miR-124 inhibits proliferation and invasion of human retinoblastoma cells by targeting STAT3

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Abstract. A growing body of evidence suggests that microRNA-124 (miR-124) functions as tumor-suppressor, and involves in tumor initiation, development and metastasis in major classes of human cancers; however, the biological role and underlying molecular mechanism of miR-124 in retinoblastoma (RB) remain unknown. Therefore, we investigated the biological activity and underlying molecular mechanism of miR-124 in human retinoblastoma. In the present study, our results demonstrated the downregulation of miR-124 in RB tissues and RB cell lines compared with normal retinal tissues. The ectopic expression of miR-124 in the RB cell lines (Y79 and SO-RB50) suppresses cell proliferation, migration and invasion, induced cell apoptosis in vitro. Furthermore, signal transducer and activator of transcription 3 (STAT3) was identified as a new target of miR-124, and overexpression of miR-124 decreased STAT3 expression on mRNA level and protein level in human RB cells. We also found that STAT3 mRNA expression was upregulated and inversely correlated with miR-124 expression in the RB tissues (r=-0.683; P<0.001). Restoration of the expression of STAT3 rescues the effects induced by miR-124 in RB cells. The findings of the present study suggested that miR-124 functioned as tumor suppressor in RB, at least in part, by targeting STAT3, and that it could serve as a potential candidate for RB therapeutics.

Introduction

Retinoblastoma (RB), a deadly pediatric eye cancer, is the most common primary intraocular malignancy in children worldwide (1). The mortality rate among children with RB is 50-70% in the underdeveloped countries (2). The reason of high mortality rate of RB is mainly frequent metastasis of RB and intracranial neuroblastic malignancy (trilateral RB) (3,4). It is an urgent need to further study the biology and molecular mechanisms of RB that cause RB procession and metastasis, and identify the specific biomarkers and therapy agents for improving the therapeutic outcome of patients with RB.

MicroRNAs (miRNAs) are a novel class of short (18-25 nucleotides in length) noncoding RNAs that regulate gene expression by repressing translation and cleaving their target mRNAs through binding to complementary sites in their 3' untranslated region (3'UTR) (5,6). It has been demonstrated due to aberrant expression. An accumulating body of evidence showed that miRNAs are involved in various biological processes such as development, differentiation, invasion, proliferation, apoptosis and stress response (7). It has been demonstrated that the altered expression of miRNAs contributes to the initiation and progression of cancer, and functions as tumor suppressors and oncogene (8,9). Numerous miRNAs have been found to be involved in the development of RB (10,11), suggesting that miRNAs may potentially serve as a novel strategy of diagnosis and therapy to RBs.

There is a growing interest particularly toward microRNA-124 (miR-124) in context of various types of cancers. miR-124 has been reported to be downregulated and function as a tumor suppressor in a variety of human cancers, such as prostate cancer (12), glioma (13), lung adenocarcinoma (14), breast (15), gastric (16) and colorectal cancer (17). However, the expression of miR-124 in patients with RB, and its biological functions in human RB cells, as well as the molecular mechanisms by which miR-124 exerts its functions, remains largely unclear. Therefore, the aims of the present study were to investigate the miR-124 expression in RB tissues and cell lines, and to evaluate its role and underlying mechanisms in RB. The result of the present study showed that miR-124 expression was downregulated in RB tissues and cell lines; and that overexpression of miR-124 in RB cells inhibited cell proliferation, migration and invasion and induced cell apoptosis in vitro by targeting signal transducer and activator of transcription 3 (STAT3).

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of transcription 3 (STAT3). These findings provide a novel therapeutic strategy for treatment of RB.

Materials and methods

Human tissue samples and cell lines. Forty human RB and 20 normal retina tissues were provided by the First Hospital of Jilin University (Changchun, China). All tissue samples were harvested at surgery, immediately frozen in liquid nitrogen and stored at -80°C until RNA extraction. The present study was approved by the Ethics Committee of the First Hospital of Jilin University (Changchun, China). All of the experiments were undertaken with the understanding and written consent of each patients or family.

