

# Exosomal-miR-1184 derived from mesenchymal stem cells alleviates cisplatin-associated acute kidney injury

JINSHI ZHANG<sup>1\*</sup>, WENFANG HE<sup>1\*</sup>, DANNA ZHENG<sup>1</sup>, QIANG HE<sup>1</sup>, MINGMING TAN<sup>2</sup> and JUAN JIN<sup>2,3</sup>

Departments of <sup>1</sup>Nephrology and <sup>2</sup>Quality Management, Zhejiang Provincial People's Hospital;

<sup>3</sup>Dean's Office, Hangzhou Medical College, Hangzhou, Zhejiang 310014, P.R. China

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**Abstract.** Acute kidney injury (AKI) poses a severe threat to human health. MicroRNAs (miRNAs/miRs) are known to be involved in the progression of AKI; however, the function of miR-1184 in AKI remains unclear. Thus, the aim of the present study was to examine the role of this miRNA in kidney injury. In order to mimic AKI *in vitro*, HK-2 cells were treated with cisplatin. Bioinformatics analysis was performed to explore the differentially expressed miRNAs in AKI. A Cell Counting Kit-8 assay and flow cytometry were performed to examine cell viability and apoptosis, respectively. mRNA expression levels were detected via reverse transcription-quantitative PCR, and protein levels were investigated by western blot analysis. ELISA was performed to examine the levels of IL-1 $\beta$  and TNF- $\alpha$  in the cell supernatants. The results revealed that miR-1184 expression was downregulated in AKI. Exosomes derived from miR-1184 agomir-treated mesenchymal stem cells (MSCs) significantly reversed cisplatin-induced cell growth inhibition by inhibiting apoptosis. Moreover, forkhead box O4 (FOXO4) was found to be the direct target of miR-1184, and exosomes expressing miR-1184 notably inhibited cisplatin-induced inflammatory responses in HK-2 cells via the mediation of IL-1 $\beta$  and TNF- $\alpha$ . Furthermore, exosomes derived from miR-1184 agomir-treated MSCs significantly induced G<sub>1</sub> phase arrest in HK-2 cells via the regulation of FOXO4, p27 Kip1 and CDK2. In conclusion, the present study demonstrated that exosomal-miR-1184 derived from MSCs alleviates cisplatin-associated AKI. Thus, the findings presented herein may shed new light onto the exploration of novel strategies for the treatment of AKI.

## Introduction

Acute kidney injury (AKI) is known to be associated with a decline in kidney function within 48 h (1). It is often associated with a high mortality rate in patients, specifically in 1% of the affected population (2). Ischemia-reperfusion injury, sepsis and nephrotoxic insults are the major risk factors for AKI (3-5). Meanwhile, renal tubular epithelial cells are the main cell type in kidney tissue; therefore, the dysfunction of these cells is the main pathophysiological process of AKI (6). Studies have shown that the injury of renal tubular epithelial cells plays an important role in the pathogenesis of AKI (7,8). Moreover, the main features of AKI include inflammatory cell infiltration and the massive secretion of inflammatory factors, necrosis, apoptosis and delayed renal resident cell proliferation (9,10). Although significant efforts have been made in the treatment of AKI, patient outcomes remain dismal. Thus, there is an urgent need for the development of novel strategies for the treatment of AKI.

MicroRNAs (miRNAs/miRs) are a novel class of non-coding small ribonucleic acids, which can regulate gene expression by suppression of mRNA translation or degradation of mRNAs (11,12). Moreover, miRNAs participate in cellular process, including the growth of renal tubular epithelial cells (13,14). Meanwhile, miRNAs have been reported to be associated with AKI progression (15-17). For instance, miR-21 has been shown to inhibit the progression of AKI (18); Jiang *et al* (19) found that miR-500a-3p alleviated kidney injury by targeting mixed lineage kinase domain like pseudokinase. However, other miRNAs associated with the progression of AKI warrant further investigation.

Exosomes are microvesicles ranging from 70 to 120 nm in diameter and are derived from multivesicular bodies (20). In addition, exosomes participate in cell communication by transferring proteins and nucleic acids, and this function can lead to the mediation of intercellular communication (21,22). In recent studies, a number of exosomal proteins, miRNAs and lncRNAs have been reported to promote the progression of AKI. For example, Cao *et al* (14) found that exosomal miR-125b-5p deriving from mesenchymal stem cells (MSCs) could promote tubular repair by suppression of p53 in ischemic AKI; Zhang *et al* (23) indicated that endothelial progenitor cells-derived exosomal miR-21-5p could alleviate sepsis-induced AKI by inhibiting runt-related transcription

*Correspondence to:* Dr Mingming Tan or Dr Juan Jin, Department of Quality Management, Zhejiang Provincial People's Hospital, 158 Shangtang Road, Hangzhou, Zhejiang 310014, P.R. China  
E-mail: mingmingtan@yandex.com  
E-mail: jinjuan\_12@126.com

\*Contributed equally

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factor 1 expression. However, the function of exosomes in AKI needs to be further explored. On the other hand, among the potential targets of miR-1184, forkhead box O4 (FOXO4) has been found to be associated with cell growth (24). Thus, the present study focused on the relationship between miR-1184 and the FOXO signaling pathway.

On the other hand, MSCs, characterized by the abilities of self-renewal, differentiation, immunomodulation and trophic support, are essential in regenerative medicine owing to the capacity to create a microenvironment conducive to the repair of injured tissues (14). Previous studies indicated that exosomes derived from MSCs have been proposed as an alternative to MSC-based therapy for several diseases (14,25). In addition, MSC-derived exosomes are known to be involved in AKI progression. For instance, Cao *et al* (26) found that exosomes derived from MSCs (MSC-exos) could significantly attenuate cisplatin-induced murine AKI through inhibiting inflammation; Ji *et al* (25) demonstrated that platelet-rich plasma could promote MSC-derived exosome paracrine signaling to repair AKI via the AKT/Rab27 pathway. Thus, exosomes derived from MSCs can play a vital role in AKI progression.

Based on this background, the present study aimed to detect the differentially expressed miRNAs closely associated with the progression of AKI. The findings presented herein may provide new insight on the role of miRNAs in AKI and may aid in the development of novel treatment methods for AKI.

## Materials and methods

**Cell culture and treatment.** HK-2 cell lines (American Type Culture Collection) and MSCs (American Type Culture Collection) were maintained in RPMI-1640 medium (Thermo Fisher Scientific, Inc.), supplemented with 10% FBS (Gibco; Thermo Fisher Scientific, Inc.), 1% penicillin (Thermo Fisher Scientific, Inc.) and 1% streptomycin (Thermo Fisher Scientific, Inc.) in a condition with 5% CO<sub>2</sub> and 37°C. To mimic AKI *in vitro*, HK-2 cells were treated with 20 µM cisplatin (MedChemExpress) for 48 h according to previous refs. (8,27).

