MicroRNA-489-3p plays a significant role in congenital hypothyroidism through regulating neuronal cell apoptosis via targeting translationally controlled tumor protein 1

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Abstract. Accumulating reports have indicated that congenital hypothyroidism (CH) is an endocrine disorder caused by underdeveloped thyroid gland or thyroid dyshormonogenesis. It has been also reported that certain microRNAs (miRNAs) may exert protective effects against the development of CH. However, whether miR-489-3p regulates CH progression remains unclear. The aim of the present study was to investigate the effects of miR-489-3p on CH and elucidate the underlying mechanisms. Therefore, Sprague Dawley rats were injected with propylthiouracil (50 mg/day) to establish a CH model. Reverse transcription-quantitative PCR (RT-qPCR) assay demonstrated that miR-489-3p was upregulated in the hippocampal tissues of CH rats. Furthermore, the TargetScan software was employed to predict the target gene of miR-489-3p, and a dual luciferase reporter assay revealed that translationally controlled tumor protein 1 (TPT1) was directly targeted by miR-489-3p. Additionally, RT-qPCR and western blot assays suggested that TPT1 was markedly downregulated in the hippocampal tissues of CH rats compared with control rats. In addition, inhibitor control, miR-489-3p inhibitor, control-shRNA or TPT1-shRNA were injected into CH rats. The results of the open-field and forced swimming tests revealed that miR-489-3p inhibitor notably improved the behavior of CH rats. Flow cytometry was applied to explore the effects of miR-489-3p inhibitor on neuronal cell apoptosis, and the findings indicated that miR-489-3p inhibitor attenuated CH-induced neuronal cell apoptosis, whereas these effects were reversed by treatment with miR-489-3p inhibitor and TPT1-shRNA. Finally, the function of miR-489-3p in neuronal cells was investigated in vitro. Neuronal cell viability, apoptosis and the expression of apoptosis-related proteins were determined using MTT assay, flow cytometry and western blot analysis, respectively. The results demonstrated that miR-489-3p inhibitor enhanced cell viability, suppressed apoptosis and upregulated Pim-3, phosphorylated (p)-Bad (Ser112) and Bcl-XL expression. Rescue experiments indicated that these effects were reversed following silencing of TPT1. Taken together, the findings of the present study demonstrated that miR-489-3p inhibitor could relieve CH-induced neurological damage through regulating TPT1 expression.

Introduction

Previous reports have revealed that thyroid hormone deficiency may lead to functional injuries, including congenital hypothyroidism (CH) (1,2). Several developmental processes may be induced by CH, such as disrupted neurogenesis and abnormal hippocampal neuron apoptosis (3-5). It was previously demonstrated that the majority of newborn infants with CH did not have significant clinical manifestations (6). It is widely accepted that the hippocampus is involved in cognitive activities in humans. Accumulating evidence has suggested that hypothyroidism promotes hippocampal neuron apoptosis (4,7-9). Moreover, a previous study suggested that perinatal hypothyroidism affects behavioral development and may lead to a decrease in spatial learning ability and memory, and these changes are closely associated with the increasing number of apoptotic neurons in the hippocampus (10). Therefore, early diagnosis and timely treatment are crucial for patients with CH.

MicroRNAs (miRNAs) are highly conserved small RNAs, 20-22 nucleotides in length, which modulate protein function via directly binding to the 3'-untranslated region (3'-UTR) of their target mRNAs (11). Recent studies have demonstrated that changes in miRNA levels are involved in numerous diseases, including myocardial ischemia (12), inflammation (13), diabetes (14) and CH (4,8,9). In addition, miRNAs serve important functions in physiological processes, such as post-transcriptional regulation, cell proliferation and apoptosis. For example, You et al (15) found that miR-498 suppressed gastric cancer cell proliferation, migration and invasion via targeting B lymphoma Mo-MLV insertion region 1 homolog and inactivating the AKT pathway. Zhou et al (16) reported that miR-429 attenuated neuroblastoma cell viability, migration and...
invasion via the nuclear factor-κB pathway. Furthermore, it has been reported that miR-489-3p serves key functions in several diseases. For example, Chen et al.(17) demonstrated that increased expression of miR-489-3p and miR-630 inhibited oxaliplatin uptake in renal cell carcinoma via targeting octamer-binding protein 2. Additionally, Kuppa et al (18) reported that autotaxin accelerated cancer development via enhancing the expression of mitogen-activated protein kinase kinase 1 and overriding the role of miR-489-3p. However, the role of miR-489-3p in neuronal cells and CH remains elusive. Therefore, the present study was undertaken to investigate the role of miR-489-3p in CH and elucidate the potential underlying mechanisms.

