Research Article

miR-101-3p Serves as a Tumor Suppressor for Renal Cell Carcinoma and Inhibits Its Invasion and Metastasis by Targeting EZH2

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Received 31 March 2021; Accepted 24 June 2021; Published 8 July 2021

Academic Editor: Tao Huang

Background. The role of miRNAs in renal cell carcinoma (RCC) is not certain. We wanted to study the biological functions and potential mechanisms of miR-101-3p in RCC. Methods. miR-101-3p was inhibited in A498 and OSRC-2 (two RCC cell lines). We studied its effect on cell invasion and proliferation. Target EZH2 of miR-101-3p was designated by different methods, including luciferase functional analysis and Western blotting. The expression level of the target gene in treated cells was quantitatively analyzed by quantitative real-time polymerase chain reaction. In addition, induction of miR-101-3p to prevent tumor formation of A498 cells in mice was further studied. Results. The overexpression of miR-101-3p significantly inhibited the proliferation, migration, and invasion in two RCC cells. Western blotting and luciferase functional analysis indicated that miR-101-3p regulated the expression of EZH2 in two cell lines. Mice inoculated with A498 and OSRC-2 cells transfected with miR-101-3p mimics showed significantly smaller xenografts and weaker EZH2 expression levels than the control group. Conclusions. miR-101-3p inhibited RCC cell proliferation, migration, and invasion by targeting EZH2.

1. Introduction

RCC is one of the most common malignancies of the urinary system [1], accounting for about 2%-3% of adult malignancies [2]. For localized and early-stage RCC, the best treatment is radical and partial nephrectomy, which has a 92.6% 5-year survival rate [3]. However, for metastatic RCC, the 5-year survival rate is expected to be very poor due to its resistance to radiotherapy and chemotherapy [4]. Although the advent of antiangiogenic drugs with tyrosine kinase inhibitors (TKI) has improved the prognosis of RCC, the associated resistance remains elusive [5]. It is important to uncover the mechanism of renal cancer and find potential prognostic biomarkers and therapeutic targets for renal cancer. Thus, we sought to find the molecular mechanism underlying RCC pathogenesis.

miRNA is a small nontranslated RNA that inhibits the expression of target genes through translation inhibition or transcriptional silencing [6]. Current studies have shown that miRNAs can act on at least 30% of protein-coding genes [7]. With the continuous development of technology, various miRNAs have been identified and have been widely proven to play an important role in tumorigenesis. Among them, miR-101-3p has been declared to regulate cancer development by acting as a tumor suppressor or an oncogene in tumorigenesis. For example, Wang et al. found that miR-101-3p promotes apoptosis of oral cancer cells by targeting BICC1 [8]. Wu et al. reported that it diametrically targets SRF to inhibit HOX transcript antisense RNA- (HOTAIR-) induced proliferation of gastric cancer cells [9]. In hepatocellular carcinoma, the LINC00052/miR-101-3p axis inhibited cell proliferation and metastasis by targeting SOX9 [10]. In non-small-cell lung cancer (NSCLC), it inhibits the growth and metastasis of NSCLC by blocking the PI3K/Akt signaling pathway by targeting MALAT-1 [11]. However, until now, its roles and underlying molecular mechanisms in RCC progression still remain unknown and need to be elucidated.
Figure 1: Continued.
Thus, we focused on the role of miR-101-3p in RCC and found its low expression in RCC tissue based on the starBase database and clinical tumor specimen analysis. Its overexpression inhibited the proliferation, invasion, and migration of RCC cells. In addition, we found that the enhancer of zeste 2 polycomb repressive complex 2 subunit (EZH2) is a direct downstream target of miR-101-3p. Then, this miRNA in RCC was studied using A498 and OSRC-2 cells as models. These two cell lines have been widely used in the study of RCC. Therefore, we used the functional gain/loss method to demonstrate that it affected the proliferation and invasion of RCC cells in vitro and in vivo, respectively. Our findings suggest the oncogenic role of miR-101-3p in RCC and may provide a new potential target for RCC therapy.

