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Force, Hangzhou 310021, Zhejiang Province, People's Republic of China ORIGINAL RESEARCH

The microRNA miR-3174 Suppresses the Expression of ADAM15 and Inhibits the Proliferation of Patient-Derived Bladder Cancer Cells

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Background: Bladder cancer is a major urinary system cancer, and its mechanism of action regarding its progression is unclear. The goal of this study was to examine the expression of ADAM panel in the clinical specimens of bladder cancer and to investigate the role of miR-3174/ADAM15 (a disintegrin and metalloprotease 15) axis in the regulation of bladder cancer cell proliferation.

Methods: The expression of an ADAM gene panel (including ADAM8, 9, 10, 11, 12, 15, 17, 19, 22, 23, 28, and 33), including 30 pairs of bladder tumor and non-tumor specimens, was examined by Ion AmpliSeq Targeted Sequencing. A microRNA (miRNA) that could potentially target the ADAM with the highest expression level in the tumor tissue was identified using the online tool miRDB. Next, the interaction between the miRNA and ADAM15 was identified by Western blot. Finally, the proliferation of bladder cancer cells was examined using MTT (3-(4,5-dimethyl-2-thiazolyl)-2,5-diphenyl-2-H-tetrazolium bromide) experiments (cell proliferation examining) and subcutaneous tumor models by using nude mice.

Results: The expression of ADAM15 in tumor tissue was found statistically significant when compared to its expression in non-tumor tissue. Additionally, ADAM15's expression in tumor tissue was found the highest of all other tested ADAMs. Next, by using the online tool miRDB, a microRNA termed miR-3174 was identified that targets ADAM15 and inhibits its expression by binding to its 3'-untranslated region. Finally, we found that overexpression of miR-3174 in bladder cancer cells inhibited the proliferation of cells due to the inhibition of ADAM15.

Conclusion: In the present work, the data highlight that miR-3174 inhibits the proliferation of bladder cancer cells by targeting ADAM15.

Keywords: ADAM, bladder cancer, metastasis, microRNA, proliferation

Introduction

Bladder cancer is one of the major urinary system cancers; however, the details of its mechanisms, such as cell proliferation and metastasis, remain scarce.¹⁻³ ADAMs (a disintegrin and metalloproteases), which belong to a protein family of de-integrating metalloproteinases, are type I transmembrane proteins (zinc-containing proteases) and play important roles in regulating cell-cell and cell-matrix interactions.⁴ So far, more than 30 types of ADAMs have been identified.^{4,5} The structure of an ADAM is featured by a conserved RGD (Arg-Gly-Asp) motif in its integrin domain that regulates cell-cell adhesion.^{5,6} ADAMs have also been found to regulate various physical processes of the cell, including proteolysis, cell fusion, migration, signal transduction, and release of

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biological activity factors.⁶ Recently, ADAMs have started to be considered as important regulators of human cancers, including lung, gastric, colon, breast, and pancreatic cancer.⁷ In the present study, we demonstrate that among 12 types of ADAMs, the expression of ADAM15 was found the highest in bladder cancer tissue. Here, miR-3174 was found as a microRNA that targets ADAM15. Moreover, the expression of miR-3174 was found to inhibit the expression of ADAM15 in bladder cancer tissue. As a result, overexpression of miR-3174 inhibited the cell proliferation, metastasis, and subcutaneous growth in vivo of patientderived bladder cancer cells.

Materials and Methods

Patient-Derived Bladder Cancer Cell Lines The paired tissues (the bladder cancer tissues or the non-tumor tissues obtained in the same patient) of bladder tumor and nontumor specimens were obtained by surgery. The collection of clinical specimens was in compliance with the Declaration of Helsinki, and the pathological stages (Class 0, A, B, C and D) of bladder cancer tissues were measured using the Jewett-Marshall methods.^{8–10} Furthermore, this study and its applied methods were approved by the ethics committee of the Beijing Rehabilitation Hospital, which is affiliated to the Capital Medical University (Beijing, China). Written informed consent was obtained from all the patients. The surgically resected bladder cancer specimens were preserved in Dulbecco's Modified Eagle Medium (DMEM, Thermo Fisher Scientific, Waltham, MA, USA) supplemented with 20% fetal bovine serum (FBS, Thermo Fisher Scientific). Then, to obtain a patient-derived tumor cell line, the tissue was ground with a 200-mesh steel sieve, and the cell suspension washed with PBS. Next, the single cells derived from clinical specimens of bladder cancer tissues were cultured in DMEM with 10% FBS. There were five lines of PDCs including in the present work, and the PDCs included two lines of Class C (No. 1 and No. 2) and three lines of Class D (No. 3, 4 and 5).

