Research Paper

Novel biomarkers for post-contrast acute kidney injury identified from long non-coding RNA expression profiles

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Abstract

Background: Post-contrast acute kidney injury (PC-AKI) is a severe complication of cardiac catheterization. Emerging evidence indicated that long non-coding RNAs (lncRNAs) could serve as biomarkers for various diseases. However, the lncRNA expression profile and potential biomarkers in PC-AKI remain unclear. This study aimed to investigate novel lncRNA biomarkers for the early detection of PC-AKI.

Methods: lncRNA profile in the kidney tissues of PC-AKI rats was evaluated through RNA sequencing. Potential lncRNA biomarkers were identified through human-rat homology analysis, kidney and blood filtering in rats and verified in 112 clinical samples. The expression patterns of the candidate lncRNAs were detected in HK-2 cells and rat models to evaluate their potential for early detection.

Results: In total, 357 lncRNAs were found to be differentially expressed in PC-AKI. We identified lnc-HILPDA and lnc-PRND were conservative and remarkably upregulated in both kidneys and blood from rats and the blood of PC-AKI patients; these lncRNAs can precisely distinguish PC-AKI patients (area under the curve (AUC) values of 0.885 and 0.875, respectively). The combination of these two lncRNAs exhibited improved accuracy for predicting PC-AKI, with 100% sensitivity and 83.93% specificity. Time-course experiments showed that the significant difference was first noted in the blood of PC-AKI rats at 12 h for lnc-HILPDA and 24 h for lnc-PRND.

Conclusion: Our study revealed that lnc-HILPDA and lnc-PRND may serve as the novel biomarkers for early detection and profoundly affect the clinical stratification and strategy guidance of PC-AKI.

Key words: post-contrast acute kidney injury; long non-coding RNA; biomarker; bioinformatics

Introduction

Post-contrast acute kidney injury (PC-AKI), also called contrast-induced acute kidney injury (CI-AKI), is defined as acute kidney injury that occurs after intravascular administration of iodine-based contrast media [1, 2]. PC-AKI, which is the third leading cause of hospital-acquired AKI, occurs in 5% to 35.8% of the 80 million people worldwide who receive an iodinated contrast agent and even up to 50% in high-risk patients [3–5]. Besides, after a cardiac catheterization, PC-AKI confers an unfavorable prognosis, resulting in 3-fold and 7-fold increases in short-term and long-term mortality, respectively [6, 7]. Therefore, accurate diagnosis, which is dependent on identifying an increase in serum creatinine (SCr), is

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the first step in PC-AKI intervention [8]. However, the insensitivity of and delay in SCr testing and the absence of accurate novel biomarkers render early detection of PC-AKI very challenging [9, 10]. Additionally, because the key molecular mechanisms and regulatory networks underlying PC-AKI are still unclear, the effectiveness of classical prevention strategies, such as hydration, is controversial and limited, and specific targeted drugs are lacking [11-13]. Therefore, screening potential biomarkers and revealing key aspects of the molecular pathogenesis may provide the cornerstone for managing PC-AKI.

Long non-coding RNAs (lncRNAs), a group of single-stranded, non-coding RNAs containing more than 200 nucleotides and having broad regulatory functions across a wide range of biological processes and human ailments, can serve as novel biomarkers to predict various diseases due to their high specificity and detectability [14-16]. Recent studies have identified that MALAT1 can effectively regulate proliferation and inflammatory responses in ischemia/reperfusion-induced AKI, while TapSAKI and RP11-354P17.15-001 can sensitively predict the occurrence of AKI in renal allograft recipients [17-19]. However, lncRNA expression profiles in PC-AKI and potential lncRNA biomarkers for its detection are still largely unknown.

In this study, we explored the lncRNA expression profile in rat models. Then we conducted bioinformatics analysis and clinical screening to identify lncRNA biomarkers for the early detection of PC-AKI.

**Results**

**Establishment of the PC-AKI rat model**

As shown in Figure 1A, we randomly divided 6 rats into PC-AKI and control groups. The PC-AKI rat model was established through the classic combination of furosemide and contrast medium (CM). All rats in the PC-AKI group met the PC-AKI diagnostic criteria, SCr and blood urea nitrogen (BUN) concentrations were significantly increased after CM exposure (1.56- and 1.82-fold, respectively, Figure 1B). Haematoxylin-eosin (HE) staining was performed to detect the histopathological changes in PC-AKI. HE staining revealed that the kidney tissues of the control group were basically normal, while those of the PC-AKI group demonstrated typical pathological features of AKI, including severe renal tubular occlusion, extensive tubular cell detachment and infiltration of numerous inflammatory cells. The semiquantitative tubular damage scores in the PC-AKI group (histological score: 3.10 ± 0.73) were significantly increased compared with those in the control group (histological score: 0.60 ± 0.51) (Figures 1C, D). Transmission electron microscope (TEM) examination showed that ultrastructural changes of mitochondria and significant accumulation of lysosomes and autophagosomes were examined in the PC-AKI group (Figures 1E). The elevated SCr concentration, increased kidney injury marker levels, HE staining and TEM examination results indicated the successful establishment of the PC-AKI rat model.

