

lncRNA UCA1 Contributes to 5-Fluorouracil Resistance of Colorectal Cancer Cells Through miR-23b-3p/ZNF281 Axis

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Purpose: The chemoresistance of 5-fluorouracil (5-FU) limited the application of chemotherapy in colorectal cancer (CRC) treatment. Herein, we aimed to uncover the potential mechanism behind the 5-FU resistance of CRC cells.

Methods: The abundance of long noncoding RNA urothelial carcinoma associated 1 (lncRNA UCA1), microRNA-23b-3p (miR-23b-3p) and zinc finger protein 281 (ZNF281) was measured by quantitative real-time polymerase chain reaction (qRT-PCR) in CRC tissues and cells. Western blot was conducted to examine autophagy-related proteins, apoptosis-associated proteins and ZNF281 in CRC tissues and cells. Cell counting kit-8 (CCK8) assay was performed to detect the viability and inhibitory concentration 50% (IC50) value of 5-FU of CRC cells. The apoptosis of CRC cells was measured by flow cytometry. The binding sites between miR-23b-3p and UCA1 or ZNF281 were predicted by miRcode and Starbase software, respectively, and the combination was confirmed by dual-luciferase reporter assay and RIP assay. Murine xenograft model was established to verify the role of UCA1 on the 5-FU resistance of CRC in vivo.

Results: The 5-FU resistance of CRC was positively related to the level of UCA1 and autophagy. UCA1 accelerated the 5-FU resistance of CRC cells through facilitating autophagy and suppressing apoptosis. MiR-23b-3p was a target of UCA1 in 293T and CRC cells. The knockdown of miR-23b-3p reversed the inhibitory effects of UCA1 interference on the 5-FU resistance and autophagy and the promoting impact on the apoptosis of CRC cells. ZNF281 could bind to miR-23b-3p in 293T cells. MiR-23b-3p elevated the 5-FU sensitivity through down-regulating ZNF281 in CRC cells. UCA1 interference enhanced the 5-FU sensitivity of CRC through miR-23b-3p/ZNF281 axis in vivo.

Conclusion: UCA1 mediated 5-FU resistance of CRC cells through facilitating autophagy and inhibiting apoptosis via miR-23b-3p/ZNF281 axis in vivo and in vitro.

Keywords: colorectal cancer, 5-FU, UCA1, miR-23b-3p, ZNF281, autophagy, apoptosis

Introduction

Chemotherapy and surgical resection are the primary treatment methods of colorectal cancer (CRC).¹ 5-fluorouracil (5-FU) is a common chemotherapeutic drug for CRC therapy. However, chemoresistance is an enormous challenge for the effective application of chemotherapeutic agents in CRC.

Long noncoding RNAs (lncRNAs) are long-chain RNAs, and they have been reported to regulate the levels of microRNAs (miRNAs) through serving as miRNAs sponges.²⁻⁶ Wang et al demonstrated that lncRNA urothelial carcinoma

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associated 1 (UCA1) facilitated the progression of bladder cancer, suggesting that UCA1 played an oncogenic role in bladder cancer.⁷ Accumulating articles have also confirmed the oncogenic functions of UCA1 in multiple cancers.^{8–11} For instance, Fang et al claimed that UCA1 contributed to multi-drug resistance of gastric cancer through sponging miR-27b.⁹ Han et al reported that the enrichment of UCA1 was positively related to tumor size and depth.¹¹ Besides, Bian et al found that UCA1 facilitated CRC cell proliferation, and it also contributed to 5-FU resistance through miR-204-5p/CREB1.¹² Nevertheless, the precise signal pathway of UCA1 in the chemoresistance of CRC is not fully addressed.

miRNAs participated in the proliferation, metastasis and apoptosis by negatively modulating the levels of their target messenger RNAs (mRNAs) or inhibiting their translation.^{13–16} Accumulating articles reported that miR-23b-3p played an anti-tumor role in a variety of cancers.^{17–20} For example, Kou et al demonstrated that the low expression of miR-23b was associated with the poor prognosis of CRC patients.²⁰ Herein, we aimed to assess the function of miR-23b-3p in the chemoresistance of CRC.

Zinc finger protein 281 (ZNF281) mediates both transcriptional activation and suppression of its target genes.²¹ Hahn et al claimed that ZNF281 promoted the epithelial-mesenchymal transition (EMT), and it was positively regulated by SNAIL and negatively modulated by miR-34a.²² Besides, ZNF281 has been reported to accelerate the proliferation and metastasis of pancreatic cancer cells and CRC cells through Wnt/ β -catenin signal pathway.^{23,24} However, the potential role of ZNF281 in the chemoresistance of CRC remains poorly understood.

In this study, we assessed the role of UCA1 in the chemoresistance of CRC. And then we explored the molecular mechanism by which UCA1 contributed to the 5-FU chemoresistance of CRC cells.

Patients and Methods

Patients

5-FU-resistant CRC patients (n=25) and 5-FU-sensitive CRC patients (n=25) who had not received surgery, chemotherapy or radiotherapy in The Sixth Affiliated Hospital of Sun Yat-sen University were recruited in this study. According to RECIST 1.1 criteria, CRC patients were divided into Sensitive-CRC group (containing Complete Response, Partial Response and Stable Disease) and Resistant-CRC group (Progressive Disease).²⁵ This

research had granted approval by the Ethics Committee of The Sixth Affiliated Hospital of Sun Yat-sen University. Patients whose tissues used in this study had provided written informed consents.

Cell Culture and Drug

Human CRC cell line SW480 and SW620 and human embryonic kidney cell line 293T were obtained from Bena Culture Collection (Beijing, China). All cells were cultivated with Dulbecco's Modified Eagle Medium (DMEM; Gibco, Carlsbad, CA, USA) added with 10% fetal bovine serum (FBS; Gibco) and 10% penicillin (100 U/mL)-streptomycin (100 μ g/mL) mixed solution in a 37°C, 5% CO₂ humidified incubator.

5-FU was obtained from Sigma (St. Louis, MO, USA). The 5-FU-resistant CRC cell lines were built by treating normal SW480 and SW620 cells with increased concentrations (1 μ g/mL, 2 μ g/mL, 5 μ g/mL, 10 μ g/mL, 20 μ g/mL, 40 μ g/mL) of 5-FU. 1 μ g/mL 5-FU was used for the maintenance of drug resistance in SW480/5-FU and SW620/5-FU cells.

Quantitative Real-Time Polymerase Chain Reaction (qRT-PCR)

qRT-PCR was performed through using SYBR Green PCR Kit (Takara, Dalian, China) and the following primers (GeneCopoeia, Rockville, MD, USA). U6 or β -actin acted as the internal reference of UCA1, miR-23b-3p and ZNF281. Relative expression of the above genes was analyzed by 2^{- $\Delta\Delta$ Ct} method.²⁶ The primers were listed as follows: UCA1 (Forward, 5'-CTCTCCATTGGG TTCACCATTC-3'; Reverse, 5'-GCGGCAGGTCTTAAG AGATGAG-3'), miR-23b-3p (Forward, 5'-NBS]? >GAGCATCACATTGCCAGGG-3'; Reverse, 5'-GTGCA GGGTCCGAGGT-3'), ZNF281 (Forward, 5'-GAGG ACACATAGTGGAGAAAAGCC-3'; Reverse, 5'-GTGG ATCTAGACTAGGGCTGGAGTT-3'), U6 (Forward, 5'-CTCGCTTCGGCAGCACA-3'; Reverse, 5'-AACGCT TCACGAATTTGCGT-3'), β -actin (Forward, 5'-TGGAAT CCTGTGGCATCCATGAAAC-3'; Reverse, 5'-ACGCA GCTCAGTAACAGTCCG-3').

