miR-509-3p Suppresses Migration, Invasion, and Epithelial–Mesenchymal Transition in Melanoma Cells by Targeting Collagen Triple Helix Repeat Containing 1

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Background: microRNAs (miRNAs) are ubiquitously dysregulated in numerous tumor cell types, including melanoma cells. The anti-tumor effect of miR-509-3p was widely evaluated in various cancers. Aims: To determine the functional role of miR-509-3p in melanoma. Study Design: Cell culture study. Methods: Expression of miR-509-3p in melanoma cell models were assessed by qRT-PCR. Cell migration and invasion were analyzed by wound healing and transwell assays, respectively. Expression levels of biomarkers of epithelial–mesenchymal transition were determined by Western blot. Luciferase vectors containing wildtype or mutant miR-509-3p binding site were constructed, and then dual-luciferase reporter assay. Results: Dysregulated miR-509-3p level was found in melanoma cells. Elevated miR-509-3p expression suppressed melanoma cell migration (P < .001) and invasion (P < .001) capacities. Epithelial–mesenchymal transition of melanoma cells was repressed by miR-509-3p, along with increased α-catenin/E-cadherin (P < .001) and decreased vimentin/fibronectin (P < .001). CTHRC1 (collagen triple helix repeat containing 1) contained a potential binding site for miR-509-3p, and miR-509-3p decreased protein expression of CTHRC1 in melanoma cells (P < .001). CTHRC1 promoted melanoma cell migration and invasion (P < .001), as well as contributed to epithelial–mesenchymal transition. Increased CTHRC1 expression attenuated miR-509-3p-induced inhibition of melanoma cell migration (P < .001), invasion, and epithelial–mesenchymal transition. Conclusion: miR-509-3p suppressed the biological function of melanoma cells through negatively regulating CTHRC1, shedding light on miR-509-3p as a potential candidate for melanoma therapeutics and treatments.

INTRODUCTION

Considered the most serious type of skin cancer, melanoma is mainly derived from melanocytes in the skin, mucous membranes, nerve centers, and other regions.1 Melanoma causes high mortality due to its strong invasiveness, easy metastasis, and easy tolerance to the treatment.2 The current strategies targeting melanoma mainly rely on early surgical removal, and there are no sustained effective treatments for the advanced stages of melanoma.3 Investigation of the molecular mechanism underlying the pathogenesis, as well as the development of novel therapeutic strategies, are urgent for the prevention of malignant metastasis of melanocytes.

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miR-509-3p was differentially expressed in A375 with resistance to vemurafenib\textsuperscript{10} and melanoma patients.\textsuperscript{11} However, the functional role of miR-509-3p in melanoma has not been thoroughly elucidated yet.

CTHRC1 retards collagen I synthesis and promotes cell migration of fibroblasts, facilitating vascular remodeling and tissue repair.\textsuperscript{12} Considering the tight link between tissue repair and carcinogenesis, CTHRC1 was regarded as an oncogene in tumours.\textsuperscript{13} CTHRC1 has been found to be overexpressed in melanoma cells, thus contributing to melanoma metastasis.\textsuperscript{14} We explored whether miR-509-3p regulated melanoma progression through targeting CTHRC1.

**MATERIAL AND METHODS**

**Cell Culture**

A375, SK-MEL-28, WM115, and WM35 (Chinese Academy of Sciences, Shanghai, China) were cultured in RPMI-1640 medium containing 10\% fetal bovine serum (Lonza, Basel, Switzerland) in a 37°C humidified incubator.

**Plasmid Construction and Cell Transfection**

CTHRC1 ORF was inserted into pcDNA3.1 backbone (Invitrogen, Waltham, MA, USA) to generate pcDNA-CTHRC1. SK-MEL-28 and WM115 were transfected with pcDNA vectors, miR-509-3p mimics, and an inhibitor (GenePharma, Shanghai, China) via Lipofectamine\textsuperscript{TM} 2000 (Invitrogen) for 48 h.

**Cell Migration**

Transfected SK-MEL-28 and WM115 cells with 80-90\% confluence were scratched, and a wound field in each well was generated. After 24 h of post-wound scratching, the mobilized cells were evaluated by a microscope (Olympus CY3, Tokyo, Japan).

**Cell Invasion**

Upper chambers (Corning, NY, USA) were seeded with transfected SK-MEL-28, and WM115 cells were supplemented with a serum-free medium. Medium containing serum was added into the lower chambers. After 48 h post-seeding, the cells in the lower chambers were fixed with 4\% methanol and stained with crystal violet before counting using a microscope.

**Western Blot**

Protein samples were extracted from transfected SK-MEL-28, and WM115 cells were extracted using RIPA lysis buffer (BioTeke, Beijing, China). Proteins (30 µg) were separated using SDS-PAGE and transferred onto a polyvinylidene difluoride membrane. The membrane was blocked in 5\% skim milk and incubated with anti-CTHRC1 (1:1000; Proteintech Group, Chicago, IL, USA), anti-α-catenin (1:2000), anti-E-cadherin (1:2000), anti-vimentin (1:2500), or anti-β-actin (1:3000) antibodies. The membrane was incubated with horseradish peroxidase-conjugated secondary antibodies (1:5000) by using Super Signal West Pico Chemiluminescent Substrate kit (Thermo Fisher; Waltham, MA, USA) as per the manufacturer’s instruction.