**Overview of lncRNA and mRNA profiles in PC-AKI**

To systematically reveal dynamic changes in RNAs in PC-AKI, 6 rat kidney tissues from the PC-AKI and control groups were subjected to next-generation RNA sequencing. A total of 30829 lncRNAs and 11962 mRNAs were detected in PC-AKI tissues. Among these, 135 lncRNAs and 169 mRNAs were significantly upregulated while 222 lncRNAs and 181 mRNAs were significantly downregulated according to the following criteria: |log2 (fold change vs. control)| >1 and P<0.001 (Figures 1F, Figure S1A). Analysis of the lncRNA locations showed that the differentially expressed lncRNAs in PC-AKI were widely distributed across all chromosomes except chromosomes 21 and 22. Chromosomes 1 and 2 had the highest number of differentially expressed lncRNAs (Figure 1G). The differentially expressed lncRNAs were classified into the following 6 categories: 18.8% were exonic sense, 14.8% were exonic antisense, 16.2% were bidirectional, 23.0% were intergenic, 12.9% were intronic sense and 14.6% were intronic antisense [20, 21] (Figure 1H).

**Functional analyses and experiments revealed the potential biological processes and pathways mediating PC-AKI**

Functional annotation analyses of the differentially expressed genes were performed to predict the roles of RNAs in PC-AKI. Gene ontology (GO) analysis of differentially expressed mRNAs revealed that several cellular processes were enriched, including apoptosis (GO terms: 0097194, execution phase of apoptosis) and autophagy (GO terms: 0006914, autophagy). To validate the apoptosis and autophagy levels in PC-AKI, we conducted terminal deoxynucleotidyl transferase dUTP nick end labeling (TUNEL) assays, Western blotting and TEM in kidney tissues from the rat model (Figure S1B-D, Figure 1E). The results indicated that apoptosis and autophagy might be the potential cell processes to mediate PC-AKI [22, 23]. However, further experiments are needed to determine the exact role of these cellular processes in PC-AKI.
Figure 1. Establishment and expression profiles of the PC-AKI rat model. (A) Experimental grouping and operative procedures for establishing the PC-AKI rat model (PC-AKI group) and the control group. (B) The concentration of SCr and BUN levels among two groups. (C) Representative pathological changes in the kidney tissues of PC-AKI (HE staining; 40×, scale bar = 50 µm). (D) Tubular damage score. Tissue damage was scored according to the percentage of damaged tubules. (E) TEM examination of kidney tissues. (F) Heat map and volcano plot of differentially expressed lncRNAs in kidney tissues of PC-AKI rat, as determined by RNA sequencing. (G) Circos plot showed the chromosomal locations of the differentially expressed mRNAs. The first (outermost) layer of the Circos plot was a chromosomal map of the human genome, and the black and white bars indicated chromosome cytobands. In the second layer (next innermost), the red lines indicated lncRNA expression levels on the chromosomes. In the third layer, the green lines indicated mRNA expression levels. The innermost circle indicated lncRNA-mRNA interaction relationships. (H) Types and counts of differentially regulated lncRNAs were classified into six categories according to the genomic loci of their neighboring genes. The measurement data are expressed as the means ± SD. *: P<0.05, **: P<0.01, ***: P<0.001.
Kyoto Encyclopedia of Genes Genomes (KEGG) analysis and Ingenuity pathway analysis (IPA) were conducted to reveal the PI3K/Akt and HIF-1 pathways may mediate the occurrence of PC-AKI (Figure S2A-C). In kidney tissues from the PC-AKI group, HIF-1α was significantly upregulated. As for PI3K/Akt pathway, the ratio p-Akt(S473)/Akt was significantly decreased, while p-Akt(T308)/Akt has no significant changes, which indicates that the renal injury caused by CM was probably due to the suppression effects of the PI3K/Akt pathway via phosphorylation at Ser473 but not at Thr308 [24] (Figure S2D). Immunohistochemical staining of kidney tissues further verified that the HIF-1 pathway was activated in PC-AKI (Figure S2E).

**Selection of candidate lncRNAs as potential biomarkers for PC-AKI**

Candidate lncRNAs were defined as human-rat homology and significantly upregulated in both kidney tissues and blood samples of PC-AKI rats, and blood samples of human (Figure 2A). First, 135 lncRNAs with greater than 2-fold upregulation and P value <0.001 in PC-AKI rat kidney tissues were selected. Then, 44 lncRNAs were excluded because they were not included in the NONCODE database. Additionally, 77 of these lncRNAs were excluded because of the lack of human-rat homology. The homology was determined by nearby gene anchoring and sequence similarity using the UCSC Genome Browser (http://genome.ucsc.edu/) [25, 26]. Some Inc-HILPDA and Inc-PRND sequences conserved across *Homo sapiens*, *Mus musculus* and *Rattus norvegicus*, as determined by the UCSC Genome Browser, are shown in Figure 2C. We validated the expression of the 14 human-rat homologous lncRNAs in PC-AKI rat kidney tissues via qPCR and excluded 3 lncRNAs with an increase <2-fold or a P value >0.01. Then, the expression levels of the remaining 11 lncRNAs were measured in the blood of PC-AKI rats, and 6 were excluded due to an increase of <2-fold or a P value >0.01 in the blood. Ultimately, 5 lncRNAs, namely, Inc-HILPDA, Inc-PRND, Inc-CDK6, Inc-CDC42SE1, and NEAT1, were identified as candidate lncRNA biomarkers for PC-AKI. The expression levels of these 5 candidate lncRNAs in rat kidney tissues and blood are shown in Figure 2B.

