SRSF3 recruits DROSHA to the basal junction of primary microRNAs

KIJUN KIM,1,2,5 TRUNG DUC NGUYEN,3,4,5 SHAOHUA LI,3 and TUAN ANH NGUYEN3,4

1Center for RNA Research, Institute for Basic Science, Seoul 08826, Korea
2School of Biological Sciences, Seoul National University, Seoul 08826, Korea
3Division of Life Science, Hong Kong University of Science and Technology, Hong Kong, China
4HKUST Shenzhen Research Institute, Shenzhen 518057, China

ABSTRACT

The Microprocessor complex, consisting of an RNase III DROSHA and the DGCR8 dimer, cleaves primary microRNA transcripts (pri-miRNAs) to initiate microRNA (miRNA) maturation. Pri-miRNAs are stem–loop RNAs, and ~79% of them contain at least one of the three major and conserved RNA motifs, UG, UGU, and CNNC. We recently demonstrated that the basal UG and apical UGU motifs of pri-miRNAs interact with DROSHA and DGCR8, respectively. They help orient Microprocessor on pri-miRNA in a proper direction in which DROSHA and DGCR8 localize to the basal and apical pri-miRNA junctions, respectively. In addition, CNNC, located at ~17 nucleotides (nt) from the Microprocessor cleavage site, interacts with SRSF3 (SRp20) to stimulate Microprocessor to process pri-miRNAs. The mechanism underlying this stimulation, however, is unknown. In this study, we discovered that SRSF3 recruits DROSHA to the basal junction in a CNNC-dependent manner, thereby enhancing Microprocessor activity. Furthermore, by generating various pri-miRNA substrates containing CNNC at different locations, we demonstrated that such stimulation only occurs when CNNC is located at ~17 nt from the Microprocessor cleavage site. Our findings reveal the molecular mechanism of SRSF3 in pri-miRNA processing and support the previously proposed explanation for the highly conserved position of CNNC in SRSF3-enhanced pri-miRNA processing.

Keywords: SRSF3; microprocessor; microRNA; DROSHA; DGCR8

INTRODUCTION

MicroRNAs (miRNAs), short and single-stranded RNAs of ~22 nucleotides (nt), are mostly produced via the canonical microRNA biogenesis pathway (Ha and Kim 2014). This process involves two consecutive cleavages by RNase III-type enzymes. The primary miRNA transcript (pri-miRNA) synthesized in the nucleus is initially cleaved by the Microprocessor complex composed of nuclear RNase III DROSHA and its cofactor DGCR8 (Denli et al. 2004; Gregory et al. 2004; Han et al. 2004, 2006; Landthaler et al. 2004; Auyeung et al. 2013; Nguyen et al. 2015; Fareh et al. 2016; Herbert et al. 2016; Kwon et al. 2016). The resulting product, precursor miRNA (pre-miRNA) is exported to the cytoplasm and is further cleaved by another RNase III enzyme, Dicer, that produces a miRNA duplex of ~22 nt (Ma et al. 2004; MacRae et al. 2006a,b; Park et al. 2011; Gu et al. 2012; Tian et al. 2014). Only one strand of each duplex is loaded into an Argonaute (Ago) protein, and the other strand is eliminated (Kawamata and Tomari 2010). The Ago-miRNA interaction forms the core of an RNA-induced silencing complex that plays a critical role in posttranscriptional gene regulation in various essential biological processes (Bartel 2009; Ameres and Zamore 2013; Ha and Kim 2014).

Pri-miRNA (Fig. 1A), a substrate of Microprocessor, has a local stem–loop, consisting of a three-helix stem flanked by the basal and apical junctions at two ends (Han et al. 2006; Ha and Kim 2014). DROSHA, a catalytic subunit of the Microprocessor complex, must localize at the pri-miRNA basal junction to cleave pri-miRNA at the productive sites (~13 nt from the basal junction) and generates pre-miRNA (Fig. 1A). However, the somewhat symmetrical nature of pri-miRNA secondary structure often leads Microprocessor to interact with the apical junction, thereby cleaving pri-miRNAs at the unproductive sites ~13 nt from the apical junction (Han et al. 2006; Nguyen et al. 2015). Multiple primary sequence motifs that have been recently identified (Auyeung et al. 2013; Fang and Bartel 2015) create