Translationally controlled tumor protein 1 (TPT1), a multifunctional protein, has been evidenced to be highly expressed in various diseases, including polycystic ovary syndrome (19) and breast cancer (20), and is involved in cell proliferation, invasion, cell cycle progression and apoptosis (21). Furthermore, accumulating evidence has suggested that miR-489-3p inhibits glioblastoma progression via downregulating TPT1 (22).

The aim of the present study was to investigate the effects of miR-489-3p on hippocampal neuronal cell apoptosis in CH in vivo and vitro and to determine the role of the TPT1/Pim-3 signaling pathway in CH, thereby improving our understanding of the molecular biology of the CH.

Materials and methods

Animals. A total of 50 female pregnant Sprague Dawley rats (age, 4–6 weeks; weight, 200±5 g) were purchased from the Experimental Animal Center of Shanghai and kept in a controlled environment (temperature, 22±1˚C; humidity, 50-60%; 12-h light/dark cycle). All animal experiments were carried out according to the guidelines provided by the National Institutes of Health (NIH) for the Care and Use of Laboratory Animals. The study protocol was approved by the Animal Ethics Committee of the Experimental Animal Center of Yancheng Maternal and Child Health Hospital.

Establishment of the CH model. To establish the CH rat model, pregnant rats were intraperitoneally injected with propylthiouracil (50 mg/kg) starting on gestational day 15 and then daily thereafter until parturition, in order to generate pups with CH (4,23). For CH therapy, the newborn rats (12 days old) were anesthetized with intraperitoneal injection of 2% pentobarbital sodium (40 mg/kg), and then their skulls were opened as previously described (9). Subsequently, inhibitor control (5’-CAG UAU UUU UGU GUA CAA-3’; GenePharma Co., Ltd.), miR-489-3p inhibitor (5’-CAG UAC UUU UGU GUA CAA-3’; GenePharma Co., Ltd.), miR-489-3p inhibitor + control-shRNA (cat. no. sc-108060; Santa Cruz Biotechnology, Inc.) or miR-489-3p inhibitor + TPT1-shRNA (cat. no. sc-40675-SH; Santa Cruz Biotechnology, Inc.) containing 2% B27, 100 U/ml penicillin/streptomycin and 0.5 mmol/l glutamine (Gibco; Thermo Fisher Scientific, Inc.) were synthesized by Shanghai GenePharma Co., Ltd. and transfected into neurons using the Lipofectamine 2000 reagent (Invitrogen; Thermo Fisher Scientific, Inc.) according to the manufacturer's protocol. Following incubation for 48 h, the transfection efficiency was confirmed using reverse transcription-quantitative PCR (RT-qPCR) and western blot analyses.

MTT assay. Cell proliferation was assessed using an MTT assay. Briefly, neurons (5x10⁴ cells per well) were seeded into 96-well plates in triplicate, transfected with inhibitor control, miR-489-3p inhibitor, control-shRNA or TPT1-shRNA, and cultured at 37˚C for 48 h. Following transfection, each well was supplemented with 20 µl MTT solution (Sigma-Aldrich; Merck KGaA) and the cells were cultured for an additional 4 h. Subsequently, the medium was discarded, and 100 µl DMSO was added to dissolve the formazan. The absorbance at 490 nm was determined using a microplate reader (BioTek Instruments, Inc.).

Cell apoptosis assay. Treated neurons were digested, cleaned and collected at 4˚C overnight. For the detection of cell apoptosis, the Annexin V-FITC/propidium iodide apoptosis detection kit (Beyotime Institute of Biotechnology) was used according to the manufacturer's protocol. Subsequently, apoptotic cells were identified using the FACScan flow cytometry system (BD Biosciences), and their number was measured using the FlowJo 7.6.1 software (BD Biosciences).