**Cell transfection.** HK-2 cells or MSCs (5x10<sup>3</sup> cells/well) were transfected with miR-1184 agomir or agomir-negative control (NC) using Lipofectamine® 2000 (Thermo Fisher Scientific, Inc.) at 37°C for 48 h. miR-1184 agomir (50 nM) and agomir-NC (50 nM) were obtained from Shanghai GenePharma Co., Ltd. The sequences were as follows: miR-1184 agomir, 5'-CCU UCGGUAGUUCAGCGACGUCC-3' and agomir-NC, 5'-UUC UCCGAACGUGUCACGUUU-3'. After 48 h of transfection, cells were used in subsequent experiments.

For FOXO4 overexpression, MSCs (5x10<sup>3</sup> cells/well) were transfected with pcDNA3.1 (1 µg/µl; Shanghai GenePharma Co., Ltd.) or pcDNA3.1-FOXO4 (1 µg/µl; Shanghai GenePharma Co., Ltd.) using Lipofectamine 2000 for 48 h at 37°C, according to the manufacturer's instructions. After 48 h of transfection, cells were used in subsequent experiments.

**Bioinformatics analysis.** The differentially expressed miRNAs were presented in a volcano plot and a heatmap using the

GSE53771 dataset (28) from the Gene Expression Omnibus (GEO) database (<https://www.ncbi.nlm.nih.gov/geo>). In addition, the differentially expressed miRNAs were screened using R analysis (29). Using a t-test, P<0.05 and fold-change >1.5 or <0.667 were considered as differentially expressed miRNAs. On the other hand, the functions of miRNA-targeted mRNAs in terms of 'Cellular Components' and 'Biological Processes' were investigated by Gene Ontology (GO) analysis (<http://www.geneontology.org>). Pathway analysis was performed to define biological pathways by using the KEGG orthology-based annotation system (KOBAS; version 3.0; <http://kobas.cbi.pku.edu.cn/index.php>).

**Prediction of miRNA downstream target.** The downstream target of miR-1184 was predicted using TargetScan (version 7.2; [http://www.targetscan.org/vert\\_72/](http://www.targetscan.org/vert_72/)) and miRDB (version 2.0; <http://www.mirdb.org/>).

**Exosome extraction and identification.** Briefly, MSCs were maintained in RPMI-1640 medium until they reached 80% confluence. The medium was then replaced with the serum-free medium. The supernatants were centrifuged for 1 h (300 x g for 15 min at 4°C, 2,000 x g for 15 min at 4°C and 10,000 x g for 30 min at 4°C) following 48 h of culture. Subsequently, the supernatants were filtrated and collected to extract exosomes using ultracentrifugation (2,000 x g for 15 min at 4°C). Western blot analysis was used to detect the exosome isolation, and the detailed protocol was in accordance with a previous study (25). The particle sizes were detected by Nanoparticle tracking analysis (NTA).

**NTA.** A total of ~0.3 ml supernatant was loaded into the sample chamber of an LM10 NanoSight unit (NanoSight, Ltd.) and three videos of either 30 or 60 sec were recorded of each sample. Data analysis was performed using NTA 2.1 software (NanoSight, Ltd.). In NTA, the paths of unlabeled particles acting as point scatterers, undergoing Brownian motion in a 0.25 ml chamber through which a 635-nm laser beam was passed, was determined from a video recording, with the mean squared displacement determined for each possible particle. The diffusion coefficient and sphere-equivalent hydrodynamic radius were subsequently determined using the Stokes-Einstein equation (25).

**Fluorescence staining.** MSCs (5x10<sup>4</sup> per well) were seeded overnight. Subsequently, cells were labeled for 24 h at 4°C with phalloidin (1:1,000; Abcam; cat. no. ab176753) or PKH26 red membrane dye (1:1,000; Biolab Co., Ltd.; cat. no. HR9070). The nuclei were stained with 5 µl/ml DAPI (Beyotime Institute of Biotechnology). The results were observed under a fluorescence microscope (magnification, x200; Olympus Corporation).

**Western blot analysis.** RIPA lysis buffer (Beyotime Institute of Biotechnology) was used to extract total protein from the HK-2 cells. A BCA protein kit (Thermo Fisher Scientific, Inc.) was used to quantify the total protein. SDS-PAGE (10%) was used to separate the protein (40 µg per lane), and the protein was then transferred to PVDF membranes (Thermo Fisher Scientific, Inc.). After blocking with 5% skimmed milk at room

temperature for 1 h, the membranes were incubated overnight at 4°C with anti-CD63 (cat. no. ab134045; 1:1,000), anti-CD81 (cat. no. ab109201; 1:1,000), anti-Bax (cat. no. ab32503; 1:1,000), anti-Bcl-2 (cat. no. ab32124; 1:1,000), anti-cleaved caspase-3 (cat. no. ab32042; 1:1,000), anti-TSG101 (cat. no. ab125011; 1:1,000), anti-FOXO4 (cat. no. ab128908; 1:1,000), anti-p27 Kip1 (cat. no. ab32034; 1:1,000), anti-CDK2 (cat. no. ab32147; 1:1,000) and anti- $\beta$ -actin (cat. no. ab8227; 1:1,000) primary antibodies (all Abcam). Subsequently, the membranes were incubated with secondary anti-rabbit antibodies (HRP-conjugated; Abcam; cat. no. ab7090; 1:5,000) for 1 h at room temperature. Protein bands were visualized using an ECL kit (Thermo Fisher Scientific, Inc.).  $\beta$ -actin was used as the loading control. All antibodies were purchased from Abcam. ImageJ software (version 6.0; National Institutes of Health) was used for densitometry.

**Reverse transcription-quantitative PCR (RT-qPCR).** TRIzol<sup>®</sup> reagent (Invitrogen; Thermo Fisher Scientific, Inc.) was used to isolate total RNA from HK-2 cells or MSCs. The PrimeScript RT Reagent Kit (Takara Bio, Inc.) was used to reverse transcribe total RNA into cDNA, according to the manufacturer's protocol. Subsequently, qPCR was performed using the SYBR Premix Ex Taq II kit (ELK Biotechnology). RT-qPCR reactions were performed under the following protocol: Initial denaturation for 2 min at 94°C, followed by 35 cycles (30 sec at 94°C and 45 sec at 55°C). The following primer pairs were used for RT-qPCR: miR-1184 forward, 5'-CTGGACTGAGCC GTGCTAC-3' and reverse, 5'-CTCAACTGGTGTCGTGGA GTC-3'; and U6 forward, 5'-CTCGCTTCGGCAGCACAT-3' and reverse, 5'-AACGCTTCACGAATTTGCGT-3'. The  $2^{-\Delta\Delta C_q}$  method (30) was used to quantify the data. U6 was used as an internal control.

**Cell counting kit-8 (CCK-8) assay.** HK-2 cells ( $5 \times 10^3$  cells/well) were seeded and treated with cisplatin (20  $\mu$ M), MSC-Exo<sup>miR-1184 agomir</sup> or cisplatin + MSC-Exo<sup>miR-1184 agomir</sup> for 72 h at 37°C. Subsequently, CCK-8 reagent (10  $\mu$ l; Beyotime Institute of Biotechnology) was added to the cells for 2 h at 37°C. The absorbance (450 nm) was measured using a microplate reader (Thermo Fisher Scientific, Inc.).

