MicroRNA-365a-3p promotes tumor growth and metastasis in laryngeal squamous cell carcinoma

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Abstract. MicroRNAs (miRNAs) are increasingly recognized as oncogenes or tumor suppressors in laryngeal squamous cell carcinoma (LSCC). In this study, we analyzed the roles of miR-365a-3p, miR-143-5p, and miR-494-3p in LSCC using Annexin V/propidium iodide double staining and flow cytometry, along with a Transwell migration and invasion assay. The results showed that miR-365a-3p inhibitor significantly facilitated cell apoptosis and suppressed cell cycle progression, migration, and invasion in Hep-2 cells. However, miR-143-5p and miR-494-3p had no such influences. We then investigated the role of miR-365a-3p in LSCC in vivo and found that miR-365a-3p inhibitor suppressed LSCC xenograft tumor growth and metastasis in xenograft mouse models. Moreover, miR-365a-3p inhibitor significantly decreased the expression of p-AKT (Ser473), which indicated that miR-365a-3p can mediate PI3K/AKT signaling pathway transduction via p-AKT (Ser473) in LSCC. The data suggest that miR-365a-3p may act as an oncomiR and may promote growth and metastasis in LSCC via the PI3K/AKT signaling pathway, and thus miR-365a-3p may be a potential therapeutic target for treatment of LSCC.

Introduction

Laryngeal squamous cell carcinoma (LSCC) is the most common type of laryngeal carcinoma (LC) and accounts for more than 90% of LC cases. In 2015, the estimated number of new cases of LC was 13,560 in the United States, and about 3,640 Americans were likely to die from LC (1). During the past 2 decades, clinicians have seen a worrying decline in the survival rate of LC patients, partly due to lymph node metastases (LNM). Recent studies showed that the 5-year overall survival was 39-44% in patients with advanced LC, and the 2-year local relapse rate was 27.5% in stage III LC (2,3). Although a variety of molecules associated with the oncogenesis and the treatment of LSCC have been identified, the potential therapeutic targets and the precise molecular mechanisms of LSCC proliferation and metastasis remain to be elucidated.

In the past few years, miRNAs have been identified in most carcinomas and have become potential biomarkers for cancer diagnosis and treatment (4-6). miRNAs are small, single-stranded non-coding RNA molecules that regulate gene expression by binding to miRNA at the 3'-untranslated region (7). As of May 2014, 2,578 miRNAs have been identified in the human genome (4). Deregulated miRNAs have been shown to play pivotal roles in the multistep process of carcinogenesis via regulated oncogenes or tumor suppressor genes. For example, miR-10b and miR-320b enhance cell proliferation and invasion in pancreatic cancer and colorectal cancer, respectively (8,9), and miR-218 and miR-195 inhibit cell proliferation, migration, and invasion in gliomas and non-small cell lung cancer (10,11).

miRNAs also play important roles in LSCC (12,13). miR-27a was significantly upregulated in LSCC, promoted cell viability and colony formation, and repressed apoptosis by targeting PLK2 in Hep-2 cells (13). Our previous study (14) reported the identification via miRNA microarray and verification by real-time PCR (RT-PCR) of 10 miRNAs that were expressed differentially in LSCC patients with vs. those without LNM, including 9 upregulated miRNAs (miR-365a-3p, miR-143-5p, miR-634, miR-223-3p, miR-409-5p, miR-1224-3p, miR-192-5p, miR-30d-5p, and miR-1249-3p) and 1 downregulated miRNA (miR-494-3p).

Key words: miR-365a-3p, miR-143-5p, miR-494-3p, laryngeal squamous cell carcinoma, AKT
Based in part on findings from our previous study, we analyzed the effects of miR-365a-3p, miR-143-5p, and miR-494-3p in LSCC in vitro. However, only miR-365a-3p can affect cell apoptosis, mitosis, migration, and invasion. We also found that miR-365a-3p promoted LSCC xenograft tumor growth and metastases in liver and hepatic lymph nodes in mouse models. Moreover, miR-365a-3p was demonstrated to mediate the PI3K/AKT signaling pathway via p-AKT (Ser473) in LSCC.

