Abstract. miR-212 as a tumor suppressor has been reported to be downregulated in multiple cancer cells lines and tumor tissues. However, its role in thyroid cancer has nor been investigated. Therefore, the present study aimed to investigate the role of miR-212 in human thyroid cancer and the underlying mechanisms. In the present study, we demonstrated that miR-212 expression was significantly decreased in thyroid cancer specimens and cell lines compared with adjacent normal tissues and normal thyroid cell lines. In addition, we demonstrated that miR-212 downregulation in thyroid cancer tissues was negatively associated with lymph node metastasis and advanced clinical stage. Functionally, ectopic expression of miR-212 by transfection with miR-212 mimic significantly inhibited proliferation, colony formation, migration and invasion in TPC-1 cells. In addition, Sirtuin 1 (SIRT1) was identified as a direct target of miR-212 and its expression was inversely correlated with miR-212 expression in thyroid cancer tissues. Overexpression of SIRT1 could effectively rescue miR-212 mimic-induced suppression of cell proliferation, migration and invasion in TPC-1 cells. In vivo, miR-212 overexpression significantly inhibited tumor growth in a nude mice model. In light of these findings, miR-212 may function as a tumor suppressor in thyroid cancer by targeting SIRT1.

Introduction

Thyroid cancer is the most common endocrine malignancy worldwide with a rapidly increasing incidence and prevalence over the past 20 years (1). Despite the fact that thyroid cancer patients in early stages have a favorable prognosis with high overall 5-year survival rate, patients with advanced thyroid cancer have a 5-year survival rate of only ~59% (2,3). Therefore, elucidating the molecular mechanisms underlying thyroid cancer is required in order to develop novel therapeutic strategies for this disease.

MicroRNAs (miRNAs) are a class of endogenous small RNAs containing ~22 nucleotides that disturb the protein translation process by targeting the sequences on their 3' untranslated region (3'UTR) (4-6). By negatively regulating the protein expression levels of their target genes, miRNAs play central roles in tumor initiation, development and progression (7,8). Since protein expression is dysregulated in various human cancers, including thyroid cancers and some miRNAs act as oncogenes or tumor suppressors in thyroid cancer, miRNAs may serve as potential molecular targets or candidates for the treatment of thyroid cancer (9,10).

miR-212, a tumor associated miRNA, has been reported to play a suppressive role in many types of cancer, including gastric (11), hepatocellular carcinoma (12), non-small cell lung (14), ovarian (15) and cervical cancer (16). However, to the best of our knowledge, the role and molecular mechanism of miR-212 in thyroid cancer has not been determined. The present study aimed to investigate the expression levels of miR-212 in thyroid cancer tissues and cell lines, as well as its clinical significance. Furthermore, the present study examined the role of miR-212 in the regulation of thyroid cancer cell growth and invasion, as well as the underlying regulatory mechanism of its action by in vitro and in vivo experiments.

Materials and methods

Patients and tissue samples. Primary thyroid cancer samples and their matched non-cancerous tissues (normal) were obtained from 42 patients who underwent surgery of thyroid cancer at China-Japan Union Hospital of Jilin University (Changchun, China). Following surgery all samples were snap-frozen in liquid nitrogen and stored at -80°C until RNA extraction. Patients who underwent chemotherapy, radiotherapy or other treatment before the operation were excluded. The study was approved by the Research Ethics Committee of...
Cell lines and transfection. Three human thyroid cancer cell lines (TPC-1, BCPAP and SW1736) and human thyroid follicular epithelial cells (Nthy-ori3-1) were obtained from the Type Culture Collection of the Chinese Academy of Sciences (Shanghai, China) and were grown in Dulbecco's modified Eagle's medium (DMEM; Gibco, Grand Island, NY, USA) supplemented with 10% fetal bovine serum (FBS; Gibco) at 37°C in a humidified chamber supplemented with 5% CO₂.

miR-212 mimic or negative control mimic (miR-NC) were designed and synthesized by Shanghai GenePharma Co., Ltd. (Shanghai, China). Overexpression SIRT1 plasmid (pcDNA3.1-SIRT1) was granted by Dr Peng Li (Jilin University, Changchun, China). These molecules were transiently transfected into TPC-1 cell using Lipofectamine™ 2000 (Invitrogen, Waltham, MA, USA) according to the manufacturer's protocol. After 24-72 h transfection, the cells were harvested for further analysis.

