Serum miR-21 may be a Potential Diagnostic Biomarker for Diabetic Nephropathy

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Abstract

MiRNAs play important roles in initiation and progress of many pathologic processes. MiR-21 was closely associated with diabetic nephropathy (DN). However, whether serum miR-21 was as a potential diagnostic biomarker for DN and the relationship between serum miR-21 and tissue miR-21 remained unclear. In this study, real-time RT-PCR, cell transfection, luciferase reporter gene assays, western blot and confocal microscope were used, respectively. Here, we found that serum and renal tissue miR-21 was significantly elevated with the progress of DN. Moreover, luciferase reporter gene assays showed that smad7 was a validated miR-21 target, cell transfection showed that miR-21 overexpression downregulated target smad7 expression. Interestingly, serum miR-21 was significantly consistent with the alterations of tissue miR-21 with the development of DN. In addition, serum miR-21 was also positively correlated with GBM, GA, ACR and CCF, while negatively correlated with Ccr. Importantly, antagomiR-21 not only alleviated GBM, GA, ACR and CCF, but also ameliorated Ccr by increasing target smad7. In conclusion, our data demonstrated that serum miR-21 was closely associated with renal structure and function, and serum miR-21 may be regarded as a potential diagnostic biomarker of DN.

Abbreviations

DN                  Diabetic nephropathy
ACR                urine albumin creatinine ratio
Ccr                creatinine clearance ratio
AntagomiR-21   miR-21 antagonist
GBM                glomerular basement membrane
GA                 glomerular area
CCF                content of collagen fibers

Introduction

Diabetic nephropathy (DN) is a chronic, progressive process that ultimately leads to renal fibrosis and end-stage renal failure, a devastating disorder that requires dialysis or kidney transplantation [1–3]. KK-Ay mouse was considered suitable as a polygenic model for human type-2 diabetes mellitus and was produced by transferring the yellow obese gene (Ay allele) into the KK-Ay/Ta mice. Renal lesions in KK-Ay mice closely resembled human DN, which develop marked high glucose, albuminuria and renal fibrosis [4]. MicroRNAs (miRs) are endogenous non-coding small RNA, 20–22 nucleotides in length, which bind to the 3'-UTR of target genes, thereby, repress translation and/or induce degradation of target gene mRNAs [5]. Increasing evidences have indicated that miRNAs play significant roles in many diseases including diabetes mellitus and its complications [6]. For example, miR-29 aggravates insulin resistance and inhibits fibrosis [7,8]. miR-377 can lead to increased fibronectin production in DN [9]. miR-451 regulates p38 MAPK signaling by targeting of Ywhaz and suppresses the mesangial hypertrophy in early DN [10]. miR-503 contributes to diabetes mellitus-induced impairment of endothelial function and reparative angiogenesis after limb ischemia [11]. miR-192 may be a critical downstream mediator of TGF-β/Smad3 signaling in the development of renal fibrosis [12]. Recent reports have documented that miR-21 plays a crucial role in DN and kidney fibrotic diseases [13,14]. For example, miR-21 expression was increased in kidney biopsies from diabetic patients and DN mice [15]. miR-21 expression increases rapidly in cultured

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were fed by common forage (12% fat, 60% carbohydrate, and bio, China) for 8 weeks. Antagomir is a miRNA antagonist, which binds with mature miRNA in the body. Antagomir prevents TGF-β1-induced EMT [22, 23]. However, whether serum miR-21 was as a potential diagnostic biomarker for DN and relationship between serum miR-21 and tissue miR-21 in DN remained unclear.

The aim of this study was to identify the relationship between serum miR-21 and tissue miR-21, and to explore whether serum miR-21 was a novel diagnostic biomarker for DN. The results suggest that both serum miR-21 was closely associated with renal structure and function, and serum miR-21 may be as a novel diagnostic biomarker for DN.