Dual luciferase reporter assay. TargetScan bioinformatics software (version 7.2; http://www.targetscan.org/vert_72/) was employed to identify the potential targets of miR-489-3p. Subsequently, the 3’-UTR of TPT1 containing miR-I-489-3p-binding was sub-cloned into pMIR vectors (Ambion; Thermo Fisher
Scientific, Inc.) to generate the TPT1 wild-type (TPT1-WT) and TPT1 mutant (TPT1-MUT) plasmids. TPT1 was identified as a possible target of miR-489-3p. A QuikChange Site-Directed Mutagenesis Kit (Stratagene; Agilent Technologies, Inc.) was applied according to the manufacturer's instructions to point-mutate the miR-489-3p-binding domain in the 3'-UTR of TPT1. For luciferase reporter activity analysis, 293T cells were co-transfected with 1 ng TPT1-WT or 1 ng TPT1-MUT and 100 nM miR-489-3p mimic (5'-GUGACAUCAUAUAACGG CAGC-3'; Suzhou GenePharma Co., Ltd.) or 100 nM mimic control (5'-UUUGCCACAGUGUCAGCGT-3'; Suzhou GenePharma Co., Ltd.) with Lipofectamine® 2000 (Invitrogen; Thermo Fisher Scientific, Inc.) according to the manufacturer's instructions. After 24 h, the luciferase activity was determined using the Dual-Luciferase Reporter Assay System (Promega Corporation) and normalized to Renilla luciferase activity.

Behavioral tests
Open-field test (OFT). This test was conducted to assess the anxiety and physical activity of experimental rats (21 days old). OFT was carried out into a 48x48x36 cm opaque apparatus partitioned into 16 equal squares by white lines, including center squares and a peripheral area. Mice were left in the center squares for 5 min and their activity was tracked using a digital camcorder (Ethovision 2.0; Noldus Information Technology). In this test, the following four parameters were analyzed: Ambulation distance, referring to the total distance of the grid lines crossed; center square entries, referring to the frequency of squares crossed with all four paws; center area duration, referring to the accumulated time of rat in central square; and rearing, referring to the frequency of each rat standing on the hind paws. The maze was wiped with 75% ethanol prior to testing another rat.

Forced swimming test (FST). This test was conducted to assess depression-like behavior in rats (21 days old). Briefly, the rats were placed into a transparent cylinder container (diameter, 22 cm; depth, 40 cm) filled with water to a height of 30 cm at 25˚C for 6 min. Subsequently, the swimming duration of each rat was recorded. During this experiment, immobility time was defined as the time during which the rats were floating motionless or keeping their head above water for 4 min. Swimming activity was indicated as a non-depressive behavior.

RT-qPCR analysis. Total RNA from cultured neurons or hippocampal tissues was harvested using TRIzol® reagent (Takara Biotechnology Co., Ltd.) according to the manufacturer's instructions. Subsequently, RNA was reverse-transcribed into cDNA using the cDNA Reverse Transcription kit (Takara Biotechnology Co., Ltd.) according to the manufacturer's instructions. The expression of miR-489-3p in the hippocampal tissue of CH rats was determined by RT-qPCR analysis. The results revealed that the miR-489-3p expression levels were higher in the hippocampal tissues of CH rats compared with those in the control group (Fig. 1A). To verify the mechanisms underlying the effects of miR-489-3p, bioinformatics analysis was performed to predict its target genes. The analysis identified a potential binding site of miR-489-3p on TPT1 3'-UTR (Fig. 1B). Previous reports have indicated that TPT1 is a target gene of miR-489-3p. Therefore, a dual luciferase reporter assay was performed to verify the association between miR-489-3p and TPT1. It was first confirmed that, compared with the mimic control group, miR-489-3p mimic significantly enhanced miR-489-3p expression in 293T cells (Fig. 1C). As shown in Fig. 1D, miR-489-3p notably decreased luciferase activity in the TPT1-WT group, but not in the TPT1-MUT group, compared with the control group. These findings suggested that miR-489-3p directly targets TPT1. Furthermore, RT-qPCR and western blot assays were performed to assess whether miR-489-3p regulates TPT1 expression. RT-qPCR analysis indicated that TPT1 mRNA was

Western blot analysis. Total proteins were extracted from neurons and hippocampi using a lysis buffer (Beyotime Institute of Biotechnology), and protein concentration was measured using a BCA assay (Solarbio Science and Technology Co., Ltd.). Subsequently, the protein samples were resolved on 10% SDS-PAGE and transferred onto PVDF membranes. The membranes were blocked with 5% skimmed milk in TBS containing 0.1% Tween at room temperature for 1.5 h. The membranes were then incubated with primary antibodies against TPT1 (cat. no. 5128; dilution, 1:1,000; Cell Signaling Technology, Inc.), Pim-3 (cat. no. 4165; dilution, 1:1,000; Cell Signaling Technology, Inc.), Bcl-xL (cat. no. 2764; dilution, 1:1,000; Cell Signaling Technology, Inc.) and GAPDH (cat. no. 5174; dilution, 1:1,000; Cell Signaling Technology, Inc.) at 4˚C overnight. After washing with PBS-T, the membranes were incubated with corresponding secondary antibody (cat. no. 7074; dilution, 1:1,000; Cell Signaling Technology, Inc.) for 2 h at room temperature. Finally, the protein bands were quantified using an ECL reagent (Cytiva) according to the manufacturer's instructions.