Western Blot Assay

The proteins were isolated from CRC cells using RIPA lysis solution (Beyotime, Shanghai, China). The proteins were quantified through using a BCA detecting kit (Beyotime), and then the samples were subjected to

sodium dodecyl sulfate polyacrylamide gel electrophoresis (SDS-PAGE) gel and transferred onto a polyvinylidene fluoride (PVDF) membrane. After incubating with 5% non-fat milk for 1 h, the PVDF membrane was incubated with specific antibodies overnight. The antibodies against Bcl-2 associated X, apoptosis regulator (Bax; ab32503), Caspase-3 (ab13847), light chain 3 I/II (LC3I/II; ab51520), Beclin-1 (ab210498), p62 (ab155686), ZNF281 (ab101318) and β -actin (ab8226) were obtained from Abcam (Cambridge, MA, USA). The membrane was then incubated with a secondary antibody (ab205718, Abcam) for 1 h. The protein images were obtained through the enhanced chemiluminescent (ECL) system (Beyotime).

Cell Transfection

When CRC cells grew to 7080% confluence, transfection was carried out using Lipofectamine 2000 (Invitrogen, Carlsbad, CA, USA). Small interfering RNA negative control (si-NC), siRNA against UCA1 (si-UCA1), siRNA against ZNF281 (si-ZNF281), short hairpin RNA negative control (sh-NC) and shRNA against UCA1 (sh-UCA1) were purchased from Genepharma (Shanghai, China). miRNA-NC (miR-NC), miR-23b-3p, anti-miR-NC and miR-23b-3p inhibitor were obtained from Ribobio (Guangzhou, China).

Cell Counting Kit-8 (CCK8) Assay

CRC cells were cultured in 96-well plates to assess cell viability and inhibitory concentration 50% (IC50) value of 5-FU with Cell Counting Kit-8 (Sigma). Ten microliter CCK8 reagent was added to the wells of 96-well plates, and the CRC cells were incubated with CCK8 reagent for an additional 2 h. The optical density was measured by a microplate reader at 450 nm.

Cell Apoptosis Analysis

CRC cells were harvested using cold phosphate buffer saline (PBS) buffer and re-suspended in binding buffer. The cells were mixed with Annexin V and propidium iodide (PI; Solarbio, Beijing, China) for 10 min. Then the samples were analyzed by the flow cytometer (BD Biosciences, San Jose, CA, USA).

Dual-Luciferase Reporter Assay

miRcode online software was used for predicting the targets of lncRNA UCA1. The miR-23b-3p binding sites in UCA1 sequences (wild-type or mutant type) were embedded into the luciferase reporter vector. The

luciferase activity was analyzed through the dual-luciferase reporter assay system (Promega, Madison, WI, USA). The targets of miR-23b-3p were predicted by Starbase software. The confirmation of the combination between miR-23b-3p and ZNF281 was conducted following the same approach.

RNA Immunoprecipitation (RIP) Assay

RIP buffer (Millipore, Billerica, MA, USA) was used to lyse CRC cells, and the cell extract was incubated with anti-Argonaute-2 (anti-Ago2; Millipore) or anti-Immunoglobulin G (anti-IgG; Millipore) beads. The immunoprecipitated RNAs were identified by qRT-PCR.

Murine Xenograft Assay

This protocol was permitted by the Animal Care and Use committee of The Sixth Affiliated Hospital of Sun Yat-sen University. All animal procedures were performed in accordance with the Guidelines for Care and Use of Laboratory Animals of "National Institutes of Health". Nude mice were obtained from Orient Bio Inc (Seongnam, South Korea). SW480/5-FU cells stably expressing sh-NC or sh-UCA1 were inoculated to the flank of nude mice. When the average tumor size reached about 50 mm³, 30 mg/kg 5-FU was intraperitoneally injected into murine three times a week. The dimension of tumor was measured every 5 d. After 37-d inoculation, the nude mice were sacrificed and the tumors were excised and weighed.

Statistical Analysis

All data were displayed as mean \pm standard deviation (S.D.). Student's *t*-test was used to assess the differences between two groups. The correlation between the expression of miR-23b-3p and UCA1 or ZNF281 in 5-FU resistant-CRC patients was calculated by Spearman correlation coefficient. $P < 0.05$ was considered to be statistically significant.

Results

5-FU Resistance of CRC is Related to the Abundance of UCA1 and Autophagy

To illustrate the chemoresistance mechanism of CRC, we aimed to find crucial molecules involved in 5-FU resistance. As showed in [Figure 1A](#), UCA1 was up-regulated in 5-FU resistant-CRC tissues compared with that in 5-FU sensitive-CRC tissues. Besides, we found that autophagy was promoted in 5-FU resistant-CRC tissues ([Figure 1B](#)). These findings demonstrated that the 5-FU resistance of CRC

