strong and canonical Pol II promoter. Therefore, the miRNA promoter of the t-WT constructs was replaced by the constitutive CMV promoter (Fig. 5A, left panel). The histogram in Fig. 5A (right panel) shows a reduced attenuation effect when transcription was driven from the CMV promoter. Due to the effect of the Microprocessor complex on transcriptional attenuation, we tested whether various Pol II-dependent promoters could differentially contribute to Drosha recruitment to the downstream hairpin. The p23, PGK, U1 snRNA, and CMV promoters were cloned upstream to pri-miR-223, and Drosha loading was tested in vivo by ChIP analysis. Figure 5B shows that among the promoters, the miRNA promoter displays the most efficient Drosha recruitment. The other promoters have reduced activity which reaches the lowest level in the case of the CMV promoter. Notably, compared to the CMV promoter, the miRNA promoter displayed also a more efficient Xrn2 recruitment (see Fig. S3B in the supplemental material).

Moreover, when a poly(A) signal was inserted downstream to the pre-miR-223 coding sequence in miR-WT and CMV-WT constructs, miRNA transcripts arising from the miR promoter were retained to the chromatin and efficiently processed, while those originating from the CMV promoter were rapidly released from the chromatin mainly as polyadenylated species (see Fig. S4 in the supplemental material).

**DISCUSSION**

The realization that mRNA transcription by RNA Pol II and processing of nascent precursors are closely connected events (3, 27) has led to a consensus view that the general “RNA factory” model can be extended to other Pol II genes (2, 22).
Coupling between transcription and processing has also been demonstrated for intron-encoded miRNAs. Thus, the Microprocessor complex was shown to be recruited to the sites of host pre-mRNA transcription and to cleave pri-miRNA transcripts during ongoing transcription (16, 23).

Here we show that for intergenic miRNAs, the cotranscriptional processing is coupled with termination. As a consequence of Drosha cleavage, Pol II release occurs in downstream regions. Searching for the mechanism of this attenuation, we establish the involvement of the 5'-Xrn2 exonuclease. Thus, we show that Xrn2 is recruited to sites of miRNA transcription and that its depletion promotes transcriptional readthrough. These data suggest that, similarly to the torpedo model proposed for mRNAs (4, 31, 32, 33), after Drosha cleavage of the pri-miRNA transcript, exposure of an unprotected 5' end offers a substrate for Xrn2 recruitment. The exonuclease acts to degrade the 3'-cutoff product resulting in Pol II template destabilization and transcription termination.

We also show that the position at which termination occurs depends on the efficiency of Drosha cleavage, since mutations in the Microprocessor cleavage site, or its deletion, allow Pol II to extend over greater distances. According to these findings, we predict that the distance between tandem miRNAs may have been appropriately selected in order to prevent a polarity effect on the production of the downstream miRNA. Short distances ensure that even if upstream Drosha cleavage occurs efficiently, the Xrn2 exonuclease will not catch the polymerase before transcription has proceeded into the downstream miRNA sequence. Alternatively, for longer distances between adjacent miRNAs, Pol II processivity, Drosha recruitment and cleavage, and Xrn2 trimming should be appropriately balanced to ensure the synthesis of all the members of the cluster. Our bioinformatics analysis reveals that 80% of endogenous clusters have spacer sequences shorter than 1.5 kb. Notably, in our artificial dicistronic constructs, the attenuation effect becomes relevant above 1.5 kb. Therefore, it can be suggested that longer spacing provides a way for nonstoichiometric production of clustered miRNAs or for imposing posttranscriptional control through Drosha activators or repressors.

In conclusion, transcriptional attenuation, as a consequence of Drosha cleavage, may only be involved in primary transcript...
termination without affecting the expression of the majority of clustered miRNAs.

This study also reveals that an miRNA promoter, in contrast to mRNA or snRNA Pol II promoters, favors the recruitment of Drosha on nascent transcripts. Possibly, the miRNA promoter acts either by favoring the recruitment of factors that in turn mediate Drosha loading or by retaining the transcript at chromatin sites where Drosha is localized. Several reports have previously demonstrated the existence of specific factors (some of which are Drosha interactors) that can modulate processing of pri-miRNAs (8, 12, 15, 28). This allows speculation that regulation mediated by auxiliary factors may operate early during transcription and affect Drosha recruitment or its affinity for the substrate. Alternatively, a miRNA promoter may drive less processive transcription providing more time for Drosha–pri-miRNA interaction to occur. Promoter structure and recruitment of transcription factors or coactivators may also affect polymerase processivity by conferring different elongation properties to the enzyme (17).

Moreover, in agreement with previous observations, our data suggest that poly(A) signals interfere with miRNA maturation, very likely by mislocalizing the pri-miRNA from the sites of Microprocessor action (7, 25, 26). Even though the utilization of poly(A) sites is detrimental for efficient miRNA biogenesis, their occurrence downstream of many miRNA transcriptional units (29) could act in a quality control mechanism. In contrast to mRNA coding genes, where poly(A) sites direct efficient export to the cytoplasm, for miRNA transcriptional units they may represent a way of preventing transcriptional readthrough to downstream sequences when correct processing has not occurred. This is resonant with snoRNA transcription where incorrect 3’-end processing leads to readthrough, unless poly(A) sites are present in the downstream regions (21, 30).

By analogy with previous findings, linking Pol II transcription with tightly coupled processing of nascent pre-mRNAs (3,