**Inc-HILPDA and Inc-PRND as new biomarkers to detect PC-AKI in patients receiving CAG/PCI**

To confirm the expression levels of the 5 candidate lncRNAs in patients, 568 consecutive patients receiving coronary angiography (CAG) or percutaneous coronary intervention (PCI) in our center from April 2016 to November 2019 were enrolled. Among the 71 patients who developed PC-AKI, blood samples were available and qualified for 56. We performed a case-control study with these 56 PC-AKI cases and another 56 controls who were matched for age, sex, presence of diabetes mellitus and stage of chronic kidney disease and were randomly selected among patients who did not develop PC-AKI. Clinical characteristics were collected, and no significant differences were observed (Table 1). The expression levels of these 5 candidate lncRNAs in human blood were evaluated via qPCR. After CM exposure, the Inc-HILPDA and Inc-PRND expression levels in blood of PC-AKI patients were significantly higher than those in the controls (mean difference in lncRNA fold changes: Inc-HILPDA, 4.490 [3.135–5.845], P<0.0001; Inc-PRND, 4.334 [2.769–5.899], P<0.0001; Figure 2D). However, the Inc-CDK6, Inc-CDC42SE1, and NEAT1 expression levels did not meet the significant differential expression criteria of fold change >2 and P value <0.01.

Receiver operating characteristic (ROC) analysis was performed to determine the sensitivity and specificity of Inc-HILPDA and Inc-PRND for predicting PC-AKI. As shown in Figure 2E, the levels of these 2 lncRNAs significantly discriminated patients with PC-AKI from those without PC-AKI (Inc-HILPDA: AUC, 0.885 [0.824-0.946]; Inc-PRND: AUC, 0.875 [0.811-0.934]). Based on ROC analysis and the Youden index, the cut-off fold change increase values for Inc-HILPDA and Inc-PRND were established as 1.855 and 1.170, respectively (Table 2). An increase in Inc-HILPDA expression >1.855-fold demonstrated relatively high efficacy for detecting PC-AKI, with a Youden index of 0.6 (sensitivity, 80.36%; specificity, 82.93%). The Youden index was 0.63 when the Inc-PRND cut-off fold change value was 1.170 (sensitivity, 85.71%; specificity, 76.79%). To obtain the optimal diagnostic sensitivity and specificity, we combined these two lncRNAs to predict PC-AKI. This analysis showed that any positive lncRNA combination (Inc-HILPDA increase >1.855-fold or Inc-PRND increase >1.170-fold) yielded the maximized Youden index of 0.84 (sensitivity, 100%; specificity, 83.93%, Table 2). Therefore, Inc-HILPDA and Inc-PRND may be novel biomarkers for detecting PC-AKI in patients receiving CAG/PCI.
Figure 2. Selection of candidate IncRNAs and clinical validation of PC-AKI biomarkers. (A) Flow chart of the candidate IncRNA screening process. (B) Expression levels of 5 selected candidate IncRNAs in PC-AKI rat kidney tissues and blood. (C) Sequence similarity of Inc-HILPDA and Inc-PRND across species. (D) Blood levels of the 5 candidate IncRNAs in 56 PC-AKI patients and 56 matched controls, determined by RT-qPCR. (E) The discriminatory potential of Inc-HILPDA and Inc-PRND. Receiver operating characteristic curves were drawn with the fold change data of blood IncRNAs from 56 PC-AKI patients and 56 matched controls. The dashed line indicates the “random guess” diagonal line. AUC indicates the area under the curve. The measurement data are expressed as the means ± SD. *, P<0.05; **, P<0.01; ***, P<0.001; ****, P<0.0001.
To investigate the potential of Inc-HILPDA and Inc-PRND as early biomarkers of PC-AKI, we conducted fluorescence in situ hybridization (FISH) assays and time-course expression experiments of these two lncRNAs. FISH assays showed that both Inc-HILPDA and Inc-PRND were mainly localized in the tubules (Figure 3A). We treated HK-2 cells with CM to induce cell damage. MTS assays and flow cytometry analysis indicated that HK-2 cells showed damage in a time-dependent manner, with decreased cell viability and an increased apoptotic cell ratio (Figure 3B-C). RT-qPCR was performed on HK-2 cells treated with iopromide at baseline, 6 h, 12 h and 24 h. The temporal trends are shown in Figure 3D. Furthermore, the time-course expression patterns of Inc-HILPDA and Inc-PRND were simultaneously detected in kidney tissues, blood and urine from PC-AKI rat models. The results indicated that both the Inc-HILPDA and Inc-PRND expression levels were significantly different from those in the control group at 6 h in kidney tissues, while a significant difference was first noted in blood at 12 h for Inc-HILPDA and 24 h for Inc-PRND (Figure 3E). However, due to their low abundance in urine, neither Inc-HILPDA nor Inc-PRND could be stably detected in urine (data not shown).

To further evaluate the biomarker potential of Inc-HILPDA and Inc-PRND, we then calculated the correlation of these two lncRNAs with previously validated kidney injury biomarkers, including KIM-1, IL-6, TNF-α, and TIMP2×IGBFP7. As shown in Figure S3A, we found that Inc-HILPDA was positively correlated with TNF-α (r=0.26, P<0.001) and TIMP2×IGBFP7 (r=0.49, P<0.001). Meanwhile, Inc-PRND was positively correlated with TNF-α (r=0.21, P<0.05) and TIMP2×IGBFP7 (r=0.22, P<0.001). The results of the ROC curve analysis for the novel potential biomarkers (KIM-1, IL-6, TNF-α, and TIMP2×IGBFP7) are shown in Figure S3B.

