PCAF-mediated acetylation of Lin28B increases let-7 biogenesis in lung adenocarcinoma H1299 cells

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Abstract

Background: Lin28B and its paralog Lin28A are small RNA binding proteins that have similar inhibitory effects, although they target separate steps in the maturation of let-7 miRNAs in mammalian cells. Because Lin28B participates in the promotion and development of tumors mostly by blocking the let-7 tumor suppressor family members, we sought to explore the associated mechanisms to gain insights into how Lin28B might be decreased in human cancer cells to increase let-7 levels and reverse malignancy.

Results: We demonstrated that the histone acetyltransferase PCAF, via its cold shock domain, directly interacts with and subsequently acetylates Lin28B in lung adenocarcinoma-derived H1299 cells. RT-qPCR assays showed that both let-7a-1 and let-7g were increased in PCAF-transfected H1299 cells. Lin28B is acetylated by ectopic PCAF and translocates from the nucleus to the cytoplasm in H1299 cells.

Conclusions: The effects of acetylated Lin28B on let-7a-1 and let-7g are similar to that of stable knockdown of Lin28B in H1299 cells. The new role of PCAF in mediating Lin28B acetylation and the specific release of its target microRNAs in H1299 cells may shed light on the potential application of let-7 in the clinical treatment of lung cancer patients.

Keywords: Lin28B, PCAF, let-7, Acetylation

Background

Lin28 and the microRNA let-7 were first discovered in C. elegans as heterochronic genes that regulate developmental timing [1–3]. In eukaryotes including worms and mammals, Lin28 blocks let-7 expression, whereas let-7 negatively regulates Lin28 expression by binding to the 3'UTR of Lin28 mRNA, thereby establishing a double negative feedback loop. The Lin28/let-7 axis plays a pivotal role in stem cell biology and the development and control of glucose metabolism, as well as in human diseases [4, 5]. In mammals, there are two Lin28 paralogs: Lin28A and Lin28B. Although it is structurally similar to Lin28A, Lin28B contains a cold shock domain (CSD) and a retroviral-type CCHC zinc finger (ZF) motif. Lin28B has a coding extended C terminus that contains a nuclear localization signal (NLS) in addition to a nucleolus localization signal (NoLS) between the CSD and ZF domains, both of which participate in the subcellular localization of Lin28B in human cells [6–10]. The expression of Lin28A in the cytoplasm blocks let-7 processing by Dicer and uridylation of pre-let-7 by TUTase [11], whereas Lin28B primarily accumulates in the nucleus, where it binds pri-let-7 miRNAs and blocks the activity of the microprocessor complex [5, 8, 11]. However, the subcellular localization of Lin28B is controversial [4].

Lin28B was first cloned and identified as an overexpressed factor in hepatocellular carcinoma cells [6]. Lin28B is currently known to be involved in the promotion and development of tumors, thus indicating that it may be a potential target in human cancer therapy [7, 12–15]. A high Lin28A or Lin28B and low let-7 expression pattern is found in approximately 15% of human cancers [16]. The expression of Lin28B in cancer cells can be activated by transcription factors and epigenetic modifiers, such as Myc, NF-κB and Sirt6 [17–20]; however, much of the underlying mechanism remains unclear.
Acetylation is an important modification pattern that has been widely investigated in recent years. Protein acetylation is known to participate in regulating multiple cellular processes in normal and cancer cells [21–23]. As a bona fide cancer-related protein, Lin28B is subject to polyubiquitination that leads to the enhancement of let-7 biogenesis [24, 25]. However, whether the acetylation of Lin28B affects the let-7 biogenesis involved in tumorigenesis is not yet fully understood.

In this study, we found that knockdown of Lin28B in the human lung adenocarcinoma cell line H1299 abrogated the inhibition of let-7 miRNA. The histone acetyltransferase PCAF was found to directly interact with Lin28B via its CSD, and this interaction facilitated Lin28B acetylation by the HAT domain of PCAF. Most importantly, we demonstrated that the PCAF-mediated acetylation of Lin28B might de-repress the processing of let-7a-1 and let-7g, and these findings shed light on the potential application of acetylated Lin28B for future cancer therapy.