Ion AmpliSeq Targeted Sequencing

Amplification primers for the ADAM8, ADAM9, ADAM10, ADAM11, ADAM12, ADAM15, ADAM17, ADAM19, ADAM22, ADAM23, ADAM28, and ADAM33 genes were designed using the Ion AmpliSeq Designer tool from Thermo Fisher Scientific.¹¹ Before use, the primers were mixed together into one primer pool. For testing samples, the total RNA was extracted from each bladder cancer cell line using an RNA-extraction Kit (Applied Biosystems instruments, from

Thermo Fisher Scientific). These samples were then subjected to reverse transcription by using the RNeasy Mini Kit (Qiagen, Valencia, CA, USA). Next, the library construction of cDNA was performed using the Ion AmpliSeq Library Kit and the Ion Xpress Barcode Adapter reagent (Life Technologies, USA). The DNA targets-including the ADAM gene panel —were amplified using a 2 \times primer pool buffer with 22 cycles' amplify (Ion PGM Hi-Q View Sequencing Kit, Catalog number: A30044, Life Technologiesy Corporation, USA). The Ion PGM Hi-Q sequencing kit and the Ion Torrent 318 chip (Ion 318 Chip Kit v2 BC, Catalog number: 4488150, Life Technologiesy Corporation, USA) were used for sequencing on the Ion Torrent PGM platform (Ion Personal Genome Machine [PGM] System, Catalog number: 4462921, Life Technologiesy Corporation, USA). Finally, the Ion Torrent data were analyzed using the Ion Torrent Suite v3.0 software (Supplemental Table 1).

Quantitative Polymerase Chain Reaction (qPCR)

Total RNA samples were extracted from bladder cancer cells by using an RNA-extraction Kit (Applied Biosystems Instruments, from Thermo Fisher Scientific), and the RNA samples were reverse transcribed into cDNA by using an RNeasy Mini Kit (Qiagen) according to the manufacturer's instructions. The kit (miRNA examination kit) to examine the expression level of miR-3174 was purchased from Thermo Fisher Scientific. Next, the qPCR experiments were performed following the manufacturer's instructions (Applied Biosystems Instruments, from Thermo Fisher Scientific) and the primers from Table 1. The level of GAPDH mRNA was used as an internal control. The expression of ADAM15 and miR-3174 in bladder cancer

 Table I
 The Inhibitory Rates of miR-3174 on Five Patient-Derived Bladder Cancer Cells' Proliferation

PDCs	Control mi RNA	mi R -3174	miR-3174 + inhibitor	miR-3174 + ADAM15 ^{mut}
	Inhibitory Rates on the Proliferation of Bladder Cancer Cells (%)			ladder Cancer
No. I	-	75.33±5.25	25.42±2.88	5.45±0.56
No. 2	-	61.47±3.33	14.22±1.78	3.21±0.44
No. 3	-	70.92±4.05	22.36±3.01	6.35±0.73
No. 4	-	57.44±4.55	10.21±1.11	4.67±0.88
No. 5	-	66.67±3.33	16.81±2.23	5.26±0.75

Abbreviations: PDCs, patient-derived cells included two lines of Class C (No. I and 2) and three lines of Class D (No. 3–5); ADAM15^{Mut}, ADAM15 with the mutated miR-3174's targeting sites.

specimens was processed into a scatter plot image: The x-axis represented the mRNA level of ADAM15 and the y-axis the miR-3174 level, both from the same sample.