Materials and methods

Cell culture and transient infection. Human Hep-2 cell lines (American Type Culture Collection, ATCC; Manassas, VA, USA) were cultured in RPMI-1640 medium (Gibco-Life Technologies, Carlsbad, CA, USA) with 10% fetal bovine serum at 37°C and 5% CO₂ in a humidified-atmosphere incubator. For overexpression and inhibition of miRNAs (all from GE Healthcare Dharmacon, Lafayette, CO, USA), miR-365a-3p inhibitor (IH-300666-05-0002), miR-143-5p inhibitor (IH-301057-02-0002), miR-494-3p mimic (C-300761-05-0002), normal control (NC)-inhibitor (IN-001005-01-05), and NC-mimic (CN-001000-01-05) were transiently transfected in Hep-2 cells by Lipofectamine 2000 (Invitrogen-Life Technologies, Carlsbad, CA, USA) following the manufacturer's protocol.

RNA extraction and real-time PCR analysis. Total RNAs of the transfected Hep-2 cells were extracted with TRIzol (Invitrogen-Life Technologies) according to the manufacturer's protocol. Single-stranded cDNA of the miRNAs was synthesized by reverse transcription using the miScript Reverse Transcription kit (Qiagen, Valencia, CA, USA). The expression level of transfected miRNAs was assessed with RT-PCR (TaqMan MicroRNA Assay; Life Technologies). The PCR primers for miR-365a-3p (sense, GCCCCTAAAAATCCTT and antisense, GTGCAGGCTTCCAGGT); miR-143-5p (sense, AAAAAAGAGAAAAACACCC and antisense, GTGCAGGTCAGGT); miR-494-3p (sense, GAAACATACACGAGGAAACC and antisense, GTGCAGGCGTCCAGGT); and U6 (sense, TGCAGGTTGCTCCTTCGCAGC and antisense, CCAGTGAGGTCGAGAGGT) were obtained from Life Technologies. The Hep-2 cells were treated according to the procedures for cell apoptosis assay. NC-mimic and NC-inhibitor were used as controls. Subsequently, the transfected cells were resuspended with 400 µl PBS and fixed with prechilled 80% ethanol at 4°C for 24 to 48 h for cell cycle analysis. Then, the fixed cells were washed with PBS and resuspended in staining solution (Boehringer Mannheim, Mannheim, Germany) containing 50 µg/ml of RNase A and 65 µg/ml of PI solution and incubated at 37°C for 30 min. Cell cycle flow cytometric analysis was performed with a FACSCalibur flow cytometer.

Cell apoptosis assay. Hep-2 cells were seeded in 6-well plates, grown to about 80% confluence, and transiently transfected with the prepared miRNA inhibitors and mimics. NC-mimic and NC-inhibitor were used as controls. The transfected cells were trypsinized, washed twice in prechilled phosphate-buffered saline (PBS), and collected after transfection at 48 and 72 h. Subsequently, the cells were resuspended with 300 µl of 1X binding buffer. Then, 5 µl of Annexin V-FITC conjugate and 5 µl of propidium iodide (PI) solution (both from BD Biosciences, San Diego, CA, USA) were added, and cells were incubated at room temperature for 15 min in the dark. Flow cytometric analysis for cell apoptosis was performed with a FACSCalibur flow cytometer (BD Biosciences) according to the manufacturer's instructions.

Cell cycle assay. The Hep-2 cells were treated according to the procedures for cell apoptosis assay. NC-mimic and NC-inhibitor were used as controls. Subsequently, the transfected cells were resuspended with 400 µl PBS and fixed with prechilled 80% ethanol at 4°C for 24 to 48 h for cell cycle analysis. Then, the fixed cells were washed with PBS and resuspended in staining solution (Boehringer Mannheim, Mannheim, Germany) containing 50 µg/ml of RNase A and 65 µg/ml of PI solution and incubated at 37°C for 30 min. Cell cycle flow cytometric analysis was performed with a FACSCalibur flow cytometer.