Two human RB cell lines (Y79 and SO-RB50) were purchased from the Type Culture Collection of the Chinese Academy of Sciences (Shanghai, China), and were grown in RPMI-1640 medium (Gibco, Grand Island, NY, USA) supplemented with 10% heat-inactivated fetal bovine serum (FBS; HyClone, Logan, UT, USA), 100 U/ml penicillin or 100 mg/ml streptomycin in a humidified atmosphere of 95% air and 5% CO₂ at 37°C.

Transfection experiments. miR-124 mimic (miR-124) and corresponding miRNA-negative control (miR-NC) were purchased from GenePharma Co., Ltd. (Shanghai, China). The STAT3 sequence was amplified from human genomic DNA using a standard PCR protocol, and inserted into the pcDNA3.1 (Invitrogen, Shanghai, China). For the transfection experiments, 2x10⁵ RB cells were seeded in a 6-cm dish in antibiotic-free RPMI-1640 medium with 10% FBS. Twenty-four hours later, the specific molecular production was transfected into the RB cells using Lipofectamine 2000 (Invitrogen, Carlsbad, CA, USA) according to the manufacturer's instructions.

Quantitative reverse transcription polymerase chain reaction (qRT-PCR). Total RNA from the cultured cells and frozen tissues was isolated using TRIzol reagent (Invitrogen) following the manufacturer's instructions. To quantify miR-124, total RNA was reversely transcribed into cDNA using One Step PrimeScript miRNA cDNA Synthesis kit (Qiagen, Valencia, CA, USA). Then, miR-124 level was detected using the TaqMan miRNA assay kit (Takara, Dalian, China). For qRT-PCR, the STAT3 mRNA level was detected using the real-time PCR system. The primer and GAPDH was used as previously described (18). Relative expression of miRNA and mRNA was calculated using the 2^ΔΔCt method following normalization to U6 or GAPDH expression, respectively.

Cell proliferation and apoptosis analyses. Cell proliferation was determined using 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT; Sigma, St. Louis, MO, USA) assays. Briefly, 5x10⁴ transfected cells/well were seeded into a 96-well plate and cultured for 72 h. Then, 20 µl MTT solution (5 mg/ml) was added into each well and additionally cultured for 4 h, then, MTT solution was removed and 150 µl dimethyl sulfoxide (DMSO; Sigma-Aldrich, St. Louis, MO, USA) was added to each well. Optical density (OD) was detected at the wavelength of 570 nm using a Benchmark Plus™ microplate spectrometer (Bio-Rad, Hercules, CA, USA).

For cell apoptosis, the cells were collected 48 h after transfection and washed with phosphate-buffered saline (PBS). Then, 5x10⁴ cells were resuspended in 500 µl of binding buffer containing 5 µl of Annexin V-fluorescein isothiocyanate (FITC) and 5 µl of propidium iodide (PI), following the manufacturer's instructions of the Annexin V-FITC apoptosis detection kit (KeyGen, Shanghai, China). After incubation for 15 min at room temperature in the dark, all the samples were analyzed within 1 h with a BD flow cytometry system with FACSDiva software (BD Biosciences, Franklin Lakes, NJ, USA). The data were analyzed using FlowJo v5.7.2 software (BD Biosciences).

Wound healing assay. The cells were transfected and cultured to near (>80%) confluency in 24-well dishes. Then, an artificial homogeneous wound was created onto the monolayer with a sterile pipette tip, and cultured in RPMI-1640 medium containing 10% FBS for 24 h. To visualize migrating cells, wound closure was measured by photographing at five selected random fields at the time of wounding (time 0 h) and at 24 h after wounding under a light microscope (Olympus, Tokyo, Japan).