Furthermore, we examined whether Inc-HILPDA and Inc-PRND were specific for distinguishing PC-AKI. An IRI-induced AKI rat model was established by clamping the bilateral kidney pedicles (Figure S4A). The expression levels of both Inc-HILPDA (P=0.69 for kidney; P=0.81 for blood) and Inc-PRND (P=0.67 for kidney; P=0.60 for blood) in the IRI group were similar to those in the PC-AKI group (Figure S4B-C), indicating that these two lncRNAs are not specific for distinguishing PC-AKI from other types of AKI.

**Functional predictions for Inc-HILPDA and Inc-PRND**

The considerable biomarker potential of Inc-HILPDA and Inc-PRND indicates that these two lncRNAs may strongly mediate PC-AKI, but their functions in PC-AKI are still unknown. Therefore, focusing on Inc-HILPDA and Inc-PRND, we conducted interaction and functional analyses. We constructed Inc-HILPDA-mRNA and Inc-PRND-
mRNA coexpression networks and lnc-HILPDA-miRNA-mRNA and lnc-PRND-miRNA-mRNA competing endogenous RNAs (ceRNA) networks (Figure 4A, D). GO and KEGG pathway analyses of the mRNAs associated with lnc-HILPDA and lnc-PRND predicted that lnc-HILPDA and lnc-PRND may mediate PC-AKI through the HIF-1 and AGE-RAGE pathways (Figure 4B, C).

Figure 3. Localization and expression patterns of lnc-HILPDA and lnc-PRND in PC-AKI. (A) Localization of lnc-HILPDA and lnc-PRND in kidney tissue. White frames indicate the glomeruli. (B and C) Evaluation in the PC-AKI cell models. MTS assays (B) and flow cytometry measurement (C) of CM-treated HK-2 cells. (D) The expression level of lnc-HILPDA and lnc-PRND in CM-treated HK-2 cells over a 24-h time course. (E) The expression level of lnc-HILPDA and lnc-PRND in kidney tissues and blood of PC-AKI rats over a 24-h time course. The measurement data are expressed as the means ± SD. *, P<0.05; **, P<0.01; ***, P<0.001.
Figure 4. Functional predictions for Inc-HILPDA and Inc-PRND. (A) The IncRNA-mRNA coexpression networks of Inc-HILPDA and Inc-PRND. Green nodes, mRNAs; red nodes, Inc-HILPDA or Inc-PRND. (B) Biological process annotations with the top 15 enrichment scores determined by GO analysis for Inc-HILPDA- and Inc-PRND-related differentially expressed mRNAs. (C) Pathways with the top 20 enrichment scores determined by KEGG pathway analysis for Inc-HILPDA- and Inc-PRND-related differentially expressed mRNAs. (D) mRNA-miRNA-incRNA networks in PC-AKI. Circles, mRNAs; arrows, miRNAs; diamonds, Inc-HILPDA or Inc-PRND.
**Discussion**

In this study, we examined the RNA expression profile of PC-AKI. After rat kidney profiling, human-rat homologous screening and rat blood filtration, we selected 5 candidate lncRNAs for PC-AKI detection. A 112 clinical sample validation demonstrated that lnc-HILPDA and lnc-PRND were significantly increased in PC-AKI patients compared with the controls, which allowed PC-AKI patients to be precisely distinguished, with AUC values of 0.885 and 0.875, respectively. Time-course experiments showed that significant differences of lnc-HILPDA and lnc-PRND were both first noted in the blood of PC-AKI rats at 12 h for lnc-HILPDA and 24 h for lnc-PRND. These observations in this study suggest that lnc-HILPDA and lnc-PRND may serve as the new potential biomarker to distinguish PC-AKI in early phases.

We found that lnc-HILPDA and lnc-PRND may serve as novel biomarkers for PC-AKI, with AUC values of 0.885 and 0.875, respectively. Recent studies have reported that the lncRNAs TapSAKI and RP11-354P17.15-001 can effectively detect the occurrence of AKI [2, 3], which together with our results demonstrates the promising potential of lncRNAs for AKI detection and patient stratification. However, several differences exist between the previous study and ours. First, the populations are different. Previous studies focused on severe AKI patients requiring renal replacement therapy or renal allograft-induced AKI patients. In contrast, our study focused on PC-AKI, a common and specific type of AKI. Second, the screening processes were different. In the previous study, TapSAKI and RP11-354P17.15-001 were directly selected from human blood/urine samples using a two-step screening process, including a small-sample blood/urine lncRNA array analysis and larger sample clinical validation. However, we identified lnc-HILPDA and lnc-PRND through a four-step screening process that included rat kidney RNA sequencing and validation, rat blood screening, human-rat homologous screening and clinical validation. Each of the two different screening processes has advantages and disadvantages. Two-step screening in human samples is convenient and can detect more lncRNAs as candidate biomarkers but cannot identify highly abundant or kidney-specific lncRNAs in the blood and requires more statistical analysis due to the lack of lncRNA expression data in the kidney. Comparatively, four-step screening can narrow down the candidate lncRNAs and select highly abundant kidney biomarkers from combined kidney and blood lncRNA data. However, many data entries may be lost during multiple screening events. Third, prediction efficiencies are different. The previous study showed that TapSAKI and RP11-354P17.15-001 had relatively high prediction efficiencies, with AUC values of 0.80 and 0.76, respectively. In contrast, Inc-HILPD and Inc-PRND, which we identified, showed even better performance, with AUC values of 0.885 and 0.875, respectively. We think the population heterogeneity and the screening criteria may contribute to the differences. Severe AKI may be caused by various factors, such as inflammation, ischemia and medications with renal toxicity [27, 28], and the incidence of renal allograft-induced AKI is affected by many factors, such as donor kidneys, surgeons and immune state [29, 30]. In contrast, PC-AKI, which is induced by contrast media, has a similar pathophysiological process [2, 31] and may have less heterogeneity than severe AKI and renal allograft-induced AKI. Additionally, we applied strict screening criteria (fold change>2 and P<0.01) during multiple screening processes. Finally, we identified lnc-PRND and Inc-HILPD.