Methods

Cell culture

HEK293T, HCT116, MCF7, HeLa, HepG2, and H1299 cells were cultured in Dulbecco’s modified Eagle’s medium (DMEM) supplemented with 10% fetal bovine serum (FBS, HyClone) at 37 °C in 5% CO₂ atmosphere. The HEK293T cells, MCF7, and H1299 cells were stored in our Lab. The HeLa (Cat. #3111C0001CCC000011) and HepG2 (Cat. #3111C0001CCC000035) cell lines were purchased from Chinese National Infrastructure of Cell Line Resource (Beijing, China). HCT116 cell line was a gift from Dr. Depei Liu (Institute of Basic Medical Sciences, Chinese Academy of Medical Sciences, Cat.# 3111C0001CCC000158). Before the experiments, the two cell lines were authenticated on cell micrograph compared to the cell lines on ATCC. HEK293T cells showed 90% transfect efficiency with GFP-tag plasmid. H1299 cells showed the lack of p53 protein expression by western blot assay. Mycoplasma contamination was detected by the EZ-PCR Mycoplasma Test Kit (Cat. #20–700–20), a PCR-based mycoplasma test kit, in cell cultures before the experiment. The kit includes a unique reaction mix that contains all the ingredients required for PCR: nucleotides, primers, Taq Polymerase and magnesium. After performing agarose gel electrophoresis, positive samples will yield a 270 bp fragment, but HEK293T and H1299 cell lines not.

We established two stably transfected clones of H1299 cells, in which Lin28B was knocked down by co-transduction of the cells with lentivirus encoding each of the shRNA specific for Lin28B, designated shLin28B-1, shLin28B-2, and shLuc was as a negative control. The shLin28B-1, shLin28B-2, or shLuc were cloned into the pLKO.1 vector. The shRNA sequences were as follows: shLin28B-1: GCAGGCGATAAGCAGGTTA; shLin28B-2: GCCTTGAGTCAATACGGGTAA; shLuc: CGCTGAGTACTCGAAATGTC.

Antibodies

Antibodies against β-actin (sc-47778), GAPDH (sc-166545), Myc (sc-789), Lamin B (sc-6216) and PCAF (sc-13,124) were purchased from Santa Cruz Biotechnology (Santa Cruz, CA). Antibodies to FLAG (F3165) and M2 (F2426) were purchased from Sigma (St. Louis, MO). Antibodies to Lin28B (4196) and acetyl lysine (9441) were purchased from Cell Signaling Technology (Beverly, MA).

Plasmids, constructs, and transfection

The FLAG-tagged and GFP-tagged Lin28B eukaryotic expression plasmids and GST-tagged Lin28B prokaryotic expression plasmids were constructed by cloning Lin28B into the pcDNA6-FLAG, pEGFP-C1-GFP and pGEX-4 T-1 vectors, respectively, using a PCR product from a cDNA library derived from HEK293T cells. pCX-FLAG-PCAF, pGEX-PCAF-HAT and pCX-FLAG-PCAF-ΔHAT were as previously described [16]. Truncated fragments of Lin28B were cloned using the PCR product of full-length GFP-tagged Lin28B. FLAG-Lin28B-ΔCSD was constructed by deleting the CSD. Myc-tagged PCAF was cloned from FLAG-tagged PCAF into the pCMV-3tag7-Myc vector. The eukaryotic expression plasmid of PCAF containing the Cys574Ser mutation was generated by site-directed mutagenesis of Myc-PCAF. The cells were transfected with Vigofect reagent (Vigorous Biotech, Beijing, China) according to the manufacturer’s instructions. Assays were performed 3 times each in triplicate, and all results are shown as the mean ± SD.

Immunofluorescence

Cells were cultured on glass cover slips and fixed in 4% paraformaldehyde at room temperature for 10 min, washed three times with PBS for 15 min, permeabilized with 0.25% Triton X-100 in PBS for 15 min at room temperature, washed three times as described above and blocked with 1% BSA blocking solution at 37 °C for 1 h. The cells were incubated with primary anti-Lin28B antibody and anti-PCAF antibody at 1:50 dilution, anti-Myc antibody at 1:100 dilution and anti-FLAG antibody at 1:500 dilution at 4 °C overnight and washed three times with PBS. After washing, the cells were incubated with FITC- or TRITC-conjugated secondary antibodies at a dilution of 1:200 at 37 °C for 1 h and then washed three times with PBS. The coverslips were stained with DAPI, mounted and examined with a laser scanning confocal microscope.
Co-immunoprecipitation (co-IP) and immunoblot analyses

Transiently transfected HEK293T cells were homogenized in lysis buffer (50 mM HEPES pH 7.5, 150 mM NaCl, 2 mM EDTA, 2 mM EGTA, 1% Triton X-100, 50 mM NaF, 5 mM Sodium Pyrophosphate, 50 mM Sodium β-glycerophosphate, 1 mM NaVO₃, 1 mM DTT, 1 mM PMSF, 10 μg/ml leupeptin and 10 μg/ml aprostation). Cell lysates were rotated at 4 °C for 30 min and centrifuged at 12,000 rpm for 15 min to remove insoluble material and incubated with anti-GFP beads or anti-FLAG M2 beads at 4 °C overnight. The collecteduble material and incubated with anti-GFP beads or anti-FLAG M2 beads at 4 °C overnight. The collected beads were then washed three times and boiled in SDS gel-loading buffer for western blot analysis.

