miR-21 attenuates contrast-induced renal cell apoptosis by targeting PDCD4

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Abstract. Contrast medium (CM) is widely used in cardiac catheterization; however, it may induce acute kidney injury or renal failure, although the underlying mechanism remains to be elucidated. MicroRNA-21 (miR-21) is involved in renal disease and has been indicated to regulate cellular apoptosis and fibrosis, although its role in CM-induced renal cell injury is unknown. The present study examined the expression and potential targets of miR-21 in human renal proximal tubular epithelial (HK-2) cells following CM treatment. CM induced renal cell apoptosis and decreased miR-21 expression. The expression level of the apoptosis regulator protein, B-cell lymphoma 2 (Bcl-2) was upregulated, whereas that of the apoptosis regulator, Bcl-2-associated X protein (Bax) was downregulated upon transfection of miR-21 mimics; miR-21 overexpression additionally directly inhibited the expression of programmed cell death protein 4 (PDCD4), as determined by a dual luciferase reporter assay, and PDCD4 silencing reduced the rate of HK-2 cell apoptosis. The results of the present study indicated that miR-21 protected renal cells against CM-induced apoptosis by regulating PDCD4 expression.

Introduction

Contrast-induced acute kidney injury (CI-AKI) is the third most common cause of hospital-acquired renal failure, and is associated with increased cardiovascular and renal morbidity and mortality (1,2). Although efforts have focused on the treatment and prevention of the development of renal dysfunction, its incidence is increasing (3,4). Contrast medium (CM)-induced renal epithelial cell apoptosis is an important underlying cause of renal failure (5), although the mechanism remains unclear.

MicroRNAs (miRNAs/miRs) are a class of small, non-coding RNA molecules with a length of 18-25 nucleotides that serve an important role in normal biological functions, including cell growth, proliferation, differentiation and apoptosis, via post-transcriptional regulation of gene expression (6). Recent evidence has implicated miRNAs in ischemia-reperfusion injury (IRI), in addition to cisplatin-/cyclosporine-induced AKI (7-10). Specifically, miR-21 was observed to regulate renal cell apoptosis and fibrosis (11), although these findings are controversial, with certain studies demonstrating miR-21 upregulation (12,13) and others reporting a downregulation, or no change, in expression levels (14,15), depending on the tissue or disease model.

Programmed cell death protein 4 (PDCD4) is expressed in proliferating cells where it is known to suppress tumorigenesis and induce apoptosis, and is negatively regulated by miR-21 in various types of cancer (16). PDCD4 has been demonstrated to be associated with IRI (12). However, the role of PDCD4 in CM-induced renal cell injury has yet to be fully elucidated. It may be hypothesized that PDCD4 may be involved in renal tubular epithelial cell apoptosis induced by CM exposure (5). In order to test this hypothesis, the present study examined miR-21 expression levels in human renal proximal tubular epithelial (HK-2) cells under CM treatment and in gain- and loss-of-function experiments, to determine the association between miR-21 and PDCD4 expression and renal cell apoptosis. The

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Abbreviations: miR-21, microRNA-21; PDCD4, programmed cell death protein 4; Bcl-2, B-cell lymphoma 2; Bax, Bcl-2-associated X protein; CM, contrast medium; CI-AKI, contrast-induced acute kidney injury; HK-2, human renal proximal tubular epithelial; HEK, human embryonic kidney; IRI, ischemia-reperfusion injury; MUT, mutant type; WT, wild type; RT-qPCR, reverse transcription-quantitative polymerase chain reaction; siRNA, small interfering RNA; TUNEL, terminal deoxynucleotidyl transferase dUTP nick-end labeling

Key words: microRNA-21, programmed cell death protein 4, contrast medium, contrast-induced acute kidney injury, HK-2 cells
results of the present study indicated that miR-21 protected kidney cells against CM-induced apoptosis by directly targeting PDCD4.