**Cell apoptosis analysis.** HK-2 cells were trypsinized, washed with PBS, resuspended in Annexin V Binding Buffer, and stained with 5  $\mu$ l FITC and 5  $\mu$ l PI for 15 min in the dark. The cells were analyzed using a flow cytometer (FACSlyric<sup>™</sup>; BD Biosciences) to assess the incidence of cell apoptosis (early + late apoptosis). FlowJo (version 10.6.2; FlowJo LLC) was used to analyze the data.

**Dual-luciferase reporter assay.** The FOXO4 3'-untranslated region (UTR) containing putative miR-1184 binding sites were obtained from Sangon Biotech Co., Ltd., and cloned into pmirGLO Dual-Luciferase miRNA Target Expression Vectors (Promega Corporation) to construct FOXO4 wild-type (WT) or mutant (MUT) reporter vectors. The mutated 3'-UTR was generated using a site directed mutagenesis kit (Sangon Biotech Co., Ltd.). FOXO4 (WT) or FOXO4 (MUT) were transfected into HK-2 cells ( $5 \times 10^3$ ) together with NC or miR-1184 agomir using Lipofectamine 2000. After 48 h of transfection, relative

luciferase activities were then analyzed using a Dual-Glo Luciferase Assay System (Promega Corporation). *Renilla* luciferase activity was used for normalization.

**ELISA.** HK-2 cell supernatants were collected by centrifugation (500 x g, 10 min, 4°C). Subsequently, the levels of TNF- $\alpha$  (cat. no. H052-1) and IL-1 $\beta$  (cat. no. H002) were investigated using ELISA kits (Nanjing Jiancheng Bioengineering Institute), according to the manufacturers' protocols.

**Cell cycle analysis.** In brief, HK-2 cells were harvested, fixed with 75% ethanol on ice for 20 min, permeabilized with 0.25% Triton X-100 and stained with PI/RNase (BD Pharmingen; BD Biosciences). Following incubation at 4°C for 15 min, cells were analyzed using a flow cytometer (BD FACSaria III; BD Biosciences). The data were quantified using FlowJo software (version 3.0; FlowJo LLC).

**Statistical analysis.** Each group was examined in three independent experiments and all data are expressed as the mean  $\pm$  SD. Western blot analysis, RT-qPCR, flow cytometry, CCK-8 assay and immunofluorescence staining were repeated three times. An unpaired Student's t-test was used to analyze the differences between two groups, and one-way ANOVA followed by Tukey's test were used to analyze differences among multiple groups (>2 groups, using GraphPad Prism 7; GraphPad Software, Inc.).  $P < 0.05$  was considered to indicate a statistically significant difference.

## Results

**Differentially expressed miRNAs in AKI.** To detect differentially expressed miRNAs in AKI, bioinformatics analysis was performed. As indicated in Fig. 1A and B, differentially expressed miRNAs (miR-1269a, miR-1184, miR-299-5p, miR-411, miR-451 and miR-499) are presented using a volcano plot and heatmap. In addition, the six differentially expressed miRNAs (miR-1269a, miR-1184, miR-299-5p, miR-411, miR-451 and miR-499; miR-1269a and miR-1184 were downregulated, while the other four miRNAs were upregulated) in AKI are presented (Fig. 1C). Moreover, GO and pathway analyses were performed to determine the most common 'Biological Process' of miR-1184. The results indicated that miR-1184 was enriched in the following 'Cellular Component' terms: 'Brush border membrane', 'main axon', 'platelet dense granule', 'membrane region', 'pigment granule', 'melanosome', 'myelin sheath', 'membrane raft', 'membrane microdomain' and 'microfibril' (Fig. 2A). miR-1184 was enriched in the following 'Molecular Function' terms: 'Protein domain specific binding', 'nucleocytoplasmic carrier activity', 'disordered domain specific binding', ' $\beta$ -catenin binding', 'protein C-terminus binding', 'nuclear import signal receptor activity', 'RNA polymerase II general transcription initiation factor binding', 'DNA-binding transcription activator activity' and 'RNA polymerase II-specific' (Fig. 2A). miR-1184 was enriched in the following 'Biological Process' terms: 'Modulation of process of other organism involved in symbiotic interaction', 'modulation by symbiont of host process', 'melanocyte differentiation', 'modulation of process of other organism',

