

HNFI A-AS1 Regulates Cell Migration, Invasion and Glycolysis via Modulating miR-124/MYO6 in Colorectal Cancer Cells

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Background: Accumulating evidence determined that lncRNAs play multiple roles in cell progression in colorectal cancer (CRC). Long noncoding RNA (lncRNA) hepatocyte nuclear factor 1 homeobox A (HNF1A)-antisense RNA 1 (AS1) has been identified to affect cell growth and disease diagnosis in various cancers, including CRC. However, the underlying regulatory mechanism of HNF1A-AS1 in cell progression and glycolysis has not been fully explored in CRC.

Materials and Methods: The expression of HNF1A-AS1, microRNA-124 (miR-124) and Myosins of class VI (MYO6) was detected using reverse transcription-quantitative polymerase chain reaction (RT-qPCR). The analysis of glucose consumption, lactate production and hexokinase 2 (HK2) protein level was used to assess glycolysis in cells. The protein level of HK2 and MYO6 was measured with Western blot. Cell migration and invasion were evaluated using the transwell assay. The relationship among HNF1A-AS1, miR-124 and MYO6 was determined via luciferase reporter and RNA immunoprecipitation (RIP) assay.

Results: In this study, we found that HNF1A-AS1 was upregulated in CRC tissues and cell lines. Functional experiments determined that reduction of HNF1A-AS1 or promotion of miR-124 inhibited cell migration and invasion as well as glycolysis in CRC cells. What's more, luciferase reporter assay manifested that miR-124 was a target of HNF1A-AS1 and MYO6 was a target mRNA of miR-124 in CRC cells. Additionally, reverse experiments showed that the effects of si-HNF1A-AS1 on colorectal cancer cells were impaired by anti-miR-124 and the effects of high miR-124 expression on CRC cells were rescued by upregulating MYO6. HNF1A-AS1 regulated MYO6 expression via targeting miR-124 in CRC cells.

Conclusion: In this study, we first found that HNF1A-AS1 regulated cell migration, invasion and glycolysis via modulating miR-124/MYO6 in CRC cells.

Keywords: HNF1A-AS1, miR-124, MYO6, colorectal cancer

Introduction

Colorectal cancer (CRC) is the third most common cancer worldwide, with high morbidity and mortality in China.^{1,2} Because advanced colorectal cancer still lacks effective diagnosis and treatment, it is necessary to further explore and analyze the pathogenesis and regulation mechanism of CRC to determine effective biomarkers.

At present, a large number of studies have shown that long noncoding RNAs (lncRNAs) are important regulators in cells and are widely involved in the occurrence and development of human diseases.³⁻⁹ lncRNAs are noncoding RNAs with a length of more than 200 nt, as ceRNAs, which further regulated the production of

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downstream proteins by regulating the miRNA/mRNA axis.^{10–13} Some lncRNAs have been reported to be differentially expressed in colorectal cancer tissues and cells, but their functions have been verified.¹⁴ For example, lncRNA DLEU1 promoted CRC cell progression through regulating KPNA3.¹⁵ Moreover, lncRNA CRNDE induced cell proliferation and chemoresistance in CRC.¹⁶ lncRNA BANCR has been determined to relate to CRC poor prognosis.¹⁷ However, the regulation mechanism of lncRNA in colorectal cancer has not yet been elucidated, and further exploration and research are needed.

The lncRNA HNF1A antisense RNA 1 (HNF1A-AS1), as an oncogene, was involved in cell progression and metabolism in many cancers, such as lung adenocarcinoma, colorectal carcinoma, oesophageal adenocarcinoma, nonsmall cell lung cancer, bladder cancer and hepatocellular carcinoma.^{18–23} Zhao et al demonstrated that induction of HNF1A-AS contributed to cell proliferation and metastasis through activating Wnt/ β -catenin signaling pathway in osteosarcoma.²⁴ In addition, lncRNA HNF1A-AS1 has been determined to be a novel diagnostic predictor for prognosis of cancers.^{25,26} However, the effect of HNF1A-AS1 on cell glycolysis has not been explained and the regulatory network HNF1A-AS1 remains to be further explored and analyzed.

In the present study, we investigated the functional effects of lncRNA HNF1A-AS1 on the cell metabolism and glycolysis in CRC cells, as well as the potential regulatory mechanism.

Materials and Methods

Tissues, Cell Culture and Transfection

Forty pairs of CRC tissues and adjacent normal tissues were obtained from patients who were diagnosed at Xiangya Hospital Central South University. This experiment has been approved by the Ethics Committee of Xiangya Hospital Central South University. Informed written consents were acquired from patients.

CRC cell lines (SW620, HT-29, HCT116 and SW480) were purchased from American Type Culture Collection (ATCC, Manassas, VA, USA). The immortalized colon cell (NCM460) was purchased from Rongbai (Shanghai, China). All cells were cultured in Dulbecco's modified Eagle's medium (DMEM; Thermo Fisher Scientific, Inc., Waltham, MA, USA) containing 10% fetal bovine serum (FBS, Sciencell Research, Carlsbad, CA, USA) and 1% penicillin at 37°C in a humidified atmosphere of 5% CO₂.

Small interfering RNA (siRNA) targeting HNF1A-AS1 (si-HNF1A-AS1), siRNA scrambled control (si-NC), pcDNA-HNF1A-AS1 (HNF1A-AS1), pcDNA-MYO6 (MYO6), pcDNA, miR-124 mimic (miR-212-3p), miRNA negative control (miR-NC), miR-124 inhibitor (anti-miR-212-3p) and inhibitor negative control (anti-miR-NC) were purchased from GenePharma Co. Ltd. (Suzhou, China). These vectors and oligos were transfected into SW620 and HCT116 cells using Lipofectamine 2000 reagent (Invitrogen, Carlsbad, CA, USA) according to the manufacturer's instructions.