As the third leading cause of hospital-acquired AKI [3], PC-AKI was also named CI-AKI [32]. The 2018 ESUR guidelines recommended that the term CI-AKI should only be used when a causal relationship can be shown, while PC-AKI is recommended to be used when AKI occurs after intravascular administration of iodine-based contrast media. Although CI-AKI may be better known than PC-AKI, recent studies have suggested that the risk of AKI due to contrast material is overestimated [33]. Therefore, in our study, it is better to use “PC-AKI” to indicate the included patients. Previous studies have verified that LNC_000343, LNC_000985 and LNC_002216 may serve as novel biomarkers in a rat PC-AKI model [34]. Cheng et al. screened and validated the biomarker potential in a rat model, and their screening process was a relatively simple two-step screening process. In some sense, our four-step screening and human sample validation may have a better chance of identifying potential PC-AKI biomarkers. In this study, we found that lnc-HILPDA and lnc-PRND can distinguish PC-AKI patients with relatively high sensitivity and specificity. Furthermore, the expression of lnc-HILPDA and lnc-PRND in kidney tissues and blood samples was time-dependent, indicating that these two lncRNAs could likely serve as biomarkers in the early stage of PC-AKI. Although showing high sensitivity and specificity, neither Inc-HILPD nor Inc-PRND could distinguish PC-AKI from other types of AKI, such as IRI-induced AKI. This might be because the pathophysiological mechanisms of
PC-AKI are similar to those of IRI-AKI. These two AKI types are both related to oxidative stress [35], vascular endothelial cell damage [36], and imbalanced secretion of vasomotor substances [37], among other factors.

Our study has several limitations. First, the IncRNA profiles were assessed in a rat model rather than in human samples, and the number of rats in each group was limited. As PC-AKI is not a clinical indication for renal biopsy, it would be difficult to acquire PC-AKI patient samples. Although we validated the IncRNA biomarkers in human blood, our profiles and analysis should still be considered when applied to humans. Second, the IncRNA screening process can be further improved. Due to the poor conservation of IncRNAs between humans and rats, the human–rat homologous screening may result in the unavailability of RNA data in humans. Third, the IncRNA biomarker validation was performed on monocentric, small-scale clinical samples, and the efficiency of these biomarkers requires further validation in large-scale, multicentric clinical trials. Moreover, the biological mechanisms by which Inc-HILPDA and Inc-PRND contribute to PC-AKI remain unclear, and further investigations into their functions may provide novel targets and strategies.

In conclusion, this study offers the first analysis of IncRNAs in the blood as potential markers of PC-AKI. We found that Inc-HILPDA and Inc-PRND can effectively predict PC-AKI risk after a cardiac catheterization, therefore increasing the probability that IncRNAs may serve as novel biomarkers for PC-AKI in early phases. Due to the potential clinical application of these IncRNAs, PC-AKI patients may receive precise stratification, early detection and better interventions.

Materials and methods

Rat model establishment

Male Sprague-Dawley rats (200-220 g) were purchased from Guangdong Animal Center (Guangzhou, China). For the PC-AKI model, rats were deprived of water from 24 h before the experiment until sacrifice. Six hours before CM exposure, rats were administered 15 mL/kg furosemide via intraperitoneal injection. Rats in the PC-AKI group were injected with iopromide (Ultravist, 300 mg iodine/mL, 330 mOsm/kg H₂O and 9.5 MPa at 37 °C; Bayer AG, Leverkusen, Germany) at a dose of 15 mL/kg body weight via the tail vein over 5 min. Control group rats received an equal volume of normal saline. PC-AKI rats were sacrificed, and the kidneys were excised after CM exposure for 6, 12, or 24 h. Blood samples were collected from the orbital vein and aorta ventralis at baseline and after sacrifice. SCr concentrations were measured, and AKI was defined as a 50% increase in the SCr concentration above baseline after contrast administration [38].

For the ischemia-reperfusion injury model, bilateral kidney pedicles were identified through two small paramedial dorsal incisions and clamped for 45 min. The clamps were then released for reperfusion. The wound was then sutured after good renal reperfusion. Control animals underwent the same surgical procedure, except the renal pedicles were not clamped [39]. All rats were sacrificed at 24 h after reperfusion. The AKI definition was the same as that for the PC-AKI group.

All experimental procedures were performed in compliance with the Guide for the Care and Use of Laboratory Animals (National Research Council Publication, 8th Edition, 2011) and in accordance with the Chinese Regulations for the Administration of Affairs Concerning Experimental Animals. The animal experimental protocol was approved by the Guangdong General Hospital Ethics Research Committee (No. 2015253). The model was established at the Department of Experimental Animal Research Centre, Southern Medical University (Guangzhou, China).