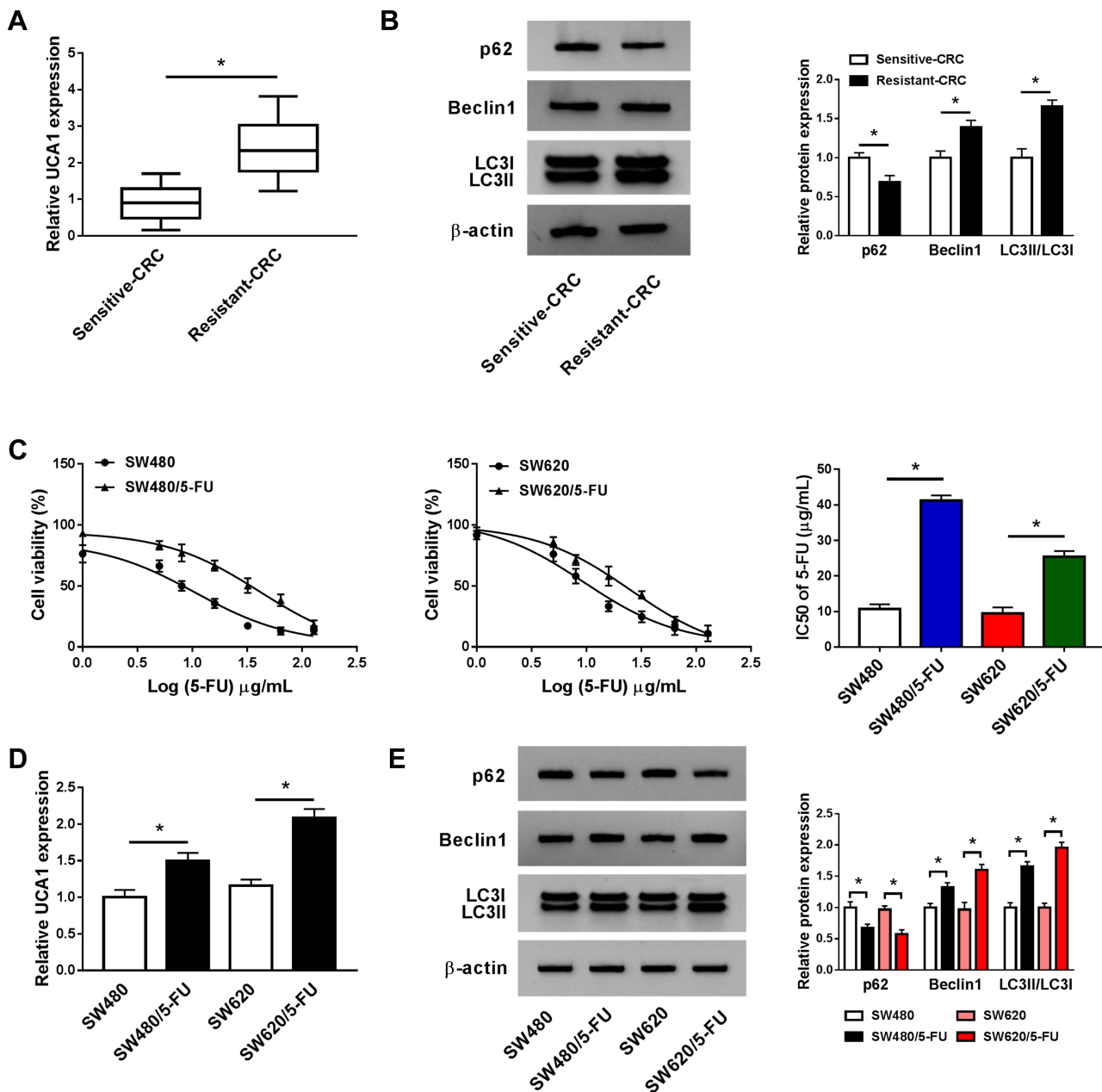


Figure 1 5-FU resistance of CRC is related to the abundance of UCA1 and autophagy. **(A)** The abundance of UCA1 was measured in 5-FU sensitive-CRC and resistant-CRC tissues by qRT-PCR. **(B)** Western blot was conducted to detect autophagy-associated proteins (LC3I/II, Beclin1 and p62) in 5-FU sensitive-CRC and resistant-CRC tissues. **(C)** Cell viability in SW480, SW620 and their corresponding 5-FU-resistant cell lines was assessed by CCK8 assay. **(D)** qRT-PCR was performed to examine the enrichment of UCA1 in SW480, SW620 and their 5-FU-resistant subclones (SW480/5-FU and SW620/5-FU). **(E)** Western blot was carried out to determine autophagy-associated proteins in SW480, SW620, SW480/5-FU and SW620/5-FU cells. * $P < 0.05$.

was associated with the level of UCA and autophagy. We established two 5-FU-resistant CRC cell lines (SW480/5-FU and SW620/5-FU) to further confirm our results. The IC50 values of 5-FU in SW480/5-FU and SW620/5-FU cell lines were notably elevated compared with their parental cell lines (SW480 and SW620) (Figure 1C), suggested that the

establishment of 5-FU-resistant CRC cell lines was successful. As indicated in Figure 1D, the expression of UCA1 was elevated in the two 5-FU-resistant subclones compared with that in their parental cells. Additionally, the autophagy-related proteins (LC3II and Beclin1) were up-regulated in the two 5-FU-resistant subclones (Figure 1E), while p62 was

down-regulated in SW480/5-FU and SW620/5-FU cells, suggesting that autophagy was accelerated in two 5-FU-resistant subclones compared with that in their parental cells.

UCA1 Contributes to 5-FU Resistance of CRC Cells Through Promoting Autophagy and Inhibiting Apoptosis

To explore the role of UCA1 in 5-FU resistance of CRC cells, we conducted loss-of-function experiments. We evaluated the knockdown efficiency of si-UCA1 in SW480/5-FU and SW620/5-FU cells. As mentioned in Figure 2A, the expression of UCA1 in SW480/5-FU and SW620/5-FU was prominently

reduced by the transfection of si-UCA1. As indicated in Figure 2B and C, UCA1 knockdown decreased the viability and IC₅₀ of 5-FU in SW480/5-FU and SW620/5-FU cells, demonstrating that UCA1 depletion elevated the 5-FU sensitivity of CRC cells. To investigate the mechanism by which UCA1 contributing to 5-FU resistance of CRC cells, we performed flow cytometry and Western blot assay. The apoptosis was notably facilitated in si-UCA1 transfected SW480/5-FU and SW620/5-FU cells compared with that in si-NC group (Figure 2D). Besides, Western blot assay showed that the abundance of Bax, Caspase-3 and p62 was enhanced (Figure 2E), while the enrichment of Beclin1 and LC3II was conspicuously decreased, suggesting that UCA1 intervention promoted the

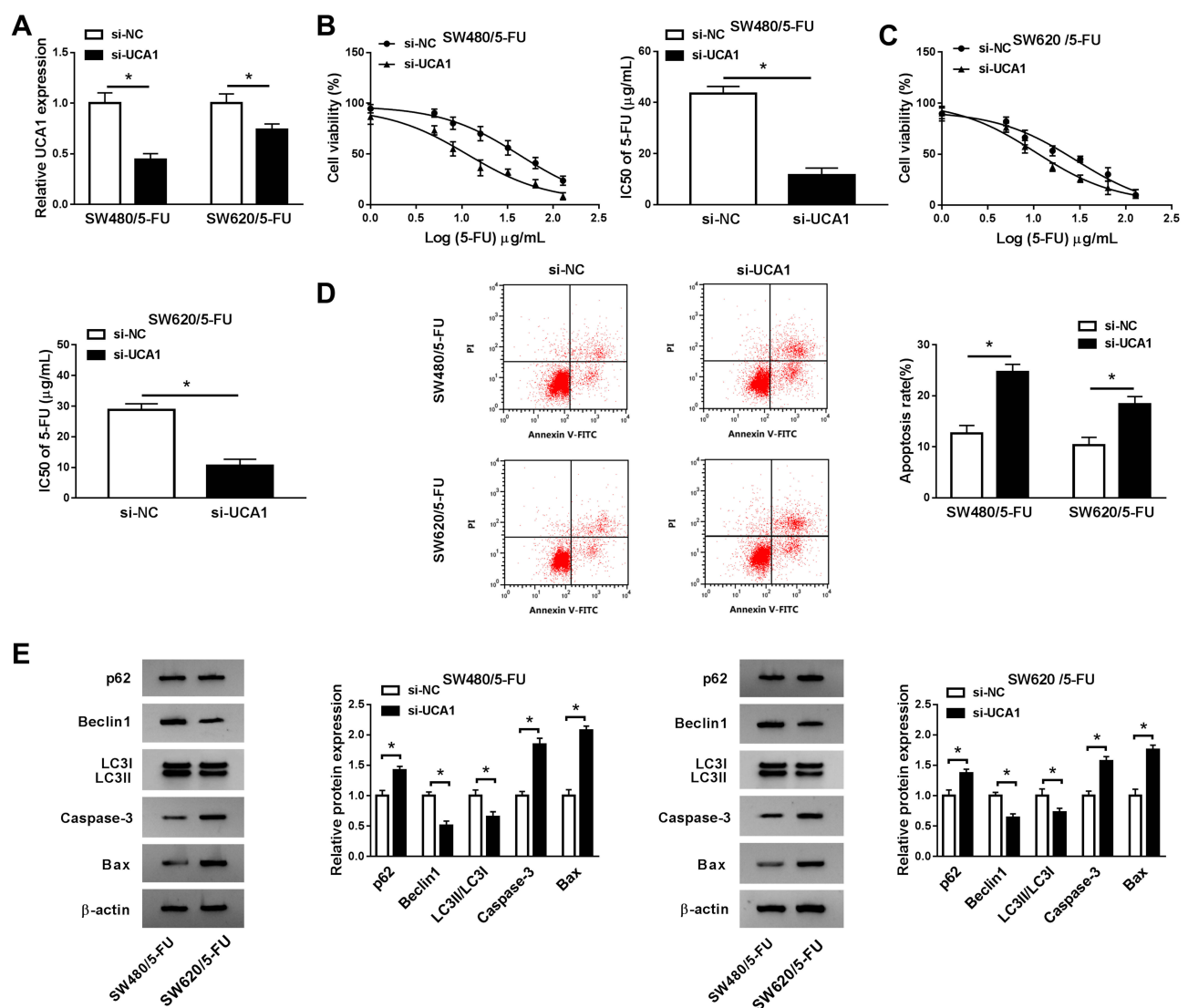


Figure 2 UCA1 contributes to 5-FU resistance of CRC cells through promoting autophagy and inhibiting apoptosis. (A) The level of UCA1 was examined in SW480/5-FU and SW620/5-FU cells transfected with si-NC or si-UCA1 by qRT-PCR. (B and C) CCK8 assay was conducted to measure the viability and IC₅₀ of 5-FU in SW480/5-FU and SW620/5-FU cells transfected with si-NC or si-UCA1. (D) Flow cytometry was applied to detect the apoptosis of si-NC or si-UCA1 transfected SW480/5-FU and SW620/5-FU cells. (E) Western blot was conducted to measure the expression of apoptosis-related proteins (Bax and Caspase-3) and autophagy-associated proteins (LC3I/II, Beclin1 and p62) in SW480/5-FU and SW620/5-FU cells transfected with si-NC or si-UCA1. **P*<0.05.