Reverse Transcription-Quantitative Polymerase Chain Reaction (RT-qPCR)

Total RNA was extracted from CRC cells and tissues using TRIzol reagent (Invitrogen) referring to the protocols supplied by the manufacturer. 1 ng of RNA was reversely transcribed into cDNA using PrimeScript RT reagent kit (Takara Biotechnology Co., Ltd., Dalian, China). For HNF1A-AS1 and MYO6 detection, One Step SYBR PrimeScript™ RT-PCR Kit (Takara) was used to measure the relative expression of HNF1A-AS1 and MYO6. For miR-124 detection, TaqMan miRNA assays (Applied Biosystems, Carlsbad, CA, USA) were used to measure the relative expression of miR-124. All samples were performed on a CFX96 Touch™ Real-Time PCR system (Bio-Rad, Hercules, CA, USA). Glyceraldehyde phosphate dehydrogenase (GAPDH) and U6 were employed to normalize for mRNA and miRNA, respectively. The primer sequences were as follows: the primers for U6: forward, 5'-CTCGCTTCGGCAGCAC-3' and reverse 5'-AACGCTTCACGAATTTGCGT-3'. The primers for miR-124, forward, 5'-GCCTAAGGCACGCGGTG-3' and reverse, 5'-GTCGTATCCAGTGCAGGGTCCGAGGTA TTCGACTGGATACGACGGCATT -3'. The primers for HNF1A-AS1: forward 5'-TCAAGAAATGGTGGCTAT-3' and reverse, 5'-GATCTGAGACTGGCTGAA-3'. The primers for MYO6: forward 5'-GATGGAGCTGCACCCTGACA-3' and reverse, 5'-GCTCTCAATGGCGCTCTGAAG-3'. The primers for GAPDH: forward 5'-GGAGCGAGATCCCTCCAAAAT-3' and reverse, 5'-GGCTGTTGTCATAC TTCTCATGG-3'.

Western Blot Assay

Cells were lysed with RIPA Lysis Buffer (Thermo Fisher Scientific) and the protein concentrations were quantified by a bicinchoninic acid (BCA) Protein Assay Kit (Thermo Fisher Scientific). Protein samples were added onto sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE) and

transferred to polyvinylidene difluoride (PVDF) membranes (Millipore, Bedford, MA, USA). The membranes were incubated with primary antibodies HK2 (1:2000; Santa Cruz Biotechnology, Inc., Santa Cruz, CA, USA), MYO6 (1:2000 dilution; Santa Cruz Biotechnology) or GAPDH (1:1000 dilution; Santa Cruz Biotechnology) at 4°C overnight. After being washed in TBS, the membrane was incubated with horseradish peroxidase (HRP)-conjugated goat anti-rabbit secondary antibody (Santa Cruz Biotechnology) for 1 h at room temperature. The blots were detected by Clarity Western ECL substrate (Bio-Rad) and visualized with a ChemiDoc™ MP Imaging System (Bio-Rad).

Glucose Consumption and Lactate Production Assay

The dose of cloxiquine (CLQ, 50 μM) with/without GW9662 (10 μM, PPAR gamma antagonist) was used to treat SW620 and HCT116 cells for 24 h. Glucose content was determined using a Glucose Assay kit (Rongsheng Biotechnology, Shanghai, China) according to the supplier's instructions. The concentrations of intracellular lactate were analyzed with a lactate assay kit (Sigma, St Louis, MO, USA) following the protocols of the manufacturer.

Transwell Assay

For invasion assay, 2×10^3 cells were seeded in the upper chambers with 8-μm pores Matrigel-coated (BD Biosciences, San Jose, CA, USA) and cultured in concentration in DMEM. The lower chamber was added with DMEM containing 10% FBS as a chemoattractant. After incubation for 48 h at room temperature, cells in the upper chamber were scraped off with cotton swabs. Cells in the lower chamber were fixed with methanol and then stained by crystal violet for 30 min. For migration assay, cells were added onto the upper chambers without Matrigel-coated, other steps were similar to invasion assay. The migrated and invasive cells were calculated using a Countess automatic cell counter (Invitrogen).

Luciferase Reporter Assay

The fragments of HNF1A-AS1 containing the wild-type or mutated miR-124 binding sites were amplified and cloned into a pRL-TK (Promega, Madison, WI, USA), namely HNF1A-AS1-WT-Luc or HNF1A-AS1-MUT-Luc. In addition, the fragments of MYO6 containing the wild-type or mutated miR-124 binding sites were amplified and inserted into pRL-TK, namely MYO6-

WT-Luc or MYO6-MUT-Luc. Then, these vectors were co-transfected with miR-124 or miR-NC into SW620 and HCT116 cells using Lipofectamine 2000 reagent (Invitrogen). The renilla luciferase control vector pRL-ubi-63E was a control vector using transfection reagent Lipofectamine 2000 (Invitrogen). After transfection for 48 h, cells were collected and the luciferase activity was measured using the Dual-Luciferase Reporter Assay System (Promega).

RNA Immunoprecipitation (RIP) Assay

The EZ-Magna RIP Kit (Millipore) was used for RIP experiments according to the manual. Briefly, cells were collected by centrifugation and lysed in RIP buffer. Then, cells were incubated with human anti-Argonaute 2 (Ago2) antibody (Abcam) or normal mouse IgG (negative control, Sigma-Aldrich) overnight. Nonspecific binding was removed with proteinase K buffer (Abcam), and then the immunoprecipitated RNA was isolated. RT-qPCR was used to analyze the enrichment of HNF1A-AS1 or MYO6.

Statistical Analysis

Statistical analyses were performed using GraphPad Prism 7.0 (GraphPad Software, San Diego, CA, USA). Comparison between the two groups was analyzed using Student's *t*-test and among three or more sets were calculated using LSD. All data were presented as means ± standard deviation (SD). $P < 0.05$ was considered to be statistically significant.

Results

HNF1A-AS1 Was Upregulated in Colorectal Cancer Tissues and Cell Lines

To explore the expression of HNF1A-AS1 in CRC tissues and cell lines, we collected 40 pairs of colorectal cancer tissues and adjacent tissues as well as purchased four CRC cell lines (SW620, HT-29, HCT116 and SW480) and immortalized colon cells (NCM460). As shown in [Figure 1A](#), the expression of HNF1A-AS1 in cancer tissues was significantly higher than that in normal tissues. In addition, we also found HNF1A-AS1 expression was upregulated in SW620, HT-29, HCT116 and SW480 cells compared with that in NCM460 cells ([Figure 1B](#)). Moreover, we selected two CRC lines (SW620 and HCT116), which have a relative higher expression of HNF1A-AS1 for the following experiments. Collectively, these results showed that HNF1A-AS1 was upregulated in CRC tissues and cell lines.