GST pull-down assay

The GST-tagged Lin28B fusion protein was expressed in Escherichia coli and purified using anti-GST beads. GST or GST-tagged Lin28B beads were individually incubated with cell lysates at 4 °C overnight. Finally, the beads were washed three times, and the bound proteins were analyzed by western blotting.

Quantitative real-time RT-PCR assays (RT-qPCR)

RT-qPCR assays were carried out as described previously [26, 27]. Specific primers for Lin28B (forward: AGCCCCCTTGATATCAGTTC and reverse: AATGTGAATTCACGTGTTCTCCT). The relative expression of let-7a-1 and let-7g was normalized to that of U6 snRNA using the comparative CT method according to the manufacturer’s instructions (Bio-Rad CFX Connect Real-Time System). The primer sequences used for let-7 are listed below (F: forward; R: reverse; RT: reverse transcription).

- Mature let-7a-1 (RT: GTGCTAATCCCTG CAGGGTTCGGATATTGCAGTAGATGTGTGA and R: GTGCAGGGTGCCAGGT), mature let-7g (RT: GTGCTATCCAGTG CAGGGTTCGGATATTGCAGTAGATGTGTGA and R: GTGCAGGGTGCCAGGT), mature miR-16-1 (RT: GTGCTATCCAGTG CAGGGTTCGGATATTGCAGTAGATGTGTGA and R: GTGCAGGGTGCCAGGT), mature mir-16-1 (RT: GTGCTATCCAGTG CAGGGTTCGGATATTGCAGTAGATGTGTGA and R: GTGCAGGGTGCCAGGT), mature mir-16-1 (RT: GTGCTATCCAGTG CAGGGTTCGGATATTGCAGTAGATGTGTGA and R: GTGCAGGGTGCCAGGT).

- F: GCCGCTAGCAGCACGTAAATAT and R: GTGCA
- GCCGCTAGCAGCACGTAAATAT and R: GTGCA
- GCCGCTAGCAGCACGTAAATAT and R: GTGCA

Preparation of cell fractions

The cell fractions were prepared as previously described [28], and the prepared cell fractions were separated with 10% SDS-PAGE and detected by western blotting.

Statistical analysis

Statistical analysis was performed using two-tailed Student’s t-test. All data were shown as mean with standard deviations (SD). Probabilities of $P > 0.05$ were considered as no significant (#), $P \leq 0.05$ as significant (*) and $P \leq 0.01$ as highly significant (**).

Results

The effect of Lin28B knockdown on let-7 expression in H1299 cells

Consistent with the literature [8], Lin28B is mainly found to be distributed in the nucleus (Fig. 1a), where it abrogates the expression of let-7 miRNA. To explore the functions of Lin28B in cancer cells, we first established two stably transfected clones of H1299 cells, in which Lin28B was knocked down by co-transduction of the cells with lentivirus encoding each of the shRNA specific for Lin28B, designated sh-Lin28B-1 and sh-Lin28B-2. We found that the expression of the Lin28B protein (Fig. 1b) and its mRNA (Fig. 1c) was lower in the Lin28B knockdown cells (Lin28B-K/D) than in cells mock transduced with shRNA against luciferase (sh-Luc). We then determined the expression of let-7 in H1299 cells and their Lin28B-K/D counterparts and found that knockdown of Lin28B enhanced the biogenesis of let-7 family members detected (let-7a-1, b, c, d, e, f, and g). The results of two representatives of let-7 miRNAs were shown in Fig. 1d, the let-7a-1 was increased and let-7g was drastically induced in Lin28B-K/D cells relative to their basal levels in the wild-type H1299 cells but mir-16-1 was not induced at all.