Transwell migration and invasion assay. For the migration assay, the transiently transfected Hep-2 cells with the miRNA inhibitors and mimics described above were seeded in the upper Transwell chamber (Sigma-Aldrich, St. Louis, MO, USA) coated with 8-µm pore Transwells (Millipore, Billerica, MA, USA). After incubation for 24 h, the non-migrated cells in the upper chamber were removed. The invaded cells were fixed with methanal, stained with Giemsa staining solution, and then counted by an inverted light microscope. The Hep-2 cells invasion assays were performed using the Matrigel invasion chamber (Sigma-Aldrich) according to the manufacturer's protocol. The seeding, staining, and counting of Hep-2 cells were performed as the migration assay. In the above two assays, the medium of upper Transwell chamber contains serum-free culture, and of the lower chamber contains 10% fetal bovine serum served as the chemotactrant.

Lentivirus-mediated inhibition of miR-365a-3p. To establish the stable miR-365a-3p and miR-365a-3p inhibitor cell lines, Hep-2 cells were transfected with NC-pLenti6/TR inhibitor (Lv-NC-inhibitor) and pLenti6/TR miR-365a-3p inhibitor (Lv-miR-365a-3p inhibitor), respectively, by selection for 42 days in fresh complete medium supplemented with 3 µg/ml puromycin (Invitrogen-Life Technologies). Single colonies were selected and amplified, and the expression level of miR-365a-3p was detected by RT-PCR.

Analysis of tumorigenesis and metastatic potential of miR-365a-3p in vivo. For the in vivo tumorigenesis assay, 2x10⁶ stable transfected Hep-2 cells with Lv-NC-inhibitor were injected into the enterococelia of 10 nude mice (Laboratory Animal Center of the Capital Medical University, Beijing, China), aged 6 weeks, so were the 2x10⁶ stable transfected Hep-2 cells with Lv-miR-365a-3p inhibitor. The mice were sacrificed on the day 42 after injection, and the final tumors were isolated. Tumor size was measured with a digital caliper, and tumor volume was calculated with the formula: tumor volume = (length x width²) x 0.5. For the tumor-metastasis assay in vivo, all the organs of the sacrificed mice were examined at necropsy. Lungs, livers, and the corresponding lymph nodes were stained with hematoxylin and eosin and were examined by two pathologists.

All mice used in this experiment were bred and maintained in sterile cages and were handled according to the National Institutes of Health Animal Care and Use Committee Regulations. All experimental procedures were approved by the Animal Care and Use Committee of Capital Medical University (Beijing, China).
Protein extraction and western blot analysis. Cells were lysed in RIPA buffer with a protease inhibitor cocktail (both from Millipore). Protein was extracted and analyzed by standard western blot analyses. The primary antibodies, including total ERK, p-ERK1/2, total AKT, p-Akt (Ser473), and p-Akt (Thr308) were obtained from Life Technologies. GAPDH was used as an endogenous normalizer. The membrane was incubated with alkaline phosphatase secondary antibodies (Thermo Fisher Scientific, Waltham, MA, USA). Then the intensities of enhanced bands of immunoreactive protein was quantified (Image-Pro Plus 6.0; Media Cybernetics, Rockville, MD, USA).

Statistical analyses. All of the experiments were repeated in triplicate, and one representative experiment was selected for data analysis. The statistical analyses were performed with...
SPSS 17.0 (IBM, Chicago, IL, USA). The statistical significance was tested using two-tailed Student's t-test. A p-value of <0.05 was considered statistically significant.

Results

miR-365a-3p inhibitor facilitates apoptosis and suppresses mitosis in Hep-2 cells. Our previous study determined that miR-365a-3p and miR-143-5p were upregulated and miR-494-3p was downregulated in LSCC patients with LNM. In the present study, we successfully transfected Hep-2 cells with miR-365a-3p inhibitor (P<0.001, Fig. 1A), miR-143-5p inhibitor (P<0.001, Fig. 1B), and miR-494-3p mimics (P<0.001, Fig. 1C) for 48 and 72 h.