Transwell invasion assay. Cell invasion was performed using Transwell chamber assay (8.0-µm pore size; Corning Inc., Corning, NY, USA). Briefly, the 2x10⁵ transfected cells were seeded onto the upper chamber coated with Matrigel (BD Biosciences, Bedford, MA, USA) in serum-free medium, while the lower chamber was added with 20% FBS (600 µl). After the cells were incubated for 48 h at 37°C with 5% CO₂, the upper chamber was removed and the cells that had migrated to the lower chamber through the membrane were fixed in 90% alcohol and stained with 0.1% crystal violet for 5 min, then, photographed under a light microscope (magnification, x200; Olympus). The number of invaded cells was counted at five randomly selected fields.

Luciferase reporter assay. Regions of the 3’ untranslated region (3’UTR) for STAT3 containing the miR-124 binding sites was amplified from human genomic DNA, and inserted into the pGL3-control vector (Ambion, Austin, TX, USA) at the Nhel and Xhoi restriction sites, named as: Wt-STAT3-3’UTR. Mutant STAT3 containing four-point mutations in the putative miR-124 seed recognition motif were generated using overlapping PCR, and then inserted into the pGL3-control vector (Ambion) at the Nhel and Xhoi restriction sites, named as: Mut-STAT3-3’UTR. For the luciferase assay, RB cells were seeded onto 12-well plates at ~70% confluency and co-transfected with 50 ng of plasmid DNA and 50 nM miR-124 or miR-NC using Lipofectamine 2000 according to the manufacturer's protocol. Measurements of firefly and Renilla luciferase activity were performed 48 h after transfection using the Dual-Luciferase Reporter Assay System (Promega, Madison, WI, USA). Renilla luciferase was used for normalization.
Western blot analysis. Total proteins of the cell lines and tissues were extracted using a RIPA buffer with 0.5% sodium dodecyl sulfate (SDS) in the presence of a proteinase inhibitor cocktail (Complete Mini; Roche Diagnostics, Basel, Switzerland). The concentration of protein was quantified using a Bradford assay protein assay kit (Beyotime Biotech, Shanghai, China). Equal amounts of protein (20 µg) were separated using 10% SDS-polyacrylamide gels (SDS-PAGE) and transferred to polyvinylidene fluoride (PVDF) membranes (Millipore, Bedford, MA, USA) for 1 h at 100 V at 4˚C. The membranes were blocked with 5% non-fat milk/Tris-buffered saline with Tween-20 (TBST) for 1 h at room temperature, and then, incubated at room temperature with primary antibodies against STAT3 (1:1,000) and GAPDH (1:3,000) (both from Santa Cruz Biotechnology, Inc., Santa Cruz, CA, USA). GAPDH was used as control. Membranes were incubated with corresponding horseradish peroxidase (HRP)-conjugated secondary antibody (1:0000; Santa Cruz Biotechnology, Inc.) for 2 h at room temperature. Proteins were detected using an enhanced chemiluminescence luminol-based reagent, and visualized on X-ray film under an ECL detection system (both from Thermo Fisher Scientific, Inc., Waltham, MA, USA).

Statistical analysis. Data from at least three independent experiments are expressed as mean ± standard deviation (SD). Statistical analysis was performed using IBM SPSS 19.0 statistical software (version 19.0; SPSS, Inc., Chicago, IL, USA). Statistical analysis was performed using Student’s t-test or one-way ANOVA. Statistical significance was considered to indicate a statistically significant result at a P-value of <0.05.

Results

miR-124 expression is downregulated in primary RB tissues and cell lines. Quantitative real-time polymerase chain reaction (qRT-PCR) was performed to examine the expression levels of miR-124 in 20 normal retina and 40 RB tumor samples, and normalized against endogenous U6 controls. As shown in Fig. 1A, the expression level of miR-124 in RB tissues was significantly downregulated when compared to the normal retina tissues. Next, we evaluated the expression of miR-204 in two human RB cell lines (Y79 and SO-RB50). In comparison to normal retina tissues, miR-204 was downregulated in two cell lines (Fig. 1B). These data suggested that miR-124 may play crucial roles in the RB process.