Statistical analysis. Data are expressed as the mean ± standard deviation from three independent experiments. GraphPad Prism 6 (GraphPad Software, Inc.) and SPSS 21.0 software (IBM Corp.) were employed for statistical analyses. The variables were analyzed using Student's t-test or one-way ANOVA followed by Tukey's post hoc test. P<0.05 was considered to indicate a statistically significant difference.

Results
miR-489-3p is upregulated in hippocampal tissues of CH rats via targeting TPT1. The expression of miR-489-3p in the hippocampal tissue of CH rats was determined by RT-qPCR analysis. The results revealed that the miR-489-3p expression levels were higher in the hippocampal tissues of CH rats compared with those in the control group (Fig. 1A). To verify the mechanisms underlying the effects of miR-489-3p, bioinformatics analysis was performed to predict its target genes. The analysis identified a potential binding site of miR-489-3p on TPT1 3'-UTR (Fig. 1B). Previous reports have indicated that TPT1 is a target gene of miR-489-3p. Therefore, a dual luciferase reporter assay was performed to verify the association between miR-489-3p and TPT1. It was first confirmed that, compared with the mimic control group, miR-489-3p mimic significantly enhanced miR-489-3p expression in 293T cells (Fig. 1C). As shown in Fig. 1D, miR-489-3p notably decreased luciferase activity in the TPT1-WT group, but not in the TPT1-MUT group, compared with the control group. These findings suggested that miR-489-3p directly targets TPT1. Furthermore, RT-qPCR and western blot assays were performed to assess whether miR-489-3p regulates TPT1 expression. RT-qPCR analysis indicated that TPT1 mRNA was
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miR-489-3p inhibitor improves CH rat behavior and reduces CH-mediated neuronal cell apoptosis. To determine whether miR-489-3p is involved in CH, inhibitor control, miR-489-3p inhibitor, miR-489-3p inhibitor + control-shRNA or miR-489-3p inhibitor + TPT1-shRNA were injected into rats to establish CH rat models. The rats were divided into the following six groups: Control; CH; inhibitor control; miR-489-3p inhibitor; miR-489-3p inhibitor + control-shRNA; and miR-489-3p inhibitor + TPT1-shRNA groups. First, RT-qPCR analysis was performed to determine miR-489-3p and TPT1 expression in rat hippocampal tissues in different groups. Compared with the control group, miR-489-3p was upregulated in the model group, while this increase was reversed following treatment with miR-489-3p inhibitor (Fig. 2A). Furthermore, the expression of TPT1 was lower in the model group compared with that in the control group. In addition, compared to the CH + inhibitor control group, TPT1 was upregulated in CH rats in the CH + miR-489-3p inhibitor group, and this effect was significantly reversed by TPT1-shRNA (Fig. 2B).

Subsequently, behavioral tests were performed to evaluate the neuronal injury-induced behavioral variation following treatment with miR-489-3p inhibitor. miR-489-3p inhibitor attenuated rat anxiety- and depressive-like behavior, as observed in the FST (Fig. 2C) and OFT (Fig. 2D and E), respectively. These results indicated that miR-489-3p inhibitor significantly improved the behavior of CH rats. Furthermore, cell apoptosis assay was performed to assess the effect of miR-489-3p inhibitor on neuronal cell apoptosis. The data demonstrated that miR-489-3p inhibitor suppressed CH-induced neuronal cell apoptosis, while this effect was reversed by TPT1-shRNA (Fig. 2F and G). The aforementioned findings suggested that miR-489-3p inhibitor may relieve CH via regulating neuronal cell apoptosis.