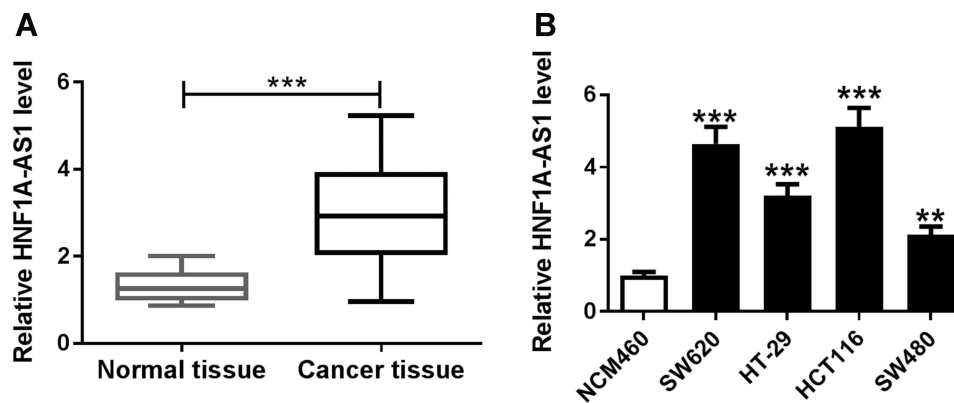


Figure 1 HNF1A-AS1 was upregulated in CRC tissues and cell lines. **(A, B)** The expression of HNF1A-AS1 in colorectal cancer tissues and normal tissues **(A)**, as well as CRC cells (SW620, HT-29, HCT116 and SW480) and immortalized colon cells (NCM460) was detected by RT-qPCR. ** $P < 0.01$. *** $P < 0.001$.

Knockdown of HNF1A-AS1 Inhibited Cell Migration and Invasion as Well as Glycolysis in Colorectal Cancer Cells

To determine the effects of HNF1A-AS1 on cell migration, invasion and glycolysis in CRC cells, we transfected two CRC cell lines (SW620 and HCT116) with siHNF1A-AS1 to construct CRC cells with lower HNF1A-AS1 expression than control cells (Figure 2A). As shown in Figure 2B and C, both relative glucose consumption and lactate production were obviously reduced in SW620 and HCT116 cells with HNF1A-AS1 knockdown compared with cells transfected with si-NC (Figure 2B and C). More than that, the expression of HK2 was also decreased in the si-HNF1A-AS1 group compared with that in the si-NC group of SW620 and HCT116 cells (Figure 2D). In addition, cell migration and invasion were significantly lower in SW620 and HCT116 transfected with si-HNF1A-AS1 than that in cells transfected with si-NC (Figure 2E and F). These data determined that knockdown of HNF1A-AS1 inhibited cell migration and invasion as well as glycolysis in CRC cells.

miR-124 Was a Target of HNF1A-AS1

To further investigate the potential regulatory mechanisms of HNF1A-AS1 in CRC cells, we sought potential target miRNAs for HNF1A-AS1 through DIANA tools and found that miR-124 had a reverse complement to 3'UTR of HNF1A-AS1 (Figure 3A). Therefore, we constructed HNF1A-AS1-WT-Luc (binding sites of wild type) or HNF1A-AS1-MUT-Luc (binding sites of mutate) vectors. Luciferase reporter assay determined that luciferase activity of HNF1A-AS1-WT-Luc in SW620 and HCT116 cells were remarkably reduced by miR-124, while the luciferase

activity of HNF1A-AS1-MUT-Luc has not changed (Figure 3B and C). Otherwise, RIP assay was applied to further validate the interaction between miR-124 and HNF1A-AS1, the results showed that HNF1A-AS1 was significantly enriched in the RIP-Ago2 than RIP-IgG in SW620 and HCT116 cells (Figure 3D and E). Meanwhile, we analyzed the expression of miR-124 in SW620 and HCT116 cells with HNF1A-AS1-overexpression or HNF1A-AS1-down-expression via RT-qPCR (Figure 3F and G). The results showed that upregulated HNF1A-AS1 inhibited miR-124 expression while downregulated HNF1A-AS1 promoted miR-124 expression in SW620 and HCT116 cells. Based on the above results, we ensured that miR-124 was a target miRNA of HNF1A-AS1.

The Effects of Downregulated HNF1A-AS1 on Colorectal Cancer Cells Were Impaired by Silenced miR-124

To further understand the function of miR-124 in CRC cells, we analyzed the expression of miR-124 in four CRC cell lines (SW620, HT-29, HCT116 and SW480) by RT-qPCR and observed that miR-124 expression was significantly reduced in CRC tissues and cells compared to immortalized colon cells (NCM460) (Figure 4A and B). As shown in Figure 4B and C, we constructed SW620 and HCT116 cells with a lower expression of miR-124 through transfection with anti-miR-124. Furthermore, the si-HNF1A-AS and anti-miR-124 were co-transfected into SW620 and HCT116 cells for functional rescue experiments. We found that si-HNF1A-AS transfection significantly induced the expression of miR-124, which the promoted impact was alleviated by anti-miR-124 transfection in SW620 and HCT116 cells. Additionally, both