Materials and methods

Cell culture. HK-2 and human embryonic kidney (HEK)-293T cells were provided by the Stem Cell Bank, Chinese Academy of Sciences (Shanghai, China), and were cultured in Dulbecco’s modified Eagle's medium/nutrient mixture F-12 (Gibco®; Thermo Fisher Scientific, Inc., Waltham, MA, USA) supplemented with 10% fetal bovine serum (Hyclone; GE Healthcare Life Sciences, Logan, UT, USA) at 37°C and 5% CO₂, and were subcultured when they reached 80-85% confluence. The cells were treated with 150 mg iodide (mgI)/ml Ultravist CM (370 mgI/ml; Bayer AG, Leverkusen, Germany) for 2 h at 37°C.

Terminal deoxynucleotidyl transferase dUTP nick-end labeling (TUNEL) assay. Cells (2x10⁶) were washed with PBS and fixed with 4% paraformaldehyde for 30 min at room temperature. Following three washes with PBS, cells were treated with 0.2% Triton X-100 for 5 min on ice, washed with PBS, and incubated with fluorescein isothiocyanate-labeled dUTP and terminal deoxynucleotidyl transferase for 1 h at 37°C. Following a series of washes with PBS and nuclear staining with DAPI at room temperature away from light, cells were analyzed with an epifluorescence microscope in five different visual fields at x400 magnification.

Target gene prediction. The Targetscan (www.targetscan.org) and microRNA.org databases (www.microrna.org/microrna/home.do) were used to predict the potential target of miR-21.

Reverse transcription-quantitative polymerase chain reaction (RT-qPCR) analysis. Total RNA was extracted using TRIzol reagent (Invitrogen; Thermo Fisher Scientific, Inc.), according to the manufacturer's protocol. Subsequent to the addition of 100% chloroform followed by 100% isopropanol, washes with 75% ethanol, and a series of centrifugation steps, RNA concentration and quality were measured by spectrophotometry at 260 and 280 nm. Isolated RNA was reverse-transcribed using M-MLV reverse transcriptase (Takara Bio, Inc., Otsu, Shiga, Japan), and the cDNA was amplified using SYBR Premix Ex Taq (Takara Bio, Inc.) on a CFX96 Touch real-time PCR detection system (Bio-Rad Laboratories, Inc., Hercules, CA, USA), according to a standard protocol. The working concentrations were 100 nM. After 48 h, cells were treated with 150 mgI/ml Ultravist CM for 2 h at 37°C. The efficiency of each mimic or inhibitor was confirmed by RT-qPCR. Proteins were detected by western blotting. The fraction of apoptotic cells was determined with the TUNEL assay. The sequences were as follows: miR-21-mimics, 5'-UAG CUU AUC AGA CUG AUG UUG A-3'; miR-21-inhibitors, 5'-UCA ACAUACUGAUGUUGA-3'; and negative control siRNA (Shanghai GenePharma Co., Ltd., Shanghai, China) using Lipofectamine 2000 (Invitrogen; Thermo Fisher Scientific, Inc.), according to the manufacturer's protocol. The working concentrations were 100 nM. After 48 h, cells were lysed by incubation with trypsin-EDTA solution (Invitrogen; Thermo Fisher Scientific, Inc.) and protein was extracted with radioimmunoprecipitation assay buffer (Thermo Fisher Scientific, Inc.) containing protease inhibitor. Protein concentration was quantified using a BCA Protein Assay kit (Thermo Fisher Scientific, Inc.). Equal amounts of protein (20 µg) were separated by SDS-PAGE on a 12% gel and electrophoretically transferred to a polyvinylidene difluoride membrane, which was incubated in blocking solution containing 5% non-fat milk in a Tris-buffered saline/Tween 20 solution (TBST) for 60 min at room temperature. Blocking was followed by overnight incubation at 4°C with rabbit primary antibodies against the following proteins: PDCD4 (cat. no. 9535; 1:1,000), the apoptosis regulators, B-cell lymphoma 2 (Bcl-2; cat. no. 4223; 1:1,000) and Bcl-2-associated X protein (Bax; cat. no. 5023; 1:500) (all Cell Signaling Technology, Inc., Danvers, MA, USA); and β-actin (cat. no. ab8227; 1:1,000; Abcam, Cambridge, UK). Following washing with TBST, bound antibodies were detected by incubation for 60 min at room temperature with horseradish peroxidase-labelled goat anti-rabbit immunoglobulin G (cat. no. 7074; 1:5,000; Cell Signaling Technology, Inc.). Enhanced chemiluminescence (Thermo Fisher Scientific, Inc.) was used to visualize protein bands on X-ray film. Protein expression levels were normalized to that of β-actin. Image J v1.48 U (National Institutes of Health, Bethesda, MD, USA) was used for densitometric analysis.