Plasmids and Lentivirus Particles

The expression vectors containing full-length sequences of pri-muR-3174 and ADAM15 were purchased from Vigene Biosciences (Jinan City, Shandong Province, China). The lentivirus particles containing full-length sequences of pri-muR-3174, wild type ADAM15, and ADAM15 with mutated miR-3174 binding sites were also prepared by Vigene.

Cell Culture and Proliferation

The patient-derived bladder cells were cultured in DMEM (Thermo Fisher Scientific) supplemented with 10% FBS (Thermo Fisher Scientific). The cells were treated with lentivirus particles to introduce the expression of miR-3174 or ADAM15. Then, for the proliferation experiments, 2000 cells were seeded per well into a 96-well cell culture plate. After 3 days of culture, the cells were analyzed by MTT analysis following the manufacturer's instructions (Ameresco, USA). The inhibition rates were calculated as [(O.D. [optical density] 490 nm of the control group at day 3 of culture -O.D. 490 nm of the control group at day 0 of culture) – (O.D. 490 nm of the administration group at day 3 of culture - O.D. 490 nm of the administration group at day 0 of culture)]/(O.D. 490 nm of the control group at day 3 of culture – O.D. 490 nm of the control group at day 0 of culture) \times 100%.^{12,13}

Transwell Experiments

The patient-derived bladder cells were cultured in DMEM (Thermo Fisher Scientific) supplemented with 10% FBS (Thermo Fisher Scientific). The cells were treated with lentivirus particles to introduce the expression of miR-3174 or ADAM15. For the cell invasion transwell experiment, the bottom of the chamber was first coated with ECM gel (extra-cellular matrix gel, Sigma-Aldrich, St. Louis, USA). For the cell migration transwell experiment, the ECM gel was not required. The cells were seeded into the transwell chambers (8000 cells per chamber). For both experiments, the chambers were cultured for 12–16 hours at 37°C with 5% CO₂ condition. Next, the chambers were fixed using absolute ethanol, followed by staining with a crystal violet dye solution (0.25% w/v). Afterward, images of

the chambers were obtained, and the chambers extracted by absolute ethanol. The absorbance value (O.D., optical density) was measured at a wavelength of 546 nm. The inhibition rates were calculated as (O.D. 546 nm of the control group – O.D. 546 nm of the administration group)/(O.D. 546 nm of the control group) \times 100%.^{14–16}

Subcutaneous Tumor Model

The patient-derived bladder cells were cultured in DMEM (Thermo Fisher Scientific) supplemented with 10% FBS (Thermo Fisher Scientific). The cells were treated with lentivirus particles to introduce the expression of miR-3174 or ADAM15. The subcutaneous in vivo tumor model experiments were reviewed and approved by the Institutional Animal Care and Use Committee of the 302nd Hospital of People's Liberation Army (Beijing, People's Republic of China). Nude mice (4-6 weeks old) were purchased from Si-Bei-Fu Biotechnology (Beijing, People's Republic of China). All animal studies were performed in accordance with the UK (United Kingdom) Animals (Scientific Procedures) Act 1986 and associated guidelines. The cells were injected into a subcutaneous position in the mice. Next, the mice were sacrificed after 6 to 8 weeks, and the subcutaneous tumors were collected (n = 10 for per group). The mass of the tumors was measured using a precise balance. The length and width of the subcutaneous tumors were measured by a vernier caliper, and the volumes of subcutaneous tumors were calculated as (tumor length × tumor width × tumor width)/ 2.17-19 The inhibition rates were calculated as (tumor volume of the control group tissue - tumor volume of the administrated group tissue)/(tumor volume of the control group tissue) \times 100% and as (tumor weight of the control group tissue - tumor weight of the administrated group tissue)/(tumor weight of the control group tissues) \times 100%.20

Statistical Analysis

Analyses to determine the statistical significance of the obtained data were performed using GraphPad Software (version 6.0, San Diego, CA. USA). The statistical significance between the two groups was analyzed using the two-way ANOVA (Analysis of Variance) with a Bonferroni correction. Additionally, the correlation analysis of the expression of miR-3174 or ADAm15 was performed using linear regression methods. Paired samples were tested using the paired-

sample *t*-test. All values with a P-value <0.05 were concerned statistically significant.

