miR-433 accelerates acquired chemoresistance of gallbladder cancer cells by targeting cyclin M

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Abstract. Resistance to chemotherapy is associated with dismal prognosis in patients with gallbladder cancer. Cyclin-dependent kinase 10 (CDK10) influences the chemosensitivity of gallbladder cancer cells, and cyclin M is the activating factor and binding partner of CDK10. To determine the effect of CDK10 or cyclin M overexpression on chemosensitivity, gemcitabine-resistant (GR) subclones were established from CDK10 or cyclin M stable transfectants. Stable overexpression of CDK10 increased the sensitivity to gemcitabine in non-resistant cells and did not further increase the sensitivity to gemcitabine in the GR subclones. GR subclones exhibited a significantly decreased expression of cyclin M while maintaining the expression levels of CDK10, compared with the non-resistant cells. MicroRNA (miR)-433 was identified as a candidate factor involved in the mechanism of the downregulation of M cyclin in GR subclones. Luciferase assays confirmed the interaction between miR-433 and the 3′ untranslated region (3′UTR) of cyclin M. Additionally, ectopic expression of miR-433 significantly decreased the expression of cyclin M. Finally, increased expression of circulating miR-433 was associated with poor outcome of chemotherapy. The results of the present study suggest that miR-433 is a potential biomarker for evaluating chemosensitivity in gallbladder cancer.

Introduction

Gallbladder cancer is a relatively rare disease, with an estimated annual incidence of between 2 and 3 individuals/100,000; however, it is the most frequently encountered malignancy of the biliary tract (1,2). Complete surgical resection is the only potentially curative approach in early stages of the disease; however, the majority of cases are diagnosed in advanced stages. Additionally, because the aggressive malignancy tolerates traditional chemotherapy and radiotherapy, these cases have a dismal prognosis with an overall 5-year survival rate of <5% (3). Therefore, improving the diagnostic accuracy during the early stages and identifying the molecular mechanisms associated with chemotherapy or radiation resistance may provide a potential therapeutic approach for the treatment of gallbladder cancer.

Cyclin M, also known as FAM58A, is a member of the cyclin family, which serves important regulatory functions in the cell cycle and transcription (4). Mutations in cyclin M cause toe syndactyly, telecanthus and anogenital and renal malformations syndrome, a human developmental anomaly (4,5). However, the function of cyclin M in other diseases remains unclear. Cyclin M has been identified as the binding partner for cyclin-dependent kinase 10 (CDK10) (5). CDK10 is involved in the regulation of cell division and function in various types of tumor (6-10). It was confirmed that CDK10 functions as a tumor suppressor gene and regulates the survivability of biliary tract cancer cells, including gallbladder cancer cells (8). These results suggest that cyclin M may also be involved in cancer development through binding to CDK10.

MicroRNAs (miRNAs) are key negative regulators of gene expression in eukaryotes and abundant evidence indicates that miRNAs are involved in human cancers (11,12). miR-433 has been identified as a cancer-associated miRNA in various types of tumor, including bladder cancer, ovarian cancer, gastric cancer, hepatocellular carcinoma and oral squamous cell carcinoma (13-17). Furthermore, aberrant expression levels of miR-433 have been identified to be associated with chemosensitivity and enhanced survival of ovarian cancer cells (15,18).

In the present study, gemcitabine-resistant (GR) subclones were established on the basis of stable transfectants of CDK10 or cyclin M overexpression. To provide a possible mechanism for cyclin M downregulation in acquired chemoresistance of gallbladder cancer cells, miR-433 was predicted to target cyclin M. Furthermore, the association of serum levels of miR-433 and the chemotherapeutic response in gallbladder cancer was evaluated. Collectively, the results of the present study suggest that aberrant expression of miR-433 affects the chemoresistance of gallbladder cancer cells by targeting cyclin M.