PACA is associated with Lin28B

In mammalian cells, PCAF acetylates numerous proteins with multiple roles in the normal growth and function of cells, and it is also associated with the occurrence of cancer [21–23]. Previously we reported that PCAF directly interacted with and acetylated Lin28A, the paralog
of Lin28B [26]. To identify whether PCAF is associated with Lin28B, we assessed the interaction between PCAF and Lin28B by co-IP assay. Anti-GFP antibody was applied to co-immunoprecipitate GFP-tagged Lin28B with ectopic FLAG-PCAF in HEK293T cells (Fig. 2a). In addition, in vitro GST-pulldown assays indicated that FLAG-PCAF was pulled-down by GST-Lin28B in the HEK293T cell lysates (Fig. 2b). These results demonstrated an interaction between Lin28B and PCAF. To further identify the binding domain of Lin28B that interacts with PCAF, the coding region of the Lin28B gene was truncated; the representative protein fragments are shown in Fig. 2c. The co-IP assay revealed that the CSD of Lin28 (Fig. 2d, L1 & L4) was essential in mediating the interaction between Lin28B and PCAF. Similar results from in vitro GST-pulldown assays also indicated that the CSD of Lin28B mediated its binding with PCAF (Fig. 2e). Additionally, endogenous Lin28B- and PCAF-expressing H1299 cells were cultured on glass cover slips and treated with antibodies for immunofluorescence staining. Our results showed that Lin28B and PCAF co-localized in the nucleus of H1299 cells (Fig. 2f).

PCAF acetylates Lin28B and involves in repression of let-7
To explore whether Lin28B is acetylated by PCAF, an in vitro acetyltransferase assay was performed, as previously reported [26]. Using anti-acetyl lysine (AcK) antibody, we found that GST-Lin28B was acetylated only in the presence of the purified GST-fused HAT2-domain of PCAF (GST-PCAF-HAT), whereas PCAF was auto-acetylated (Fig. 3a). The expression constructs for FLAG-Lin28B were co-transfected into HEK293T cells with either Myc-PCAF or HAT-domain-deleted Myc-PCAF (Myc-PCAF-ΔHAT). Ectopically expressed Lin28B was acetylated by Myc-tagged PCAF in the cells, whereas no acetylation of Lin28B was observed with the HAT-domain-deleted PCAF (Fig. 3b). To explore the functional role of PCAF compared with its HAT-domain-deleted counterpart in non-modified H1299 cells, RT-qPCR analysis was performed, and elevated expression levels of the let-7a-1 and let-7g miRNAs were found in the presence of PCAF but not the HAT-domain-deleted PCAF, whereas mir-16-1 was not induced by PCAF transfection (Fig. 3c). These results suggested that the acetylation of Lin28B by PCAF is a critical event that abrogates the inhibitory effect of native Lin28B on the biogenesis of let-7a-1 and let-7g miRNAs in H1299 cells.

We have previously reported that the protein stability of Lin28A, the paralog of Lin28B, decreases when it is acetylated by PCAF [26]; however, no apparent change was observed in the protein level of Lin28B in the cells transfected with PCAF (data not shown). Co-IP assays showed that PCAF-ΔHAT still interacted with Lin28B in cells (Fig. 3d). To reveal the mechanism underlying the PCAF-mediated acetylation of Lin28B and its effect on the
biogenesis of let-7, we first determined the expression of Lin28B in different cell types and detected it in three cell lines, H1299, HepG2, and HEK293T, but it was not detectable in other cell lines including HCT116, MCF7, and HeLa cells (Fig. 3e). However, unlike HEK293T cells, Lin28B was mainly distributed in the nucleus of H1299 cells (last group of cells in Fig. 3e). Furthermore, as compared with localization of Lin28A mostly in the cytoplasm but not in the nucleus [8, 26], Lin28B was mainly located in nucleus in these cells, and acetylation of Lin28B by ectopic PCAF enabled its translocation from the nucleus to the cytoplasm of H1299 cells, as shown by the results of immunofluorescence assays (Fig. 3f) and western blot assays on nuclear/cytoplasmic extracts (Fig. 3g). The distribution of endogenous Lin28B detected by anti-Lin28B is similar with that of ectopic Lin28B staining with anti-FLAG in H1299 cells (Fig. 2f vs Fig. 3f), which is confirmed with results of western blot for the fractions of nuclear/cytoplasmic (Fig. 1a vs the first two lines of Fig. 3g). To analyze the effects of PCAF activity on Lin28B, we used a Cys574 mutant of PCAF (M-PCAF) and Lin28B-ΔCSD and detected the cytoplasmic distribution of Lin28B in H1299 cells. The results showed that M-PCAF did not increase the cytoplasmic distribution of Lin28B and that the CSD-deleted Lin28B was mainly distributed in the cytoplasm and was not affected by M-PCAF (Fig. 3h). These findings indicated that the acetylation of Lin28B was induced by PCAF, which removed the inhibition on let-7 biogenesis in H1299 cells, as schematically shown in Fig. 3i.