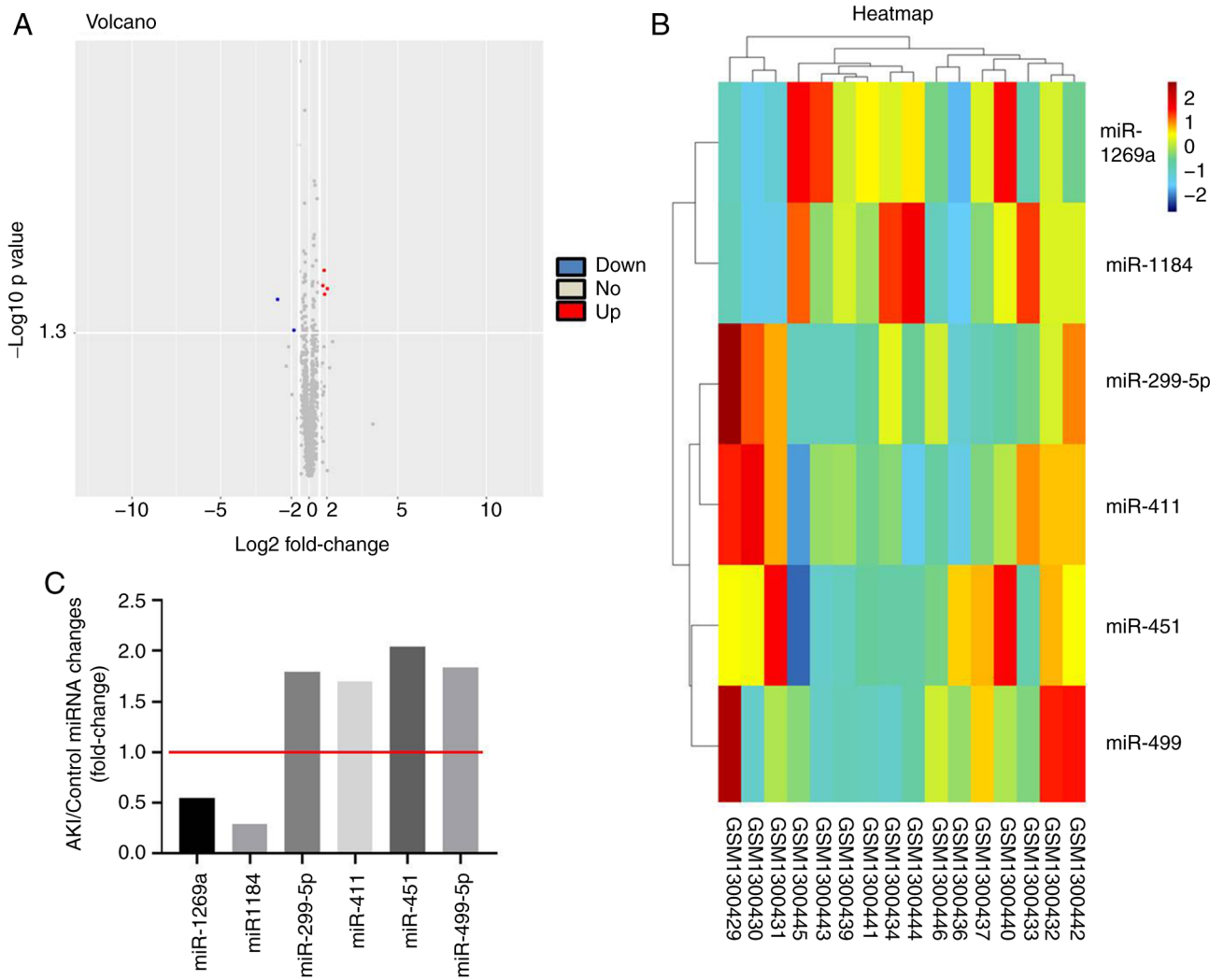


Figure 1. Differentially expressed miRNAs in AKI. (A) Volcano plots and (B) heatmap illustrates the miRNAs differentially expressed in AKI. (C) The AKI/Control ratio of miR-1184, miR-1269a, miR-299-5p, miR-411, miR-451 and miR-499-5p was calculated. miR/miRNA, microRNA; AKI, acute kidney injury.

‘gland morphogenesis’, ‘modulation by virus of host process’, ‘pigmentation’, ‘pigment cell differentiation’, ‘regulation of DNA-binding transcription factor activity’ and ‘regulation of viral process’ (Fig. 2A). Notably, the most enriched ‘Cellular Component’ was ‘brush border membrane’, the most enriched ‘Molecular Function’ was ‘protein domain specific binding’ and the most enriched ‘Biological Process’ was ‘modulation of process of other organism involved in symbiotic interaction’ (Fig. 2A). Pathway analysis revealed that the ‘Hedgehog signaling pathway’, ‘pathways in cancer’, ‘sphingolipid signaling pathway’, ‘antifolate resistance’, ‘oocyte meiosis’, ‘TGF-beta signaling pathway’, ‘FOXO signaling pathway’, ‘prostate cancer’, ‘gastric cancer’, ‘pantothenate and CoA biosynthesis’, ‘cholinergic synapse’ and ‘cysteine and methionine metabolism’ were associated with the development of AKI (Fig. 2B). Based on the aforementioned data, miR-1184 was selected for further analysis.

**Successful isolation of exosomes from MSCs.** In order to detect the efficiency of cell transfection and exosome isolation, RT-qPCR and NTA was performed. As shown in

Fig. 3A, the expression of miR-1184 in MSCs was significantly upregulated by miR-1184 agomir, and NTA revealed a similar size distribution of exosomes (Fig. 3B). Moreover, exosomal proteins (TSG101, CD63 and CD81) were highly expressed in exosomes derived from MSCs, while they were expressed at low levels in MSCs (Fig. 3C). Moreover, the expression of miR-1184 in exosomes derived from MSCs was significantly upregulated by miR-1184 agomir (Fig. 3D), and MSC-derived exosomes labeled with fluorescent PKH26 were internalized by unstained MSCs (Fig. 3E). Furthermore, the expression level of miR-1184 in MSCs was significantly increased by exosomes derived from miR-1184 agomir-treated MSCs (Fig. 3F). Taken together, the data demonstrated that exosomes were successfully isolated from MSCs.

*Exosomes derived from miR-1184 agomir-treated MSCs significantly reverse cisplatin-induced HK-2 cell growth inhibition.* In order to detect the effects of exosomes on AKI *in vitro*, a CCK-8 assay was used. As illustrated in Fig. 4A, the viability of the HK-2 cells was significantly increased when the cells were co-cultured with exosomes derived from miR-1184