apoptosis while impeding the autophagy of SW480/5-FU and SW620/5-FU cells. Taken together, UCA1 enhanced the 5-FU resistance of CRC cells through promoting autophagy and restraining apoptosis.

miR-23b-3p Could Bind to UCA1 in CRC Cells

The enrichment of miR-23b-3p was lower in 5-FU resistant-CRC tissues than that in 5-FU sensitive-CRC tissues (Figure 3A). Apart from this, the level of miR-23b-3p was declined in 5-FU-resistant CRC cells (SW480/5-FU and SW620/5-FU) compared with that in SW480 and SW620 cells (Figure 3B). The level of miR-23b-3p was negatively correlated with the abundance of UCA1 in 5-FU resistant-CRC tissues (Figure 3C). miR-23b-3p was predicted as a target of UCA1 by miRcode software, and we confirmed the combination between miR-23b-3p and UCA1 in 293T cells by dual-luciferase reporter assay. Luciferase activity was conspicuously declined in 293T cells co-transfected with miR-23b-3p mimic and UCA1 WT (Figure 3D), while it has little impact in miR-23b-3p mimic and UCA1 MUT group, suggesting that miR-23b-3p was a direct target of UCA1 in 293T cells. Meanwhile, RIP assay also showed that UCA1 could bind to the RNA-induced silencing complex (RISC) in SW480/5-FU and SW620/5-FU cells, likely through the combination with miR-23b-3p (Figure 3E).

UCA1 Promotes the 5-FU Resistance of CRC Cells Through Sponging miR-23b-3p

To clarify the modulatory relationship between UCA1 and miR-23b-3p in CRC cells and whether miR-23b-3p was involved in UCA1-mediated 5-FU resistance of CRC cells, we transfected si-NC, si-UCA1, si-UCA1 + anti-miR-NC or si-UCA1 + miR-23b-3p inhibitor into SW480/5-FU and SW620/5-FU cells. As showed in Figure 4A, the level of miR-23b-3p was dramatically increased by si-UCA1 transfection, and the addition of miR-23b-3p inhibitor reversed this promoting effect, demonstrating that miR-23b-3p was inversely modulated by UCA1 in CRC cells. miR-23b-3p depletion abolished the inhibitory effect of UCA1 interference on the 5-FU resistance of SW480/5-FU and SW620/5-FU cells (Figure 4B and C). Besides, the addition of miR-23b-3p inhibitor counteracted the promoting effect of UCA1 depletion on the apoptosis and the suppressive effect on the autophagy of SW480/5-FU and SW620/5-FU cells (Figure 4D and E). Taken together, UCA1 contributed to 5-FU resistance of CRC cells through negatively regulating miR-23b-3p.

ZNF281 is a Direct Target of miR-23b-3p in CRC Cells

To illustrate the underlying mechanism by which miR-23b-3p elevating the 5-FU sensitivity of CRC cells, we aimed to investigate the downstream component of miR-23b-3p in

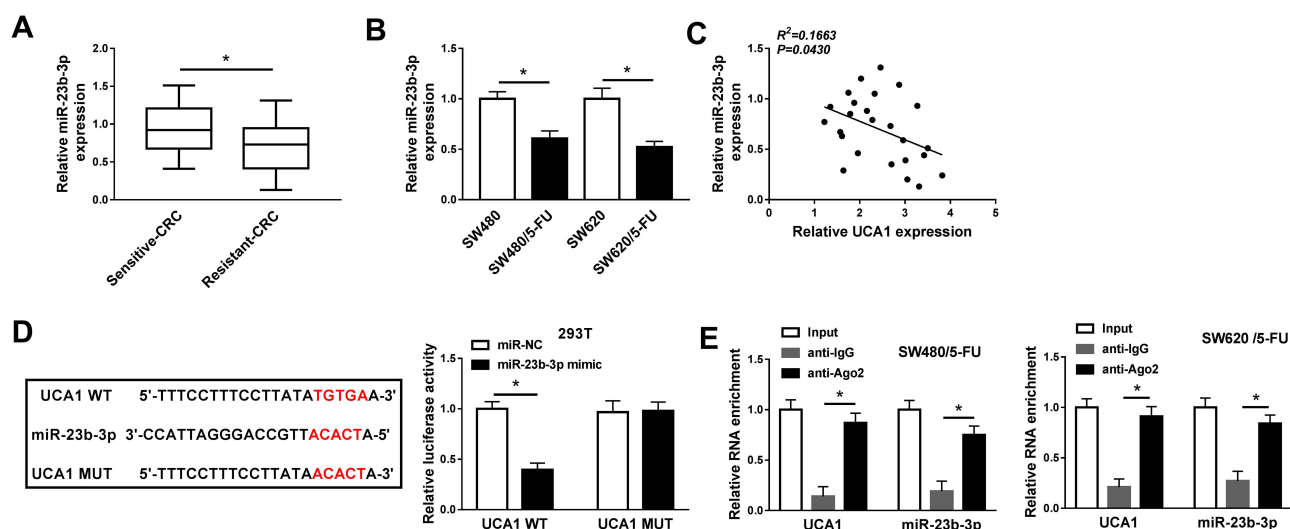


Figure 3 miR-23b-3p could bind to UCA1 in CRC cells. (A) qRT-PCR was applied to examine the abundance of miR-23b-3p in 5-FU sensitive-CRC and resistant-CRC tissues. (B) qRT-PCR was performed to measure the level of miR-23b-3p in SW480, SW620 and their 5-FU-resistant subclones (SW480/5-FU and SW620/5-FU). (C) Spearman correlation coefficient was used to analyze the liner relationship between expression of miR-23b-3p and UCA1 in 5-FU resistant-CRC tissues. (D) The binding sites between miR-23b-3p and UCA1 were predicted by miRcode software, and dual-luciferase reporter assay was applied to confirm this combination in 293T cells. (E) RIP assay was conducted to validate the combination between miR-23b-3p and UCA1 in SW480/5-FU and SW620/5-FU. * $P < 0.05$.