2. Materials and Methods

2.1. RCC Tissue Samples. Between January 2019 and June 2020, 11 RCC samples and normal samples were collected from Shanghai Tenth People’s Hospital. Fresh tumor tissue was stored in liquid nitrogen to avoid RNA degradation. The specimens were used with the consent of all patients and approved by the Ethics Committee of Shanghai Tenth People’s Hospital of Tongji University.

All procedures in this study were performed in accordance with the ethical standards of the institutional and/or national research committee and with the World Medical Association’s Declaration of Helsinki.

2.2. Bioinformatics Analysis. The GEO dataset GSE16441 was used for analysis. In this dataset, microRNAs from 17 RCC tumors and 17 corresponding nontumor samples were hybridized on a single-channel platform for miRNA expression analysis. Differentially expressed miRNAs were identified and presented by a clustered heat map and volcano plot [12, 13]. The expression of miR-101-3p and EZH2 was evaluated based on the starBase database (http://starbase.sysu.edu.cn/). The potential target genes of miR-101-3p were predicted using TargetScan (http://www.targetscan.org/), miRTarBase (http://mirtarbase.cuhk.edu.cn/), and miRDB (http://mirdb.org/) [14].

2.3. Cell Culture. Human RCC cell lines A498, OSRC-2, 786-O, and SW839 and human normal kidney tubular epithelial cell line (HK-2) were purchased from the Cell Bank of the Chinese Academy of Sciences (Shanghai, China) and were cultured, respectively, in DMEM (Invitrogen, Carlsbad, CA) supplemented with 10% fetal bovine serum (FBS), 50 U/ml of penicillin, and 50 μg/ml of streptomycin (Invitrogen).

2.4. RNA Extraction, Reverse Transcription, and Quantitative Real-Time PCR (qRT-PCR) Analyses. Total RNA was measured using a rapid RNA extraction kit (Epizyme, Shanghai). The purity and concentration of RNA were measured using a NanoDrop 2000 spectrophotometer. The cDNA was synthesized using a Bulge Loop™ miRNA RT primer kit (RiboBio, China). An SYBR green qPCR kit (Yeason, Guangzhou, China) was used for qRT-PCR. Expressions of miR-101-3p and EZH2 were normalized to GAPDH or U6 applying the 2-ΔΔCt method.

2.5. Western Blotting Analysis. The sample is lysed on ice for 30 minutes with a RIPA buffer (1:100 protease inhibitors). After measuring the protein content, calculate the sample amount. After adding enough electrophoresis solution, start to prepare for loading. After electrophoresis, the electrophoresis can be stopped just after the bromophenol blue runs out, and the membrane is transferred. The hybridization was carried out overnight at 4°C (1:10000 for GAPDH, 1:1000 for EZH2). Then, it was extensively washed and incubated in a blocking buffer for 1 h at room temperature with appropriate horseradish peroxidase combined with a
Figure 2: Continued.
secondary antibody (1:1000). After three times of washing and film scanning, we analyzed the molecular weight and net optical density value of the target band with the gel image processing system.

2.6. Cell Proliferation Assays. Cell Counting Kit-8 is used to measure cell proliferation. Briefly, 1 × 10^3 transfected RCC cells per well were seeded into 96-well plates and cultured for 24-120 h. Every 24 h, we measured optical density values using an automatic microplate reader (BioTek Solutions, Thousand Oaks, CA, USA) and incubated with CCK-8 (10 μl) in 5% CO₂ for 2 h [15].

2.7. Wound Healing Assays. First, draw horizontal lines evenly behind the 6-hole plate with a marker, cross the hole every 1 cm, and then lay the plate. On the next day, lines perpendicular to the horizontal lines were scratched with a spearhead. Then, wash the cells with PBS 3 times, and the cells were removed. After adding a serum-free medium, the cells were cultured in an incubator at 37 °C and 5% CO₂. Samples were taken at 0, 6, 12, and 24 h, and photos were taken.