We first assessed the roles of these miRNAs in apoptosis in Hep-2 cells by flow cytometric analysis. At 48 and 72 h after transient transfection, analyses based on Annexin V-FITC/PI apoptosis revealed that with the decreased expression of miR-365a-3p, the number of Hep-2 cells that expressed apoptotic protein Annexin V was elevated (representative data are shown in Fig. 2A). The result indicates that miR-365a-3p inhibitor significantly facilitated the apoptosis of Hep-2 cells at 48 and 72 h (P<0.001, Fig. 2B). In contrast, miR-143-5p inhibitor and miR-494-3p mimics had no effect on Hep-2 cells (Fig. 3).

To characterize the influences of these miRNAs on the cell cycle, we used flow cytometry with PI staining to measure the DNA content in order to analyze the cell cycle distribution in Hep-2 cells. The percentages of cells in the S1 phase infected with the NC-inhibitor at 24 and 48 h after transfection were 36.2 and 37.7% and with the miR-365a-3p inhibitor were 28.6 and 30.6%, respectively (representative data are shown in Fig. 2C). These findings suggest that the miR-365a-3p inhibitor significantly suppressed cell cycle progression in Hep-2 cells, as shown by the histogram in Fig. 2D (P=0.002). However, Hep-2 cells infected with the miR-143-5p inhibitor and the miR-494-3p mimics showed no significant differences compared with the homologous controls (Fig. 4).

miRNA-365 promotes migration and invasion of Hep-2 cells. To determine the biological functions of these miRNA mimics and inhibitors in LSCC metastasis, we performed Transwell migration and invasion assays on Hep-2 cells transfected with miR-365a-3p inhibitor, miR-143-5p inhibitor, and miR-494-3p mimics at 24 h after transfection. As shown in Fig. 5A, transfection of the miR-365a-3p inhibitor led to
Figure 4. The mitosis assay of infected Hep-2 cells with miR-143-5p inhibitor and miR-494-3p mimic based on flow cytometry analyses. (A and B) The percentage of Hep-2 cells infected with miR-143-5p inhibitor and miR-494-3p mimic in S1 phase at 24 and 48 h later. (C and D) The numbers of infected homologous Hep-2 cells in S1 phase were counted and are shown in the histograms.

Figure 5. The migration and invasion of Hep-2 cells transfected with miR-365a-3p were measured by the Transwell assay. (A) The migration and (C) invasion of infected Hep-2 cells were measured by the Transwell assay without and with Matrigel, respectively. (B and D) The numbers of Hep-2 cells migrating to and invading the lower chamber were quantified and are shown in histograms B and D, respectively. *P<0.05 vs. the control; **P<0.01 vs. the control.
significantly decreased cell migration of Hep-2 cells, as shown in the histogram in Fig. 5C (P=0.023). Similarly, Transwell invasion assays using Matrigel further demonstrated that only Hep-2 cells transfected with the miR-365a-3p inhibitor were significantly reduced compared to the parental control cells (P=0.0024) (Fig. 5B and D). As shown in Fig. 6, the miR-143-5p inhibitor and the miR-494-3p mimics had no significant influence on the cell migration and invasion of Hep-2 cells.

miR-365a-3p promotes LSCC tumor growth and metastasis in vivo. To further view the potential effects of miR-365a-3p,
we established stably transfected Hep-2 cells containing an Lv-NC-inhibitor and an Lv-miR-365a-3p inhibitor (Fig. 7A) and injected the inhibitors into the enterocoelia of nude mice. The mice were euthanized, and the tumors were harvested on day 42 after injection. Tumors formed from Hep-2 cells stably transfected with the Lv-NC-inhibitor grew much faster than tumors from the cells that stably expressed the Lv-miR-365a-3p inhibitor (Fig. 7B). Accordingly, the final tumor volumes in the two groups were calculated and compared, as shown in Fig. 7C (P<0.0001 for the difference).