Results

The Expression of ADAM15 in Bladder Tumor Tissue Was Significantly Higher Compared to the Paired Non-Tumor Tissue

First, the expression level of all ADAMs (ADAM8, 9, 10, 11, 12, 15, 17, 19, 22, 23, 28, and 33) was examined in 30 pairs of clinical specimens (one cancer and one non-cancer as one pair) from bladder cancer patients. The results show that the expression of ADAM23 in tumor

tissue was lower than that in non-tumor tissue (Figure 1A). The expression of ADAM11 (Figure 1K), ADAM28 (Figure 1B), and ADAM33 (Figure 1C) was slightly lower in tumor tissue compared to non-tumor tissue; nevertheless, no statistical difference was found. For ADAM17 (Figure 1E), ADAM15 (Figure 1F), ADAM12 (Figure 1G), and ADAM10 (Figure 1H), the expression was significantly higher in tumor tissue than in non-tumor tissue (Figure 1). This, while the expression of ADAM8 (Figure 1J), ADAM9 (Figure 1I), ADAM19 (Figure 1L), and ADAM22 (Figure 1D) was basically the same between the tumor and non-tumor tissue.



Figure I The expression of ADAMs in paired bladder cancer-nontumor specimen. Total mRNA samples from 30 paired bladder tumor and non-tumor specimens were extracted for Ion AmpliSeq Sequencing experiments. The expression of ADAMs (ADAM23 (A), ADAM28 (B), ADAM33 (C), ADAM22 (D), ADAM17 (E), ADAM15 (F), ADAM12 (G), ADAM10 (H), ADAM9 (I), ADAM8 (J), ADAM11 (K) and ADAM19 (L)) in paired bladder tumor and non-tumor specimens is represented as a scatter plot showing the median values of mRNA levels. The differences between bladder tumor and non-tumor specimens were determined (*P < 0.05).

Compared to the expression level of different ADAMs in the tumor tissue, the expression level of ADAM15 was significantly higher (Figure 2). A correlation analysis was performed of ADAM15's expression in the tumor as well as non-tumor tissue. As a result, the expression of ADAM15 was found to be inversely correlated (y = 0 - 0.0918x + 0.0252; R2 = 0.2862; P = 0.0023) in both the tumor and nontumor tissue (Figure 3A). To further examine the correlation between the expression of ADAM15 and various stages of bladder cancer, the expression level of ADAM15 was evaluated in various stages: Class 0, A, B, C, and D (Figure 3B). The results show that a high level of ADAM15 is present in more aggressive stages of bladder cancer (eg, Class C and D) than in less aggressive stages of bladder cancer (eg, Class 0



Figure 2 The expression of ADAMs in paired bladder tumor and non-tumor specimens. Total mRNA samples from 30 paired bladder tumor and non-tumor specimens were extracted for Ion AmpliSeq Sequencing experiments. The expression of ADAMs (ADAM9, ADAM10, ADAM10, ADAM15, ADAM17, ADAM22, ADAM23, ADAM28, ADAM8, ADAM11, ADAM19 and ADAM33) in paired bladder tumor and non-tumor specimens is represented as a scatter plot showing the median values of mRNA levels. The asterisk indicator (ADAM) indicates the members of ADAM (ADAM8, 9, 10, 11, 12, 15, 17, 19, 22, 23, 28 or 33). *P<0.05 versus the endogenous mRNA level of ADAMs in cancer specimens within the non-tumor tissues.

or Class A) (Figure 3B). Overall, ADAM15 was selected for further investigation.