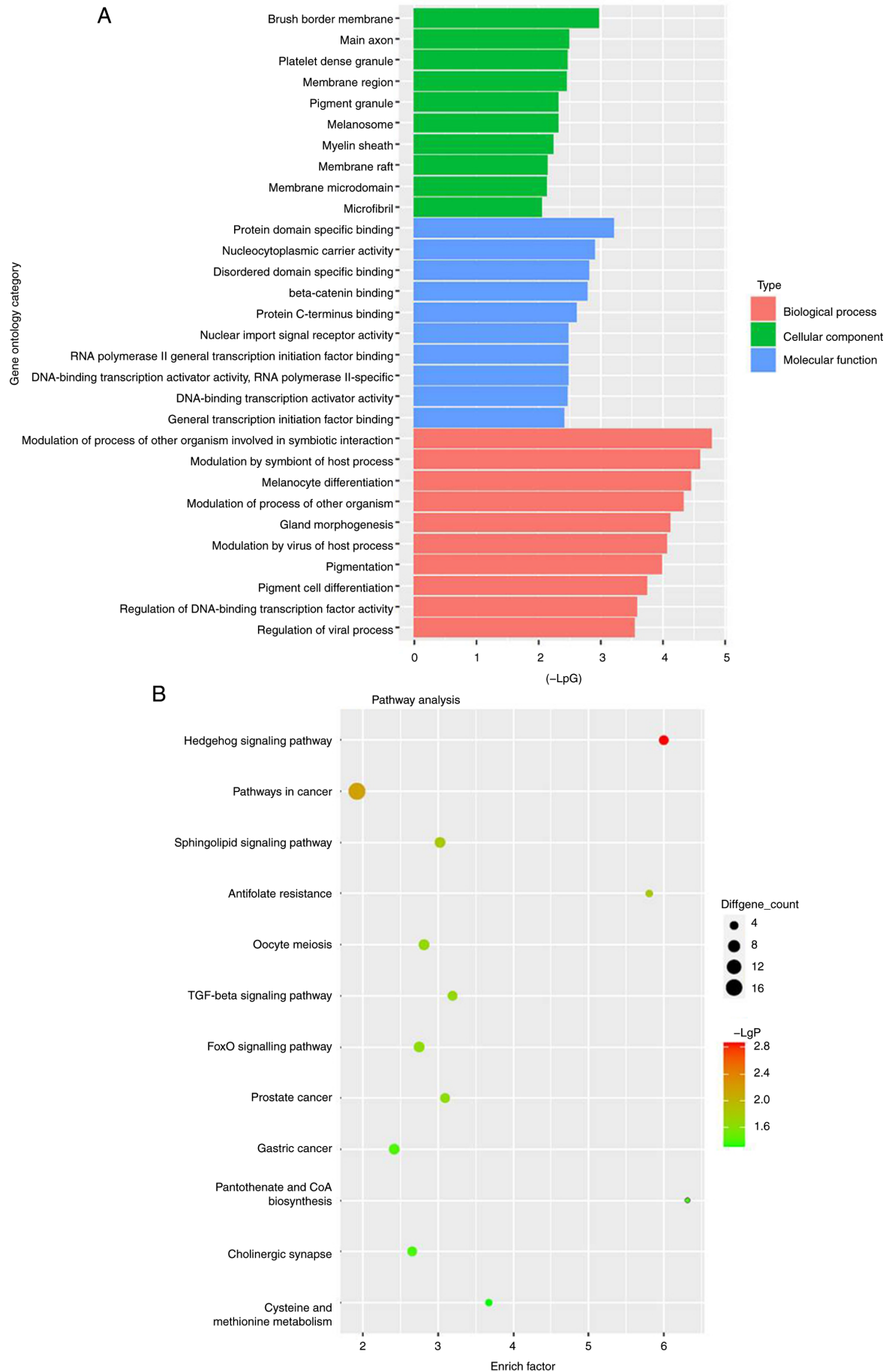


Figure 2. Profile of miR-1184 in AKI analyzed with GO and pathway analyses. (A) Go analysis exploring the potential functions of differentially expressed miRNAs. (B) Pathway analysis exploring the signaling pathways related to acute kidney injury. miRNA, microRNA; GO, Gene Ontology.

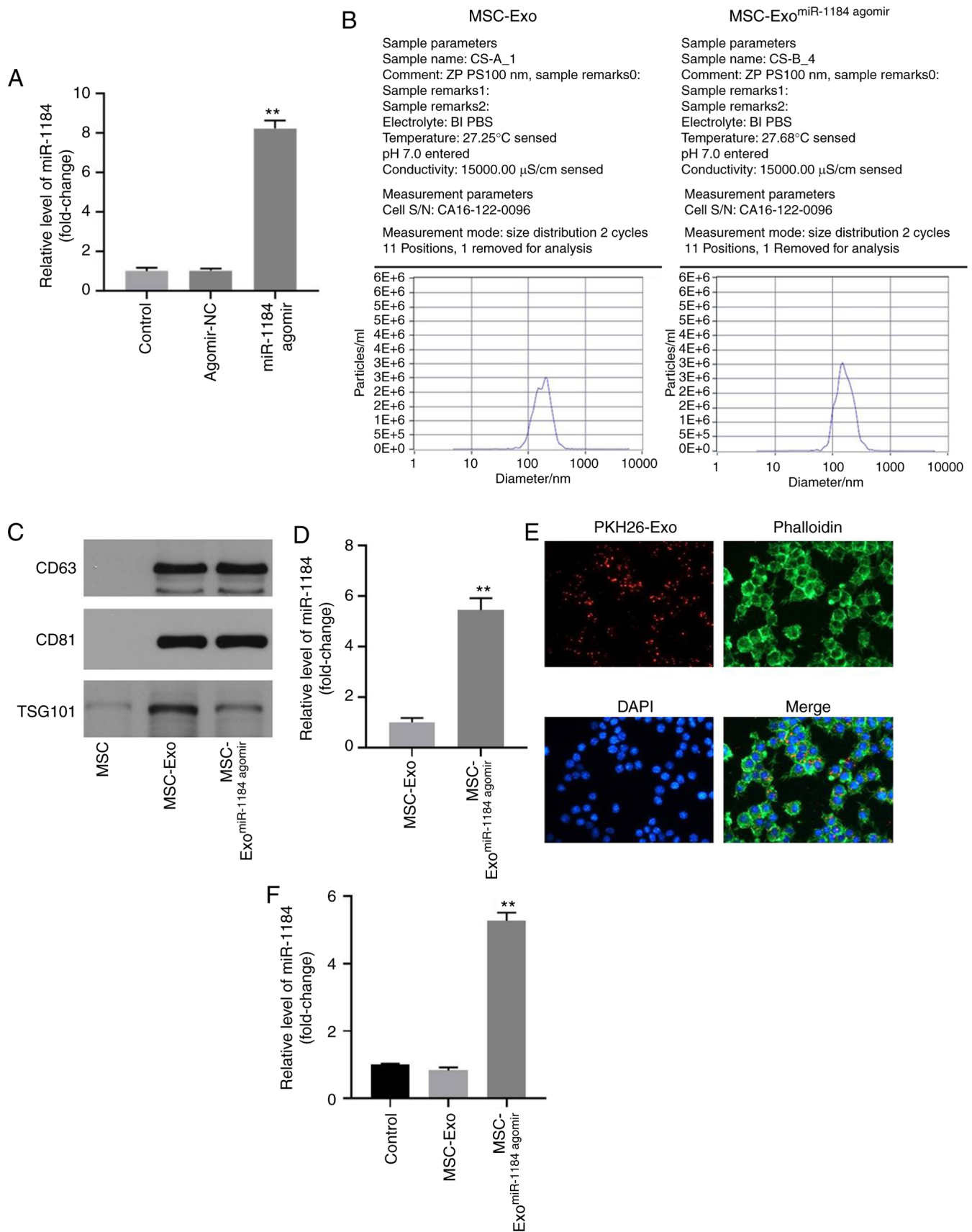


Figure 3. Exosomes were successfully isolated from MSCs. (A) MSCs were transfected with miR-1184 agomir or agomir-NC. Then, the expression of miR-1184 in MSCs was detected by RT-qPCR. \*\* $P < 0.01$  vs. control. (B) The particle sizes of exosomes were measured by Nanoparticle tracking analysis. (C) The expression levels of CD63, CD81 and TSG101 in MSCs, MSC-Exo or MSC-Exo<sup>miR-1184 agomir</sup> were detected by western blotting. (D) The expression of miR-1184 in MSC-Exo or MSC-Exo<sup>miR-1184 agomir</sup> was detected by RT-qPCR. \*\* $P < 0.01$  vs. MSC-Exo. (E) The location of exosomes was observed by immunofluorescence staining. (F) MSCs were co-cultured with MSC-Exo or MSC-Exo<sup>miR-1184 agomir</sup>. Then, the expression of miR-1184 in MSCs was tested by RT-qPCR. \*\* $P < 0.01$  vs. control. MSCs, mesenchymal stem cells; miR, microRNA; NC, negative control; RT-qPCR, reverse transcription-quantitative PCR; MSC-Exo, exosomes derived from MSCs; MSC-Exo<sup>miR-1184 agomir</sup>, exosomes derived from miR-1184 agomir-treated MSCs.