Population

Patients who were scheduled for CAG or PCI in the Cardiology Department of Guangdong Cardiovascular Institute between April 2016 and November 2019 were consecutively enrolled in this study. The inclusion criteria were as follows: (1) provision of a signed informed consent form, (2) age ≥ 18, and (3) treatment with elective CAG or PCI in the Cardiology Department of Guangdong Cardiovascular Institute. The exclusion criteria were as follows: (1) death during the procedure; (2) severe heart failure (cardiogenic shock or NYHA IV); (3) end-stage renal failure or receipt of renal transplant; (4) exposure to CM within 1 week before or 72 h after the procedure; (5) allergy to CM; (6) pregnancy, lactation or the presence of a malignant tumor resulting in a life expectancy of <1 year; (7) use of sodium bicarbonate, non-steroidal anti-inflammatory drugs (NSAIDs), metformin, aminoglycosides, cyclosporine, or cisplatin before or within 48 h after the procedure; or (8) severe valvular disease or planned surgery. Non-ionic, iso-osmolar CM and intravenous hydration were used in all patients. The duration and volume of hydration and the use of N-acetylcysteine were determined at the discretion of the physicians. Peripheral blood (5 mL) was collected 24 h before and 24-30 h after the procedure directly
into PAXgene Blood RNA tubes (BD Vacutainer, Franklin Lakes, NJ). SCr was measured at the time of admission and within 72 h after the intervention procedure. PC-AKI was defined as an absolute increase in SCr≥0.3 mg/dL or a relative increase in SCr≥50% over baseline within 72 h [1, 9]. The estimated glomerular filtration rate (eGFR) was determined as follows: (1) hypotension: score=5; (2) intra-aortic balloon pump: score=5; (3) congestive heart failure: score=5; (4) age >75 years: score=4; (5) anaemia: score=3; (6) diabetes mellitus: score=3; (7) CM volume: score=1 per 100 cc; and (8) score=2 for 40 mL/min per 1.73 m² ≤ eGFR <60 mL/min per 1.73 m², score=4 for 20 mL/min per 1.73 m² ≤ eGFR <40 mL/min per 1.73 m² or score=6 for eGFR <20 mL/min per 1.73 m² [41]. The baseline values for the enrolled patients are shown in Table 1. The study protocol strictly complied with the ethical guidelines of the 1975 Declaration of Helsinki and was approved by the Guangdong Provincial Hospital Medical Ethics Committee (No. 2015253). All patients who participated in this programme were fully informed and signed an informed consent form before enrolment.

**Blood collection and RNA isolation**

For the patients’ peripheral blood samples and rat blood samples, the PAXgene Blood RNA system was used to stabilize RNA and to store the samples for a relatively long time. Whole blood was collected directly into PAXgene Blood RNA tubes (BD Vacutainer, Franklin Lakes, NJ). Total RNA was isolated within 24 h after sample collection with a PAXgene Blood RNA kit (Qiagen, Hilden, Germany) following the manufacturer’s protocol. For rat kidney samples, total RNA was harvested and purified with the standard TRIzol reagent (Invitrogen, Carlsbad, CA) according to the manual. The concentration and purity of total RNA were assessed with a NanoDrop 2000 spectrophotometer (Thermo Fisher Scientific, Wilmington, DE). The quality of RNA samples was checked with an Agilent 2100 Bioanalyzer and RNA 6000 Nano Kit (Agilent, Santa Clara, CA, USA). Samples with an RNA integrity number ≥8 were considered qualified for use. Qualified RNA was stored at -80 °C until further analysis.

**Next-generation sequencing and data processing**

The lncRNA, microRNA and mRNA profiles in rat kidney tissues were assessed by next-generation sequencing. rRNA-depleted total RNA was purified from 6 kidney tissues from PC-AKI and control rats using a Ribo-Zero rRNA Removal Kit (Epicentre). Libraries were constructed using an Illumina Gene Expression Sample Preparation Kit and sequenced on an Illumina HiSeq™ 2000 system (Beijing Genomics Institute (BGI)) with 50-60 million 2×100 bp paired raw passing filter reads. The raw reads were filtered to obtain the high-quality reads by removing reads containing an adaptor sequence or reads with a percentage of unknown bases (N) exceeding 10%; low-quality reads contained more than 50% unknown bases and had a Q value of <5. The high-quality reads were mapped to the rat reference genome (IRGSP build 5.0) using SOAPaligner/SOAP2, allowing no more than two mismatches in the alignment. Manipulation of the false discovery rate (FDR) was used to determine the threshold P values in multiple testing and analyses.

**Reverse transcription quantitative real-time polymerase chain reaction (RT-qPCR) analysis**

RT-qPCR was used to verify the expression of differentially expressed lncRNAs screened via next-generation sequencing in rat and human blood. Reverse transcription was carried out using total RNA (800 ng from PAX). cDNA was synthesized via reverse transcription using a PrimeScript™ RT Reagent Kit (RR037) with random 6mers and oligo dT primers according to the manufacturer’s protocols (Takara, Japan), and reactions were performed in an Eppendorf 5331 PCR instrument (Eppendorf, Hamburg, Germany). Then, quantitative real-time PCR was carried out using TB Green® Premix Ex Taq™ II (Tli RNaseH Plus, RR820) according to the manufacturer’s protocols (Takara, Japan). The primers used to amplify lncRNAs in the reactions were synthesized by Generay (Shanghai, China). All RT-qPCR primer sequences are shown in Table S1. qPCR was performed in a CFX Connect Real-Time System (Bio-Rad, USA). The relative expression levels of the lncRNAs were normalized to those of the reference gene GAPDH and calculated using the $2^{\Delta\Delta CT}$ method. All reactions were repeated in triplicate.