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Materials and methods

Cell culture, plasmids, RNA oligonucleotides and transfection. The GBC-SD gallbladder carcinoma cell line was obtained from the Cell Bank of Type Culture Collection of the Chinese Academy of Sciences (Shanghai, China). Cells were cultured at 37°C in 5% CO₂ in RPMI-1640 medium with 10% fetal bovine serum (both from Gibco; Thermo Fisher Scientific, Inc., Waltham, MA, USA), 100 IU/ml penicillin and 100 µg/ml streptomycin. Human CDK10 and cyclin M coding sequences were cloned into pBudCE4.1 vector (Invitrogen; Thermo Fisher Scientific, Inc.). Total RNA was isolated from normal liver tissue using TRIZol reagent (Thermo Fisher Scientific, Inc.) and cDNA was synthesized using the PrimeScript reagent kit (Applied Biosystems; Thermo Fisher Scientific, Inc.), according to the manufacturer’s instructions. The primer sequences for PCR amplification of CDK10-open reading frame (ORF) were as follows: 5’-CCAAGCTTATGGCGG AGGCCAGATCTGGA-3’ (forward) and 5’-CCGGATCTCT CAGGTTTACAGCGCTTG-3’ (reverse). The PCR products were purified and cloned into the HindIII/BamHI sites of the vector. The primer sequences for PCR amplification of cyclin M-ORF were as follows: 5’-GGGTACCATGGA AGCCCCGAGGG-3’ (forward) and 5’-CCCTCAGT TAGGGATCTCTTGTGTTCA-3’ (reverse). The PCR products were purified and cloned into the KpnI/Xhol sites of the vector. CDK10-ORF was cloned into the pBudCE4.1 vector and cyclin M-ORF was cloned into the pBudCE4.1 vector to produce pBudCE4.1-CDK10 and pBudCE4.1-CyclinM, respectively. Finally, CDK10-ORF was cloned into the pBudCE4.1-CyclinM vector and named pBudCE4.1-Co. All the constructs were confirmed by DNA sequencing and the pBudCE4.1 empty vector was used as a vector control.

GBC-SD cells were transfected with pBudCE4.1-CDK10, pBudCE4.1-CyclinM, pBudCE4.1-Co or empty vector plasmid using LipoFectamine® 2000 (Invitrogen; Thermo Fisher Scientific, Inc.). At 48 h post-transfection, the culture medium was replaced with medium containing Zeocin (Invitrogen; Thermo Fisher Scientific, Inc.) and the medium containing Zeocin was replaced every 3-4 days. After 2 weeks, isolated colonies began to appear. After 3 weeks, stable transfectants expressing CDK10 or cyclin M were selected for further study. The control clones expressing the empty vector were isolated at the same time. The stable transfectants were named as follows: GBC-Mock, GBC-CDK10, GBC-CyclinM and GBC-Co, respectively.

Small interfering RNA (siRNA) targeting the proto-oncogene serine/threonine-protein kinase c-Raf was obtained from RiboBio (Guangzhou, China) and the sequence was 5’-GCUGCAUCUCUCCUCAAU-3’. Control siRNA was obtained from RiboBio (8). Other RNA oligonucleotides (including miR-433, anti-miR-433, miR-control and siRNA-control groups) were also obtained from RiboBio. At 24 h before transfection, cells were plated onto a 35-mm dish at 50-60% confluence. RNA oligonucleotides transfection was performed at 50 nM using LipoFectamine® 2000, according to the manufacturer’s instructions. The transfected cells were resuspended in RPMI-1640 medium and cultured for 48-72 h before analysis. Each assay was performed in triplicate at least.

Establishment of GR subclones. Stable transfectants were exposed to increasing concentrations (0.001, 0.0025, 0.005, 0.01, 0.025, 0.05, 0.1, 0.25, 0.3, 0.35, 0.4, 0.45 and 0.5 µg/ml) of gemcitabine (Lilly France S.A., Lilly, Fegersheim, France) every 4 weeks over a 9-month period. As cells adapted to the doses, the gemcitabine concentration was increased. The cell subclones were named as follows: GBC-Mock-GR, GBC-CDK10-GR, GBC-CyclinM-GR and GBC-Co-GR.