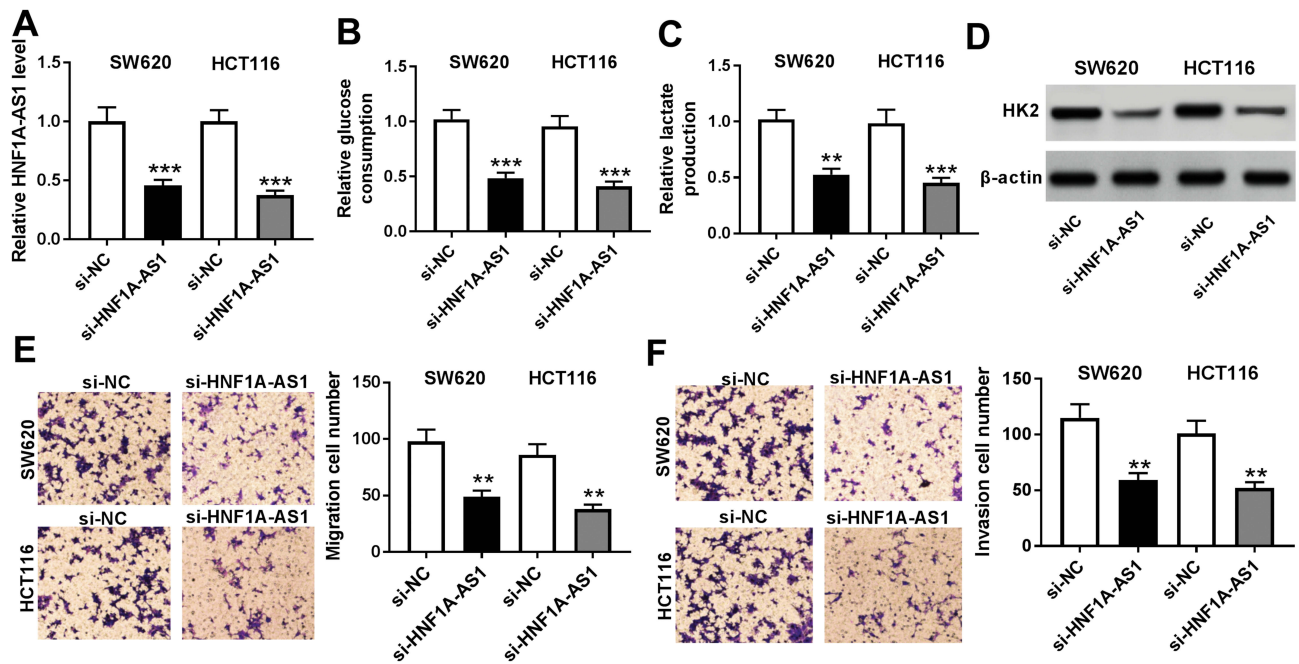


Figure 2 Knockdown of HNF1A-AS1 inhibited cell migration and invasion as well as glycolysis in CRC cells. SW620 and HCT116 cells were transfected with si-NC or si-HNF1A-AS1. (A) HNF1A-AS1 expression in transfected SW620 and HCT116 cells was monitored by RT-qPCR. (B) Glucose consumption was measured in transfected cells utilizing Glucose Assay kit. (C) Lactate production was detected in SW620 and HCT116 cells with a lactate assay kit after transfection. (D) The protein level of HK2 in transfected SW620 and HCT116 cells was detected by Western blot. (E and F) Cell migration (E) and cell invasion (F) in SW620 and HCT116 cells were examined with transwell assay. ***P* < 0.01. ****P* < 0.001.

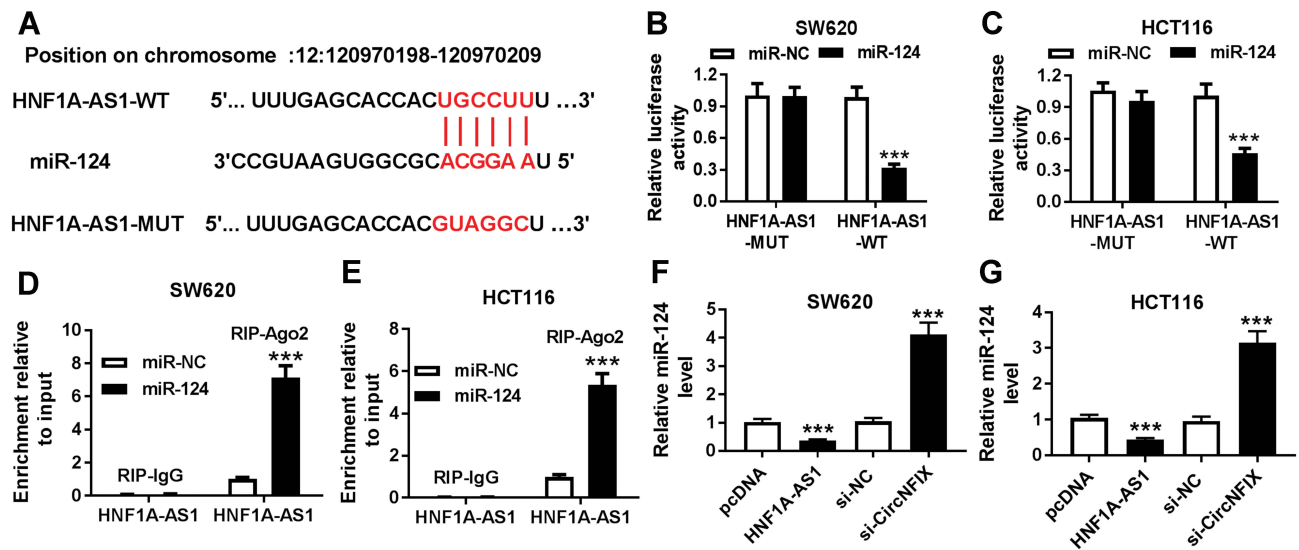


Figure 3 miR-124 was a target of HNF1A-AS1. (A) The predicted wild-type and mutated type of miR-124 binding sites in the 3'UTR of HNF1A-AS1 were exhibited. (B and C) Luciferase activity was measured in SW620 (B) and HCT116 (C) cells co-transfected with HNF1A-AS1-WT or HNF1A-AS1-MUT and miR-124 or miR-NC through luciferase reporter assay. (D and E) RIP assay was performed to ensure the relationship between HNF1A-AS1 and miR-124 in SW620 (D) and HCT116 (E) cells. (F and G) the expression of miR-124 was detected in SW620 (F) and HCT116 (G) cells transfected with pcDNA, HNF1A-AS1, si-NC or si-HNF1A-AS1. ****P* < 0.001.

relative glucose consumption and lactate production were inhibited by downregulating HNF1A-AS, while this inhibitory effect was weakened by anti-miR-124 transfection in SW620 and HCT116 cells (Figure 4D–G). In addition, the reduction of HNF1A-AS inhibited the expression of

HK2, which was rescued by anti-miR-124 transfection in SW620 and HCT116 cells (Figure 4H). Strikingly, cell migration and invasion were suppressed by reduction of HNF1A-AS, while they were relieved by inhibition of miR-124 in SW620 and HCT116 cells (Figure 4I–M).