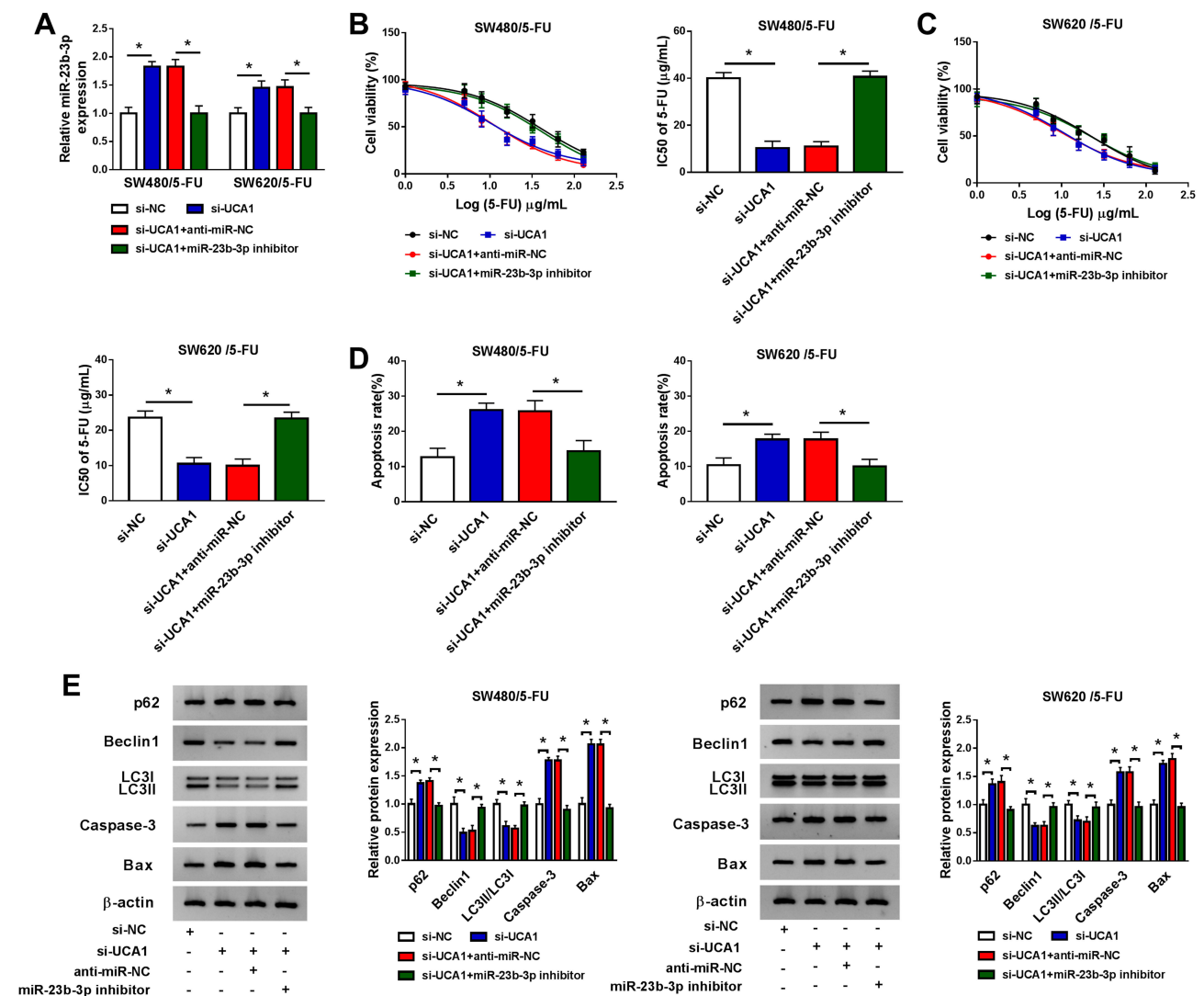


Figure 4 UCA1 promotes the 5-FU resistance of CRC cells through sponging miR-23b-3p. SW480/5-FU and SW620/5-FU cells were transfected with si-NC, si-UCA1, si-UCA1 + anti-miR-NC or si-UCA1 + miR-23b-3p inhibitor. (A) The enrichment of miR-23b-3p was measured in the above two cells by qRT-PCR. (B and C) The viability and IC50 value of 5-FU in the above SW480/5-FU and SW620/5-FU cells were detected by CCK8 assay. (D) The apoptosis of the above two cells was detected by flow cytometry. (E) Western blot was performed to measure the apoptosis-associated proteins and autophagy-related proteins in the above two cells. * $P < 0.05$.

CRC cells. As showed in Figure 5A and B, the mRNA and protein expression of ZNF281 was markedly up-regulated in 5-FU resistant-CRC tissues. Meanwhile, the abundance of ZNF281 mRNA and protein was notably increased in 5-FU-resistant CRC cells compared with that in their matching parental CRC cells (Figure 5C and D). As mentioned above, the expression of miR-23b-3p was lower in 5-FU resistant-CRC tissues, and we proposed the hypothesis that there was a significantly negative correlation between the levels of ZNF281 and miR-23b-3p in 5-FU resistant-CRC tissues. Subsequently, the result of the correlation analysis supported our hypothesis (Figure 5E). ZNF281 was predicted as a target of miR-23b-3p by Starbase software, and then dual-luciferase

reporter assay verified the combination between ZNF281 and miR-23b-3p in 293T cells (Figure 5F). The luciferase activity was markedly reduced with the accumulation of miR-23b-3p in ZNF281 3' UTR-WT group compared with that in ZNF281 3' UTR-MUT group, suggesting that ZNF281 was a direct target of miR-23b-3p in 293T cells.

miR-23b-3p Elevates the 5-FU Sensitivity Through Negatively Regulating ZNF281 in CRC Cells

To explore whether ZNF281 could regulate the 5-FU resistance of CRC cells through serving as a downstream gene of miR-23b-3p, we transfected si-NC, si-ZNF281, si-ZNF281 +

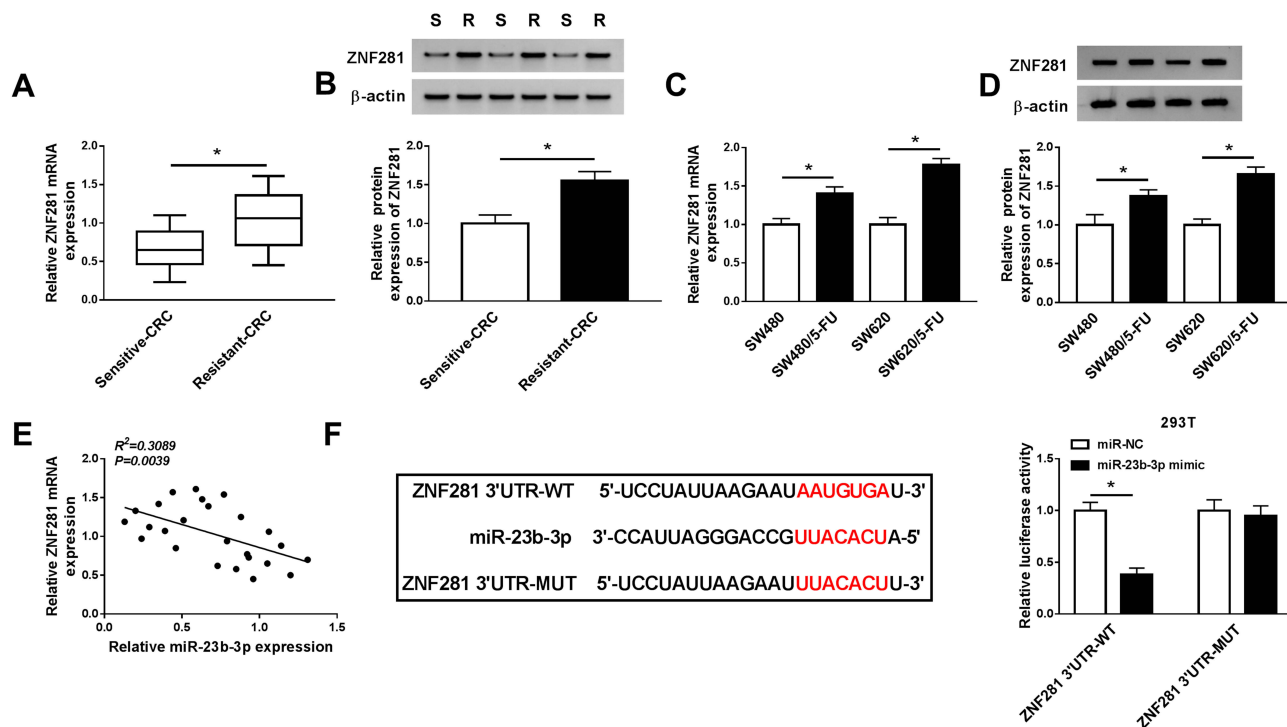


Figure 5 ZNF281 is a direct target of miR-23b-3p in CRC cells. (A and B) The abundance of ZNF281 mRNA and protein was determined in 5-FU sensitive-CRC or resistant-CRC tissues by qRT-PCR and Western blot. (C and D) qRT-PCR and Western blot were performed to measure the mRNA and protein expression of ZNF281 in SW480, SW620, SW480/5-FU and SW620/5-FU cells. (E) The correlation between the expression of ZNF281 and miR-23b-3p was analyzed in 5-FU resistance-CRC tissues. (F) ZNF-281 was predicted as a target of miR-23b-3p by Starbase software, and this combination was verified by dual-luciferase reporter assay. * $P < 0.05$.