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an asymmetry for pri-miRNAs and facilitate association of DROSHA with the basal junction (Nguyen et al. 2015; Kwon et al. 2016). The U and G at positions -14 and -13 (Fig. 1A) whose frequencies in human pri-miRNAs conserved to mice, so-called representative pri-miRNAs, are ~40% and 48%, respectively, independently contribute to pri-miRNA processing (Auyeung et al. 2013). The apical UGU motif of the apical loop that appeared at 22 nt from the cleavage site is recognized by DGCR8. The CNNC motif of the basal segment that resides at 17 nt from the cleavage site interacts with SRSF3. (B) Protein constructs used in this study. The first and end residue numbers for each construct are shown. (P-rich) Proline-rich domain, (RS) arginine/serine-rich domain, (CED) central domain, (RIIIDa and RIIIDb) RNase III domains, (dsRBD) dsRNA-binding domain, (Rhed) RNA-binding heme domain, (CTT) C-terminal tail region, (RRM) RNA recognition motif. SRSF3 is composed of 164 amino acids and its molecular weight is about 19 kDa. It has one RRM at the N terminus and one RS domain at the C terminus with extensive phosphorylation on serine residues. (C) The purity of the GFP-tagged SRSF3 on SDS–PAGE. (D) The pri-miR-30a substrates. The uppercase letters represent the pre-miRNA region. The CNNC and ΔCNNC motifs are highlighted in red. The green and purple arrowheads indicate the productive and unproductive cleavage sites of DROSHA, respectively. (E) Processing of the pri-miR-30a substrates by D3–G2 or D3–G4G4 and SRSF3. The internally labeled pri-miR-30a substrates were incubated with the indicated proteins for 1 h under the conditions described in Materials and Methods. The three products, a 5′-fragment, pre-miRNA, and a 3′-fragment were named as F1, F2, and F3, respectively.

FIGURE 1. SRSF3 recruits DROSHA to the CNNC-containing basal junction. (A) Microprocessor is a trimeric complex consisting of an RNase III DROSHA and a DGCR8 dimer. The typical pri-miRNA, a substrate of the Microprocessor complex, has a stem–loop structure. The green arrowheads indicate the DROSHA cleavage sites. The basal junction and the basal UG motif located at 14 nt from the cleavage site are recognized by DROSHA. The apical UGU motif of the apical loop that appeared at 22 nt from the cleavage site is recognized by DGCR8. The CNNC motif of the basal segment that resides at 17 nt from the cleavage site interacts with SRSF3. (B) Protein constructs used in this study. The first and end residue numbers for each construct are shown. (P-rich) Proline-rich domain, (RS) arginine/serine-rich domain, (CED) central domain, (RIIIDa and RIIIDb) RNase III domains, (dsRBD) dsRNA-binding domain, (Rhed) RNA-binding heme domain, (CTT) C-terminal tail region, (RRM) RNA recognition motif. SRSF3 is composed of 164 amino acids and its molecular weight is about 19 kDa. It has one RRM at the N terminus and one RS domain at the C terminus with extensive phosphorylation on serine residues. (C) The purity of the GFP-tagged SRSF3 on SDS–PAGE. (D) The pri-miR-30a substrates. The uppercase letters represent the pre-miRNA region. The CNNC and ΔCNNC motifs are highlighted in red. The green and purple arrowheads indicate the productive and unproductive cleavage sites of DROSHA, respectively. (E) Processing of the pri-miR-30a substrates by D3–G2 or D3–G4G4 and SRSF3. The internally labeled pri-miR-30a substrates were incubated with the indicated proteins for 1 h under the conditions described in Materials and Methods. The three products, a 5′-fragment, pre-miRNA, and a 3′-fragment were named as F1, F2, and F3, respectively.
SRSF3 fails to stimulate the Microprocessor activity for pri-mir-30c wild-type (WT), where the CNNC motif is present in the duplex RNA region (Fernandez et al. 2017). This emphasizes the physiological significance of the SRSF3–CNNC interaction in the appropriate cellular context. However, a molecular mechanism explaining how SRSF3 stimulates pri-miRNA processing and why the position of CNNC is restricted to an area within 16–18 nt from the DROSHA cleavage sites remains elusive.

In this study, we explored the mechanism of SRSF3 action using purified recombinant proteins from human cells (Fig. 1B). We discover that SRSF3 stimulates Microprocessor by recruiting DROSHA to the pri-miRNA basal junction. This recruitment is dependent on the CNNC motif of pri-miRNA at the precise position.