Proliferation assay. The proliferation of thyroid cancer cell was assessed by the Cell Counting Kit-8 (CCK-8) assay kit (Dojindo Laboratories, Kumamoto, Japan). Briefly, ~1×10⁴ transfected cells were seeded in each well of 96-well plates and cultured for 24-72 h. At indicated time-points (24, 48 and 72 h), 10 µl of CCK-8 was added to each well. After incubation at 37°C for 4 h, the absorbance was detected at 450 nm.

Colony formation assay. Transfected cells were digested with trypsin and suspended into a single cell status. A total of 1,000 cells from each group were seeded in 6-well plates and cultured in complete DMEM medium with 10% FBS for 14 days. The colonies were fixed with 4% paraformaldehyde for 20 min and stained with 0.1% crystal violet for 15 min. The total number of colonies was counted under a light microscope (Olympus, Tokyo, Japan).

Luciferase assay. The human SIRT1 3'UTR oligonucleotides containing the wild-type (WT) or mutant-type (MT) miR-212 binding site were amplified by PCR and inserted downstream of the luciferase gene in the pGL3-luciferase reporter plasmid (Ambion, Austin, TX, USA). TPC-1 cells seeded in 96-well plates in triplicate were cotransfected with WT/MT-SIRT1-3'UTR reported plasmid and miR-212 mimic or miR-NC using Lipofectamine 2000 according to the manufacturer's protocol. Forty-eight hours after transfection, the cells were harvested and luciferase activity was assessed using a Dual-Luciferase reporter assay kit (Promega).

Western blotting. The protein was extracted by lysing cells in ice-cold radioimmunoprecipitation assay buffer (Nanjing KeyGen Biotech, Co., Ltd., Nanjing, China). The protein concentration was quantified with the bicinchoninic acid (BCA) kit (Pierce, Rockford, IL, USA). Equivalent amounts of protein samples (20 µg each lane) were separated on 10% sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE) and transferred onto a polyvinylidene fluoride membrane (Millipore, Billerica, MA, USA), which was then blocked for 1 h with 5% non-fat milk in PBST. After incubation overnight with antibodies against SIRT1 and GAPDH, the membranes were incubated with horseradish peroxidase-conjugated (HRP) goat-anti-mouse secondary antibody at room temperature for 2 h. The protein bands were observed using chemiluminescence (Thermo Fisher Scientific, Waltham, MA, USA).

Wound healing assay. Wound healing assays were performed to assess the motility of indicated cells. Briefly, the transfected cells were seeded in 6-well plates and allowed to reach to 90-95% confluency. The cells were scratched with a sterile plastic micropipette tip in the cell monolayer. After wounding, the cells were cultured in complete DMEM medium with 10% FBS for 24 h. Wound closure was observed at 0 and 24 h, and photographed under a light microscope (Olympus).

Transwell invasion assay. The invasive ability of indicated cells was analyzed by Transwell (Corning Costar Corp., Cambridge, MA, USA) assay. In brief, 2×10⁴ transfected cells suspended in serum-free medium were added to each upper chamber precoated with Matrigel matrix and 500 µl of DMEM medium containing 10% FBS were added to the lower chamber as a chemoattractant. After 48-h incubation, the remaining cells in the upper chambers were removed with a cotton swab and the invasive cells on the lower membrane surface were fixed with 4% paraformaldehyde, stained with 0.1% crystal violet. Invasive cells were photographed and quantified by counting them in five random fields using a light microscope (Olympus).

In vivo tumorigenesis assay. All animal studies were approved by the Institutional Animal Care and Use Committee of Jilin University (Changchun, China). Twenty female BALB/C nude mice (18-20 g, 6-7 weeks old) were obtained from the Experimental Animal Center of Jilin University (Changchun, China) and maintained under specific pathogen-free conditions at Jilin University.
For the in vivo tumorigenesis assay, ~2x10^6 TPC-1 cells stably expressing miR-212 or miR-NC were suspended in 100 µl of phosphate buffered saline (PBS), and then injected subcutaneously in the left posterior flank of the BALB/c-nude mice (10 mice in each group). After ten days, the tumor growth was assessed and recorded by measuring tumor length (L) and width (W) every five days until the nude mice were sacrificed. Tumor volume was calculated according to the formula: V = 1/2 x L x W^2. The mice were sacrificed and photographed at 30 days post-implantation. Xenograft tumors were excised, photographed, weighed and stored at -80˚C for further analysis.