anti-miR-NC or si-ZNF281 + miR-23b-3p inhibitor into SW480/5-FU and SW620/5-FU cells. As mentioned in Figure 6A, the transfection of si-ZNF281 down-regulated the abundance of ZNF281, and the co-transfection of si-ZNF281 and miR-23b-3p inhibitor recovered the level of ZNF281 in SW480/5-FU and SW620/5-FU cells. The viability of SW480/5-FU and SW620/5-FU cells and IC₅₀ value of 5-FU were down-regulated by si-ZNF281 transfection, and the addition of miR-23b-3p inhibitor recovered the viability and IC₅₀ value of 5-FU in the two CRC cells (Figure 6B and C). Flow cytometry revealed that miR-23b-3p depletion abated the promoting effect of si-ZNF281 transfection on the apoptosis of SW480/5-FU and SW620/5-FU cells (Figure 6D). Moreover, we conducted Western blot assay to detect the proteins-associated with apoptosis and autophagy. As indicated in Figure 6E, ZNF281 interference elevated the abundance of Caspase-3, Bax and p62 and declined the expression of Beclin1 and LC3II, and the addition of miR-23b-3p inhibitor reversed the effects caused by ZNF281 depletion, suggesting that the depletion of miR-23b-3p diminished the promoting effect of ZNF281 silencing on the apoptosis and the inhibitory impact on the autophagy of CRC cells. These findings revealed that ZNF281 functioned as a downstream gene of miR-23b-3p to contribute to the 5-FU resistance of CRC cells.

ZNF281 is Modulated by UCA1/miR-23b-3p Axis in CRC Cells

To further explore the modulatory relationship among UCA1, miR-23b-3p and ZNF281 in SW480/5-FU and SW620/5-FU cells, we conducted the following experiments. SW480/5-FU and SW620/5-FU cells were transfected with si-NC, si-UCA1, si-UCA1 + anti-miR-NC or si-UCA1 + miR-23b-3p inhibitor. As showed in Figure 7A and B, UCA1 knockdown decreased the mRNA level of ZNF281 in SW480/5-FU and SW620/5-FU cells, and the addition of miR-23b-3p inhibitor recovered the abundance of ZNF281 mRNA. Meanwhile, the change of ZNF281 protein revealed the same trend to the change of ZNF281 mRNA in the above SW480/5-FU and SW620/5-FU cells (Figure 7C and D). Taken together, the level of ZNF281 was regulated by UCA1/miR-23b-3p signaling in CRC cells.

UCA1 Contributes to 5-FU Resistance of CRC in vivo

To validate the biological significance of UCA1 on the 5-FU resistance of CRC in vivo, we established the murine xenograft model using SW480/5-FU cells stably transfected with sh-NC or sh-UCA1. As mentioned in Figure 8A and B,

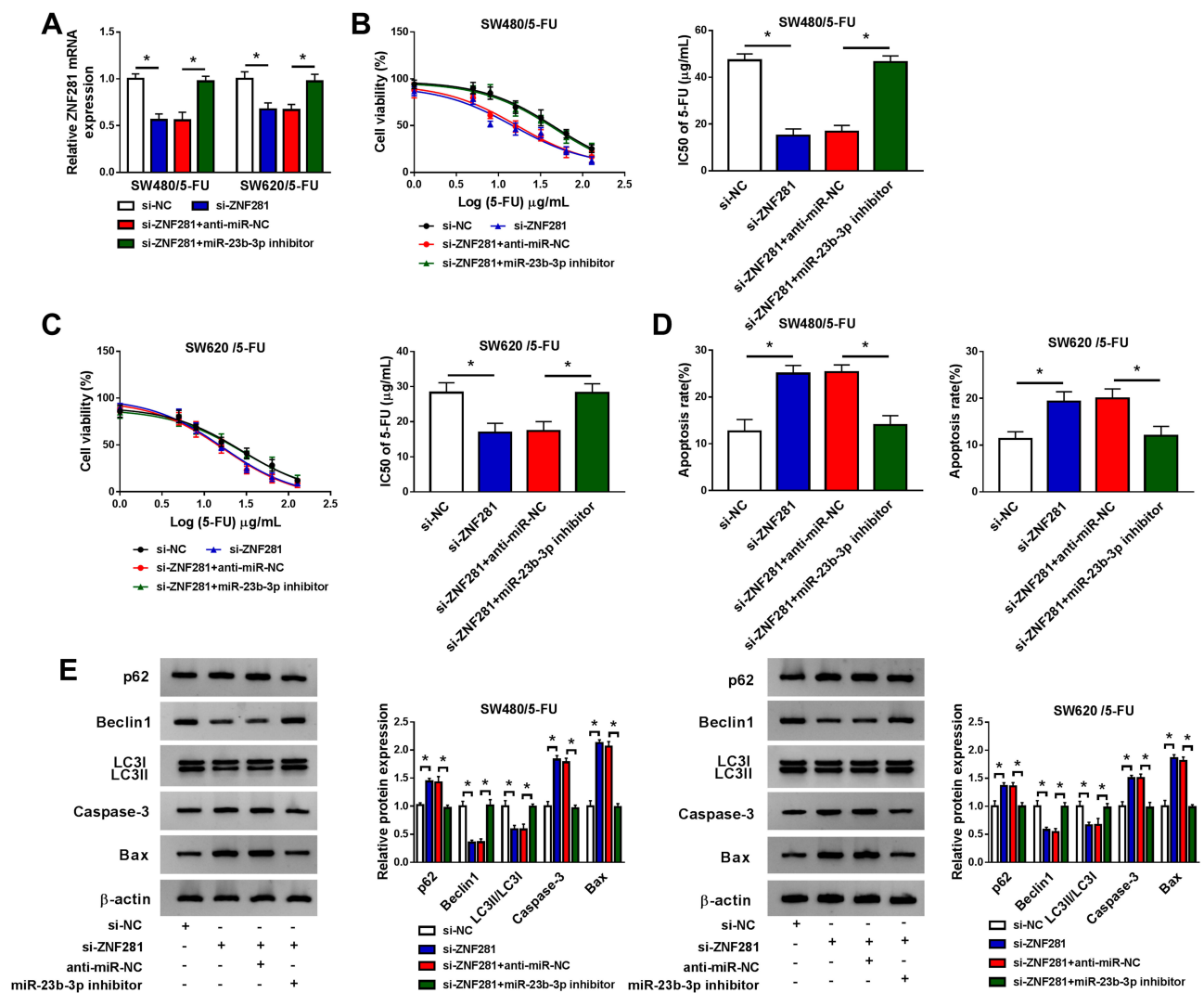


Figure 6 miR-23b-3p elevates the 5-FU sensitivity through negatively regulating ZNF281 in CRC cells. SW480/5-FU and SW620/5-FU cells were transfected with si-NC, si-ZNF281, si-ZNF281 + anti-miR-NC or si-ZNF281 + miR-23b-3p inhibitor. (A) The enrichment of ZNF281 was examined in the above two cells by qRT-PCR. (B and C) CCK8 assay was applied to measure the viability and IC50 value of 5-FU in the above SW480/5-FU and SW620/5-FU cells. (D) Flow cytometry was performed to detect the apoptosis of the above two cells. (E) The expression of apoptosis-related proteins and autophagy-associated proteins in the above two cells was determined by Western blot assay. * $P < 0.05$.