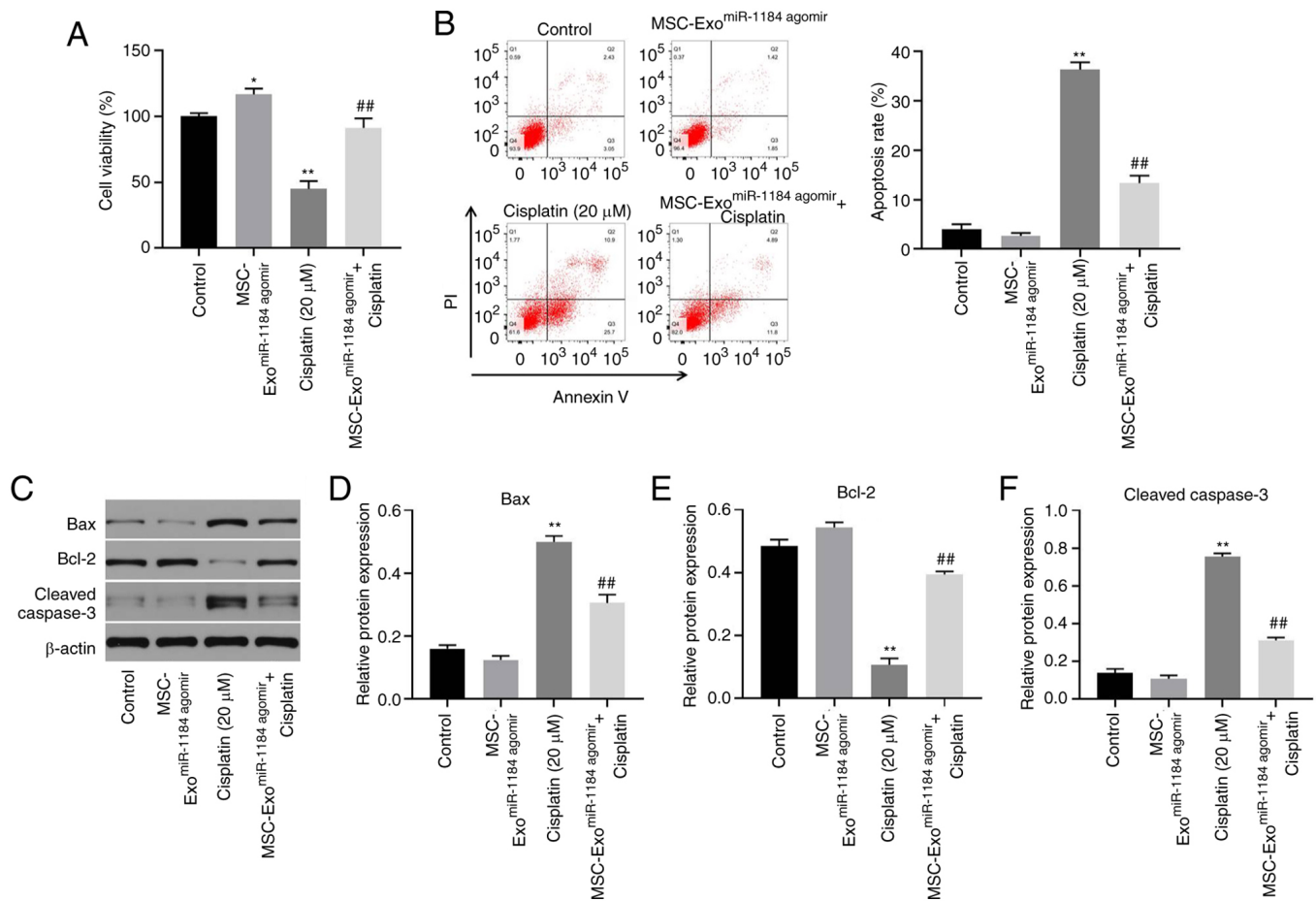


Figure 4. MSC-Exo<sup>miR-1184 agomir</sup> significantly reverses cisplatin-induced HK-2 cell growth inhibition. (A) HK-2 cells were treated with cisplatin, MSC-Exo<sup>miR-1184 agomir</sup> or cisplatin + MSC-Exo<sup>miR-1184 agomir</sup>. The viability of HK-2 cells was tested by Cell Counting Kit-8 assay. (B) The apoptosis of HK-2 cells was tested by flow cytometry. (C) The protein levels of Bax, Bcl-2 and cleaved caspase-3 in HK-2 cells were detected by western blotting. (D-F) The relative expression levels were semi-quantified by normalization to β-actin. \*P<0.05, \*\*P<0.01 vs. control; ##P<0.01 vs. cisplatin. MSCs, mesenchymal stem cells; miR, microRNA; MSC-Exo<sup>miR-1184 agomir</sup>, exosomes derived from miR-1184 agomir-treated MSCs.

agomir-treated MSCs, and exosomes expressing miR-1184 significantly reversed the cisplatin-induced inhibition of cell viability. In addition, cisplatin significantly induced HK-2 cell apoptosis, while this phenomenon was reversed in the presence of exosomes derived from miR-1184 agomir-treated MSCs (Fig. 4B). Moreover, cisplatin significantly upregulated the expression levels of Bax and cleaved caspase-3 and downregulated the protein expression level of Bcl-2, while this phenomenon was reversed by exosomes derived from miR-1184 agomir-treated MSCs (Fig. 4C-F). Taken together, the results demonstrated that exosomes derived from miR-1184 agomir-treated MSCs significantly reversed cisplatin-induced HK-2 cell growth inhibition.

**miR-1184 directly targets FOXO4 in HK-2 cells.** To explore the potential target of miR-1184, TargetScan and miRDB databases were used. As shown in Fig. 5A, FOXO4 may be the target of miR-1184, and the relative luciferase activity in the WT-FOXO4 group was significantly decreased by miR-1184 agomir (Fig. 5B). Moreover, the expression level of FOXO4 in HK-2 cells was significantly inhibited by the overexpression of miR-1184 (Fig. 5C). Furthermore, the levels of IL-1β and TNF-α in the supernatants of HK-2 cells were significantly upregulated by cisplatin, while this phenomenon was partially

reversed by MSC-Exo<sup>miR-1184 agomir</sup> (Fig. 5D and E). Therefore, miR-1184 directly targeted FOXO4 in HK-2 cells.

**Exosomes derived from miR-1184 agomir-treated MSCs significantly induce G<sub>1</sub> arrest in HK-2 cells via the mediation of FOXO1, p27 Kip1 and CDK2.** In order to further explore the underlying mechanisms through which exosomes or miR-1184 agomir regulate cisplatin-induced HK-2 cell injury, western blot analysis was performed. As indicated in Figs. 6A-C and S1A-C the protein expression levels of FOXO1 and p27 Kip1 in HK-2 cells were significantly upregulated by cisplatin. By contrast, cisplatin significantly inhibited the expression level of CDK2 (Figs. 6D and S1D). Furthermore, the effect of cisplatin on these three proteins was reversed by MSC-Exo<sup>miR-1184 agomir</sup> (Figs. 6A-D and S1A-D). In addition, cisplatin-induced G<sub>1</sub> arrest was reversed by MSC-Exo<sup>miR-1184 agomir</sup> (Fig. 6E). The expression of FOXO4 in MSCs was significantly upregulated by transfection with the FOXO4-overexpression vector (Fig. 6F). Meanwhile, the anti-inflammatory effect of exosomes derived from miR-1184 agomir-treated MSCs on cisplatin-treated HK-2 cells was rescued in the presence of exosomes derived from MSCs co-treated with miR-1184 agomir and FOXO4 overexpression (Fig. 6G). Collectively, the results demonstrated that exosomes