**Discussion**

In this report, we demonstrated that knockdown of Lin28B in H1299 cells elevated the level of let-7a-1/g, although this level of let-7a-1 was lower than that of let-7g in Lin28B-knockdown cells. The cell line H1299 was used as a model of lung adenocarcinoma and was derived from a lymph node metastasis of the lung. H1299
cells express high levels of Lin28B and low levels of let-7 [8, 16]. Lin28B, after acetylation by PCAF, might translocate from the nucleus to cytoplasm in H1299 cells and it may be further involved in the de-repression of let-7 biogenesis. These findings provide new insights into the potential application of acetylated Lin28B in mediating let-7 biogenesis, which may serve as a novel therapeutic approach for lung cancer.

In humans, there are twelve let-7 family members located at eight different chromosomal loci. Lin28A/B selectively represses the expression of let-7 miRNAs [11, 29–31]. It is well known that cytoplasmic Lin28A first binds to the conserved terminal loop of pre-let-7 and then recruits TUT4 for polyuridylation, thereby blocking Dicer cleavage [4, 5]. Although the precise mechanism underlying the Lin28B-mediated inhibition...
of let-7 remains controversial, it has been demonstrated that Lin28B expression and let-7 loss almost invariably correlate with poor prognosis [4]. Activation of Lin28B expression in cancer cells can be triggered by upstream transcriptional factors, such as c-Myc [18, 19] and NF-kB [20]. In addition, it has been reported that Merlin/NF2 is a key regulator of Lin28B localization and let-7 biogenesis in response to cell-cell contact [32]. Our data suggested that acetylation of Lin28B by PCAF may affect Lin28B localization and let-7 biogenesis.

Although we have previously reported that the protein level of Lin28A had decreases in PCAF-transfected cells [26], Lin28B was not substantially changed at the protein level (data not shown). Both Lin28A and Lin28B interact with PCAF via their CSD, and let-7 binds to the 3'UTR of both Lin28 paralogs, thereby repressing Lin28 translation in a negative feedback loop; however, TRIM71, a specific E3 ubiquitin ligase, negatively regulates Lin28B (but not Lin28A) protein levels by ubiquitinating its C-terminal unique region, which is absent in the Lin28A paralog [24, 25]. It might be interesting to study whether the acetylation of Lin28B by PCAF affects the polyubiquitination of Lin28B at the C-terminal region and may alter its stability.

**Conclusion**

In summary, we demonstrated that Lin28B interacts directly with PCAF via its cold shock domain and is acetylated by PACF. The HAT domain of PCAF is indispensable for PCAF mediated acetylation of Lin28B and de-repression of let-7a-1 and let-7g. Lin28B, after acetylation by PCAF, might translocate from the nucleus to cytoplasm in H1299 cells and it may be further involved in the de-repression of let-7 biogenesis. The discovery of a new role of PCAF in mediating Lin28B acetylation and, particularly, in elevating the level of microRNAs in lung adenocarcinoma-derived H1299 cells may shed new light on the potential application of let-7a-1 and/or let-7g in the clinical treatment of lung cancer.

**Abbreviations**

AcK: Acetyl lysine; Co-IP: Co-immunoprecipitation; CSD: Cold shock domain; DMEM: Dulbecco’s modified Eagle’s medium; Lin28B-K/D: Lin28B-knockdown cell lines; NLS: Nucleus localization signal; NoLS: Nucleolus localization signal; RT-qPCR: Quantitative real-time RT-PCR assays; ZF: CCHC zinc finger motif

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**Availability of data and materials**

The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

**Authors’ contributions**

TQ conceived, designed, performed, and analyzed experiments and also wrote the manuscript. FC, JW, YaZ, MC, and WS performed and analyzed experiments and revised the manuscript. YS conceived and designed experiments and wrote the manuscript. YeZ conceived, designed, and analyzed experiments and wrote the manuscript. All authors have read and approved the final version of this manuscript.

**Ethics approval and consent to participate**

Not applicable.

**Consent for publication**

Not applicable.

**Competing interests**

The authors declare that they have no competing interests.

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