Materials and Methods

Animals and diabetic nephropathy models

Male C57BL/6J (12 weeks of age, 24 mice) and KK-Ay mice (12 weeks of age, 48 mice) from Chinese Academy of Medical Sciences (Beijing, China) were individually housed in plastic cages with free access to food and water throughout the experiment. All mice were maintained in the same room under conventional conditions with a regular 12-h light/dark cycle with the temperature controlled at 24°C ± 1°C. To induce DN, C57BL/6J mice were fed by common forage (12% fat, 60% carbohydrate, and 28% protein), KK-Ay mice received research diets (58% fat, 25.6% carbohydrate, and 16.4% protein) for 4 weeks, random blood glucose (RBG) was checked by a portable glucometer from tail vein of each animal, C57BL/6J mice were classified as normal control group (NC group, n = 20), KK-Ay mice were considered DN when their RBG was ≥ 300 mg/dl (16.7 mmol/l) and ACR (urine albumin creatinine ratio) was ≥ 300 µg/mg were detected, KK-Ay mice were considered as DN model suitable for human type-2 diabetes mellitus [4]. KK-Ay mice were randomly divided into DN model group (DN group, n = 24), which were injected intraperitoneally with non-targeting negative control sequences (urine) expressing plasmid into 40–50% confluent T293 cells (Genechem, Shanghai, China), which grown in a 24-well plate. The cells were harvested 48 h after transfection, and luciferase activity was measured with a dual luciferase reporter assay kit (Promega, Madison, WI, USA) on a luminometer (Lumat LB9507).

Biochemical assays

Serum creatinine (SCR) and body weight was measured at 12, 16, 20 and 24 weeks of age. Urinary albumin and creatinine were measured by immunoassay (DCA 2000 system, Germany). Urinary albumin creatinine ratio (ACR) was calculated as: ACR = urinary albumin (µg)/urinary creatinine (mg). Creatinine clearance (CCR) ratio was calculated using the following equation [1]: CCR (mL/min·kg⁻¹) = [urinary Cr (mg dL⁻¹) × urinary volume (mL)/serum Cr (mg dL⁻¹)] × [1000/body weight (g)] × [1/1.440 × (min)] [25].

Light and electron microscopy

Tissues for light microscopy were fixed in 10% phosphate-buffered formalin and then embedded in paraffin. 4-micrometer-thick sections were processed for hematoxylin-eosin staining by light microscopy. Tissues for electron microscopy were fixed with 2% glutaraldehyde in 0.1 mol/L phosphate buffer at 4°C for 120 min. Ultrathin sections were collected on 100-mesh copper grids and double stained with 4% uranyl acetate and lead citrate. The sections were examined with a Hitachi 7100 transmission electron microscope. Morphological analyses were performed by an experienced pathologist who was blinded to the source of the tissue.

Transfection of cultured HKCs

To investigate the role of miR-21 in normal human kidney tubular epithelial cells (HKCs), obtained from Chinese Type Culture Collection (CTCC), we performed miR-21 transfection experiments, and cells were seeded at a density of 2 × 10⁴ cells/cm² in serum-free DMEM/F12. In this study, cells were divided into the following groups: cells without transfection were used as blank control group. Cells transfected with miR-control lentivirus vector were used as miR-control group. Cells transfected with miR-21 over-expression (pre-miR-21) lentivirus vector were used as miR-21 over-expression group (pre-miR-21 group). Cells transfected with miR-21 inhibitor lentivirus vector were used as miR-21 inhibitor group. After 12 h transfection, the medium was changed and the HKCs were incubated with fresh serum-containing medium for another 48 h. In our experiment, the most appropriate multiplicity of infection (MOI) for HKCs equals to 30, all the transfected cells were measured and sorted 48 h later according to the green fluorescent protein (GFP) intensity by flow cytometry, and the transfection efficiency was above 97%. The entire abovementioned lentivirus vector was custom-synthesized by Shanghai Genechem Co., Ltd, China. After 3 days of culturing, cells were harvested for RNA or protein isolation.

Luciferase reporter gene assays

To examine whether miR-21 regulates the expression of smad7, we transiently transfected miR-control plasmid, wild-type or mutant luciferase-smad7-3’UTR reporter) and miR-21 over-expressing plasmid into 40–50% confluent T293 cells (Genechem, Shanghai, China), which grown in a 24-well plate. The cells were harvested 48 h after transfection, and luciferase activity was measured with a dual luciferase reporter assay kit (Promega, Madison, WI, USA) on a luminometer (Lumat LB9507).

Real-time RT-PCR analysis

For analysis of serum and renal tissue miR-21 expression, TaqMan miRNA assays (Applied Biosystems, California, and USA) were used for quantitative determination of miR-21 expression according to the manufacturer’s instructions. The relative expression was normalized to the expression of U6 RNA (Applied
Western blot analysis
Renal tissues were lysed in RIPA buffer with protease inhibitors (Roche). Protein concentrations were determined by bicinchoninic assay (Pierce Biotechnology, Rockford, IL, USA). Proteins were separated on 10% SDS-PAGE gels under reducing conditions and transferred to polyvinylidene difluoride membranes. The membranes were blocked with rabbit polyclonal to col-IV (Abcam) and polyclonal to Col-I (Abcam), and then with secondary antibodies (1:5000, Roche). Membranes were stripped and reprobed with β-actin (Sigma) and secondary antibody for data normalization.