tumor volume and weight were less in sh-UCA1 + 5-FU group than that in sh-NC + 5-FU group, suggesting that UCA1 depletion elevated the 5-FU sensitivity of CRC in vivo. The abundance of UCA1 was notably decreased in sh-UCA1 + 5-FU group (Figure 8C). UCA1 intervention enhanced the level of miR-23b-3p. Meanwhile, the abundance of ZNF281 mRNA and protein was prominently reduced in sh-UCA1 + 5-FU group compared with that in sh-NC + 5-FU group (Figure 8C and D). Additionally, proapoptotic proteins (Bax and Caspase-3) and p62 were up-regulated while the enrichment of Beclin1 and LC3II was conspicuous declined in sh-UCA1 + 5-FU group compared with that in sh-NC + 5-FU group (Figure 8E). Taken

together, UCA1 promoted the 5-FU resistance of CRC through accelerating autophagy and inhibiting apoptosis via miR-23b-3p/ZNF281 axis in vivo.

Discussion

Accumulating studies have revealed the crucial role of UCA1 in multiple cancers, including bladder cancer, CRC and HCC.^{11,27,28} Han et al found that high expression of UCA1 was closely correlated with poor prognosis of CRC patients, and UCA1 was also found to accelerate the proliferation and cell cycle and suppress the apoptosis of CRC cells.¹¹ UCA1 was also reported to promote the development of HCC via suppressing miR-216b and

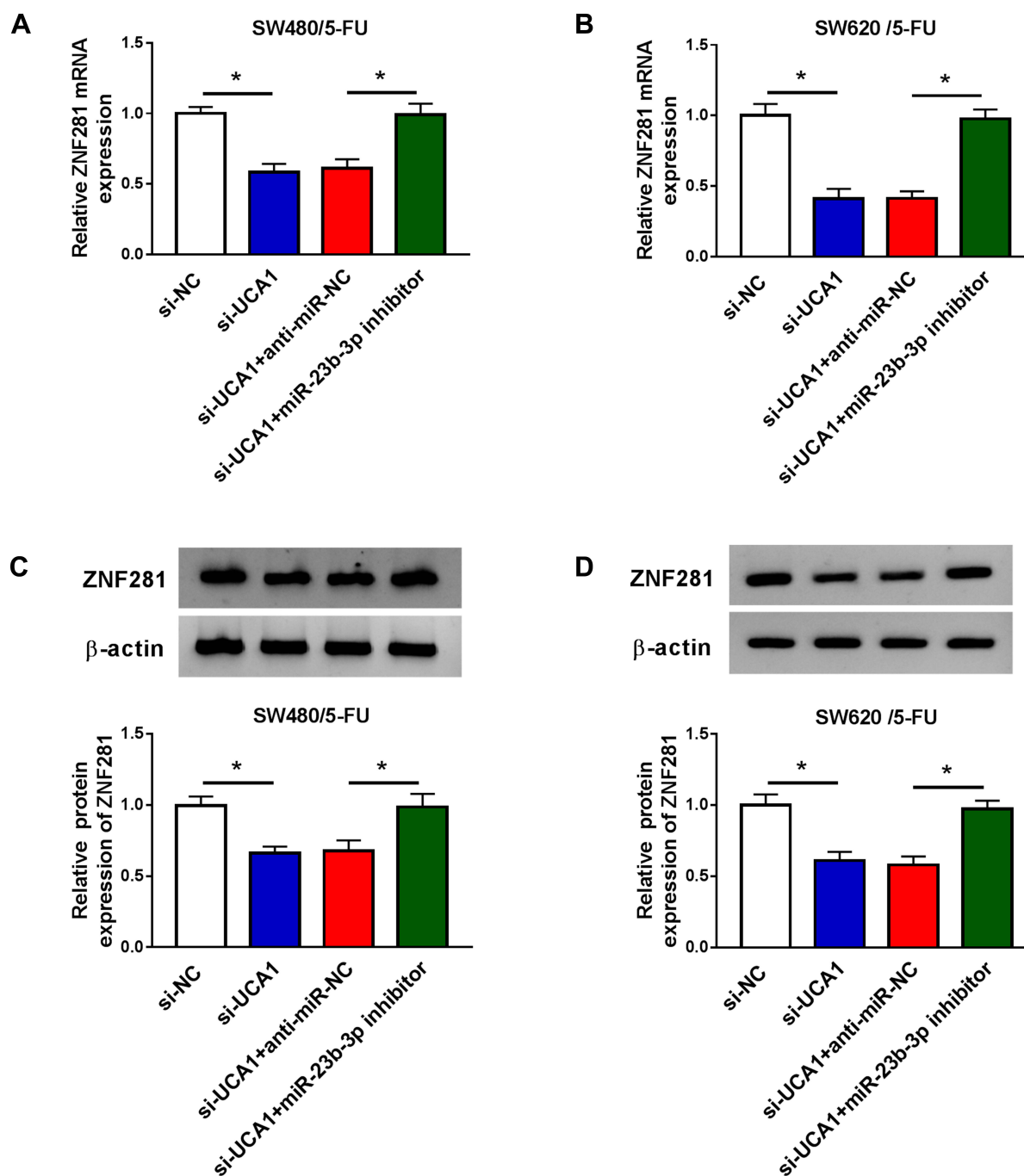


Figure 7 ZNF281 is modulated by UCA1/miR-23b-3p axis in CRC cells. (A and B) qRT-PCR was carried out to measure the expression of ZNF281 in SW480/5-FU and SW620/5-FU cells transfected with si-NC, si-UCA1, si-UCA1 + anti-miR-NC or si-UCA1 + miR-23b-3p inhibitor. (C and D) Western blot assay was performed to detect the protein expression of ZNF281 in si-NC, si-UCA1, si-UCA1 + anti-miR-NC or si-UCA1 + miR-23b-3p inhibitor transfected SW480/5-FU and SW620/5-FU cells. * $p < 0.05$.

activating FGFR1/ERK signal axis.²⁸ Furthermore, accumulating works have pointed out the functional association between UCA1 and chemoresistance in many cancers, including CRC.^{12,29,30} Zhang et al demonstrated that UCA1 enhanced the adriamycin (ADR) resistance of pediatric acute myeloid leukemia (AML) through

promoting glycolysis via regulating miR-125a/hexokinase 2 signaling.³⁰ Sun et al demonstrated that UCA1, CRNDE, H19 and HOTAIR were associated with Oxaliplatin or Irinotecan resistance in CRC cells according to the results of integrative bioinformatics analysis.³¹ Bian et al reported that UCA1 elevated the 5-FU resistance and proliferation