We further explored the metastasis in vivo using a body vision microscope. We found that tumors were metastasized in the livers (Fig. 8A) and hepatic lymph nodes (Fig. 8B). In particular, metastases to the livers in the Lv-NC-inhibitor group occurred at a significantly higher frequency than those in the Lv-miR-365a-3p inhibitor group. As shown in the stacked bars of Fig. 8C, 9 of 10 mice in the Lv-NC-inhibitor group showed liver metastases, but in the Lv-miR-365a-3p inhibitor group only 2 of 10 mice showed liver metastases. As expected, there was a similar trend of metastases in the hepatic lymph nodes in the Lv-NC-inhibitor and the Lv-miR-365a-3p inhibitor groups: 7 in 10 mice and only 1 in 10 mice showed hepatic lymph node metastases, respectively (Fig. 8D). These findings indicate that miR-365a-3p promotes LSCC tumor growth and metastases in vivo.

miR-365a-3p inhibitor downregulates p-AKT (Ser473). ERK and AKT play key roles in proliferation, migration, and invasion in malignant tumor cells. Accordingly, we performed western blotting to examine the protein levels of p-ERK1/2, p-AKT (Thr308), and p-AKT (Ser473). The results showed that the levels of p-ERK1/2 have no significant difference after miR-365a-3p inhibitor transfection compared with the levels of NC-inhibitor (P=0.082, Fig. 9A). Interestingly, miR-365a-3p inhibitor significantly decreased the expression of p-AKT (Ser473) but not p-AKT (Thr308) compared to the expression in the Hep-2 cells transfected with matched NC-inhibitor miRNA (Ser473, P<0.001; Thr308, P<0.245, Fig. 9B).

Discussion

Aberrant expression of miRNAs has been demonstrated to contribute to LSCC tumorigenesis and progression (13,15). For the first time, this study analyzed the roles of miR-365a-3p, miR-143-5p, and miR-494-3p in Hep-2 cells. Because of the overexpression of miR-365a-3p and miR-143-5p and the lower expression of miR-494-3p in LSCC patients with LNM (14), we transfected Hep-2 cells with miR-365a-3p inhibitor, miR-143-5p inhibitor, and miR-494-3p mimics. Interestingly, the loss-of-function results demonstrated that only the miR-365a-3p inhibitor significantly facilitated apoptosis and suppressed mitosis, migration, and invasion of Hep-2 cells, and we found that miR-365a-3p promotes LSCC xenograft tumor growth and metastasis in vivo. These data suggest that miR-365a-3p may be an oncomiR in LCSS, and miR-365a-3p inhibition may have potential in the treatment of laryngeal carcinoma.
The mature miR-365 is derived from two separate RNA precursors, hsa-miR-365a (previously named hsa-miR-365-1, MI0000767) and hsa-miR-365b (previously named hsa-miR-365-2, MI0000769), and mature hsa-miR-365a-3p (MIMAT0000710) is sheared from hsa-miR-365a (www.mirbase.org). miR-365 has been confirmed to act as an oncogene or tumor suppressor in different carcinomas (16,17), although this finding has not previously been reported in LSCC. Gastaldi et al (18) found that the expression of miR-365a was decreased in chemically-induced mouse skin carcinomas, which suggests a tumor suppressor role of miR-365a although miR-365a-3p has been found to be an oncogene (19). A recent study revealed that miR-365b was upregulated in cutaneous squamous cell carcinoma and induced subcutaneous tumors in vivo (16). The study also found that anti-miR-365b oligonucleotide inhibited cutaneous tumor formation in nude mice, along with apoptosis and G1 phase arrest in cancer cells. Hamada et al also found that miR-365 was highly expressed in ductal adenocarcinoma of the pancreas and contributed to the epithelial-mesenchymal transition. miR-365 can target apoptosis-promoting protein BAX and adaptor protein Src homology 2 domain containing 1, and then influences the survival of pancreatic cancer cells (20). Obviously, the expression levels of miR-365 are diverse in different malignant tumors. For example, in human gastric cancer the low expression of miR-365 correlated with poorly differentiated histology, advanced stage, and deep invasion, as well as the deregulation of phosphorylated Akt, p53, and cyclin D1. In mouse gastric cancers, the activation of Akt led to downregulated transcription of miR-365 and promoted gastric cancer cell proliferation (17). These data demonstrate that miR-365 was downregulated in gastric cancer. In addition, miR-365 was also found to be downregulated and accompanied the overexpression of NKX2-1 protein in lung cancer tissues. Furthermore, the miR-365 mimic significantly reduced the proliferation of lung cancer cells (21). The discrepancy of these findings may be due to the dynamic tumor microenvironment, stages of progression, and different types of cancers studied.