RESULTS

SRSF3 recruits DROSHA to the basal junction of pri-miRNA in a CNNC-dependent manner

To determine how SRSF3 affects the pri-miRNA processing, we purified the SRSF3 proteins tagged with GFP from suspension HEK293E cells (Fig. 1C) and selected pri-mir-30a (Fig. 1D) harboring a CNNC motif (CUUC) 17 nt from the DROSHA cleavage sites. The D3–G2 complex contains all necessary domains for DROSHA activity (D3, Fig. 1B) and a stabilizing fragment of DGCR8 (G2, Fig. 1B). The D3–G4G4 complex consists of the D3 fragment of DROSHA and the G4G4 dimer of DGCR8 (G4, Fig. 1B). These two complexes were purified as described previously (Nguyen et al. 2015). First, we tested the effect of SRSF3 on the D3–G2 complex. Consistent with previous studies (Nguyen et al. 2015; Kwon et al. 2016), D3–G2 alone primarily cleaved pri-mir-30a at the unproductive sites (Fig. 1E, lane 3). Interestingly, however, the addition of SRSF3 caused D3–G2 to successfully cleave pri-mir-30a at the productive sites (Fig. 1E, lanes 4,5). As a result, SRSF3 strongly stimulated the trimeric D3–G4G4 complex to cleave at the productive sites (Fig. 1E, lanes 6–8; Supplemental Fig. S1A, lanes 3–5). In contrast, when the CUUC motif was replaced with the UUUU motif, SRSF3 did not enhance the productive activity of the D3–G2 and D3–G4G4 complexes (Fig. 1E, lanes 11–16; Supplemental Fig. S1A, lanes 8–10). These data suggest that by interacting with the CNNC motif at the 3′-segment of pri-miRNAs, SRSF3 might recruit DROSHA to the basal junction, thereby stimulating the occurrence of productive cleavages. The other proteins, SRSF7 (9G8), DDX5, and DDX17 that were proposed to interact with the 3′-segment of pri-miRNA were also tested on the pri-miRNA processing of DROSHA (Wang et al. 2012a; Auyeung et al. 2013; Dardenne et al. 2014; Mori et al. 2014; Moy et al. 2014). Unlike SRSF3, they did not enhance the DROSHA activity (Supplemental Fig. S1B,C), suggesting that they might use different mechanisms in the pri-miRNA processing.

SRSF3 and DROSHA form a complex with the CNNC-containing pri-miRNAs

To confirm that SRSF3 recruits DROSHA to the basal junction, we examined if SRSF3 and DROSHA can physically interact with each other. We used purified SRSF3 and DROSHA proteins in immunoprecipitation assays but did not identify a direct protein–protein interaction (data not shown). Thus, their interaction is likely to be either transient or RNA-dependent. Therefore, we used an electrophoretic mobility shift assay (EMSA) to test whether SRSF3 and DROSHA can form a stable complex on RNA. The resulting EMSA data showed that individual DROSHA and SRSF3 proteins interacted with pri-mir-30a and distinctly shifted the electrophoretic mobility of RNA bands (Fig. 2, lane 2 for DROSHA and lane 8 for SRSF3; Supplemental Fig. S2A). DROSHA seemed to have a slightly higher affinity to the pri-mir-30a than pri-mir-30aΔCNNC (Fig. 2, cf. lane 2 with 10; Supplemental Fig. S2B,C). When two proteins were simultaneously incubated with pri-mir-30a, the shifted RNA bands were denser than those observed using any one of the proteins and RNA (Fig. 2, cf. lane 5 with 2 or 8; Supplemental Fig. S2A). This indicates that SRSF3 and DROSHA coordinately interact with RNA. Furthermore, the SRSF3-bound RNA band was further shifted by the addition of DROSHA (Fig. 2, cf. lane 8 with 5), suggesting that SRSF3 and DROSHA associate together on the same pri-miRNA.
molecule. In contrast with CUUC-containing pri-mir-30a, SRSF3 inefficiently stimulated the complex formation of DROSHA and UUUU-containing pri-mir-30a (Fig. 2, cf. lanes 12–13 with 4–5). This declined stimulatory effect of SRSF3 could be due to its weak interaction with the UUUU motif (Schaal and Maniatis 1999; Hargous et al. 2006; Änkö et al. 2012). In addition, the band of UUUU-containing pri-mir-30a bound with SRSF3 was not efficiently shifted by the addition of DROSHA (Fig. 2, cf. lane 16 with 13). The data from EMSA experiments support a model that SRSF3 interacts with CNNC and recruits DROSHA to the basal junction.