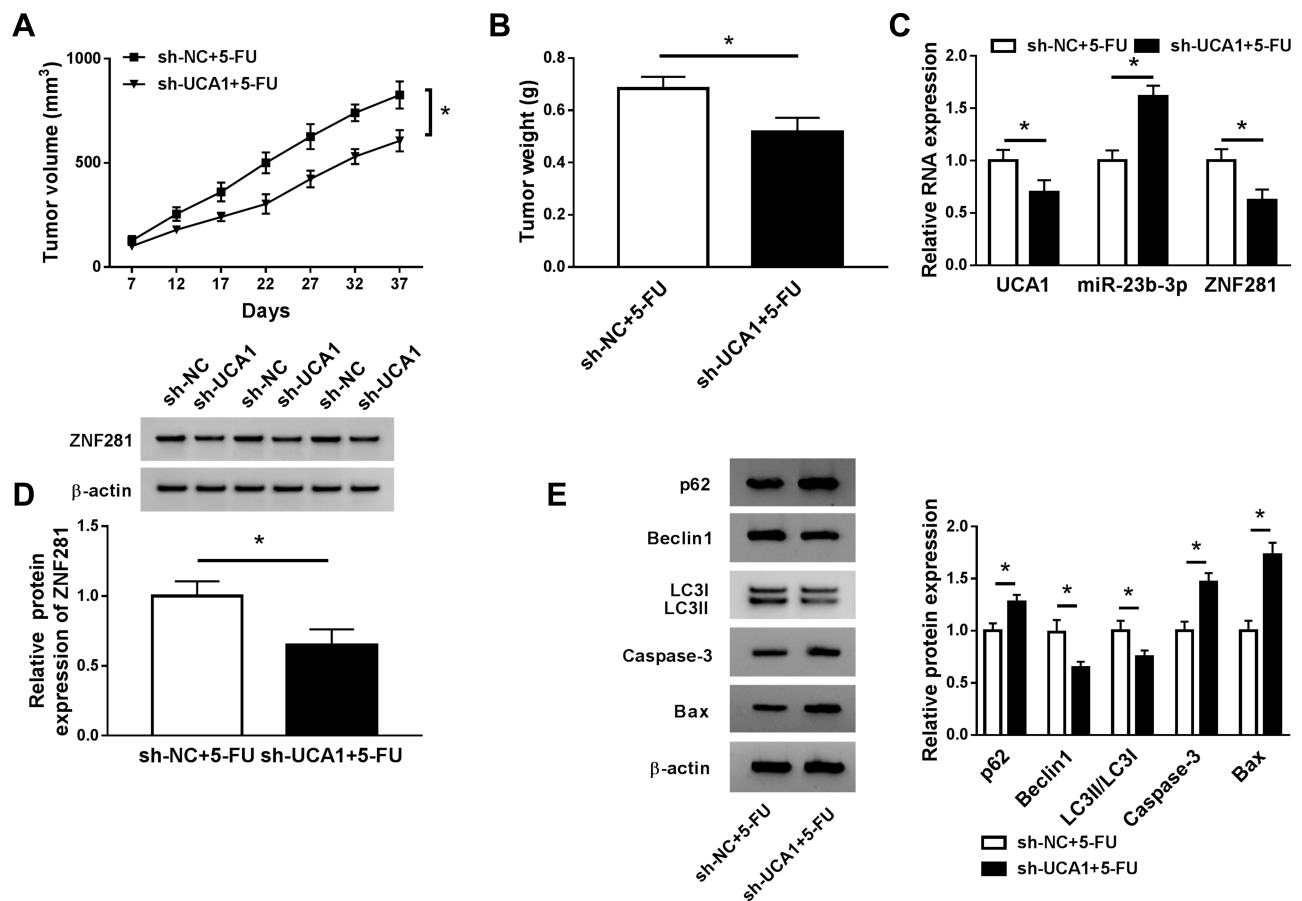


Figure 8 UCA1 contributes to 5-FU resistance of CRC in vivo. **(A)** Tumor volume was recorded every 5 d. **(B)** Tumor was weighed after 37-day inoculation. **(C)** qRT-PCR was conducted to measure the expression of UCA1, miR-23b-3p and ZNF281 in resected tumor tissues. **(D)** The protein expression of ZNF281 was determined in resected tumor tissues by Western blot assay. **(E)** Western blot assay was applied to measure the abundance of Bax, Caspase-3, LC3I/II, Beclin1 and p62 in resected tumor tissues. * $P < 0.05$.

of CRC cells through sponging miR-204-5p.¹² Herein, we found that the abundance of UCA1 was enhanced in 5-FU resistant-CRC tissues and cells, suggesting that UCA1 played a pivotal role in the chemoresistance of CRC. Besides, autophagy was enhanced in 5-FU resistant-CRC tissues and cells. Autophagy has been reported to accelerate chemoresistance and the viability of cancer cells under diverse stresses.^{32–34} lncRNA H19 was found to enhance 5-FU resistance of CRC cells through accelerating SIRT1-mediated autophagy.³² Zhang et al demonstrated that miR-22 elevated the sensitivity of CRC cells to 5-FU through promoting apoptosis and restraining the autophagy.³³ We proposed the hypothesis that UCA1 contributed to the 5-FU resistance of CRC through promoting autophagy.

We conducted knockdown experiments to assess the function of UCA1 in the 5-FU resistance of CRC cells. UCA1 depletion inhibited the viability of CRC cells treated with different concentrations of 5-FU compared with

that in the control group. And the IC₅₀ value was notably reduced by si-UCA1 transfection in CRC cells. Besides, UCA1 interference also accelerated the apoptosis and inhibited the autophagy of CRC cells. These findings suggested that UCA1 facilitated the 5-FU resistance of CRC cells through promoting autophagy and inhibiting apoptosis.

The target relationship between UCA1 and miR-23b-3p was predicted by miRcode software, and we confirmed the combination between UCA1 and miR-23b-3p in 293T cells and CRC cells by dual-luciferase reporter assay and RIP assay. MiR-23b-3p played an anti-tumor role in diverse cancers.^{20,35,36} For instance, miR-23b-3p hampered the proliferation and motility of cervical cancer cells through targeting c-Met.³⁵ Kou et al reported that low abundance of miR-23b in plasma of CRC patients was related to the poor prognosis.²⁰ Besides, Hu et al found that miR-23b-3p acted as the downstream target of MALAT1 to suppress autophagy-related chemoresistance.³⁷ The knockdown of miR-23b-3p reversed

the inhibitory effects of UCA1 depletion on the 5-FU resistance and autophagy and the promoting impact on the apoptosis of CRC cells, suggesting that miR-23b-3p functioned as a downstream component of UCA1 to elevate the 5-FU sensitivity of CRC cells.

ZNF281 has been reported to play an oncogenic role in non-small-cell lung cancer and CRC.^{24,38} ZNF281 promoted the proliferation and metastasis of CRC cells through regulating Wnt/ β -Catenin signal pathway.²⁴ Consistent with the above findings, we found that ZNF281 contributed to the 5-FU resistance of CRC cells. We found that the mRNA and protein expression of ZNF281 was conspicuously elevated in 5-FU resistant-CRC tissues and cells. Starbase software predicted that ZNF281 was a target of miR-23b-3p, and we verified the combination between ZNF281 and miR-23b-3p by dual-luciferase reporter assay. The addition of miR-23b-3p inhibitor counteracted the suppressive effects of ZNF281 depletion on the 5-FU resistance, viability and autophagy, and the promoting impact on the apoptosis of CRC cells. To further clarify the modulatory relationship between UCA1, miR-23b-3p and ZNF281 in CRC cells, we conducted the following experiments. The level of ZNF281 was declined with UCA1 depletion, and the addition of miR-23b-3p inhibitor recovered the expression of ZNF281 in SW480/5-FU and SW620/5-FU cells, demonstrating that ZNF281 was modulated by UCA1/miR-23b-3p axis in CRC cells.

We built the murine xenograft model using SW480/5-FU cells stably transfected with sh-UCA1 or sh-NC to assess the function of UCA1 in the 5-FU resistance of CRC in vivo. UCA1 depletion enhanced the 5-FU sensitivity of CRC tumor through promoting apoptosis and suppressing autophagy via miR-23b-3p/ZNF281 signaling in vivo.

In conclusion, UCA1/miR-23b-3p/ZNF281 axis was identified in CRC cells for the first time in this study. UCA1 contributed to the 5-FU resistance of CRC through promoting autophagy and suppressing apoptosis via miR-23b-3p/ZNF281 axis. Restoration of miR-23b-3p level might be an effective therapeutic strategy for CRC.

Disclosure

The authors declare that they have no conflicts of interest in this work.

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