The microRNA miR-3174 Targets ADAM15

Next, miRDB was used to identify a microRNA (miR) that potentially binds to ADAM15. As shown in Figure 4A, miR-3174 was identified to have a potential binding site in the 3'UTR (3' un-transcription region) of ADAM15. To confirm, Western blot analysis was used to determine the effect of miR-3174 on ADAM15's expression in bladder cancer cells (Figure 4B). The results showed that the transduction of lentivirus containing miR-3174 significantly decreased the expression of ADAM15 (Figure 4B). Transfection of miR-3174's inhibitor or the expression vector of ADAM15 with a mutated miR-3174 targeting site in 3'UTR almost fully blocked the inhibiting effect of miR-3174 on ADAM15' expression bladder in cancer cells (Figure 4B). Furthermore, in agreement with the inhibitory effect, expression of miR-3174 was negatively associated with the expression of the ADAM15 protein in the bladder cancer samples (y = 0 - 0.0659x + 0.0225, R2 = 0.1474, P = 0.0362, Figure 4C). Therefore, miR-3174 was found to be a microRNA that targets ADAM15.

miR-3174 Reduced the Proliferation of Bladder Cancer Cells by Targeting ADAM15

The effect of miR-3174 on the proliferation of bladder cancer cells was determined using MTT experiments. As shown in Table 1, the transduction of lentivirus containing



Figure 3 The expression of ADAM15 in clinical specimens of bladder cancer. Total mRNA samples from 30 paired bladder tumor and non-tumor specimens were extracted for Ion AmpliSeq Sequencing experiments. (A) The expression of ADAM15 in paired bladder tumor and non-tumor specimens is represented as a scatter plot: The x-axis represents the mRNA level of ADAM15 in bladder tumor tissue, whereas the y-axis represents the mRNA level of ADAM15 in the respective paired non-tumor tissue. (B) The expression of ADAM15 in bladder tumor tissue with indicated stages (Class 0, Class A, Class B, Class C, and Class D) is represented as a scatter plot showing the median values of mRNA levels. The significance levels were of the following comparisons: Class A versus Class 0 (*P < 0.05), Class B versus Class 0 (*P < 0.05), Class C versus Class 0 (*P < 0.05), and Class D versus Class 0 (*P < 0.05).



Figure 4 miR-3714 is a microRNA that potentially targets the 3'UTR of ADAM15. (A) The sequences of miR-3174's binding site and the mutated sequences of miR-3174's binding site in ADAM15's 3'UTR are shown. (B) Patient-derived bladder cancer cell lines – transducted with the control, miR-3174, miR-3174 + inhibitor, and miR-3174 + ADAM15^{mut} (ADAM15 with the mutated binding site of miR-3174 in 3'UTR)—were harvested for Western blot. GAPDH was used as a loading control. The results are shown as Western blot images. (C) The expression of ADAM15/miR-3174 in bladder cancer specimens is represented as a scatter plot: The x-axis represents the mRNA level of ADAM15 in bladder tumor tissue, whereas the y-axis represents the level of miR-3174 in the same bladder tumor tissue.

miR-3174 inhibited the proliferation of bladder cancer cells. The transfection of the miR-3174 inhibitor or the expression vector of ADAM15 with a mutated miR-3174 targeting site in 3'UTR almost fully stopped the inhibiting effect of miR-3174. For these results, the mean \pm SD is shown of the inhibition rates of miR-3174 on five patient-derived bladder cancer cell lines (Table 1). Together, these results indicate that miR-3174 reduced the proliferation of bladder cancer cells by targeting ADAM15.

miR-3174 Inhibits the in vitro Invasion and Migration of Bladder Cancer Cells by Targeting ADAM15

The effect of miR-3174 on the proliferation of bladder cancer cells in vitro was determined using MTT experiments. As shown in Figures 5 and 6, Tables 2 and 3, the transduction of lentivirus containing miR-3174 inhibited the in vitro invasion (Figure 5 and Table 2) and migration (Figure 6 and Table 3) of bladder cancer cells. Similarly, the transfection of the miR-3174 inhibitor or the expression vector of ADAM15 with a mutated miR-3174 targeting site in 3'UTR almost fully stopped the inhibiting effect of miR-3174. The results were found as typical images (Figures 5 and 6) and the mean

 \pm SD of the inhibition rates (Tables 2 and 3) of miR-3174 on five patient-derived bladder cancer cell lines. In sum, miR-3174 inhibited the in vitro invasion and migration of the bladder cancer cells by targeting ADAM15.

miR-3174 Inhibits the in vivo Invasion and Migration in Mice of Bladder Cancer Cells by Targeting ADAM15