Statistical analysis
Statistical analyses were performed by one-way ANOVA followed by the Bonferroni multiple comparison test (for comparison of more than two groups) or Student t test (for comparison of two groups) and correlated analysis by Pearson’s correlation test. A probability value of <0.05 was considered significant.

Results

Relationship between serum miR-21 and tissue miR-21
To explore the relationship between serum miR-21 and renal tissue miR-21, we measured serum miR-21 and renal tissue miR-21 by real-time quantitative RT-PCR at different weeks of age. At 12 weeks of age, serum miR-21 was initially increased (p < 0.05). With the progress of DN, serum miR-21 level was further elevated until 24 weeks of age (Fig. 1a, p < 0.05). Interestingly, the alterations of serum miR-21 were significantly consistent with the expressions of renal tissue miR-21 (Fig. 1b, c, p < 0.05). More importantly, serum miR-21 was positively correlated with tissue miR-21 expression by Pearson correlation analysis (Fig. 1d, r = 0.894, p < 0.05). Next, to clarify whether antagomiR-21 decreased the expressions of serum and tissue miR-21, KK-Ay DN mice were injected intraperitoneally with antagomiR-21, we found that serum and tissue miR-21 were significantly decreased in antagomiR-21 group compared with DN group (Fig. 1a, b, p < 0.05). Taken together, our results suggested that circulating serum miR-21 may reflect the changes of tissue miR-21.

Relationship between serum miR-21 and renal function
To evaluate the relationship between serum miR-21 and ACR/Ccr, and to further elucidate whether serum miR-21 could be as a novel diagnostic biomarker for DN, the changes of ACR and Ccr were examined. With the progress of DN (from 16 to 24 weeks of age), ACR began to increase and Ccr began to decline (Fig. 2a, b). Additionally, antagomiR-21 was able to significantly decrease ACR and increase Ccr (Fig. 2a, b, p < 0.05). Interestingly, serum miR-21 expression was positively correlated with ACR (Fig. 2c, r = 0.970, P = 0.006) and negatively correlated with Ccr (Fig. 2d, r = −0.950, P < 0.01). Importantly, we found that antagomiR-21 can decrease the levels of ACR and increase the levels of Ccr (Fig. 2a, b, p < 0.05). Thus, our results suggested that serum miR-21 was closely associated with the markers of renal function (ACR and Ccr), and that serum miR-21 may be as a novel diagnostic biomarker for DN.
serum miR-21 may be associated with the morphological changes of DN, whereas, antagomiR-21 ameliorated renal morphology in diabetic kidneys.

Serum miR-21 was closely associated with collagen fibers

Collagen fibers (CFs) are the most important ingredients comprised of the glomerulus such GBM and mesangial matrix, but the deposition of excessive CFs altered renal structure, and adversely affects renal function [26]. To determine the relationship between serum miR-21 and CFs, we examined CFs by Picrosirius red and Masson staining, respectively. Picrosirius red staining by polarized light microscopy showed that CFs was bright yellow or orange (col-I for red or yellow, col-IV for light yellow (Fig. 4a). Similarly, Masson staining showed that a large number of CFs were primarily deposited in GBM, mesangial region and renal interstitial region in DN group compared with NC group (Fig. 4a). Next, to discriminate the collagen type, the expression of col-I and col-IV were examined by western blot, we found that col-IV and col-I was remarkably increased in DN group compared with NC group (Fig. 4c, d). Interestingly, serum miR-21 was positively correlated with the protein of col-IV rather than correlated with col-I (Pearson correlation, \( r_{\text{col-IV}} = 0.870, P < 0.05 \), \( r_{\text{col-I}} = 0.39, P > 0.05 \)). Importantly, after KK-Ay DN mice were injected intraperitoneally with antagomiR-21, the content of collagen fibers (CCF) and col-IV were decreased, instead of col-I. (Fig. 4a–d). Overall, these results suggested serum miR-21 was closely associated with excessive deposition of collagen fibers, especially for col-IV.

miR-21 overexpression decreased target smad7 expression

As described above, serum miR-21 was closely associated with renal structure and function of DN. then, miR-21 was how to
Serum miR-21 was closely associated with collagen fibers at 24 weeks of age. a Yellow or orange staining represented for CFs by Picrosirius red staining and blue staining represented for CFs by Masson staining. b Quantitative analysis of CCF by Picrosirius red and Masson staining, mean optical density (MOD) value of CFs was markedly increased in DN group compared with NC group. After the treatment of antagomiR-21, CFs was slightly but significantly decreased. c Western blotting band of col-IV and col-I protein. d Quantification of col-IV and col-I for western blotting results. The grey value of col-IV and col-I was markedly increased in DN group compared with NC group. After the treatment of antagomiR-21, col-IV was significantly decreased, whereas col-I was not changed. Light microscopy × 400. (Color figure available online only).