Transfection of oligonucleotides and small interfering (si) RNA. Cells were seeded at a density of 2x10⁵ in 6-well plates and grown to 60-70% confluence in DMEM-F12, and subsequently transfected with miR-21 mimic or miR-21 inhibitor, scrambled control miR-21, siRNA against PDCD4 or negative control siRNA (Shanghai GenePharma Co., Ltd., Shanghai, China) using Lipofectamine 2000 (Invitrogen; Thermo Fisher Scientific, Inc.), according to the manufacturer's protocol. The working concentrations were 100 nM. After 48 h, cells were seeded in 24-well plates 24 h prior to transfection. Cells were co-transfected with wild-type (WT) or mutant (MUT) PDCD4 untranslated region (UTR) constructs (pGL3-PDCD4-3’UTR-WT and pGL3-PDCD4-3’UTR-MUT, respectively) or the empty control vector pGL3-promoter pRL-TK (Promega Corporation, Madison, WI, USA) or miR-21 mimic (Guangzhou Ribobio Co., Ltd, Guangzhou, China) using Lipofectamine 2000. Relative luciferase activity was measured 48 h after transfection on a GloMax luminometer (Promega Corporation) and normalized to the firefly/Renilla luciferase signal in HEK-293T cells.

Western blotting. Cells were lysed by incubation with trypsin-EDTA solution (Invitrogen; Thermo Fisher Scientific, Inc.) and protein was extracted with radioimmunoprecipitation assay buffer (Thermo Fisher Scientific, Inc.) containing protease inhibitor. Protein concentration was quantified using a BCA Protein Assay kit (Thermo Fisher Scientific, Inc.). Equal amounts of protein (20 µg) were separated by SDS-PAGE on a 12% gel and electrophoretically transferred to a polyvinylidene difluoride membrane, which was incubated in blocking solution containing 5% non-fat milk in a Tris-buffered saline/Tween 20 solution (TBST) for 60 min at room temperature. Blocking was followed by overnight incubation at 4°C with rabbit primary antibodies against the following proteins: PDCD4 (cat. no. 9535; 1:1,000), the apoptosis regulators, B-cell lymphoma 2 (Bcl-2; cat. no. 4223; 1:1,000) and Bcl-2-associated X protein (Bax; cat. no. 5023; 1:500) (all Cell Signaling Technology, Inc., Danvers, MA, USA); and β-actin (cat. no. ab8227; 1:1,000; Abcam, Cambridge, UK). Following washing with TBST, bound antibodies were detected by incubation for 60 min at room temperature with horseradish peroxidase-labelled goat anti-rabbit immunoglobulin G (cat. no. 7074; 1:5,000; Cell Signaling Technology, Inc.). Enhanced chemiluminescence (Thermo Fisher Scientific, Inc.) was used to visualize protein bands on X-ray film. Protein expression levels were normalized to that of β-actin. Image J v1.48 U (National Institutes of Health, Bethesda, MD, USA) was used for densitometric analysis.

Dual luciferase reporter assay. HEK-293T (2x10⁵) cells were seeded in 24-well plates 24 h prior to transfection. Cells were co-transfected with wild-type (WT) or mutant (MUT) PDCD4 untranslated region (UTR) constructs (pGL3-PDCD4-3’UTR-WT and pGL3-PDCD4-3’UTR-MUT, respectively) or the empty control vector pGL3-promoter pRL-TK (Promega Corporation, Madison, WI, USA) or miR-21 mimic (Guangzhou Ribobio Co., Ltd, Guangzhou, China) using Lipofectamine 2000. Relative luciferase activity was measured 48 h after transfection on a GloMax luminometer (Promega Corporation) and normalized to the firefly/Renilla luciferase signal in HEK-293T cells.
Statistical analysis. Data were described as mean ± standard deviation. The determinations were performed at least in triplicate. An unpaired t-test and one way-analysis of variance (Bonferroni post hoc test for equal variances assumed; Tambane’s T2 post hoc test for equal variances not assumed) were used to compare the groups using GraphPad Prism version 5.0 software (GraphPad Software, Inc., La Jolla, CA, USA) and SPSS software version 22.0 (IBM Corp., Armonk, NY, USA). Two-tailed P<0.05 was considered to indicate a statistically significant difference.