2.8. Transwell Assays. Dilute Matrigel gel with a serum-free cell culture medium or PBS buffer at the ratio of 1:8 at 4°C, take 100 μl and evenly smear on the surface of the polycarbonate membrane in the upper chamber, place at 37°C for 0.5-1 h, and polymerize it into a gel. Add a medium containing 10% FBS or chemokines to the lower chamber of the 6-well plate, take cell suspension to the upper chamber, and finally put it into the incubator for 12-48 h culture, cell fixation, cell staining, and counting.

2.9. Cell Transfection. The miRNA negative control (miR-NC), miR-101-3p mimics, and inhibitors were transfected instantaneously using Lipofectamine 2000 according to the manufacturer’s protocol. In the overexpressed EZH2, the EZH2 vector and the overexpressed plasmid (oeEZH2) were synthesized by IBSBIO Biotechnology. We transfected the plasmid into HEK293T cells and packaged lentivirus. After 24h, the lentivirus supernatant was collected and used to infect cells.

2.10. Luciferase Reporter Assays. The binding sites of miR-101-3p and EZH2 were obtained from TargetScan, and vectors containing the mutant-type (Mut) or wild-type (WT) sequence were synthesized by Genomeditech (China). Cultivate HEK293T cells, inoculate them in 24-well plates, and grow them for 10-24 hours. The reporter gene plasmid and transcription factor expression plasmid are cotransfected into cells. The protein is extracted and used for luciferase detection. Add the substrate and measure the luciferase activity. Calculate relative fluorescence intensity and compare with the no-load control.

2.11. Immunohistochemistry (IHC). IHC was performed to assess the expression level of EZH2 in the tissues, as described earlier [16]. Frozen sections 4-8 μm were placed at room temperature for 30 minutes, fixed in acetone at 4°C for 10 minutes, washed with PBS for 5 minutes 3x, and incubated with hydrogen peroxide for 5-10 minutes to eliminate endogenous peroxidase activity.

2.12. Statistical Analysis. Statistical analysis was performed using GraphPad Prism 7 (GraphPad Software, CA). All data were mean ± standard deviation (SD) of three independent experiments. Statistical significance between groups was analyzed using Student’s t-test [17, 18]. Correlations between miR-101-3p and EZH2 were tested using Pearson’s correlation coefficient analysis. All data were examined and statistically significant at P < 0.05 [19].

3. Results

3.1. miR-101-3p Was Upregulated in RCC Tissues and Cell Lines. To detect its expression in RCC tissues, we searched...
Figure 3: Continued.
the GEO dataset and identified GSE95384, which conducted miRNA expression profiling array analyses in 8 RCC tumor samples compared with matched nontumor samples. It was considerably upregulated in RCC samples (fold changes > 2, \( P \text{ value} < 0.05 \)) (Figures 1(a) and 1(b)). Based on the starBase database, we also found overexpression of miR-101-3p in RCC tissues compared with normal samples \( (P < 0.01) \), and Kaplan-Meier survival analysis revealed that high expression of miR-101-3p was closely associated with poor survival \( (P < 0.005 \) (Figure 1(c)). And it was highly expressed compared with normal tissues and HK-2 (Figure 1(d)). Additionally, we also validated its expression in RCC cell lines and clinical specimens (Figures 1(e) and 1(f)). Because OSRC-2 and A498 showed higher expressions of miR-101-3p, we chose these two cell lines for further studies.

3.2. Overexpressing miR-101-3p Attenuated Proliferation, Migration, and Invasion of RCC Cells. miR-101-3p mimics were transfected into OSRC-2 and A498 cells, and the expression efficiency was detected by qRT-PCR (Figure 2(a)). CCK-8 assays showed that the viability of RCC cells was attenuated after transfection with the mimics (Figures 2(b) and 2(c)). In addition, cell migration of OSRC-2 and A498 cells was significantly suppressed after transfection of the mimics (Figures 2(d) and 2(e)). Transwell assay also showed that overexpression of miR-101-3p mimics inhibited the migration and invasion of RCC cells (Figures 2(g) and 2(h)).