miR-124 inhibits cell proliferation and induces cell apoptosis in RB cells. To assess the role of miR-124 in the growth of RB, miR-124 mimic and miR-NC were transiently transfected into human RB cell lines (Y79 and SO-RB50). qRT-PCR was used to confirm miR-124 overexpression in two RB cells (Fig. 2A). The MTT assay was performed to investigate the effect of miR-124 in RB cell proliferation. The results showed that restoration of miR-124 significantly inhibited the proliferation of Y79 and SO-RB50 cells (Fig. 2B). In addition, cell apoptosis assay was performed in RB cells transfected with miR-124 or miR-NC to investigate the effect of miR-124 on apoptosis in RB cells. Our results showed that restoration of miR-124 significantly increased the apoptosis rate in Y79 cells and SO-RB50 cells (Fig. 2C).

miR-124 inhibits cell proliferation in RB cells. To examine the effect of miR-124 on cell metastasis, cell migration and invasion were determined in RB cells transfected with miR-124 mimic or miR-NC by wound healing and Transwell invasion assays, respectively. It was found that restoration of miR-124 significantly inhibited migration (Fig. 3A) and invasion (Fig. 3B) capacities in Y79 and SO-RB50 cells.

miR-124 is a direct target of miR-124 in RB cells. It has been confirmed that STAT3 is the direct target of miR-124 in many cancer cells (18-21). Considering the tissue-specific and developmental stage-specific manner of miRNAs, we wondered whether STAT3 also is a direct target of miR-124 expression in RB cell lines, thus, luciferase activity was assessed in RB cells co-transfected with miR-124 or miR-NC and Wt-STAT3-3’UTR or Mut-STAT3-3’UTR. As shown in Fig. 4A and B, miR-124 significantly inhibited the luciferase activity of the Wt-STAT3-3’UTR, but not that of the Mut-STAT3-3’UTR in Y79 and SO-RB50 cells. To directly assess the effect of miR-124 on
expression, we transfected miR-124 or miR-NC into RB cells, and found that overexpression of miR-124 reduced the STAT3 mRNA level (Fig. 4C) and protein expression (Fig. 4D) in Y79 and SO-RB50 cells. These results demonstrated that STAT3 is a direct target of miR-124 in RB cells.

STAT3 expression is inversely correlated with miR-124 expression in RB tissues. The above results proved that STAT3 is the direct target of miR-124 in RB cells, we investigated therefore STAT3 mRNA expression in RB tissues and normal retina samples by qRT-PCR. Our results showed that STAT3 expression on mRNA level was upregulated in RB tissues compared to normal retina samples (Fig. 5A). In addition, a statistically significant inverse correlation was revealed by Spearman’s correlation analysis between miR-124 and STAT3 mRNA levels in RB tissues (r=-0.638; P<0.001; Fig. 5B).

STAT3 reverses the inhibitory effect of miR-124 on cell proliferation, migration and invasion in RB cells. To investigate whether miR-124 mediates its inhibition effects in RB cells through STAT3, we increased the miR-124 level in human
RB cells using a miR-124 mimic and rescued the expression of STAT3 (using the STAT3 overexpression vector without its 3'UTR) through the transfection of the miR-124 mimic in RB cells. The result of western blot analysis showed that the miR-124 mimic obviously inhibited STAT3 protein expression compared with the miR-NC in Y79 cells and SO-RB50 cells, while the overexpression of STAT3 abolished the inhibition caused by the miR-124 mimic (Fig. 6A). Notably, overexpression of STAT3 rescues the suppressive effects on proliferation (Fig. 6B), migration (Fig. 6C) and invasion (Fig. 6D) in RB cells caused by miR-124 expression in RB cells. These results suggested that miR-124 exerted its suppressive effects in RB cells, at least in part, by regulating STAT3.