Western blot analysis. Total proteins were extracted from tissue samples or cells using lysis buffer containing phenyl-methyl sulfonylfluoride (both from Beyotime Institute of Biotechnology, Haimen, China) at 25°C and protein concentration was determined by BCA Protein Assay kit (Beyotime Institute of Biotechnology). A total of 20 µg protein was separated by SDS-PAGE (10% gel) and transferred onto a polyvinylidene fluoride membrane. Membranes were blocked with 5% non-fat milk and incubated with anti-CDK10 (cat. no. 17182-1-AP, 1:500; ProteinTech Group, Inc., Chicago, IL, USA), anti-c-Raf (cat. no. AP8716d, 1:500; Abgent, Inc., San Diego, CA, USA) and anti-cyclin M (cat. no. ab81062, 1:1000; Abcam, Cambridge, UK) primary antibodies at 4°C for 12 h. Anti-β-actin (cat. no. sc-47778, 1:1000; Santa Cruz Biotechnology, Inc., Dallas, TX, USA) was used as a loading control. Membranes were then washed and incubated with horseradish peroxidase-conjugated secondary antibody (cat. no. A21020 and A21010, 1:8000; Abbkine, Wuhan, China). Immunoreactive bands were visualized using a chemiluminescence solution with an ECL kit (Beyotime Institute of Biotechnology). Quantity One (Bio-Rad Laboratories, Inc., Hercules, CA, USA) was used as the software for quantification.

Cell counting Kit-8 (CCK-8) assay. Cells were seeded into a 96-well plate at a density of 4x10³ cells/well, and allowed to attach at 37°C overnight. Cells were treated with various concentrations (0, 0.01, 0.05, 0.1, 0.5, 1 and 10 µg/ml) of gemcitabine in ≥6 replicate wells and incubated at 37°C for 48 h. Cell proliferation was determined using the CCK-8 assay (Beyotime Institute of Biotechnology). In brief, a 1/10 volume of the CCK-8 solution was added to each well, and the cells were cultured for another 2 h. Cell viability was determined by measuring the absorbance at 450 nm using a plate reader (BioTek Instruments, Inc., Winooski, VT, USA).

Colony formation assay. Cells were seeded for colony formation in 24-well dishes at a density of 500 viable cells/well. The medium contained 1 µg/ml gemcitabine and was exchanged every 2 days. After 14 days, the cells were fixed in 4% paraformaldehyde for 15 min. Following washing, the cells were stained with 0.005% crystal violet solution for 1 h. The plates were aspirated, washed and allowed to air dry. The dishes were photographed with a camera and the colonies were counted (a single colony contained ≥20 cells) using a light microscope (magnification, x40). Each assay was performed in triplicate.

Clinical tissue and blood samples. Gallbladder cancer tissue and paired normal tissue specimens were obtained from 8 patients at the Shaoxing People’s Hospital (Shaoxing, China) from September 2012 to March 2016. All participants...
provided written informed consent and the tissue acquisition protocol was approved by the Zhejiang University Institutional Review Board (Hangzhou, China). Blood samples were collected from 23 patients with gallbladder cancer and the survival time was ≥3 months. Among the patients, 17 patients underwent chemotherapy with gemcitabine or 5-fluouracil and were assigned to the chemotherapy group. The remaining 6 patients did not receive chemotherapy and were assigned to the non-chemotherapy group. Blood samples from the patients were collected and serum was separated by centrifugation (1,000 x g, for 5 min at 4°C) collected into RNase/DNase-free tubes and stored at -80°C until use. Fresh tissues were stored at -80°C.