Results

CM induces apoptosis and inhibits miR-21 expression in HK-2 cells. HK-2 cells were treated with 150 mg/I ml Ultravist (370 mg/I/ml) for 2 h and subsequently harvested for analysis. The rate of apoptosis was increased following CM treatment, as determined by the TUNEL assay (Fig. 1A). Consistent with this observation, the expression of the pro-apoptotic factor Bax was increased, whereas that of the anti-apoptotic factor Bcl-2 was decreased under these conditions (Fig. 1B). Additionally, compared with untreated cells, the miR-21 level was downregulated by treatment with CM, as determined by RT-qPCR analysis (Fig. 1C), suggesting a negative association between miR-21 expression and HK-2 cell apoptosis in the presence of CM.

miR-21 overexpression inhibits CM-induced apoptosis in HK-2 cells. In order to investigate the effect of miR-21 on HK-2 cell apoptosis under CM treatment, cells were transfected with miR-21 mimic or inhibitor, or a negative control miRNA. The miR-21 level was increased in cells transfected with mimic and reduced in inhibitor-treated cells, demonstrating a successful transfection (Fig. 2A). Western blot analysis revealed that Bax expression was downregulated, whereas that of Bcl-2 was upregulated, following transfection of the miR-21 mimic; the converse was observed in cells transfected with miR-21 inhibitor (Fig. 2B). Additionally, overexpression of miR-21 mimic decreased CM-induced apoptosis, whereas miR-21 inhibitor exerted the opposite effect, as determined by TUNEL assay (Fig. 2C). The results of the present study demonstrated that miR-21 may protect HK-2 cell against CM-induced apoptosis.

miR-21 inhibits HK-2 cell apoptosis by binding to the PDCD4 3’ UTR. Target gene prediction indicated that PDCD4 may be a potential target of miR-21, since the PDCD4 3’ UTR harbored a miR-21 binding site (Fig. 3A). In order to test the possibility of a miR-21 interaction with PDCD4, PDCD4 expression was evaluated in HK-2 cells transfected with miR-21 under CM treatment, using RT-qPCR analysis and western blotting. PDCD4 expression was upregulated in cells in the presence of CM (Fig. 3B and C); however, this effect was reversed by overexpression of miR-21 mimic, compared with cells transfected with negative control miR-21 mimic or those that were untransfected (Fig. 3D and E). Additionally, PDCD4 expression was increased in cells transfected with miR-21 inhibitor compared with the CM-only group, whereas the level was reduced upon transfection of miR-21 mimic (Fig. 3D and E), suggesting that miR-21 may attenuate apoptosis by inhibiting PDCD4 expression.

In order to confirm this hypothesis, constructs containing WT or MUT PDCD4 3’ UTR were generated.
miR-21 attenuates contrast-induced renal cell apoptosis via PDCD4

Figure 2. Effect of miR-21 on HK-2 cell apoptosis under CM treatment. (A) MiR-21 expression in cells transfected with miR-21 mimic, inhibitor, or negative control miR was detected using the reverse transcription-quantitative polymerase chain reaction. (B) Bcl-2 and Bax protein expression in cells transfected with miR-21 mimic, inhibitor or negative control miR was measured by western blotting. (C) Detection of apoptosis (green cells) with the TUNEL assay. Magnification, x400. Cells were treated with 150 mg/ml Ultravist in the CM groups. *P<0.05, **P<0.01 vs. CM group (n=3). CM, contrast medium; miR, microRNA; HK-2, human renal proximal tubular epithelial; TUNEL, terminal deoxynucleotidyl transferase dUTP nick-end labeling; Bcl-2, B-cell lymphoma 2; Bax, Bcl-2-associated X protein.