PI3K/AKT and MAPK signaling pathways are often hyperactivated in many malignant tumors, including LSCC, and the dependence of carcinoma cells on the two activated pathways has been used successfully in the clinic (22-25). AKT and ERK play key roles in the PI3K/AKT pathway and MAPK pathway, respectively (26,27). Recent studies have also found that AKT and ERK are highly correlated with miRNAs in various human malignancies (28-30). Therefore, to explore the underlying signaling pathways of miR-365a-3p-induced LCSS cell proliferation, migration, and invasion, we examined the expression of p-AKT (Thr308), p-AKT (Ser473), and p-ERK1/2. The data showed that only the expression of p-AKT (Ser473) was significantly influenced by the miR-365a-3p inhibitor. The serine/threonine kinase Akt encoded by the protein kinase B gene is a downstream effector of
P3K. Studies have shown that activation of Akt signaling is responsible for cancer cell proliferation, invasiveness, and metastasis (17,31). Akt activation is initiated by docking of the PH domain of Akt to PIP3 on the cellular membrane, exposing two critical amino acid residues for phosphorylation (32). Both phosphorylation events, of Ser473 by the protein kinase PDK1 and of Thr308 by the mTORC2 complex, are required for full activation of Akt (33). However, the levels of Akt phosphorylation on either Ser473 or Thr308 correlate differently with tumor cell growth and proliferation in distinct carcinomas (34-37). In our study, p-AKT (Ser473) was significantly decreased by the miR-365a-3p inhibitor, but p-AKT (Ser473) had no such influence. The data suggested that miR-365a-3p can mediate the P3K/AKT signaling pathway by upregulating p-AKT (Ser473) to promote growth and metastasis in LSCC. This result is consistent with a recent study that showed that patients with head and neck squamous cell carcinomas with higher levels of p-AKT (Ser473) had worse survival, but the survival of patients with higher levels of p-AKT (Thr308) was not affected (25).

Dysregulated miR-143 and miR-494 have important roles in many carcinomas (38,39), although this association has not been reported in LSCC patients. A recent study of epithelial cancers, including esophagus and lung cancer, indicated that downregulation of miR-143 contributes to epithelial cancer development, and its re-expression suppresses cellular proliferation and triggers apoptosis of epithelial cancer cells (38). miR-494 was reported to be significantly upregulated and to negatively modulate the expression of its target gene PTEN, in human cervical cancer cell lines and tumor samples (39). These findings identified the essential roles that miR-143 and miR-494 play in many carcinomas. However, our experiment shows that the two miRNAs did not have significant influence on Hep-2 cell apoptosis, mitosis, migration, and invasion in vitro. This is partly explained by the observation that miRNAs may have different roles cancer type-dependently.

In view of the low 5-year survival rate and the limited improvements in the treatment of LSCC during the past 20 years (1,2), finding novel molecular therapeutic targets for treating LSCC is essential. Because miRNAs play pivotal roles in the development of carcinoma, it is conceivable that miRNA mimics or inhibitors may become a new class of molecular target-based therapies for various cancers (40-42). Their effects and regulatory mechanisms in LSCC remain uncertain, but the present study revealed that miR-365a-3p, a novel oncomiR, activates the PI3K/AKT signaling pathway via p-AKT (Ser473) and can promote LSCC growth and metastasis. Therefore, miR-365a-3p may become a potential therapeutic target for the treatment of LSCC.

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