Reverse transcription-quantitative polymerase chain reaction (RT-qPCR). Total RNA was isolated from cells, tissues or serum using TRIzol reagent, and RNA was reverse-transcribed into cDNA using a High Capacity cDNA Reverse Transcription kit (Takara Biotechnology Co., Ltd., Dalian, China), according to the manufacturer’s instructions. qPCR was performed using SYBR Green PCR Master Mix (Takara Biotechnology Co., Ltd.) and the ABI 7500 Real-time PCR system (Applied Biosystems; Thermo Fisher Scientific, Inc.), according to the manufacturer’s instructions. Specific primers were designed for cyclin M, CDK10 and β-actin (Table I). The PCR conditions included 95°C for 5 min, and a total of 40 cycles of 95, 60 and 72°C for 30 sec and then a final extension at 72°C for 5 min. β-actin was used as an endogenous control. miR-433 normalization was performed according to a protocol described previously (19). Expression of miR-433 was normalized using U6 RNA (for RNA from cells and tissues) and with Caenorhabditis elegans miR-39 mimic (for RNA from culture supernatant and serum samples) which was used as an RNA spike-in. Specific primers for miR-433, U6 and C. elegans miR-39 RNA were obtained from RiboBio (15,19).

Table I. Primers designed for cyclin M, CDK10 and β-actin.

<table>
<thead>
<tr>
<th>Name</th>
<th>Symbol</th>
<th>Forward (5'-3')</th>
<th>Reverse (5'-3')</th>
</tr>
</thead>
<tbody>
<tr>
<td>β-actin</td>
<td>ACTB</td>
<td>AGAAGGAGATCAGCTGCCCTGGCACC</td>
<td>CCGCTTTGCTGTACCCATCTGCTG</td>
</tr>
<tr>
<td>CDK10-ORF</td>
<td>CDK10</td>
<td>TGGCAAGGAAGGAGAGATG</td>
<td>CTGCTCAGATACCACATCC</td>
</tr>
<tr>
<td>CDK10-3'UTR</td>
<td>CDK10</td>
<td>GTGCAACACTGCTTCCTTGG</td>
<td>AGCGTCCACACAACCTCAA</td>
</tr>
<tr>
<td>Cyclin M-ORF</td>
<td>FAM58A</td>
<td>AACTTCCGAGTGGGAGGGTGCTCAT</td>
<td>TCAAGGGGTCCAGGTGGTCT</td>
</tr>
<tr>
<td>Cyclin M-3'UTR</td>
<td>FAM58A</td>
<td>CGCTTTGCCAGTCCCTGAA</td>
<td>GACAAAAACGAGATGGGCTG</td>
</tr>
</tbody>
</table>

CDK10, cyclin-dependent kinase 10; ORF, open reading frame; 3'UTR, 3' untranslated region.

Statistical analysis. Data were analyzed using SPSS software (version 13.0; SPSS, Inc., Chicago, IL, USA). The relevant data are expressed as the mean ± standard deviation (SD). Statistical significance between two groups was determined using Student's t-test. One-way analysis of variance followed by a Tukey-Kramer test was used to examine differences among multiple groups. P<0.05 was considered to indicate a statistically significant difference.

Results

Decreasing expression of cyclin M contributes to acquired chemoresistance of GR subclones. To generate stable cell lines for experimental analyses, GBC-SD cells were transfected with pBudCE4.1-CDK10, pBudCE4.1-CycM, pBudCE4.1-Co or with the empty vector plasmid. After 4 weeks, stable transfectants were obtained and named GBC-CDK10, GBC-CycM, GBC-SD-β-actin. For luciferase assays, 293T cells were cultured in 6-well plates and transfected with plasmids (including pMIR-CycM-3'UTR, mut-pMIR-CycM-3'UTR and β-galactosidase reporter plasmids) and RNA oligonucleotides using Lipofectamine® 2000. A β-galactosidase reporter plasmid was designed to normalize transfection efficiency. At 48 h after transfection, the cells were lysed, luciferase activity was determined using a Dual-Luciferase Reporter assay system and β-galactosidase activity was determined using the β-Galactosidase Enzyme assay system (both from Promega Corporation, Madison, WI, USA).