**Haematoxylin-eosin staining**

Rat kidney tissues were fixed in 10% neutral buffered formalin (NBF) for a minimum of 24 h and embedded in paraffin, after which 4-mm-thick tissue sections were cut using a microtome. Histopathological evaluation of the tissue sections was conducted with HE staining. Finally, tissue morphology was observed. Tubular damage was scored according to the percentage of damaged tubules [42]: 0, no damage; 1, <25% damaged; 2,
25-50% damaged; 3, 50-75% damaged; and 4, >75% damaged.

**TdT-mediated dUTP nick end labeling assay**

TUNEL assay for cell apoptosis was used on paraffin sections of kidney tissues from rats and was performed according to the instructions for the In-Situ Cell Death Detection Kit (Roche, Basel, Switzerland). Paraffin sections were cut with a rotary microtome (Leica RM2255, Germany). Sections were washed with xylene two times for 5 min each and then soaked one time in each component of a graded ethanol series. Then, the tissues were treated with Proteinase K working fluid for 15-30 min at 21 –37 °C. After two washes with PBS, the TUNEL-converter-POD solution reaction mixture was added and then reacted with 3,3'-diaminobenzidine (DAB) for color development. The cells were subsequently analysed under a fluorescence microscope. Positive cells were counted in 5 randomly selected visual fields at 400× magnification.

**Cell Culture**

HK-2 cells were provided by the Stem Cell Bank (Chinese Academy of Sciences, Shanghai, China). HK-2 cells were cultured in low glucose DMEM/F12 medium (Gibco™; Thermo Fisher Scientific, Waltham, MA, USA) containing 10% foetal bovine serum (FBS), 0.1% penicillin and streptomycin at 37°C with 5% CO2 and sub-cultured when cells reached 80% confluence. In the contrast medium-treated groups, the cells were treated with 150 mg iodide (mg I)/ml Ultravist Contrast Media Injection (370 mg I/ml; Bayer AG, Leverkusen, Germany) for 0, 6, 12, or 24 h in DMEM/F12 with 10% FBS.

**MTS Assay of Cell Viability**

HK-2 cell viability was assessed with a CellTiter96™ AQueous One Solution Cell Proliferation Assay (Promega, Madison, WI, USA) according to the manufacturer’s protocol. Cells were cultured in 96-well plates containing a final volume of 100 μl/well. Approximately 20 μl of MTS Reagent was added to each well, and the cells were incubated for 2 h at 37 °C. Cell viability percentages were assessed via spectrophotometry at 490 nm.

**Flow Cytometry Analysis of Apoptosis**

Cell apoptosis was examined with an APC-Annexin V and 7-amino-actino-mycin D (7-AAD) staining kit (BD Biosciences, Jersey, USA). Briefly, cells were washed twice with cold PBS and then resuspended in 1× Binding Buffer. After the addition of 5 μl of APC-Annexin V and 5 μl of 7-AAD, the cells were incubated for 15 min at 25°C in the dark. Finally, 400 μL of 1× Binding Buffer was added, and the cells were then analysed via flow cytometry within 1 h. Flow cytometry analysis was performed on a BD Biosciences Accuri C6 flow cytometer (Becton, Dickinson and Company, San Jose, CA, USA) with the 488-nm spectral line of an argon-ion laser.

**Western blot analysis**

Rat kidney tissues from the PC-AKI group and control group were homogenized and lysed in ice-cold RIPA lysis buffer (Beyotime, Shanghai, China) supplemented with 1% PMSF using a High Flux Tissue Grinder (Next Advance, New York, USA) for 3 min and then centrifuged for 20 min (16300×g, 4 °C). The supernatant was collected, and the protein concentration was evaluated using an Enhanced BCA Protein Assay Kit (Beyotime, Shanghai, China). Equal amounts of protein (30 μg) were size-fractionated in 15% and 12% SDS-PAGE gels and electrophoretically transferred to PVDF membranes. Membranes were blocked in TBST containing 5% non-fat milk for 2 h at room temperature and were then incubated with primary antibodies overnight at 4 °C. The following antibodies were used: anti-Bax (Proteintech, #50599-2, 1:1000), anti-Bcl-2 (Abcam, ab196495, 1:1000), anti-procaspase-3 (Abcam, ab32150, 1:1000), anti-Cleaved Caspase-3 (Abcam, ab32042, 1:2000), anti-P53 (Proteintech, #10442, 1:1000), anti-p-P53 (Cell Signaling Technology, #9284, 1:1000), anti-LC3 (Sigma-Aldrich, L7543, 1:2000), anti-p-LC3 (Cell Signaling Technology, #2920, 1:2000), anti-p-Akt (Cell Signaling Technology, #4060, 1:1000), and anti-p-Akt (Cell Signaling Technology, #13038, 1:1000) antibodies. Expression levels were normalized by probing the same blots with an anti-GAPDH antibody (Proteintech, #60009-1, 1:8000).