**RESULTS**

**miR-509-3p Suppressed Malignant Metastasis of Melanoma**

To elucidate the role of miR-509-3p in melanoma, A375, SK-MEL-28, WM115, and WM35 cells were processed using qRT-PCR analysis. WM35 cells showed the highest miR-509-3p expression, while WM115 cells showed the lowest expression (Figure 1A). miR-509-3p expression was higher in A375 than that in SK-MEL-28 (Figure 1A). Therefore, SK-MEL-28 (P = .001) and WM115 (P < .001) were then processed using gain-of-functional assays by overexpressing miR-509-3p mimic (Figure 1B). The migratory (Figure 1C) capacity of SK-MEL-28 (P = .032) and WM115 (P < .001) was repressed by miR-509-3p transfection, and the invasive (Figure 1D) capacity of SK-MEL-28 (P = .002) and WM115 (P = .003) cells were also repressed by miR-509-3p overexpression, suggesting that miR-509-3p repressed malignant metastasis of melanoma.

**miR-509-3p Repressed Epithelial–Mesenchymal Transition in Melanoma**

To elucidate the role of miR-509-3p in epithelial–mesenchymal transition, Western blot analysis of biomarkers was performed in SK-MEL-28 and WM115 cell models. Epithelial markers, including α-catenin (P < .001 and P = .0001) and E-cadherin (P < .001 and P < .001), were increased followed by miR-509-3p overexpression in SK-MEL-28 and WM115 cells (Figure 2). Whereas mesenchymal markers, such as vimentin (P < .001 and P < .001) and fibronectin (P < .001 and P < .001), were decreased following miR-509-3p treatment (Figure 2), suggesting that miR-509-3p suppressed epithelial–mesenchymal transition in melanoma cells.
FIG. 1. (A-D). miR-509-3p suppressed malignant metastasis of melanoma cells. (A) Differentially expressed miR-509-3p in melanoma cell lines (A375, SK-MEL-28, WM115, and WM35) detected by qRT-PCR. (B) miR-509-3p mimic was transfected into SK-MEL-28 and WM115 cells, and the expression of miR-509-3p was detected by qRT-PCR after 48 h of transfection. (C) miR-509-3p mimic was transfected into SK-MEL-28 and WM115, and cells and cell migration ability were detected by wound healing after 48 h of transfection. (D) miR-509-3p mimic was transfected into SK-MEL-28 and WM115 cells, and the cell invasion ability was detected by transwell after 48 h of transfection. **P < .01.

FIG. 2. (A,B). miR-509-3p suppressed epithelial–mesenchymal transition in melanoma cells. miR-509-3p mimic was transfected into SK-MEL-28 and WM115 cells, and protein expression levels of α-catenin, E-cadherin, vimentin, and fibronectin were detected by Western blot after 48 h of transfection. **P < .01.
CTHRC1 Was Targeted by miR-509-3p in Melanoma Cells

Target genes of miR-509-3p associated with biological effects in melanoma were screened and identified. CTHRC1 contains potential binding sites for miR-509-3p (Figure 3A). The results demonstrated that luciferase activity of CTHRC1-WT was decreased upon miR-509-3p mimic treatment compared to that of NC mimic treatment in SK-MEL-28 (\(P < .001\)) and WM115 (\(P = .004\)) cells (Figure 3B), while CTHRC1 mutation abolished such a suppressive effect of miR-509-3p (Figure 3B), suggesting that miR-509-3p directly binds to CTHRC1. Moreover, miR-509-3p overexpression directly decreased the CTHRC1 level, whereas miR-509-3p inhibition increased the expression of SK-MEL-28 (\(P < .001\) and \(P < .001\)) and WM115 (\(P < .001\) and \(P < .001\)) cells (Figure 3C). Taken together, data obtained so far illustrated that CTHRC1 was targeted by miR-509-3p in melanoma cells.

miR-509-3p Suppressed Malignant Metastasis of Melanoma Cells Through Regulation of CTHRC1

To determine the effect of miR-509-3p/CTHRC1 axis in the metastasis of melanoma, pcDNA-CTHRC1 and miR-509-3p mimic were cotransfected into SK-MEL-28 and WM115 cells. Western blot analysis showed that CTHRC1 overexpression decreased protein levels of α-catenin (\(P < .001\)) and E-cadherin (\(P < .001\)), whereas elevated vimentin (\(P < .001\)) and fibronectin (\(P < .001\)) (Figure 4A). However, cotransfection using pcDNA-CTHRC1 and miR-509-3p mimic reversed CTHRC1-induced decrease of α-catenin (\(P < .001\)) and E-cadherin (\(P < .001\)) levels, as well as restored the CTHRC1-mediated increase of vimentin (\(P < .001\)) and fibronectin (\(P < .001\)) (Figure 4A). Moreover, elevated CTHRC1 expression promoted migration (Figure 4B) and invasion (Figure 4B) of SK-MEL-28 and WM115 (\(P < .001\)) cells. miR-509-3p could attenuate the promotive effects of CTHRC1 on cell migration (Figure 4B) and invasion (Figure 4C) in SK-MEL-28 and WM115 (\(P < .001\)) cells. Collectively, miR-509-3p suppressed the malignant metastasis of melanoma cells through the regulation of CTHRC1.