Dual-luciferase reporter assay. Predictions for the potential binding sites of miR-433 in the 3'UTR of cyclin M was performed using 2 different online programs: TargetScan 6.2 (http://www.targetscan.org/vert_61/) and microRNA.org (http://www.microrna.org). The 3'UTR of cyclin M was amplified by PCR using human genomic DNA as a template. The primers used were as follows: 5'-CGACGC GTTTGACTACAATACAGGCATGACA-3' (forward) and 5'-CCAAGCTTTCTCCGATTAGATCTCAGTCAA-3' (reverse). The PCR product, which contained a potential binding site for miR-433, was digested with MluI and HindIII, and was inserted downstream of the luciferase gene in the pMIR-REPORT system (Applied Biosystems; Thermo Fisher Scientific, Inc.). The constructs were verified by DNA sequencing. The verified construct was named pMIR-CycM-3'UTR. The mutant reporter plasmids were constructed in accordance with the aforementioned criteria by annealing the following oligonucleotides: 5'-CGACGCCTTTGACTACAATACAGGCATGACATCAATGAAGGAAGATCT ataaAAATCGATGAGGCTGAAATCCCTACGGGAAAAGTTTG-3' and 5'-CCAAGCTTTGACTACAATACAGGCATGACA-3'.
GBC-Co and GBC-Mock, respectively. Western blot analysis was performed to confirm positive clones (Fig. 1A). Next, the chemosensitivity of the four stable cell lines to gemcitabine was examined. The results demonstrated that GBC-CDK10 and GBC-CycM cells exhibited a significantly increased sensitivity to gemcitabine (Fig. 1B). As expected, GBC-Co cells were also more sensitive to gemcitabine than were GBC-Mock cells (Fig. 1B).

Next, the four stable cell lines (GBC-CDK10, GBC-CycM, GBC-Co and GBC-Mock) were cultured for 9 months in the presence of gemcitabine to establish GR subclones (Fig. 1B and C). The dose-response curves indicated that GBC-CDK10-GR cells exhibited a similar chemosensitivity to that of the GBC-Mock-GR cells and were more resistant to gemcitabine compared with the GBC-Co-GR or GBC-CycM-GR cells (Fig. 1C). Following extended exposure to gemcitabine, although GBC-CDK10-GR cells maintained high levels of CDK10 expression (Fig. 1A), GBC-CDK10-GR cells did not exhibit the increased sensitivity to gemcitabine that was exhibited by GBC-CDK10 cells (Fig. 1C).

Additionally, colony formation assays were performed in all four-cell lines, which were cultured with 1 µg/ml gemcitabine. After 14 days, GBC-CDK10, GBC-CycM and GBC-Co cells were more sensitive to gemcitabine compared with GBC-Mock cells, and GBC-Co-GR cells were more sensitive to gemcitabine compared with GBC-Mock-GR and GBC-CDK10-GR cells (Fig. 1D and E). These results confirm the results obtained from the dose-response curves (Fig. 1B and C).

Cyclin M is a binding partner for CDK10 (5). It was demonstrated that cyclin M protein levels were too low to detect by western blot in the GBC-CDK10-GR cells (Fig. 1A), different from the GBC-CDK10 cells. It suggested that decreasing expression of cyclin M may contribute to the decreased chemosensitivity of CDK10-overexpressing GBC-SD cells.
Because of the transfection, both the transcripts from plasmid and genome could influence the protein levels of CDK10 and cyclin M. The exogenous transcription product (from plasmid) could cover up the practical change of endogenous expression. Considering that only the open reading frames (ORFs), without 3'UTRs, were cloned into plasmid in our study, the expression of the 3'UTRs may more accurately reflect the endogenous expression of CDK10 and cyclin M (Fig. 1F-G). To perform PCR, specific primers for the 3'UTRs of CDK10 and cyclin M were designed. The results demonstrated that the 3'UTR of cyclin M was significantly decreased in GR subclones (Fig. 1H). However, the 3'UTR of CDK10 did not exhibit a similar change (Fig. 1I). This result indicates that downregulating the expression of cyclin M may be an important mechanism for the acquired resistance in GR subclones.