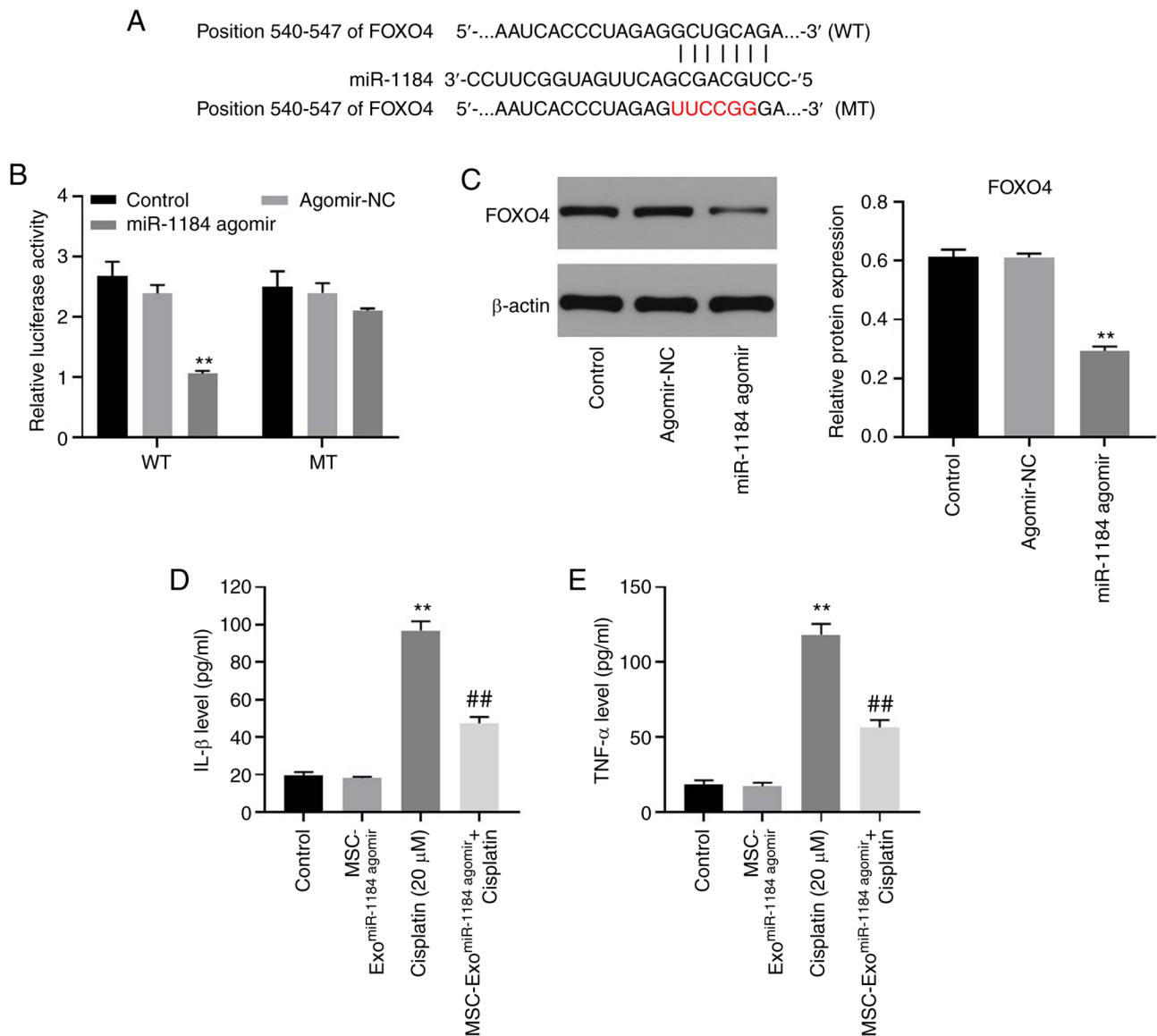


Figure 5. miR-1184 directly targets FOXO4 in HK-2 cells. (A) The target of miR-1184 was predicted by TargetScan and miRDB. (B) The relative luciferase activity was detected by dual-luciferase reporter assay. (C) HK-2 cells were transfected with NC or miR-1184 agomir. The expression of FOXO4 in HK-2 cells was detected by western blotting. The relative expression was semi-quantified by normalization to  $\beta$ -actin. (D) The level of IL- $\beta$  in supernatants of HK-2 cells was tested by ELISA. (E) The level of TNF- $\alpha$  in supernatants of HK-2 cells was tested by ELISA. \*\* $P < 0.01$  vs. control; ## $P < 0.01$  vs. cisplatin. miR, microRNA; FOXO4, forkhead box O4; NC, negative control; WT, wild-type; MT, mutant; MSCs, mesenchymal stem cells; MSC-Exo<sup>miR-1184 agomir</sup>, exosomes derived from miR-1184 agomir-treated MSCs.

derived from miR-1184 agomir-treated MSCs significantly induced G<sub>1</sub> arrest in HK-2 cells via the mediation of FOXO1, p27 Kip1 and CDK2.

## Discussion

It has been reported that miRNAs are involved in the progression of AKI (19,31,32). In the present study, it was found that miR-1184 was downregulated in cisplatin-treated HK-2 cells, and that exosomes derived from miR-1184 agomir-treated HK-2 cells significantly reversed cisplatin-induced HK-2 cell injury. Thus, the present study firstly explored the function of miR-1184 in AKI, suggesting that miR-1184 may function as an inhibitor in AKI. Moreover, miR-1184 is known to be involved in other diseases. For example, Wang *et al* (33) indicated that hsa\_circ\_0128846 promoted the tumorigenesis of colorectal

cancer by binding to miR-1184 and releasing ajuba LIM protein, and inactivating Hippo/Yes1 associated transcriptional regulator signaling. Chen *et al* (34) found that miR-1184 mediated the proliferation of colon cancer cells by targeting casein kinase 2  $\alpha$  1. Thus, other functions of miR-1184 (for example its role in renal cancer) also need to be explored in the future.

It has been reported that exosomes derived from MSCs play important roles in the progression of AKI. For example, Alzahrani (35) found that exosomes derived from MSCs could promote the progression of AKI; Cao *et al* (14) indicated that exosomal miR-125b-5p derived from MSCs could promote tubular repair by suppression of p53 in ischemic AKI. In the present study, it was found that exosomal miR-1184 derived from MSCs could inhibit cisplatin-induced HK-2 cell injury. Thus, this research first explored the relationship between miR-1184 and exosomes derived from MSCs.