Discussion

In recent years, many studies have established many miRs are closely associated with DN [9,10,20,22], mostly focusing on their actions inside the cell from the tissues samples [29]. Because of the presence of potent ribonucleases, most investigators doubted that extracellular RNA could survive in the blood [30]. Up to now, many studies have documented circulating a large number of serum miRs remain stable and consistent in severe conditions [31–34]. It is very possible to detect serum miRs which help to diagnose this disease. Therefore, to indentify the relationship between serum miR-21 and tissue miR-21 expression, we examined serum and tissue miR-21 expression by real-time RT-PCR with the course of DN. We found that serum miR-21 expressions were shown to be increased in KK-Ay DN mice at 12–24 weeks of age. More surprisingly, the altered trends of serum miR-21 levels were significantly consistent with the expressions of renal tissues miR-21. Importantly, antagomiR-21 can decrease the expressions of serum and tissue miR-21. These results suggested that directly detecting serum miR-21 was the best substitute for tissue miR-21. Albumin creatinine ratio (ACR) has been considered as a good clinical predictor of renal lesions in DN [35]. Creatinine clearance ratio (Ccr) is generally considered as marker of renal filtration function [25]. We found that serum miR-21 expression was positively correlated with ACR and negatively correlated with Ccr. Moreover, the change trend of miR-21 was consistent with ACR, suggesting that miR-21 may be a biomarker reflecting for
the volume of urine protein excretion. More importantly, antagomir-21 can not only decrease serum miR-21 and ACR but also increase Ccr. Taken together; we concluded that the changes of circulating serum miR-21 can indirectly reflect renal function, which may be as a potential diagnostic biomarker for DN.

A well-organized collagen fiber was necessary to maintain structural and functional integrity of renal tissue. Excessive CFs accumulate in the kidney, which adversely affects the structure of the kidney and further lead to the loss of renal function [36]. Additionally, CFs was the most important ingredients comprised of GBM and mesangial matrix. Moreover, GBM and glomerular area (GA) were sensitive markers of DN [37]. Accumulating studies showed that smad7 may be an effective therapy for DN and renal fibrosis via altering expression of TGF-β1/Smad3-regulated miRs. Moreover, miR-21 participates in fibrogenic events in kidneys, lungs, heart, or other organs by regulating a unique array of targets [14, 38, 39]. Interestingly, our luciferase report gene assays suggested that smad7 was a validated miR-21 target, miR-21 over-expression significantly decreased smad7 expression. All these results showed that miR-21 was involved in the pathogenesis of DN by downregulating target smad7, due to the decrease of smad7, led to the deposition of col-IV and col-I, further resulted in GBM thickened and mesangial matrix hyperplasia. Additionally, we found that serum miR-21 was positively correlated with GBM, GA, CFC and col-IV, whereas, unrelated to col-I. Antagomir-21 can decrease GBM, GA, CFC and col-IV but not col-I. Thus, we speculated that both circulating serum miR-21 and tissue local miR-21 were closely associated with CFs formation of DN, especially for col-IV rather than col-I. In summary, our study suggested that serum miR-21 may be as a potential diagnostic biomarker for DN, both circulating serum and tissue local miR-21 attenuated renal function and structure by inhibiting target smad7. Although our results suggested serum miR-21 may be a new possible marker for the detection of DN, up to now, the measure of urinary albumin excretion rate is the most reliable indicator.

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Fig. 5 miR-21 overexpression decreased target smad7 expression. a Alignment of hsa-miR-21 and mmu-miR-21 with smad7-3’-UTR based on targetScan software from (http://www.targetscan.org/), several nucleotides in the 5’-region of miR-21 contain a perfect match with the 3’-UTR sequence of smad7 genes. b The results of luciferase report gene assays. c Representative photograph of smad7 protein by ICC. d The fluorescence intensity of smad7 proteins (p<0.05). (Color figure available online only).
References


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