Cyclin M is a target of miR-433 in gallbladder cancer cells. The interactions between miRNAs and the 3'UTRs of target genes are key regulatory mechanisms that are involved in multiple biological processes and human diseases, including cancer (21-23). To examine the molecular mechanism underlying cyclin M downregulation in gallbladder cancer cells, miRNAs targeting cyclin M using the TargetScan 6.2 and microRNA.org software were employed. Considering the association between candidate miRNAs and chemoresistance (15,18), miR-433 was hypothesized to act as a candidate target for cyclin M. miR-433-binding sequences in the 3'UTR of cyclin M mRNA are indicated in Fig. 2A. Examination of miR-433 expression levels in the GBC subclones revealed that all four GR subclones, i.e., GBC-Mock-GR, GBC-CDK10-GR, GBC-CycM-GR and GBC-Co-GR, expressed significantly increased levels of miR-433 compared with their respective non-GR subclones (Fig. 2B).

Luciferase activity assays indicated that the miR-433 mimic repressed the luciferase activity of pMIR-CycM-3'UTR reporter plasmid, but did not repress the activity of mut-pMIR-CycM-3'UTR (Fig. 2C). The miR-control had no impact on the luciferase activity of pMIR-CycM-3'UTR (Fig. 2C). To further confirm the results, cyclin M expression levels were investigated following transfection with miR-433 in GBC-SD cells. Western blot analysis was performed at 72 h post-transfection. It was revealed that cyclin M was downregulated in GBC-SD cells following miR-433 transfection, whereas cyclin M was upregulated following transfection with anti-miR-433 (Fig. 2D).

miR-433 expression enhances the chemoresistance of gallbladder cancer cells. Considering that miR-433 was expressed at a high level in GR subclones and targets cyclin M, the function of miR-433 in promoting resistance to gemcitabine in GBC-SD cells was examined in vitro. The results indicated that the miR-433 mimic enhanced the resistance of GBC-SD cells in response to a moderate dose of gemcitabine (0.5 µg/ml), whereas anti-miR-433 treatment reversed this change (Fig. 2E). However, in response to a high gemcitabine dose (10 µg/ml),
the function of the miR-433 mimic or inhibitor did not induce any significant effect.

Circulating miR-433 levels are associated with chemotherapy in gallbladder cancer. A previous study demonstrated that ovarian cancer cells with increased miR-433 expression are able to release miR-433 into the growth medium (15). Therefore, the relative miR-433 expression was examined in the culture supernatant of the subclones. The results indicated that the culture supernatant from GR subclones exhibited significantly increased relative miR-433 expression compared with that from non-GR subclones (Fig. 2F).

Serum expression levels of miR-433 were examined in patients with gallbladder cancer, from the chemotherapy group and the non-chemotherapy group. The chemotherapy group comprised 17 patients with a median survival time of 9.8 months. Among these patients, 9 patients had a shorter survival time than the median (48.6 months) and the remaining 8 patients had a longer survival time than the median (12.0 months). The first serum samples were collected from the patients prior to receiving chemotherapy. The results indicated no marked difference in miR-433 levels between the non-chemotherapy group and the chemotherapy group (Fig. 1E). However, the expression of c-Raf was at low levels under conditions of increased expression of CDK10 and cyclin M, such as in the case of GBC-Co cells and GBC-Co-GR cells (Fig. 1E).

Subsequently, the involvement of c-Raf to chemoresistance was assessed. All GR subclones, including GBC-Mock-GR, GBC-CDK10-GR, GBC-CycM-GR and GBC-Co-GR, were transfected with siRNA against c-Raf and c-Raf down-regulation was confirmed by western blot analysis (Fig. 3A). The results demonstrated that c-Raf silencing significantly increased the sensitivity to gemcitabine in GBC-Mock-GR, GBC-CDK10-GR and GBC-CycM-GR cells (Fig. 3B). Therefore, increased expression of c-Raf is also an important factor that is involved in the regulation of chemoresistance in cancer.