Statistical analysis. All statistical analyses were performed using the SPSS 19.0 software (SPSS, Inc., Chicago, IL, USA). Experimental data are presented as the mean ± standard error (SE) at least from three independent experiments. The two-tailed Student's t-test was adopted for comparison between the two groups and one-way ANOVA was used for comparisons of more than two groups. A P-value of 0.05 was considered to indicate a statistically significant difference.

Results

miR-212 is downregulated in thyroid cancer tissues and cell lines. To determine the expression of miR-212 in thyroid cancer, we first detected miR-212 expression level by qRT-PCR in human thyroid cancer tissues and adjacent normal thyroid tissues. As depicted in Fig. 1A, miR-212 expression was significantly downregulated in human thyroid cancer tissues compared with adjacent normal tissues. In addition, we found that the expression levels of miR-212 in advanced clinical stage (III-IV) were significantly downregulated compared with those in low clinical stage (TNM stage I and II) (Fig. 1B). Consistent with the above mentioned results, miR-212 levels in tissues with lymph node metastases were markedly decreased compared to the tissues without lymph node metastases (Fig. 1C). In addition, we investigated the expression of miR-212 in three thyroid cancer cell lines (TPC-1, BCPAP and SW1736), using the human thyroid epithelial cell line Nthy-ori3-1 as a control. We found that miR-212 was downregulated in thyroid cancer cell lines compared with the human thyroid epithelial cell line (Fig. 1D). In particular, TPC-1 cell line exhibited the lowest levels of miR-212 expression and was used for subsequent studies (Fig. 1D). These results indicated that low miR-212 may be associated with thyroid cancer progression.

miR-212 inhibits thyroid cancer cell proliferation, migration and invasion. To explore the possible biological functions of miR-212 in thyroid cancer cells, we transfected TPC-1 cells with miR-212 mimic or negative controls (miR-NC) to enhance miR-212 expression. As shown in Fig. 2A, cells transfected with miR-212 mimic significantly increased miR-212 expression levels compared to cells transfected with miR-NC. CCK-8 assays demonstrated that miR-212 overexpression significantly inhibited thyroid cancer cell proliferation (Fig. 2B). In addition, examined the colony formation capacity of TPC-1 cells and observed that miR-212 overexpression significantly inhibited thyroid cancer cell colony formation (Fig. 2C). To investigate the effect of miR-212 on cellular motility, the migration and invasion ability of TPC-1 cells after modification of miR-212 expression were determined by wound healing and Transwell invasion assays, respectively. It was observed that miR-212 overexpression significantly inhibited the migration and invasion of TPC-1 cells (Fig. 2D and E). Collectively, these results indicated that miR-212 may impede thyroid cancer cell proliferation, migration and invasion in vitro.

SIRT1 is a direct target of miR-212 in thyroid cancer cells. To fully understand the mechanism of miR-212 inhibition of human thyroid cancer progression, three bioinformatic databases (TargetScan, miRanda and PicTar) were used to predict the targets of miR-212. We identified the 3'-UTR of SIRT1 that were able to bind to the ‘seed region’ of miR-212 (Fig. 3A). To
Figure 2. miR-212 inhibits thyroid cancer cell proliferation, migration and invasion. (A) Relative miR-212 expression was determined in TPC-1 cells after transfected with miR-212 mimic or miR-NC. (B-E) Cell proliferation, colony formation, migration and invasion were determined in TPC-1 cells after transfection with miR-212 mimic or miR-NC. *P<0.05, **P<0.01.