DISCUSSION

Distant metastasis is a major obstacle for the effective treatment of melanoma, metastasis, and invasion, which are the most important reasons for the death of patients due to melanoma.\(^1\) miRNAs, due to their ability to modulate melanoma cell growth, have been considered as new therapeutic strategies for melanoma.\(^7\) Although miR-509-3p was shown to be responsible for suppressing tumor cell metastasis, the exact role of miR-509-3p in melanoma progression remained to be investigated.

Melanoma is highly curable in the early stages; however, melanoma often develops metastasis devoid of effective diagnostic biomarkers.\(^5\) Therefore, it is important to discover diagnostic or prognostic biomarkers for melanoma. miRNAs have been reported to be related to the diagnosis and prognosis of melanoma, which could contribute to personalized treatment.\(^6\) Dysregulation of the miR-506-514 cluster, including miR-506-3p and miR-509-3p,
was found in metastatic cutaneous melanoma.\textsuperscript{17} miR-509-3p was reduced in melanoma patients, and such different expression levels could discriminate between non-melanoma controls and melanoma patients at different stages.\textsuperscript{18} Moreover, glypican 6 was upregulated in melanoma cell lines and considered as a biomarker for cutaneous melanoma, whose expression was suppressed by miR-509-3p.\textsuperscript{18} We observed that miR-509-3p was dysregulated in melanoma cells, and in this study, the relationship between overall survival or clinico-pathological parameters of patients with melanoma and miR-509-3p expression levels should be further investigated, with the aim to provide diagnostic or prognostic roles of miR-509-3p in melanoma.

Functional assays revealed that miR-509-3p could suppress melanoma cell migration and invasion, suggesting the anticancer role in melanoma. Epithelial–mesenchymal transition in melanoma cells is often accompanied by the transformation of epithelial cells into invasive mesenchymal cells, and promotion of epithelial–mesenchymal transition could contribute to metastasis of melanoma.\textsuperscript{19} Suppression of epithelial–mesenchymal transition facilitated the repressive role of miR-204-3p on melanoma cells.\textsuperscript{20} This study indicated that miR-509-3p decreased the expressions of vimentin and fibronectin, whereas increased the expression of α-catenin and E-cadherin in melanoma cells, thus suppressing metastasis of melanoma cells. Transcription factors, including TWIST1, Slug, and Snail, have been shown to promote epithelial–mesenchymal transition in tumors.\textsuperscript{21} Since miR-509-3p could target epithelial–mesenchymal transition regulator, TWIST1, in ovarian cancer,\textsuperscript{9} whether TWIST1 was involved in miR-509-3p-mediated
epithelial–mesenchymal transition in melanoma cells should be investigated in future studies.

CTHRC1 was found to be enhanced in metastatic melanoma specimens compared to nonmetastatic lesions\textsuperscript{22} and promoted melanoma cell survival.\textsuperscript{22} CTHRC1 knockdown retarded the progression of melanoma cells.\textsuperscript{23} We found that CTHRC1 overexpression contributed to the progression of melanoma cells. Previous research reported that the transforming growth factor could stimulate CTHRC1 to modulate Smad2/3, Wnt or Src, and Erk pathways, thus enhancing epithelial–mesenchymal transition to facilitate cancer metastasis.\textsuperscript{24} Elevated CTHRC1 expression in melanoma cells also promotes epithelial–mesenchymal transition with increased vimentin and fibronectin and decreased α-catenin and E-cadherin. CTHRC1 was found to be involved in miR-134,\textsuperscript{25} or miR-155,\textsuperscript{26} mediated melanoma tumorigenesis. This study first confirmed that miR-509-3p could directly target CTHRC1 and negatively regulated its protein expression. Moreover, CTHRC1 overexpression counteracted the repressive effects of miR-509-3p on melanoma cell migration, invasion, and epithelial–mesenchymal transition. Furthermore, the detailed downstream signaling pathways involved in progression regulation by miR-509-3p/CTHRC1 axis need to be further investigated.

In conclusion, this study reported miR-509-3p was a tumor suppressor in melanoma cells, modulating tumor progression through inhibiting CTHRC1 expression. Taken together, our results provide insights into miR-509-3p as a promising therapeutic agent in melanoma progression from bench to clinic.

**Ethics Committee Approval:** N/A.

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