Discussion

Resistance to chemotherapy is a major concern in gallbladder cancer (3). Acquired chemoresistance of cancer cells may arise following extended exposure to chemotherapy drugs. The majority of patients with gallbladder cancer that undergo chemotherapy become resistant following consecutive treatments and fail to benefit from chemotherapy (24). Therefore, it is important to clarify the molecular mechanism underlying acquired chemoresistance for an improved treatment of gallbladder cancer.

Gemcitabine is a deoxycytidine analogue with antitumor activity and a key drug for chemotherapy of biliary tract cancer (25,26). To investigate the acquired chemoresistance of gallbladder cancer cells, GR subclones were established. Drug concentrations were gradually increased to establish subclones of GBC-SD cells that were able to grow in medium with 0.5 µg/ml gemcitabine. This strategy may simulate the process of acquired chemoresistance. Although CCK-8 or MTT assays are used to examine sensitivity to chemotherapy during a relatively short period (48 or 72 h), a colony formation assay is able to accurately determine chemosensitivity and colony forming ability under extended exposure to gemcitabine (2 weeks).

Similar to a previous study (8), the results of the present study confirmed that GBC-CDK10 cells were more sensitive to gemcitabine compared with GBC-Mock cells. Although GBC-CDK10-GR and GBC-CDK10 cells expressed high levels of CDK10, the gemcitabine resistance of GBC-CDK10-GR cells was as high as that of GBC-Mock-GR cells in contrast.
with that of GBC-CDK10 cells. This indicates that the expression level of CDK10 alone was not able to explain the acquired gemcitabine resistance of GBC-SD cells. However, GBC-Co-GR cells and GBC-Co cells, which expressed increased CDK10 and cyclin M, exhibited a relatively increased sensitivity to gemcitabine compared with that of GBC-Mock-GR and GBC-Mock cells. Additionally, the levels of cyclin M mRNA that contained the 3'UTR were significantly downregulated in GR subclones, compared with that observed in the non-GR subclones. This suggests that cyclin M serves an important function in the acquired gemcitabine resistance of gallbladder cancer cells.

miRNAs regulate the expression of target genes by interacting with the 3'UTRs of the mRNA. The results of the present study demonstrated that the 3'UTR of cyclin M was downregulated in all GR subclones and miR-433 was identified as a predicted miRNA to target cyclin M. Overexpression of miR-433 was demonstrated to promote resistance to paclitaxel in ovarian cancer cells (15). The results of the present study also demonstrated that the acquired gemcitabine resistance of gallbladder cancer cells was associated with an upregulation of miR-433. However, other studies reported that miR-433 functions as an anti-oncomiR in various tumors by regulating various targets (13,14,27,28). The results of the present study demonstrated that the expression of miR-433 was decreased in the tumor samples, but this decrease was not statistically significant. Furthermore, the expression of cyclin M between tumor samples and normal tissues also did not exhibit a significant difference. The expression of miR-433 in serum samples was also examined. No differences in the expression of miR-433 were identified in the serum of normal volunteers and patients with gallbladder cancer. Nevertheless, the patients with poor response to chemotherapy exhibited a significant increase in the expression of serum miR-433, whereas the patients with an improved response did not exhibit changes in the serum levels of miR-433. These observations indicate that circulating miR-433 expression may not be useful as a biomarker for diagnosis, but it may be a potential biomarker to evaluate the response to chemotherapy.

The CDK10-cyclin M complex is a negative regulator of c-Raf (5,6). The results of the present study also identified that downregulated cyclin M expression or downregulated CDK10 expression resulted in increased c-Raf expression. Considering that c-Raf silencing reversed the resistance to gemcitabine in GBC-Mock-GR, GBC-CDK10-GR and GBC-CycM-GR cells, the increasing gemcitabine resistance of GBC-SD cells may be attributed to upregulated c-Raf expression, which was induced by downregulation of cyclin M expression.

In conclusion, the results of the present study suggest that the miR-433/cyclin M axis is associated with the acquired chemoresistance of gallbladder cancer cells. Circulating miR-433 expression represents a potential biomarker for evaluating the chemosensitivity of gallbladder cancer cells.

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