To further examine the effect of miR-3174 on the subcutaneous growth of bladder cancer cells, nude SCID mice were used as an in vivo model. As shown in Figure 7, Tables 4 and 5, the transduction of lentivirus containing miR-3174 inhibited the subcutaneous growth (Figure 7) of bladder cancer cells. The transduction of lentivirus containing ADAM15, containing a mutated miR-3174 targeting site in 3'UTR, almost fully stopped the inhibiting effect of miR-3174. The results are shown as typical images of subcutaneous tumors (Figure 7) and the mean \pm SD of the inhibition rates (Tables 4 and 5) of miR-3174 on five patient-derived bladder cancer cell lines. As such, it can be derived that miR-3174 reduced the subcutaneous growth of bladder cancer cells in mice by targeting ADAM15.



Figure 5 miR-3714 represses the in vitro invasion of bladder cancer cells by targeting the 3'UTR of ADAM15. Patient-derived bladder cancer cell lines – transducted with the control, miR-3174 and miR-3174 + ADAM15^{mut} (ADAM15 with the mutated binding site of miR-3174 in 3'UTR) – were analyzed by transwell experiments. The results are shown as (A) images of the transwell experiments, (B) the related invasion cell number, and (C) inhibition rates calculated from the relative invasion cell number. *P < 0.05.

Discussion

Recently, bladder cancer has been considered as a fatal urinary system disease associated with high metastasis and mortality rates when no optimal treatment is received.²¹ Although the awareness of hematuria (presence of red blood cells in urine) is still the foremost indicator of bladder cancer, the current early diagnosis of bladder cancer is still not satisfactory.^{22,23} The prognosis of patients suffering from non-muscle-invasive bladder cancer is relatively good; however, the clinical outcome of bladder cancer with muscle-invasion remains poor.^{24,25} Thus, it is urgent to research the involvement of metastasis/invasion-related mechanisms in bladder cancer. In the present work, we used Ion AmpliSeq Targeted Sequencing technology to detect the expression levels of ADAMs in 30 pairs of tumor versus non-tumor tissues (Figure 8). Here, the results show that the expression of ADAM15 was significantly higher in tumor tissue than in non-tumor tissue. At the same time, among the 12 tested ADAMs, the expression of ADAM15 was significantly higher in tumor tissue compared to the other ADAMs. The expression of ADAM15 was found to be closely related to the stages of bladder cancer as well. This finding suggests that ADAM15 might play an important role in the occurrence and progression of bladder cancer. The AmpliSeq Targeted Sequencing technology used in the present work is a highly valuable technology. It has several advances: (1) AmpliSeq Targeted Sequencing allows the simultaneous PCR reactions of thousands and even tens of thousands of reactions; (2) the minimum amount of required starting DNA is only 10 ng, which is suitable for the trace specimens; and (3) it is suitable for samples with degraded DNA or RNA due to the long-term cryopreservation or multiple freeze-thaw. So, in our study, the AmpliSeq Targeted Sequencing technology was used to detect the expression of ADAMs in a series of bladder cancer tissue specimens. As a result, the findings of these analyses lay a solid foundation for further study of the role and regulation mechanism of ADAMs in bladder cancer.



Figure 6 miR-3714 represses the in vitro migration of bladder cancer cells by targeting the 3'UTR of ADAM15. Patient-derived bladder cancer cell lines – transducted with the control, miR-3174 and miR-3174 + ADAM15^{mut} (ADAM15 with the mutated binding site of miR-3174 in 3'UTR) – were analyzed by transwell experiments. The results are shown as (**A**) images of the transwell experiments, (**B**) the related invasion cell number, and (**C**) inhibition rates calculated from the relative invasion cell number. *P < 0.05.

Previous studies have confirmed that ADAM is a novel regulator of malignant tumor metastasis and invasion;²⁶ however, similar reports regarding its role in bladder cancer are few and the effects not completely clear.²⁴ In the present work, the expression of various ADAMs in bladder cancer tissue was examined for the first time. From these results, miR-3174 was identified as a miRNA that targets the 3'UTR of ADAM15.