**Immunohistochemistry**

Rat kidney tissue slices were fixed in 10% NBF overnight and then cryoprotected in 30% sucrose solution. For immunohistochemistry, after antigen retrieval in citric acid buffer (pH 6.0), the slices were incubated with primary antibodies specific for HIF-1α (purchased from Abcam (Cambridge, UK)) overnight at 37 °C. Then, the slices were incubated with secondary antibody (1:200, Abcam, UK) at 37 °C for 40 min and washed three times with PBS. DAB (Abcam, UK) was added for color development for 2 min to visualize the antibody-antigen complexes. Immunostaining images were acquired using a microscope.
Fluorescence in situ hybridization

Renal sections from the PC-AKI group and control group were incubated at 65°C for 48 h with a Cy3-labelled probe for lnc-HILPDA and lnc-PRND purchased from GeneRay Biotechnology (Shanghai, China). Then, fluorescence signals were scanned at 260 nm using a Zeiss LSM 780 confocal microscope (Carl Zeiss SAS, Germany).

Transmission Electron Microscopy

Kidney samples (1 mm thick) from each group were carefully immersed in fixative solution (2.5% glutaraldehyde and 2% paraformaldehyde in cacodylate buffer (0.1 M, pH 7.4)) for 24 h. The samples were washed with PBS three times, fixed in 10% buffered glutaraldehyde and 1% osmic acid, dehydrated, embedded, and finally sliced to a thickness of 50-90 nm. The ultrastructure of autophagosomes was observed under a transmission electron microscope.

Enzyme-linked immunosorbent assay

We used commercially available ELISA kits (Cusabio Biotech, China) to measure IL-6, TNF-α, TIMP2, IGFBP7 and KIM-1 expression levels in patient serum according to the manufacturer’s protocols. Each experiment was performed independently and repeated three times.

Bioinformatics analysis

Differentially expressed genes were screened using the following criteria: (1) fold change ≥2 for upregulation or downregulation and (2) P value <0.001. GO analysis was applied to predict the underlying biological processes, molecular functions and cellular components in PC-AKI. The KEGG database was also used to analyse the potential pathways mediating the pathogenic process. The significant GO terms and pathways were identified with Fisher’s exact test, and the FDR was utilized to correct the P values. Additionally, differentially expressed genes were loaded into the “core analysis” function included in the IPA software and included genes associated with biological processes, canonical pathways, upstream transcriptional regulators, and gene networks. Based on the significantly different expression levels of ncRNAs and mRNAs between the control group and the PC-AKI group, miRanda (http://www.microrna.org/microrna/home.do) was used to construct a ceRNA network by predicting microRNA binding sequence sites in lncRNAs and mRNAs. The coexpression analysis was based on weighted correlation network analysis [43]. Differentially expressed lncRNAs and mRNAs with fold changes ≥2 and P values <0.01 were analysed. A soft threshold value of ≥0.8 was the recommended parameter for the coexpression analysis. K-Core scoring was used to identify core genes in the coexpression networks [44]. A higher k-core score indicates a more central location of a transcript within a network.

Statistical Analysis

Continuous variables are presented as the means ± SD or medians (with 25th and 75th percentiles). Differences between Gaussian distributed variables were determined with Student’s t-test. The Mann-Whitney U test or Kruskal–Wallis test was used to determine differences between non-normally distributed variables. Pearson correlation analysis and Spearman correlation analysis were performed to analyse Gaussian and non-normally distributed variables, respectively. Categorical variables are reported as percentages and were analysed using either a chi-square test or Fisher’s exact test. Receiver operating characteristic curves were used to establish cut-off increase values for lncRNAs in the PC-AKI predictions. All P values are 2-sided, and P values less than 0.05 were considered statistically significant. All statistical analyses were performed using SPSS 16.0 and R 3.5.0 software.

Abbreviations

PC-AKI: Post-contrast acute kidney injury; lncRNA: long non-coding RNA; AUC: area under the curve; CI-AKI: contrast-induced acute kidney injury; CM: contrast medium; SCr: serum creatinine; BUN: blood urea nitrogen; HE: Haematoxylin-eosin; TEM: Transmission electron microscope; GO: Gene ontology; TUNEL: terminal deoxynucleotidyl transferase dUTP nick end labelling; KEGG: Kyoto Encyclopedia of Genes Genomes; IPA: Ingenuity pathway analysis; CAG: coronary angiography; PCI: percutaneous coronary intervention; ROC: Receiver operating characteristic; FISH: fluorescence in situ hybridization; ceRNA: competing endogenous RNAs; eGFR: estimated glomerular filtration rate; FDR: false discovery rate; RT-qPCR: reverse transcription quantitative real-time polymerase chain reaction; NBF: neutral buffered formalin; FBS: foetal bovine serum.

Supplementary Material


Acknowledgements

Ethics approval and consent to participate

The study was approved by the Ethics
Committee of Guangdong Provincial People’s Hospital. All the patients enrolled in the study signed written informed consent.

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Availability of data and materials
Data relevant to this study are available from the corresponding authors upon reasonable request.

Authors’ contributions
Conception and study design: Yong Liu. Provision of study materials and implementation of experiments: Bowen Liu. Bioinformatics and statistical analyses and biomarker screening: Bowen Liu, Guanzhong Chen, and Shiqun Chen. Patient enrolment and collection of clinical characteristics: Jin Liu, Guoli Sun, and Lei Li. Sample collection and detection: Shanyi Huang, Huanqiang Li, Ziling Mai, Liyao Zhang, and Min Li. Administrative support and cooperation: Ning Tan, Hong Li, Yulin Liao, Jia Liu, and Jiyan Chen. Manuscript writing: Bowen Liu, Guanzhong Chen, and Shiqun Chen. Final approval of manuscript: All authors.

Competing Interests
The authors have declared that no competing interest exists.

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