TPT1-shRNA reverses the effects of miR-489-3p inhibitor on TPT1 expression in neuronal cells. Subsequently, the effect of miR-489-3p inhibitor on TPT1 expression was explored in neuronal cells by knockdown assay. miR-489-3p inhibitor, inhibitor control, miR-489-3p inhibitor + control-shRNA or miR-489-3p inhibitor + TPT1-shRNA were transfected into neuronal cells, and transfection efficiency was determined using RT-qPCR and western blot assays. The RT-qPCR results suggested that miR-489-3p inhibitor successfully downregulated miR-489-3p expression in neuronal cells compared with the inhibitor control group (Fig. 3A). Furthermore, in neuronal cells, transfection with TPT1-shRNA downregulated TPT1 expression at both the mRNA and protein levels compared with the control-shRNA group (Fig. 3B and C). In addition, compared with the inhibitor control group, miR-489-3p inhibitor upregulated TPT1 mRNA and protein expression levels. On the contrary, the effects of miR-489-3p inhibitor were reversed following transfection of neuronal cells with...
TPT1-shRNA (Fig. 3D and E). Taken together, these results confirmed that miR-489-3p negatively regulated TPT1 expression in neuronal cells.

TPT1 silencing abolishes the effect of miR-489-3p inhibitor on neuronal cell viability and apoptosis via the TPT1/Pim-3 signaling pathway. To further explore the effect of miR-489-3p on CH, MTT and flow cytometry assays were conducted to assess neuronal cell viability and apoptosis, respectively. Neuronal cells were transfected with miR-489-3p inhibitor, inhibitor control, miR-489-3p inhibitor + control-shRNA or miR-489-3p inhibitor + TPT1-shRNA. As shown in Fig. 4A, miR-489-3p inhibitor markedly increased cell viability compared with that observed in the inhibitor control group. However, TPT1-shRNA reversed this effect and significantly inhibited cell viability (Fig. 4A). In addition, flow cytometry revealed that miR-489-3p inhibitor significantly reduced cell apoptosis, while TPT1-shRNA rescued the inhibitory effects of miR-489-3p inhibitor on neuronal cells (Fig. 4B).

Additionally, the mechanism underlying the effect of the miR-489-3p inhibitor on the suppression of neuronal cell apoptosis was investigated. The expression levels of the apoptosis-related proteins Pim-3, Bcl-xL, p-Bad (Ser112) and Bad were detected using western blot assay. The results demonstrated that downregulation of miR-489-3p increased the expression levels of Pim-3, Bcl-xL, p-Bad (Ser112) and the ratio of p-Bad (Ser112)/Bad in neuronal cells. However, this effect was reversed following co-transfection with TPT1-shRNA (Fig. 4C-F). These findings suggested that miR-489-3p may promote CH progression by mediating neuronal cell apoptosis by targeting TPT1.

Discussion

CH, one of the most common preventable diseases, may be associated with mental retardation, various developmental barriers and high anxiety scores (25). Alcigir et al (5) revealed the neuroprotective activity of cannabinoid receptor 2 against oxidative stress and apoptosis in rat pups with experimentally induced CH. Li et al (9) reported that miR-124-3p attenuated the progression of CH via targeting programmed cell death protein 6. miRNAs are small non-coding molecules that play key roles in myocardial diseases (26) and diseases of the nervous system, such as Alzheimer’s (27) and Parkinson’s disease (28). In addition, miRNAs play critical roles in the development of tumors, thus serving as diagnostic markers (29). Several studies have evidenced that numerous miRNAs are differentially expressed in CH tissues and may be involved in the development of CH (4,8,9). However, the expression and associated mechanisms of action of miR-489-3p in CH remain elusive. Therefore, the present study was undertaken to investigate the exact role of miR-489-3p in CH.

Propylthiouracil was injected into pregnant rats to establish a CH model, and the results revealed that miR-489-3p was markedly upregulated in the hippocampi of CH rats compared with the control group. Furthermore, the potential targets of miR-489-3p were predicted using bioinformatics assay. Therefore, a binding site for miR-489-3p was identified in the TPT1 3'-UTR. The association between miR-489-3p and TPT1 3'-UTR was confirmed by a dual luciferase reporter assay. TPT1, a highly conserved protein, has been reported to be combined with other proteins to alter protein expression levels (30). Our findings were consistent with those of
previous reports suggesting that TPT1 was directly targeted by miR-489-3p (22). The present study demonstrated that TPT1 was downregulated in the hippocampal tissues of CH rats compared with the control group. The aforementioned observations indicated that miR-489-3p may be involved in CH via negatively regulating TPT1 expression.