Table 2The Inhibitory Rates of miR-3174 on Five Patient-Derived Bladder Cancer Cells' in vitro Invasion

PDCs	Control mi RNA	mi R -3174	miR-3174 + ADAM15 ^{mut}
	Inhibitory Rates on the in vitro Invasion of Bladder Cancer Cells (%)		
No. I	-	62.89±6.73	-
No. 2	-	65.36±5.34	1.51±3.88
No. 3	-	70.75±5.62	2.55±4.33
No. 4	-	66.11±6.92	-
No. 5	-	69.47±5.74	-

Abbreviations: PDCs, patient-derived cells included two lines of Class C (No. I and 2) and three lines of Class D (No. 3–5); ADAM15^{Mut}, ADAM15 with the mutated miR-3174's targeting sites.

Table 3 The Inhibitory Rates of miR-3174 on Five Patient-Derived Bladder Cancer Cells' in vitro Migration

PDCs	Control mi RNA	mi R -3174	miR-3174 + ADAM15 ^{mut}
	Inhibitory Rates on the in vitro Migration of Bladder Cancer Cells (%)		
No. I	_	74.40±8.83	-
No. 2	-	80.75±6.67	-
No. 3	-	78.34±6.19	-
No. 4	-	68.53±7.46	-
No. 5	-	73.95±8.48	_

Abbreviations: PDCs, patient-derived cells included two lines of Class C (No. I and 2) and three lines of Class D (No. 3–5); ADAM15^{Mut}, ADAM15 with the mutated miR-3174's targeting sites.

PDCs	Control mi RNA	mi R -3174	miR-3174 + ADAM15 ^{mut}
	Inhibitory Rates on the Subcutaneous Tumor Volumes of Bladder Cancer Cells (%)		
No. I	-	85.00±4.80	2.67±16.34
No. 2	-	86.97±5.55	-
No. 3	-	84.31±7.72	3.05±10.42
No. 4	-	73.56±4.48	0.45±9.76
No. 5	-	70.22±3.96	-

Table 4 The Inhibitory Rates of miR-3174 on Five Patient-Derived Bladder Cancer Cells' Subcutaneous Tumor Volumes

Abbreviations: PDCs, patient-derived cells included two lines of Class C (No. I and 2) and three lines of Class D (No. 3–5); ADAM15^{Mut}, ADAM15 with the mutated miR-3174's targeting sites.

 Table 5
 The Inhibitory Rates of miR-3174 on Five Patient-Derived Bladder Cancer Cells' Subcutaneous Tumor Weights

PDCs	Control miRNA	mi R -3174	miR-3174 + ADAM15 ^{mut}
	Inhibitory Rates on the Subcutaneous Tumor Weights of Bladder Cancer Cells (%)		
No. I	-	83.25±5.23	5.41±16.09
No. 2	-	87.11±4.42	-
No. 3	-	80.05±6.33	-
No. 4	-	74.73±6.51	3.33±17.44
No. 5	-	82.67±4.25	2.08±9.36

Abbreviations: PDCs, patient-derived cells included two lines of Class C (No. I and 2) and three lines of Class D (No. 3–5); ADAM15^{Mut}, ADAM15 with the mutated miR-3174's targeting sites.

Interestingly, overexpression of miR-3174 inhibited the proliferation and metastasis of bladder cancer cells by targeting and inhibiting ADAM15. Based on the factor that expression of miRNAs in human cancer cells could be used to silence oncogene expression, it is a promising approach to achieve an antitumor effect.^{27–29} Thus, our data reveal not only the role of the ADAM15/miR-3174 axis in bladder cancer regulation but also provides evidence that miR-3174 could be a potential approach for bladder cancer treatment. Moreover, previous work often focused on the genetic alteration of tumorigenesis-associated genes (eg, TP53, VEGFR, and EGFR).^{1–5} For this reason, our results are highly valuable as it expands the understanding of the molecular mechanisms involved with bladder cancer.