Proper position of CNNC is critical for SRSF3 to stimulate productive processing by DROSHA

Since the CNNC motif is highly conserved at the 3′-segment of pri-miRNAs ∼16 to 18 nt from the DROSHA cleavage sites and this position is important for miRNA expression in vivo (Auyeung et al. 2013), we examined the positional effect of the CNNC motif on the pri-miRNA processing. We generated two pri-mir-30a variants with the CUUC motif at either 19 or 21 nt from the DROSHA cleavage sites (Fig. 3A) and used them in the processing assays with D3–G2 and SRSF3. The unlabeled pri-mir-30a substrate (0.5 μM) or the internally labeled pri-mir-142 substrate was incubated with the proteins that are shown in the figure for 1 h under the conditions described in Materials and Methods. F1, F2, and F3 are the productive products. The unproductive and alternative products are indicated with the green and red arrowheads, respectively.

**FIGURE 3.** SRSF3 stimulates the productive processing by DROSHA in a CNNC position-dependent manner. (A,C) The pri-mir-30a and pri-mir-142 substrates. The uppercase letters represent the pre-miRNA region. The CNNC motif and the added nucleotides are highlighted in red and blue, respectively. The green, purple, and red arrowheads indicate the productive, unproductive, and alternative cleavage sites of DROSHA, respectively. (B,D) Processing of the pri-mir-30a and pri-mir-142 substrates by D3–G2 or D3–G4G4 and SRSF3. The unlabeled pri-mir-30a substrate (0.5 μM) or the internally labeled pri-mir-142 substrate was incubated with the proteins that are shown in the figure for 1 h under the conditions described in Materials and Methods. F1, F2, and F3 are the productive products. The unproductive and alternative products are indicated with the green and red arrowheads, respectively.

Recruitment of DROSHA to primary microRNAs
(Supplemental Fig. S3C), implying that DGCR8 (G4G4) might hold DROSHA at the productive cleavage sites (Nguyen et al. 2015). This result confirms the role of SRSF3 in DROSHA recruitment and suggests that SRSF3 stimulates the productive processing by DROSHA as the CNNC motif is present at a position ~17 nt downstream from the cleavage sites.

To further confirm the importance of the CNNC position, we investigated the effect of SRSF3 on the Microprocessor processing of pri-mir-142 in which the CNNC motif is located at 15 nt downstream from the cleavage sites (Fig. 3C). SRSF3 persistently induced DROSHA (D3–G2) to cleave at the basal sites on pri-mir-142 WT (Fig. 3D, cf. lanes 4–5 with 3) but not pri-mir-142ΔCNNC (Fig. 3D, cf. lanes 12–13 with 11). However, SRSF3 stimulated DROSHA to cleave at a position 1 or 2 nt away from the productive sites cut by DROSHA alone (Fig. 3D, cf. lanes 4–5 with 3). In addition, SRSF3 failed to stimulate D3–G4G4 to cleave pri-mir-142 (Fig. 3D, cf. lanes 7–8 with 6). This further indicates that SRSF3 functions in a CNNC position-dependent manner to facilitate the pri-miRNA processing.

DISCUSSION

Previous studies demonstrated that human Microprocessor interacts with multiple cis-acting RNA elements to accurately and efficiently cleave pri-miRNAs (Han et al. 2006; Auyeung et al. 2013; Nguyen et al. 2015; Kwon et al. 2016). The basal junction and its UG motif are recognized by DROSHA (Han et al. 2006; Nguyen et al. 2015). The apical loop and the UGU motif interact with the DGCR8 dimer assisted by hemin (Nguyen et al. 2015, 2018; Partin et al. 2017). The CNNC motif at the 3′ flanking region of pri-miRNA interacts with SRSF3, a trans-acting factor that stimulates Microprocessor (Auyeung et al. 2013). In this study, we reveal a molecular mechanism in which SRSF3 binds to the CNNC motif, thereby recruiting DROSHA to the basal junction of pri-miRNAs. Consequently, SRSF3 influences the orientation of Microprocessor on the CNNC-containing pri-miRNAs (Fig. 4 for the model).