3.3. EZH2 Was the Downstream Target of miR-101-3p. We queried the TargetScan, miRTarBase, and miRDB databases to explore downstream target genes of miR-101-3p. Seventeen candidates emerged after overlapping the results from three databases (Figure 3(a)). Since the above results show that miR-101-3p acts as an anticancer gene in RCC, we focused on tumor agonists in the predicted targets. Based on starBase and GEPIA databases, we measured the expression of RCC samples (data not shown) and found that EZH2 was apparently upregulated in RCC tissues compared with normal samples (Figures 3(f) and 3(g)). Pearson’s correlation coefficient analysis also displayed that the miR-101-3p
Figure 4: Continued.
expression level was negatively correlated with EZH2 for both our clinical samples and starBase (Figure 3(e)). And Kaplan-Meier survival analyses indicated that high EZH2 expression was closely related to poor survival ($P < 0.05$) (Figures 3(h)–3(j)) to further verify that miR-101-3p targeted EZH2. We obtained the binding sites of miR-101-3p in the 3′-UTR of EZH2 and inserted the luciferase reporter into the wild-type or mutant sequence (Figures 3(b) and 3(c)). It revealed that the mimics significantly reduced relative luciferase activity in the transfected wild-type HEK293T cells compared with the mutant group (Figure 3(d)). According to interpretation, the above results imply that EZH2 is the target gene of miR-101-3p.

3.4. Overexpressing miR-101-3p Prevented RCC Proliferation In Vivo. Western blotting and qRT-PCR verified that overexpression of miR-101-3p could reduce the protein level of EZH2 (Figures 4(a) and 4(b)). A498 cells were infected with lentivirus overexpressing miR-101-3p, and inserted the luciferase reporter into the wild-type or mutant sequence (Figures 3(b) and 3(c)). It revealed that the mimics significantly reduced relative luciferase activity in the transfected wild-type HEK293T cells compared with the mutant group (Figure 3(d)). According to interpretation, the above results imply that EZH2 is the target gene of miR-101-3p.

3.5. EZH2 Restoration Reversed the Effects of miR-101-3p in RCC Cells. We further validated that miR-101-3p mimics decreased EZH2 expression and ezrin 3p decreased EZH2 protein level in RCC cells. Western blotting and qRT-PCR were implemented to assess the expression of EZH2 in RCC cells (Figures 5(a) and 5(b)). We concluded that miR-101-3p mimics inhibited the protein level of EZH2 while the effect was likewise reversed after transfection with EZH2 overexpressed plasmids. It suggested that miR-101-3p prevented the malignant progression of RCC by preventing the expression of EZH2.

4. Discussion

Currently, studies have demonstrated that miRNAs are closely related to RCC occurrence [20–23]. However, as far as we know, its expression level and function in RCC nonetheless remain to have a lot of questions. We found it was highly reflected in tumor tissues, which was consistent with data from the starBase database. Moreover, the results from miRNA expression profiling arrays (GSE16441) also revealed that it was significantly upregulated in RCC tumor specimens.

To date, miR-101-3p has been widely mentioned in other tumors; however, its role in renal carcinogenesis has not been determined. It prevents growth and metastasis of NSCLC by...
Figure 5: Continued.
miR-101-3p mimics and EZH2 overexpressed plasmids or NC. Data represent mean ± SD. The related data would be provided if required.

**Data Availability**

The related data would be provided if required.

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**Conflicts of Interest**

The authors declare that they have no conflicts of interest.

**Authors’ Contributions**

The contributions of the authors involved in this study are as follows: (i) conception and design: Yunfei Xu and Yunze Dong; (ii) administrative support: Yunfei Xu; (iii) provision of study materials or patients: Tiancheng Xie and Huan Liu; (iv) collection and assembly of data: Yuchen Gao; (v) data analysis and interpretation: Yunze Dong; (vi) manuscript writing: all authors; and (vii) final approval of manuscript: all authors. Yunze Dong and Yuchen Gao contributed equally to this work.

**Acknowledgments**

This study was funded by the National Natural Science Foundation of China (Nos. 81971371 and 81902567).

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