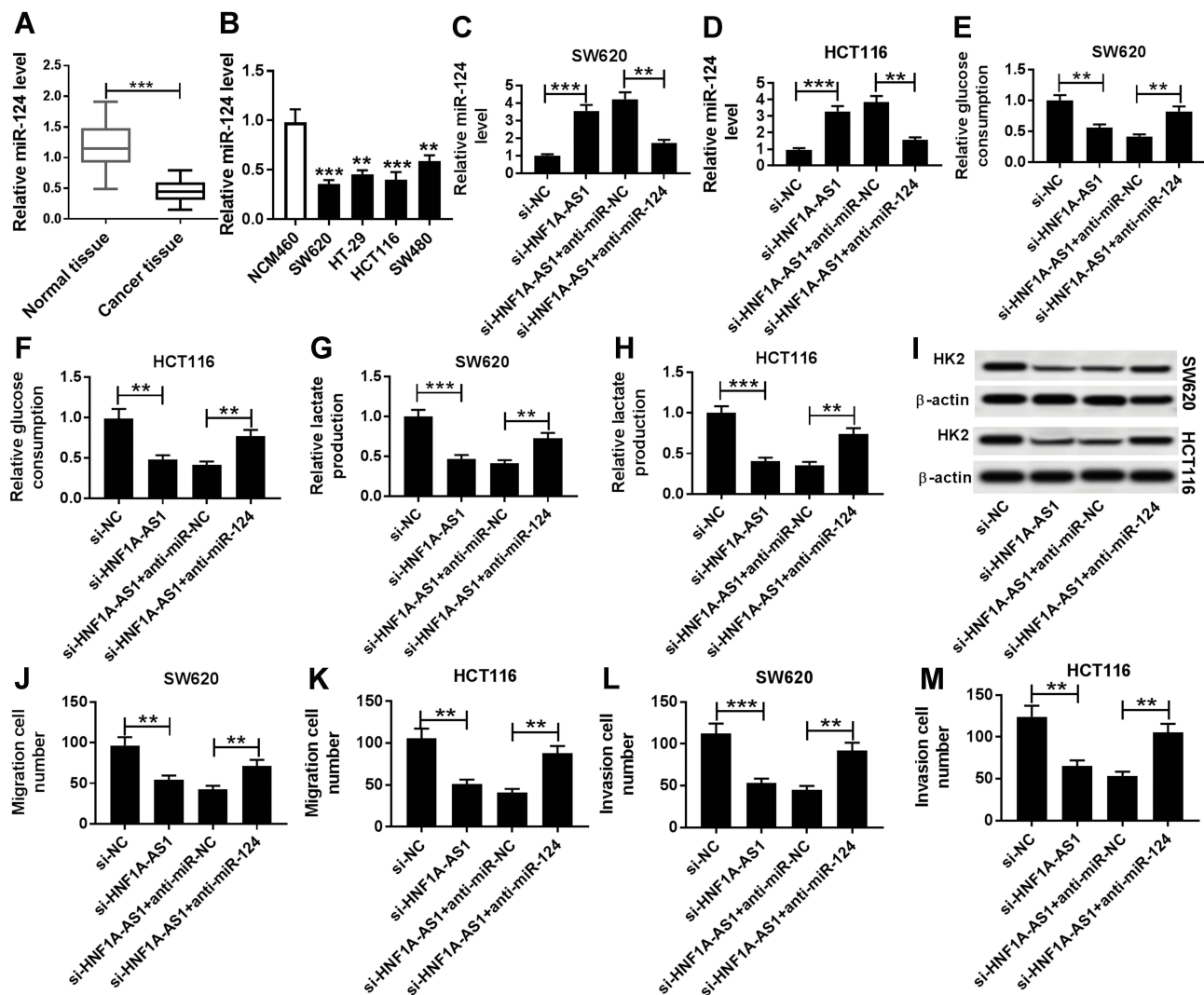


Figure 4 The repressed effects of si-HNF1A-AS1 on cell migration, invasion and glycolysis in CRC cells were impaired by anti-miR-124. **(A)** The expression of miR-124 in CRC cell tissues and normal tissues was detected using RT-qPCR. **(B)** The expression of miR-124 in CRC cell lines (SW620, HT-29, HCT116 and SW480) and immortalized colon cells (NCM460) was analyzed by RT-qPCR. **(C–M)** SW620 and HCT116 cells were transfected with si-NC, si-HNF1A-AS1, si-HNF1A-AS1 + miR-NC or si-HNF1A-AS1 + miR-124. **(C and D)** the expression of miR-124 was detected in SW620 **(C)** and HCT116 **(D)** cells using RT-qPCR. **(D and E)** Glucose consumption was measured in SW620 **(E)** and HCT116 **(F)** cells after transfection. **(G and H)** Lactate production was tested in SW620 **(G)** and HCT116 **(H)** cells after transfection. **(I)** The expression of HK2 was evaluated in transfected SW620 and HCT116 cells. **(J and K)** cell migration was detected in treated SW620 **(J)** and HCT116 **(K)** cells. **(L and M)** cell invasion was analyzed in SW620 **(L)** and HCT116 **(M)** cells after transfection. ** $P < 0.01$. *** $P < 0.001$.

Thus, these data determined that silencing of miR-124 could reverse the si-HNF1A-AS-mediated inhibited effects on cell migration, invasion and glycolysis in SW620 and HCT116 cells.

miR-124 Directly Targeted MYO6

We all know that miRNAs regulate the expression of downstream target genes. To further investigate the regulatory network of HNF1A-AS, we predicted the potential target genes of miR-124 by DIANA tools and found that MYO6 has a complementary sequence to miR-124 (Figure 5A). MYO6-WT-Luc or MYO6-MUT-Luc vector was

constructed and co-transfected into SW620 and HCT116 cells with miR-NC or miR-124. Luciferase reporter assay analysis showed that luciferase activity of MYO6-WT-Luc was apparently decreased after co-transfection with miR-124, but no significant change was observed in luciferase activity of MYO6-MUT-Luc (Figure 5B and C). RIP assay also determined that MYO6 expression was enriched in the RIP-Ago2 than RIP-IgG in SW620 and HCT116 cells (Figure 5D and E). Additionally, the expression of MYO6 was inhibited by the upregulation of miR-124, while promoted by downregulation of miR-124 in SW620 and HCT116 cells (Figure 5F and G). The above results