Figure 3. SIRT1 is a direct target of miR-212 in thyroid cancer. (A) The luciferase reporter of miR-212 binding site on the wild-type and mutant-type SIRT1 3’-UTR. WT, wide-type; MT, mutant-type. (B) Relative luciferase assay activity was analyzed in TPC-1 cells cotransfected with WT/MT SIRT1-3’-UTR reporter plasmid and miR-212 mimic or miR-NC. (C and D) The SIRT1 expression on mRNA and protein levels was measured in TPC-1 cells transfected with miR-212 mimic or miR-NC. GAPDH was used as an internal control. *P<0.05, **P<0.01.
determine whether SIRT1 is a target of miR-212, the luciferase activity assay was performed. As expected, miR-212 bound to SIRT1 3'-UTR, resulting in markedly reduced luciferase activities (Fig. 3B). In addition, we also found that miR-212 overexpression obviously decreased SIRT1 expression on mRNA expression and protein levels (Fig. 3C and D). These results indicated that miR-212 directly targets SIRT1 by binding its seed region of the 3'-UTR region in human thyroid cancer cells.

Inverse correlation between SIRT1 and miR-212 expression in thyroid cancer. Subsequently, we examined the SIRT1 mRNA in 42 pairs of thyroid cancer tissue specimens and adjacent normal tissues by qRT-PCR. The SIRT1 expression was higher in thyroid cancer specimens than that of adjacent normal thyroid tissues (Fig. 4A). The inverse correlation between miR-212 and SIRT1 mRNA levels was further confirmed by Pearson correlation analysis in 42 thyroid cancer tissues (r= -0.486, P=0.001; Fig. 4B). Furthermore, the SIRT1

Figure 4. SIRT1 is upregulated and inversely correlated with miR-212 expression in thyroid tissues. (A) The SIRT1 mRNA expression in 42 pairs of thyroid cancer tissues and adjacent normal tissues was determined by qRT-PCR. GAPDH was used as an internal control. (B) The correlation of the expression levels of SIRT1 and miR-212 was analyzed by Pearson's correlation assay in thyroid cancer tissues (n=42). (C and D) The SIRT1 expression on mRNA and protein levels was determined in three thyroid cancer cell lines (TPC-1, BCPAP and SW1736) and the human thyroid epithelial cell line Nthy-ori3-1. **P<0.01.

Figure 5. miR-212 exerts its suppressive function by targeting SIRT1 in thyroid cancer cells. (A and B) SIRT1 expression on mRNA and protein levels was determined in TPC-1 cells transfected with miR-212 mimic or miR-NC with or without SIRT1 overexpression plasmid by qPCR and western blot analysis, respectively. GAPDH was used as an internal control. (C-F) Cell proliferation, colony formation, migration and invasion were determined in TPC-1 cells transfected with miR-212 mimic or miR-NC with or without SIRT1 overexpression plasmid. *P<0.05, **P<0.01.
miR-212 expression on mRNA and protein levels was increased in thyroid cancer cell lines compared to the normal thyroid cells (Fig. 4C and D).

**miR-212 exerts its suppressive function by targeting SIRT1 in thyroid cancer cells.** To examine whether miR-212 exerted its suppressive function through its target gene SIRT1, we rescued the expression of SIRT1 in miR-212 mimic-transfected cells. qRT-PCR and western blot assays revealed that transfection of SIRT1 overexpression plasmid in miR-212 mimic-transfected cells restored the SIRT1 expression in TPC-1 cells (Fig. 5A and B). Furthermore, restoration of SIRT1 expression partially reversed the inhibition effect on cell proliferation, colony formation, migration and invasion in TPC-1 cells mediated by miR-212 (Fig. 5C-F). These results indicated that miR-212 impaired cell growth, migration and invasion of TPC-1 by targeting SIRT1.

**miR-212 suppresses tumor growth in vivo.** To determine the effects of miR-212 on tumorigenicity in vivo, TPC-1 cells transfected with miR-212 mimic or miR-NC were injected into the flanks of nude mice to form ectopic tumors. We found that tumor growth was slower in the TPC-1/miR-212 group than that in the TPC-1/miR-NC group (Fig. 6B and C). We also analyzed the expression of miR-212 and SIRT1 in xenograft tumors. In the TPC-1/miR-212 group, miR-212 expression was upregulated (Fig. 6D), whereas SIRT1 expression was downregulated both on mRNA level (Fig. 6E) and protein level (Fig. 6F). These data indicated that miR-212 suppresses tumor growth in vivo by suppressing SIRT1.