**Discussion**

A large number of microRNAs (miRNAs) has been identified to be involved in occurrence and development of retinoblastoma (RB) through regulating and inhibiting the expression of their target gene, and functioned as oncogene or tumor suppressor by regulating proliferation, cell cycle,
apoptosis, invasion and migration of RB (10,11). Wu et al reported that miR-204 was frequently downregulated in RB tissues and cell lines, and that enforced expression of miR-204 inhibited the RB cell proliferation and invasion by targeting cyclin D2 and MMP-9 (22). Sun et al demonstrated that miR-145 suppressed RB cell proliferation, migration and invasion by repressing a disintegrin and metalloproteinases 9 (ADAM9) (23). Wang et al found that miR-183 suppressed proliferation, migration and invasion of RB cells by downregulation of low-density lipoprotein receptor-related protein 6 (LRP6) (24). In the present study, our results showed that miR-124 expression is downregulated in human RB tissues and RB cell lines compared with normal retinal tissue. Our results also demonstrated that restoration of miR-124 inhibited RB cell proliferation, migration and invasion, and induced cell apoptosis by targeting STAT3. These results support the conclusion that miR-124 plays a crucial role in RB development.

miR-124, located in 8q12.3, is frequently found to be downregulated in multiple human malignancies, such as prostate cancer (12), glioma (13), lung adenocarcinoma (14), breast (15), gastric (16), colorectal (17) and bladder cancer (25), hepatocellular carcinoma (20) and ovarian cancer (26). miR-124 exerted a tumor suppressive role in various cancer cells by negative regulation of cell proliferation, apoptosis, migration and invasion through repressing multiple target genes, such as PACE4 (12), SOX9 (14), CD4 (25), STAT3 (20), SphK1 (26) and PIK3CA (27). However, at present, there is no published study regarding the biological functions of miR-124 in RB. In the present study, we found that miR-124 expression was downregulated in RB tissues and cell lines, and that miR-124 suppressed cell proliferation, migration and invasion of RB cells. These results suggested that miR-124 functioned as a tumor suppressor in RB cells.

STAT3, a member of the signal transducer and activator of transcription (STAT family), has been shown to play crucial roles in cell cycle progression, apoptosis, cellular transformation and proliferation by regulating the expression of multiple target genes such as cyclin D1, c-Myc, survivin, Bcl-xL, Bcl-2, Mcl-1, VEGF and MMP-2 and MMP-9 (28-34). It was found that downregulation of STAT3 using RNA interference targeting STAT3, and small molecule inhibitors suppressed tumor cell proliferation and invasion, induced apoptosis in vitro, and delayed tumor growth in animal models of various types of cancer (32-34). Recently, a study showed that STAT3 expression was increased in RB tissues from human patients compared to normal retinal tissues, and that inhibition of STAT3 in RB cells with targeted siRNAs resulted in impaired proliferation and downregulation of target genes in vitro, and suppressed formation of orthotopic tumors in vivo (35), suggesting STAT3 is an oncogene in RB. Although STAT3 has been reported to be a target of miR-124 in several types
of cancers, such as esophageal cancer (18), glioblastoma (19), hepatocellular carcinoma (20) and non-small cell lung cancer (21), however, the interaction between miR-124 and STAT3 has not been experimentally validated in RB. In the present study, we confirmed that STAT3 was a direct target of miR-124, and that miR-124 overexpression significantly reduced the levels of both STAT3 protein and mRNA in RB cells. We also demonstrated that miR-124 expression levels negatively correlated with STAT3 mRNA levels in human RB tissues specimens. STAT3 overexpression rescued the suppressive effect of miR-124-mediated RB cell proliferation, migration and invasion. These results displayed evidence of miR-124-mediated suppression role in RB cells, at least in part by targeting STAT3.

In summary, to the best of our knowledge, the present study is the first to provide evidence that the expression of miR-124 is downregulated in RB tissues and cell lines; and that restoration of miR-124 inhibited proliferation, migration and invasion, and induced cell apoptosis in RB cells, at least in part, by targeting STAT3. These findings suggested that miR-124 functioned as a tumor-suppressor, and may become a novel molecular therapeutic target for the treatment of RB.

References