Dysregulation of miRNA expression has been associated with several diseases. To identify whether miR-489-3p affects the progression of CH, inhibitor control, miR-489-3p inhibitor, miR-489-3p inhibitor + control-shRNA or miR-489-3p inhibitor + TPT1-shRNA were transfected into neuronal cells. (A) The levels of miR-489-3p were measured using RT-qPCR analysis in the inhibitor control and miR-489-3p inhibitor groups. (B) RT-qPCR and (C) western blot analysis of TPT1 expression in neuronal cells following treatment with control-shRNA and TPT1-shRNA. (D) RT-qPCR and (E) western blot assays were conducted to assess TPT1 expression in neuronal cells following transfection with miR-489-3p inhibitor + control-shRNA or miR-489-3p inhibitor + TPT1-shRNA. **P<0.01 vs. the inhibitor control group; ##P<0.01 vs. the control-shRNA group; ###P<0.01 vs. the miR-489-3p inhibitor + control-shRNA group. miR, microRNA; TPT1, translationally controlled tumor protein 1; RT-qPCR, reverse transcription-quantitative PCR.

The effects of miR-489 on cell viability and apoptosis have been previously reported. Wu et al (33) demonstrated that miR-489 suppressed multiple myeloma cell proliferation via inhibiting lactate dehydrogenase A-mediated aerobic glycolysis. Consistent with this finding, Gao et al (34) reported that miR-489 may suppress tumor growth and invasion via targeting histone deacetylase 7 in colorectal cancer. In addition, it has been demonstrated that TPT1 promotes cell proliferation and attenuates apoptosis (21). Hippocampal neuronal cell apoptosis is considered as the key characteristic of CH (35). Therefore, the effect of miR-489-3p on hippocampal neuronal cell apoptosis was explored. The in vitro results indicated that miR-489-3p inhibitor enhanced neuronal cell viability and inhibited apoptosis, which was alleviated by TPT1-shRNA. It has been reported that Pim-3, a member of the proto-oncogene Pim family, regulates cell viability during cancer development (36). For example, Fan et al (37) revealed that Pim-3 reduced cell proliferation and apoptosis in A549 lung adenocarcinoma cells. Additionally, Pim-3 enhanced melanoma cell migration and invasion via promoting signal transducer and activator of transcription 3 phosphorylation (38). Therefore, the expression of apoptosis-related proteins was further assessed using western blot analysis. The data demonstrated...
that the expression levels of Pim-3, p-Bad (Ser112) and Bcl-xL were significantly increased by miR-489-3p inhibitor in hippocampal neurons. Furthermore, transfection with TPT1-shRNA abolished the miR-489-3p inhibitor-mediated upregulation of Pim-3, p-Bad (Ser112) and Bcl-xL.

A limitation of the present study was that it did not include a group treated exclusively with sh-TPT1, which would further improve our understanding of the regulatory association between miR-489-3p and TPT1.

In conclusion, the results of the present study suggested that the miR-489-3p inhibitor may relieve CH-associated neurological damage via regulating the TPT1/Pim-3 signaling pathway. The findings of the study may contribute to the better understanding of the molecular biology of CH.

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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

QL and YL designed the current study, collected the data, performed statistical analysis and interpretation, and prepared the manuscript. YZ collected and analyzed the data. All authors read and approved the final manuscript. QL and YZ confirm the authenticity of all the raw data.

Ethics approval and consent to participate

All animal experiments were carried out according to the guidelines provided by the National Institutes of Health (NIH) for the Care and Use of Laboratory Animals. The study protocol was approved by the Animal Ethics Committee of the Experimental Animal Center of Yancheng Maternal and Child Health Hospital.
miRNAs as Tumor suppressor miR-145-5p sensitizes prolactin β-üä MicroRNA-copy number variations in Up-regulation of miR-489-3p and Functional omics analyses reveal only minor 16. 11. 10. 9. 6. 3. References The authors declare that they have no competing interests. Gao J, Zeng S and Yu L; Chen L, Qin Z, Lei J, Ye S, Zeng K, Wang H, Ying M, Gao J, Zeng S and Yu L: Up-regulation of mir-489-3p and mir-630 inhibits oxaliplatin uptake in renal cell carcinoma by targeting OCT2. Acta Pharmac Sin B 9: 1008-1020, 2019. 18. Kuppa SS, Jia W, Liu S, Nguyen H, Smyth SS, Mills GB, Dobkin KK, Hardman WJ and Murph MM: Autotaxin exacerbates tumor progression by enhancing MEK1 and overriding the function of mir-489-3p. Cancer Lett 432: 84-92, 2018. 19. Sun X, Su S, Zhang G, Zhang H and Yu X: MiR-204 suppresses cell proliferation and promotes apoptosis in ovarian granulosa cells via targeting TPT1 in polycystic ovary syndrome. Biochem Cell Biol 97: 554-562, 2019. 20. 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