**Discussion**

miRNAs have been found to play crucial roles in the carcinogenesis in various types of cancers (18). In line with this notion, miRNAs participation in thyroid cancer progression has been widely reported (9,10). In the present study, we found that miR-212 was downregulated in both thyroid cancer tissues and thyroid cancer cell lines and that decreased miR-212 was associated with lymph node metastasis and clinical stage. We also found that miR-212 overexpression by transfection with miR-212 mimic significantly inhibited thyroid cancer cell proliferation, colony formation, migration and invasion in vitro. In vivo, miR-212 overexpression inhibited tumor growth in nude mice model. These results may provide evidence for using miR-212 as a novel target for treating thyroid cancer.  

miR-212 has been revealed to be downregulated and function as tumor suppressor in the majority of types of cancers by regulating different oncogene (11-15). For example,
Fu et al (19) revealed that miR-212 may act as tumor suppressor in prostate cancer progression through disrupting epithelial to mesenchymal transition (EMT) process by directly targeting SOX4. Jiping et al (11) reported that miR-212 functions as a tumor suppressor involved in tumor metastasis and invasion of gastric cancer through suppressing paxillin (PXN) expression. Zhao et al (20) demonstrated that miR-212 delayed cell arrest in the G1/S phase transition and suppressed cell proliferation, as well as EMT migration and invasion in cervical cancer cell by targeting SMAD2. In contrast, recently a study revealed that miR-212 facilitated pancreatic cancer cell growth and invasion by targeting the hedgehog signaling pathway receptor patched-1 (21). Thus, the biological role of miR-212 in carcinogenesis seems to be complicated and highly tissue-specific.

In the present study, we investigated miR-212 expression in thyroid cancer tissues and cell lines by qRT-PCR and found that miR-212 was downregulated in thyroid cancer tissues and cell lines. Furthermore, miR-212 expression was significantly downregulated in patients with advanced clinical stage and lymph node metastasis. Thus, we hypothesized that miR-212 overexpression may suppress the malignant phenotypes of thyroid cancer cells. As expected, our further results revealed that ectopic miR-212 expression suppressed thyroid cancer cell growth, migration and invasion in vitro, as well as suppressed tumor growth in vivo. Altogether, both clinical and experimental data indicated a tumor suppressive role of miR-212 in thyroid cancer.

It is well known that miRNAs can act as tumor suppressors by targeting specific oncogenes (22). Thus, three bioinformatic databases (TargetScan, miRanda and PicTar) were used to predict targets of miR-212. Sirtuin 1 (SIRT1), an known oncogene, was selected as a target gene of miR-212. SIRT1 is a member of the sirtuin (SIRT) family that exerts multiple cellular functions and is conserved from bacteria to eukaryotes (23). SIRT1 expression has been reported to be higher in numerous human cancer cell lines and tissues including thyroid cancer (24). SIRT1 has been implicated in the cell cycle, as well as apoptosis and cancer metastasis by regulating its substrates such as Myc, p53, nuclear factor-κB, Ku70 and forkhead transcription factor (25,26). Recently, a study revealed that inhibition of SIRT1 expression impaired proliferation and induced cell apoptosis and cell cycle arrest in thyroid cancer cell lines (27), indicating SIRT1 as an oncogene in thyroid cancer. SIRT1 has been reported to be a target of miR-212 in prostate cancer (28), however, the interaction between miR-212 and SIRT1 has not been experimentally validated in thyroid cancer. In the present study, using luciferase reporter assays, qRT-PCR and western blot assays, we verified the SIRT1 gene as a direct target of miR-212 in thyroid cancer. In addition, SIRT1 expression was upregulated in thyroid cancer tissues and was negatively correlated with the expression level of miR-212. SIRT1 overexpression reversed the inhibition effect on cell proliferation, migration and invasion in thyroid cancer cells induced by miR-212 overexpression. In vivo, miR-212 also displayed an inhibitory role in thyroid cancer growth by suppressing SIRT1. These findings indicated that miR-212 impaired thyroid cancer development via repressing SIRT1.

In conclusion, this study first demonstrated that miR-212 is downregulated in thyroid cancer tissues and cell lines and functions as a tumor suppressor in thyroid cancer cell growth by downregulating SIRT1. Thus, miR-212/SIRT1 may provide a promising therapeutic strategy for the treatment of thyroid cancer.

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References