(Fig. 4A). The luciferase activity in HEK-293T cells transfected with pGL3-PDCD4-3’UTR-WT was reduced by ~50% relative to that in cells transfected with the MUT construct (Fig. 4B), suggesting that miR-21 may directly bind to the PDCD4 transcript, and thereby regulate its expression.

siRNA was used to knock down PDCD4 expression in HK-2 cells, and the effect of this on apoptosis was examined. PDCD4 mRNA and protein levels were decreased upon transfection of siRNA-PDCD4 (Fig. 4C and D). This decreased was accompanied by a downregulation of Bax and an upregulation of Bcl-2 compared with cells...
treated with CM only (Fig. 4D). The results of the present study demonstrated that miR-21 may protect renal cells from CM-induced apoptosis by suppressing PDCD4 expression.

Discussion

With the increasing use of CM in the diagnosis and treatment of coronary disease, CI-AKI has become the third leading cause of hospital-acquired AKI (18). Although the pathogenesis of CI-AKI is not well understood, apoptosis of renal epithelial and glomerular cells is hypothesized to be the underlying cause (19,20). Bax and Bcl-2 are two important mediators of the mitochondrial apoptosis pathway (21); Bcl-2 prevents Bax activation and thereby inhibits apoptosis. It was previously reported that Bcl-2 was downregulated in proximal renal tubular cells in AKI (22). As exhibited in the present study, Bax and Bcl-2 were increased and decreased, respectively, with a corresponding increase in the rate of apoptosis following CM treatment, consistent with previous reports of CM-induced cell apoptosis.

miRNAs contribute to kidney homeostasis and disease (23). miR-21 has been implicated in apoptosis, fibrosis, inflammation and IRI (24,25), and is considered to be a biomarker of AKI (8,26). One study demonstrated that miR-21 was upregulated 24 h subsequent to IRI in an animal model (12); this was confirmed by other investigators who reported that miR-21 expression was increased in renal proximal tubular cells 24 h following hypoxia/reoxygenation or treatment with CM (27,28). miR-21 levels have been observed to increase continuously in proliferating cells (12,25), starting at 24 h post injury (29). Therefore, its expression did not increase within 24 h reperfusion subsequent to 20 min of ischemia, or within 8 h of CM administration (13,30). It has been demonstrated that one-half of the amount of CM in the bloodstream may be eliminated quickly, in ~2 h (31). In addition, the structure of renal cells exhibited moderate changes 2 h after CM administration (32). Therefore,
Neutrophil gelatinase-associated lipocalin, a novel biomarker of CI-AKI, has been demonstrated to begin to increase at 2 h post-CM exposure (33), although the level of serum creatinine did not significantly alter. In order to observe the effect of CM on renal cells, cells were incubated with CM for 2 h in the present study, and the expression of miR-21 was downregulated. Future studies are required to investigate alterations in miR-21 expression with respect to the degree of renal injury.

miR-21 was demonstrated to be renoprotective in animal and cellular models of IRI (27,34). However, miR-21 has additionally been reported to exert deleterious effects in IRI, diabetic nephropathy and renal fibrosis (35-37). Physiological differences between models and variable times of ischemia may account for these conflicting observations. miR-21 was observed to regulate tumor cell proliferation, invasion, apoptosis and migration by targeting PDCD4 and Bcl-2 (38), while PDCD4 is thought to inhibit neoplastic transformation (39). As demonstrated in the present study, the levels of PDCD4 were reduced and enhanced upon transfection with a miR-21 mimic and inhibitor, respectively. In addition, the rate of apoptosis was decreased following PDCD4 knockdown. The results of the present study corroborated previous findings and demonstrated that PDCD4 may be negatively regulated by miR-21; additionally, these results suggested that the miR-21/PDCD4 pathway serves an important role in preventing CM-induced renal tubular cell apoptosis. To the best of our knowledge, the present study is the first to analyze the miR-21/PDCD4 pathway in vivo will be investigated in future studies.

In conclusion, the present study demonstrated that miR-21 protected renal cells against CM-induced apoptosis by directly regulating PDCD4. The results of the present study may provide a basis for the development of therapeutic strategies to treat CI-AKI.

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