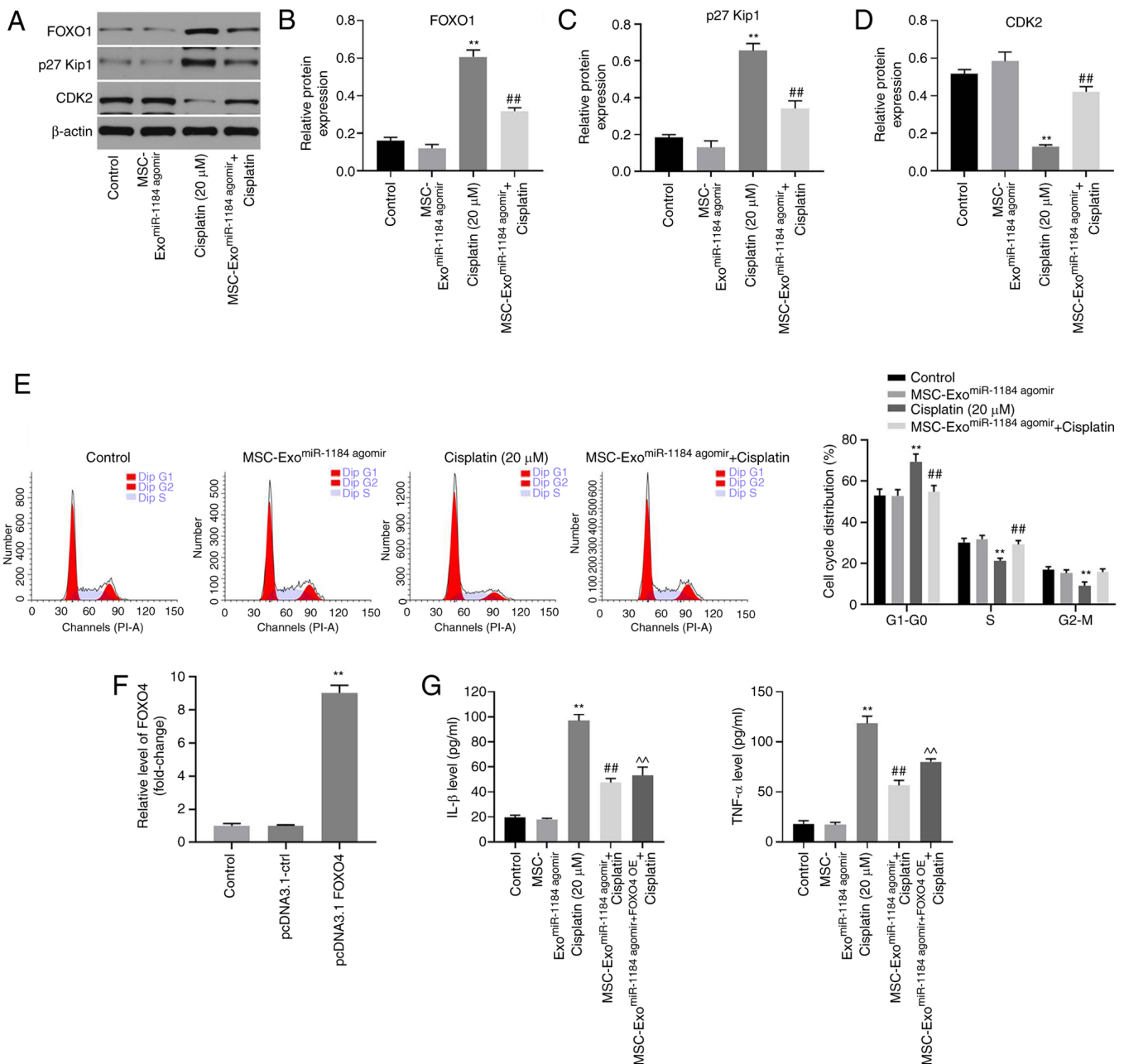


Figure 6. MSC-Exo<sup>miR-1184 agomir</sup> significantly induces G1 arrest in HK-2 cells via mediation of FOXO1, p27 Kip1 and CDK2. (A-D) The expression levels of FOXO1, p27 Kip1 and CDK2 in HK-2 cells were detected by western blotting. The relative expression levels were semi-quantified by normalization to  $\beta$ -actin. (E) The cell cycle distribution was tested by flow cytometry. (F) MSCs were transfected with pcDNA3.1-ctrl or pcDNA3.1-FOXO4. The expression of FOXO4 in MSCs was detected via reverse transcription-quantitative PCR. (G) The levels of IL-1 $\beta$  and TNF- $\alpha$  in supernatants of HK-2 cells were tested by ELISA. <sup>\*\*</sup>P<0.01 vs. control; <sup>##</sup>P<0.01 vs. cisplatin; <sup>^</sup>P<0.01 vs. MSC-Exo<sup>miR-1184 agomir</sup> + cisplatin. MSCs, mesenchymal stem cells; MSC-Exo<sup>miR-1184 agomir</sup>, exosomes derived from miR-1184 agomir-treated MSCs; FOXO1, forkhead box O1; FOXO4, forkhead box O4; OE, overexpression.

It has been confirmed that miRNAs exert their biological functions due to their target genes (36). In the present study, the results of the dual-luciferase reporter assay indicated that FOXO4 was a downstream target of miR-1184 in AKI. FOXO4 is a member of the FOXO family, and is a crucial mediator of cell proliferation (37). Consistently, the data of the present study revealed that miR-1184 may function as a mediator in AKI by targeting FOXO4. On the other hand, it has been reported that p27 Kip1 is a cell cycle regulator, firstly regarded as a cyclin-dependent kinase antagonist (38). It has been reported that p27 Kip1 expression is upregulated

during the progression of AKI (39,40). Taken together with the findings of the present study, it can be concluded that p27 Kip1 is a negative mediator in AKI. Additionally, CDK2 is a mediator of cell cycle distribution and is a downstream protein of p27 Kip1 (41). In the present study, exosomes derived from miR-1184 agomir-treated HK-2 cells reversed cisplatin-induced cell growth inhibition via the mediation of FOXO1, p27 Kip1 and CDK2. Thus, these findings are consistent with those of previous studies. Taken together, the findings presented herein revealed that miR-1184 exerted an inhibitory effect on AKI by targeting FOXO1.

There are some limitations of the present study, which are as follows: i) Other mRNAs targeted by miR-1184 in AKI need to be explored in the future; ii) rescue experiments should be performed in order to further verify the function of exosomes in AKI; iii) electron microscopy analysis is needed to further identify the exosomes; iv) the function of miR-1184 antagomir should be further confirmed; and v) the function of miR-1184 in renal cancer needs to be explored. Therefore, further investigations are required in the future.

In conclusion, the present study demonstrated that exosomal-miR-1184 derived from MSCs alleviated cisplatin-associated AKI. Thus, these findings may lead to the development of novel strategies for the treatment of AKI.

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## Availability of data and materials

All data generated or analyzed during this study are included in this published article.

## Authors' contributions

MT and JJ conceived and supervised the present study. JZ and WH designed the experiments and reviewed the results. DZ and QH performed the experiments. All authors have read and approved the final manuscript. MT and JJ confirm the authenticity of all the raw data.

## Ethics approval and consent to participate

Not applicable.

## Patient consent for publication

Not applicable.

## Competing interests

The authors declare that they have no competing interests.

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