This study demonstrates that SRSF3 can recruit DROSHA to the basal junction by interacting with the CNNC motif at the various positions (15–21 nt from the DROSHA cleavage sites). Interestingly, SRSF3 can alter the DROSHA cleavage sites at the stem of pri-miRNAs. For example, SRSF3 can shift DROSHA from the productive cleavage sites to the alternative sites toward the basal junction when the CNNC motif is farther from the basal junction (19 or 21 nt from the cleavage sites) (Fig. 3B). In contrast, SRSF3 can move DROSHA to the alternative sites away from the basal junction as the CNNC motif resides closer to the basal junction (15 nt from the cleavage sites) (Fig. 3D). These indicate that SRSF3 and DROSHA might make a protein–protein contact upon interaction with pri-miRNA and that the two proteins are likely positioned at a certain fixed distance. In addition, the EMSA results (Fig. 2; Supplemental Fig. S2) support that SRSF3 and DROSHA might form a stable complex on the CNNC-containing pri-miRNA. Of note, SRSF3 does not cause the DROSHA/RNA complex to obviously further shift in the gel, suggesting that either the DROSHA/RNA/SRSF3 complex has a similar size as the DROSHA/RNA complex or SRSF3 might dissociate from pri-miRNA after facilitating DROSHA binding. Future structural studies are expected to reveal the atomic mechanisms underlying the DROSHA/RNA/SRSF3 interaction.

Unlike DROSHA, the cleavage sites by the Microprocessor complex are less shifted by SRSF3 (pri-mir-30a, Supplemental Fig. S3C; pri-mir-142, Fig. 3D) as the CNNC motif is not located at the optimal position (16–18 nt from the productive cleavage sites). This could be explained by a role of DGCR8 in the Microprocessor complex, that strongly retains DROSHA at the productive cleavage sites (Nguyen et al. 2015) despite the effect of SRSF3. Therefore, the proper position of CNNC (16–18 nt from the DROSHA cleavage sites) is crucial for SRSF3 to coordinate with DROSHA and DGCR8 in effective stimulation of the pri-miRNA processing in vitro. However, a broader window (16–21 nt) of CNNC might be functional in vivo when a large group of pri-miRNAs was investigated (Conrad et al. 2014). It is possible that the secondary structure surrounding the CNNC motif or other cofactors may contribute to the CNNC-dependent processing of pri-miRNAs by Microprocessor.

SRSF3, the smallest member of the highly conserved SR-rich splicing factor family, regulates the splicing of numerous genes (Jumaa and Nielsen 1997, 2000; Jumaa et al. 1997; Galiana-Arnoux et al. 2003; Yu et al. 2004; de la Mata and Kornblit 2006; Gonçalves et al. 2008, 2009; Sen et al. 2009; Manley and Krainer 2010; Wang et al. 2012b; Wong et al. 2012; Ajiro et al. 2016). Furthermore, SRSF3 interacts with various CNNC-containing mRNAs (Hargous et al. 2006; Ånkö et al. 2012). Therefore, it will also be interesting
to investigate if the DROSHA/RNA/SRSF3 interaction may be involved in mRNA metabolism.

**MATERIALS AND METHODS**

**Processing assay**

The processing assay was carried out at 37°C in 20 µL assay buffer containing 50 mM Tris-hydrochloride (Tris-HCl [pH 7.5]), 100 mM sodium chloride (NaCl), 10% glycerol, 0.2 µg/µL bovine serum albumin (BSA), 1 mM dithiothreitol (DTT), and 2 mM magnesium chloride (MgCl₂). Approximately 10,000 cpm of the RNA substrates was used, and the enzyme concentrations and incubation time are indicated in the figures. The reaction was stopped by adding 20 µL TBE-urea sample buffer (Bio-Rad). Finally, the mixture was heated at 95°C for 10 min and quickly chilled on ice before loading onto a 10% Urea-polyacrylamide gel electrophoresis (Urea-PAGE) gel with the RNA size markers (Decade marker, Ambion). The RNA substrates were prepared as described previously (Nguyen et al. 2015).

**Purification of recombinant SRSF3 from human cells**

The SRSF3 fused to the C-terminal GFP and 10× His-tag was cloned into the pXG vector as used in the previous studies (Nguyen et al. 2015; Kwon et al. 2016). The plasmids were transfected into 3.6 L HEK293E suspension cell culture and the cells were harvested after 2 d. The cell pellets were resuspended in 200 mL T500 buffer (20 mM Tris-HCl [pH 7.5], 500 mM NaCl, and 4 mM β-mercaptoethanol) supplemented with 2 µg/µL RNase A and protease inhibitor cocktail. The clear lysate obtained by sonication and centrifugation was loaded on a Ni-NTA column. The column was washed with 200 mL T500 buffer (Bio-Rad). The proteins were eluted with T500 supplemented 200 mM imidazole. The eluted proteins were further purified on Superdex 75 column with the RNA size markers (Decade marker, Ambion). The RNA substrates were prepared as described previously (Nguyen et al. 2015).

**SUPPLEMENTAL MATERIAL**

Supplemental material is available for this article.

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