While the studies of ADAMs and their effects on bladder cancer are few in number, several of them can be used for comparing our data. Indeed, van Kampen et al and Jia et al already showed that miR-126 and miR-520f repress the proliferation and metastasis of bladder cancer cells by specific targeting.^{30,31} Additionally, Fu et al and Fröhlich et al reported that ADAM10 and ADAM12 are involved in regulating the proliferation and invasion process of bladder cancer cells.^{32,33} Nevertheless, in our study, the expression of a panel of various ADAMs in bladder cancer tissue was investigated for the first time. Our results also showed that the inhibited expression of ADAM15 via miR-3174 significantly reduced the proliferation and metastasis of the patient-derived bladder cancer cells. These results were consistent with previous results from Lorenzatti et al.³⁴ Interestingly, our results show that the expression of ADAM23 in the tumor tissue was lower compared to the non-tumor tissue. Also, the expression of ADAM28 and ADAM33 in the tumor tissue was slightly lower compared to the non-tumor tissue, although no statistical significance was found. Furthermore, the expression of ADAM17, ADAM15, ADAM12, and ADAM10 was significantly higher in the tumor tissue than in the non-tumor tissue. The expression of ADAM9 and ADAM22 was found similar in both the tumor and non-tumor tissue. Moreover, these results were notconsistent with the results from Tyan et al and Yang et al, who reported that ADAM28 is overexpressed in bladder transitional cell carcinoma.35,36 More clinical specimens will be collected for related testing in the future.

The biological behavior of cancer cells in tumor tissues is generally affected by the extracellular matrix (ECM) and the recognition, adhesion, and interaction between cells. In this perspective, the ECM modulates the behavior and features of solid tumors.^{37,38} Here, the collagen and osteopontin components of the ECM can modify the microenvironment in solid tumors to regulate drug-resistance, proliferation, and metastasis of tumor cells. As a membrane protein, the ADAMs can function as a key regulator of adhesion and interaction between cells in their ECM. Here, ADAM15 promotes the extracellular shedding of E-cadherin, which is involved in cell activation and proliferation. With such an effect, the down-regulation of ADAM15 could kill bladder cancer cells. Thus, the results of the present work also provide a theoretical basis for future targeted treatment of bladder cancer. Moreover, ADAMs also have other functions. For instance, Zhang et al and Dosch et al suggested that ADAM17 and ADAM10 mediate the cleaving and activation of the NOTCH (Notch Signaling pathway) protein and enhance the EMT and resistance of tumor cells.^{39,40} A such, targeting these ADAMs could be an approach for antitumor



Figure 7 miR-3714 represses the in vivo migration of bladder cancer cells by targeting the 3'UTR of ADAM15. Patient-derived bladder cancer cell lines – transducted with the control, miR-3174 and miR-3174 + ADAM15^{mut} (ADAM15 with the mutated binding site of miR-3174 in 3'UTR) – were injected into a subcutaneous position in nude SCID mice. The results are shown as (**A**) images of the tumor tissue, (**B**) tumor volumes, (**C**) inhibition rate calculated by tumor volumes, (**D**) tumor weights, and (**E**) inhibition rates calculated from the tumor weights. *P < 0.05.



Figure 8 The proposal model image of the Ion AmpliSeq Sequencing system used in the present work. The primer pool 1 includes the ADAM10, ADAM10, ADAM15, ADAM17, ADAM22 or ADAM8, whereas the primer pool 2 includes the ADAM12, ADAM19, ADAM23, ADAM28, ADAM33 and ADAM11.

strategies. Overall, thanks to the various roles that the ADAMs play in tumor biology, it is valuable to further examine their functions and mechanisms regarding the regulation of bladder cancer in the future.

The interaction between miRNAs/genes plays important roles in the tumorigenesis of human malignancies.^{41,42} The investigation on genes and miRNAs seems to be a novel diagnostic instrument in the medical field; however, the limit is that often the same gene/miRNA plays a role in different cancer types.⁴³ Besides the miRs, some new future targets of therapy, including the immune system, are the promising strategies for bladder cancer treatment.^{44–47}

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Author Contributions

All authors made substantial contributions to the design and conception, acquisition, analysis, or interpretation of data. All authors took part in either drafting or revising the manuscript. All authors also gave final approval of the version to be published and agree to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved.

Disclosure

The authors report no conflicts of interest in this work.

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