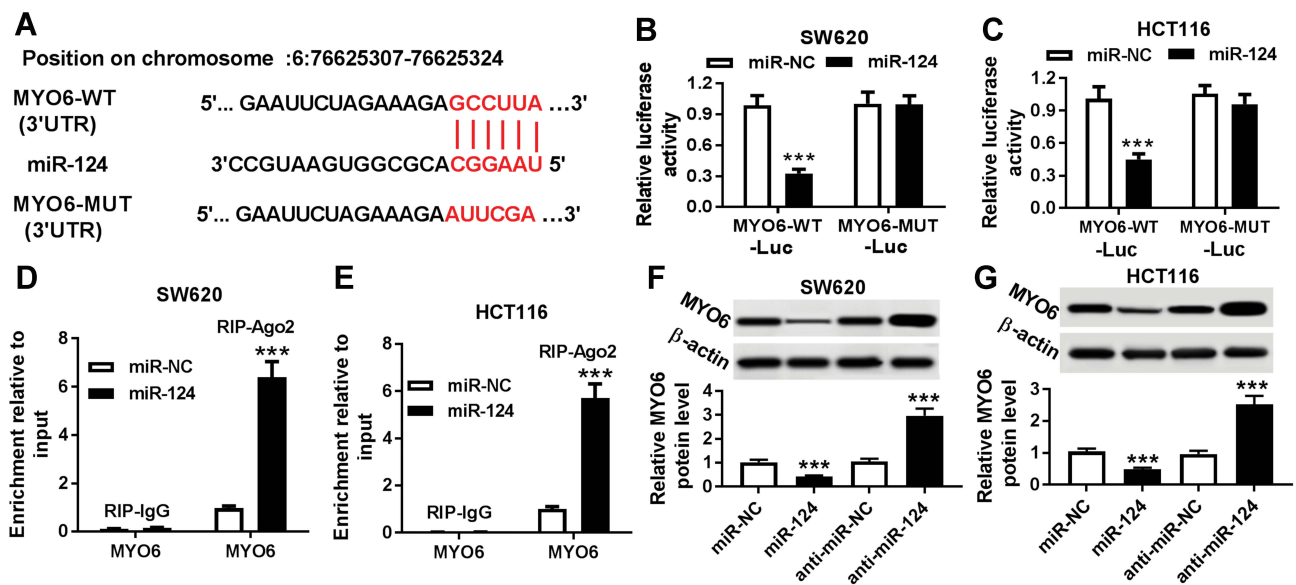


Figure 5 miR-124 directly targeted MYO6. (A) The predicted wild-type and mutated type of miR-124 binding sites in the 3'UTR of MYO6 were shown. (B and C) Luciferase activity was measured in SW620 (B) and HCT116 (C) cells co-transfected with MYO6-WT-Luc or MYO6-MUT and miR-124 or miR-NC by luciferase reporter assay. (D and E) RIP assay was performed to ensure the relationship between MYO6 and miR-124 in SW620 (D) and HCT116 (E) cells. (F and G) The protein expression of MYO6 in SW620 (F) and HCT116 (G) cells transfected with miR-NC, miR-124, anti-miR-NC or anti-miR-124 was detected with Western blot assay. *** $P < 0.001$.

indicated that MYO6 was the target gene of miR-124 in CRC cells.

The Effects Induced by Overexpression of miR-124 on Colorectal Cancer Cells Were Rescued by Upregulated MYO6

To further clarify the regulatory mechanisms of miR-124 and MYO6 in CRC, the rescue experiments were performed on SW620 and HCT116 cells. First, we measured the expression of MYO6 in CRC tissues and cell lines (SW620, HT-29, HCT116 and SW480), and the results showed that MYO6 was highly expressed in CRC tissues and cells (Figure 6A and B). In the following experiment, miR-124 mimics and pcDNA-MYO6 were co-transfected into SW620 and HCT116 cells. As depicted in Figure 6B and C, promotion of miR-124 inhibited MYO6 expression, while upregulating MYO6 could reverse this effect in SW620 and HCT116 cells. Furthermore, relative glucose consumption, lactate production and HK2 expression were obviously reduced by miR-124 transfection, which was impaired by the induction of MYO6 in SW620 and HCT116 cells (Figure 6D–H). More than that, the inhibited effects of high miR-124 expression on cell migration and invasion in SW620 and HCT116 cells were rescued by improving MYO6 expression (Figure 6I–M). These results showed that the induction of MYO6 could impair the

inhibited effects of high miR-124 expression on cell metastasis and glycolysis in CRC cells.

HNF1A-AS1 Regulated MYO6 Expression Through miR-124 in Colorectal Cancer Cells

In order to further explore the regulatory network among HNF1A-AS1, miR-124 and MYO6. Luciferase reporter methods and Western blot assay were conducted. We found that the luciferase activity was significantly reduced in MYO6-WT-Luc + miR-124 group compared with that in the MYO6-WT-Luc group. In addition, compared with MYO6-WT-Luc+miR-124 + pcDNA group, luciferase activity was notably increased in MYO6-WT-Luc + miR-124 + HNF1A-AS1 group in SW620 and HCT116 cells (Figure 7A and B). Not only that, the analysis of Western blot showed that induction of HNF1A-AS1 improved the protein level of MYO6, which was inhibited by the promotion of miR-124 in SW620 and HCT116 cells (Figure 7C and D). Therefore, these results indicated that HNF1A-AS1 regulated the expression of MYO6 through miR-124 in colorectal cancer cells.

Discussion

In this paper, we found that lncRNA HNF1A-AS1 was highly expressed in CRC tissues and cells, and further functional experiments showed that silenced HNF1A-AS1

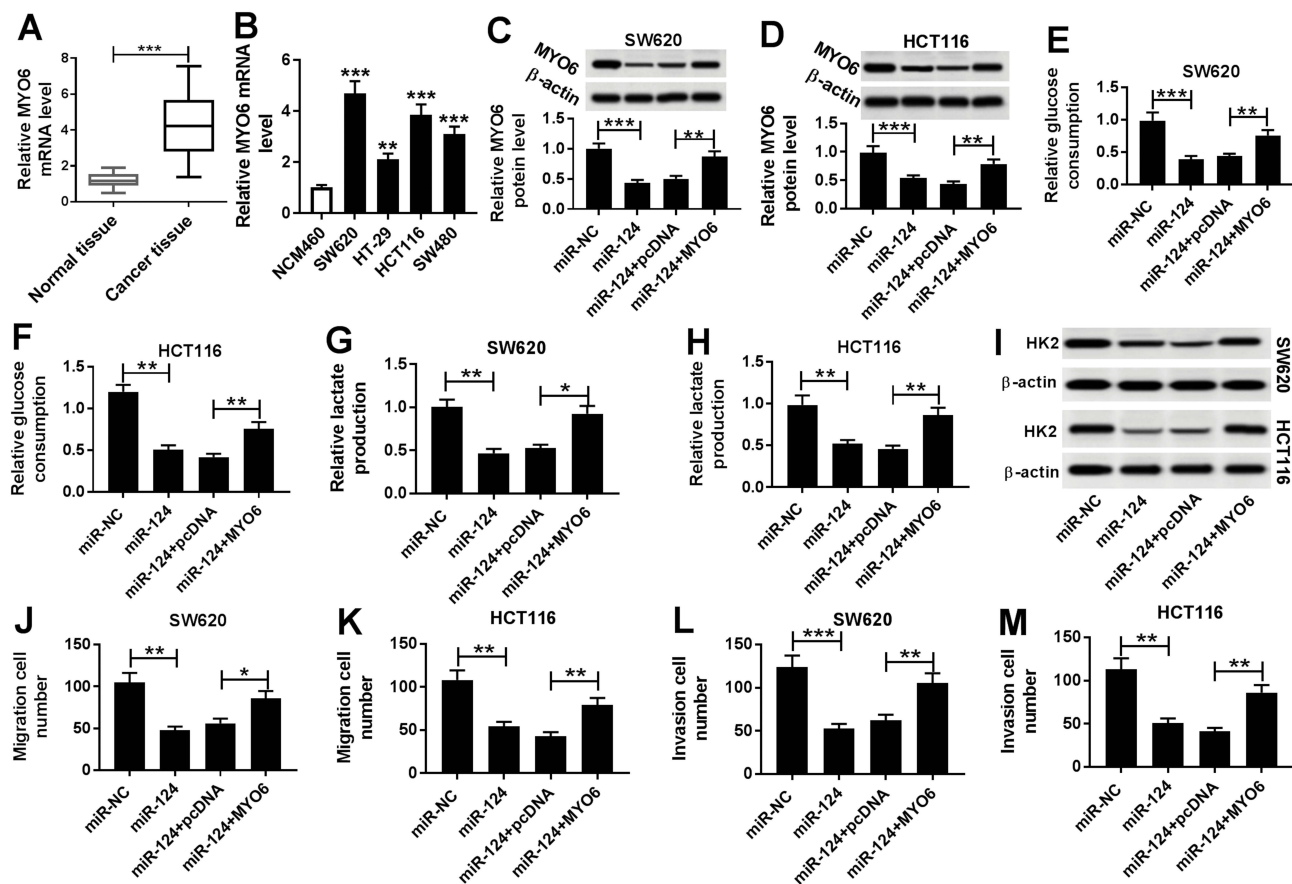


Figure 6 The inhibitory effects of miR-124 overexpression on cell migration, invasion and glycolysis in CRC cells were rescued by upregulated MYO6. (A) The expression of MYO6 was analyzed by RT-qPCR in CRC tissues and normal tissues. (B) The expression of MYO6 was analyzed by RT-qPCR in CRC cells (SW620, HT-29, HCT116 and SW480) and immortalized colon cells (NCM460). (C–M) SW620 and HCT116 cells were transfected with miR-NC, miR-124, miR-124 + pcDNA or miR-124 + MYO6. (C and D) The protein expression of MYO6 in transfected SW620 (C) and HCT116 (D) cells was detected by Western blot assay. (E and F) Glucose consumption was measured in SW620 (E) and HCT116 (F) cells after transfection. (G and H) Lactate production was examined in treated SW620 (G) and HCT116 (H) cells. (I) The expression of HK2 in transfected SW620 and HCT116 cells was analyzed via Western blot assay. (J–M) cell migration and invasion were monitored in transfected SW620 and HCT116 cells by transwell assay. * $P < 0.05$. ** $P < 0.01$. *** $P < 0.001$.

reduced the cell invasion and metastasis abilities of CRC cells. And it was found for the first time that low expression of lncRNA HNF1A-AS1 inhibited intracellular glycolysis in tumor cells. In the research on the regulation mechanism of lncRNA HNF1A-AS1, luciferase reporter assay determined that miR-124 was a target of HNF1A-AS1 and MYO6 was a target gene of miR-124. Therefore, we speculated that lncRNA HNF1A-AS1 affected cell migration, invasion and glycolysis by regulating miR-124/MYO6 axis in CRC cells. Rescue experiments further confirmed our speculation. In this paper, we first put forward lncRNA HNF1A-AS1 regulates MYO6 expression by sponging miR-124 to affect the cell glycolysis process in CRC cells, which significantly improved our understanding of the regulatory mechanisms in CRC.

Cell invasion and migration are necessary biological processes for tumor cells, marking their abilities to migrate

and spread.^{27,28} In clinical research, inhibiting and blocking cell migration and invasion are important processes in the treatment of cancer. Many noncoding RNAs, such as lncRNA and miRNA, as important regulatory factors of cell metabolism, were closely involved in cell progression for various cancers, including invasion and migration.^{29–31} For example, lncRNA UCA1 impacted cell invasion and migration through regulating FOXO3 via sponging miR-96 in pancreatic cancer.³² lncRNA ZFAS1 was upregulated in rheumatoid arthritis and contributed to cell invasion and migration through the reduction of miR-27a.³³ Moreover, lncRNA SNHG16 promoted cell migration, invasion and proliferation through modulating miR-520d-3p/STAT3 axis in hemangioma endothelial cells.³⁴ Lin et al reported that HNF1A-AS2 promoted cell progression and EMT in CRC cells via regulation of miR-129-5p/DNMP3A.³⁵ In our study, we also found that

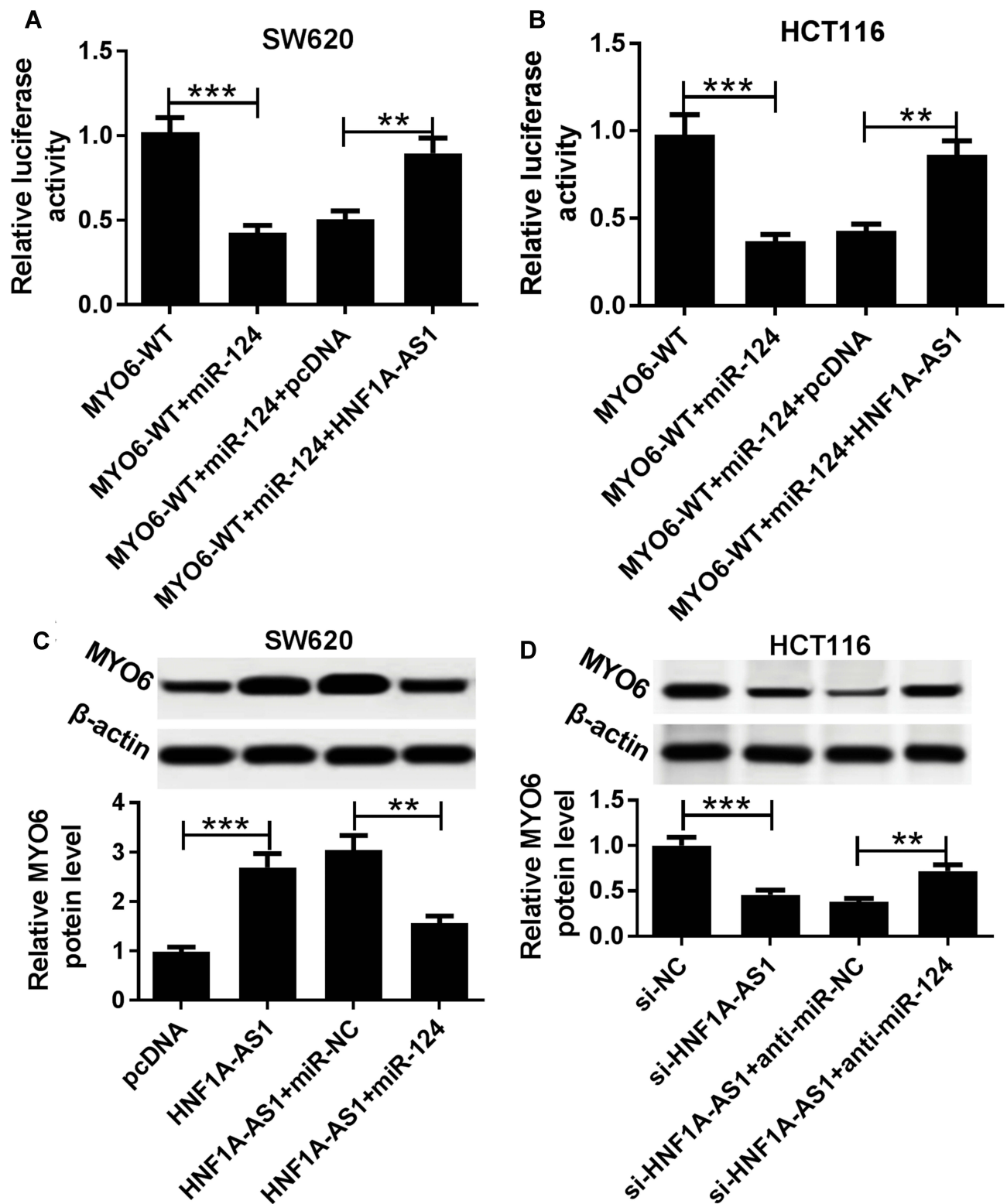


Figure 7 HNF1A-AS1 mediated MYO6 expression through miR-124 in CRC cells. (A and B) Luciferase reporter assay was performed in SW620 (A) and HCT116 (B) cells transfected with MYO6-WT-Luc following introduction with miR-124 or miR-124+ HNF1A-AS1. (C and D) The protein expression of MYO6 was detected in SW620 (C) and HCT116 (D) cells transfected with pcDNA, HNF1A-AS1, HNF1A-AS1+miR-NC and HNF1A-AS1+miR-124 utilizing Western blot assay. ** $p < 0.01$. *** $p < 0.001$.

knockdown of HNF1A-AS1 or promotion of miR-124 inhibited CRC cell migration and invasion. Furthermore, we found that miR-124 was a target of HNF1A-AS1, and MYO6 was a target of miR-124. Accumulating evidence suggested that miR-124 had multiple roles in cell progression of cancers, such as brain tumor and CRC.^{36,37} A previous study determined that miR-124 was downregulated in CRC and inhibited cell proliferation and drug resistance.^{37–39} More than that, MYO6 has been verified to highly express in many cancers, such as prostate cancer, CRC, breast cancer and gastric cancer, which also participated in cell progression.^{40–43} In our study, consistent with a previous study, miR-124 expressed low and MYO6 expressed high in CRC tissues and cells, and rescue experiments confirmed the effects of HNF1A-AS1 on cell migration and invasion through targeting miR-124/MYO6 axis.

More than that, glycolysis provides energy for the metabolism of tumor cells and guarantees the proliferation and development of tumor cells. Lactate production and glucose consumption are important indexes for the determination of glycolysis. Additionally, the glycolytic enzyme hexokinase 2 (HK2) is one of the important aerobic glycolytic mediating factors in cells.^{44–46} Increasing the expression of HK2 could promote cell glycolysis and release more energy for cell development.^{47,48} Various lncRNAs were related to glycolysis in cancers, such as lncRNA Ftx, lncRNA p23154, lncRNA MIF and lncRNA MALAT1.^{49–52} The study of Li et al suggested that lncRNA UCA1 induced glycolysis in bladder cancer cells, and promoted lactate production and glucose consumption through upregulating HK2 via activation of mTOR-STAT3/miR-143 pathway.⁴⁸ In this study, inhibition of HNF1A-AS1 or promotion of miR-124 decreased cell glycolysis. While inhibition of miR-124 or promotion of MYO6 could weaken the suppressive effect of reduction of HNF1A-AS1 or promotion of miR-124 on glycolysis, respectively. Therefore, lncRNA HNF1A-AS1 regulated cell glycolysis in CRC cells through targeting the miR-124/MYO6 axis.

Conclusion

In this study, we first found that HNF1A-AS1 regulated cell migration, invasion and glycolysis via modulating miR-124/MYO6 in CRC cells, providing a new regulatory network and a new therapeutic target of CRC.

Disclosure

The authors report no conflicts of interest in this work.

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