Radix Puerariae and Fructus Crataegi mixture inhibits renal injury in type 2 diabetes via decreasing of AKT/PI3K

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Abstract

Background: Radix puerariae (RP) is a herbal medicines for diabetes, mainly because of anti-oxidative, insulin resistance and hypoglycemic effect. Fructus crataegi (FC) also possesses strong antioxidant activity in vitro. This study focused on the effects of herbal mixture of RP and FC (RPFC) on renal protection through a diabetic rat model.

Methods: Type 2 Diabetic model was established with high fat diet followed by injecting rats a low dose of STZ (25 mg/kg body weight). Rats were randomly divided into five groups: normal, high fat diet, diabetes mellitus, high fat diet plus RPFC prevention, and RPFC prevention before diabetes mellitus. RPFC was given to rats daily by intragastric gavage. The blood bio-chemical index and renal pathological changes were examined. The later includes hematoxylin and eosin staining, periodic acid schiff staining, and Masson trichrome staining. Protein levels of were determined by Western blot and immunohistochemical staining. mRNA levels were detected by RT-PCR.

Results: Rats prevented with RPFC resulted in decreasing blood glucose with corresponding vehicle treated rats. Glomerulus mesangial matrix expansion, renal capsule constriction, and renal tubular epithelial cell edema were less severe following RPFC prevention. Moreover, RPFC prevention reduced protein levels of PI3K, AKT, α-SMA and collagen IV in the kidney of diabetic rats.

Conclusion: Combined prevention with RPFC may inhibit the PI3K/AKT pathway in the kidney, thereby prevent renal injury in diabetic rats.

Keywords: Type 2 diabetes, Radix Puerariae, Fructus Crataegi, Renal injury, PI3K/AKT

Background

Diabetic nephropathy (DN) is one of the most common microvascular complications of diabetes mellitus and is a major cause of end stage renal disease (ESRD) [1]. DN is defined as damages to the kidney caused by diabetes and mainly presents as persistent proteinuria in diabetic patients. The pathological changes mainly include glomerular mesangial expansion, basement membrane thickening, and in severe cases, glomerular sclerosis [2]. However, current therapy for DN is still limited.

Although the pathogenesis of DN is not well understood, previous studies suggest that disturbed lipid metabolism [3], abnormal hemodynamics [4], and over-produced inflammatory mediators and cytokines [5] are the major causes of DN. Moreover, a high glucose environment and changes in cytokine production can cause functional changes in mesangial cells. Increasing evidence also suggests that inflammation and fibrosis play critical roles in the development and progression of DN [6].

Traditional Chinese medicine (TCM) has been used for treating DN in China [7]. However, the exact mechanisms by which TCM relieves DN symptoms remain unknown. Radix puerariae (RP), known as 'Gegen' in Chinese, is the root of Pueraria lobata (Willd.) Ohwi or Pueraria thomsonii Benth...
Puerarin, one of the active components of RP, has various pharmacological functions, such as eliminating oxygen free radicals, protecting myocardial cells, lowering blood pressure and inhibiting platelet aggregation [9]. It is known that puerarin can improve insulin resistance of in 3 T3-L1 lipocyte induced by free fatty acids [10]. Fructus crataegi (FC) is the fruit from *Crataegus pinnatifida* Bunge Rosaceae or *Crataegus pinnatifida* var. major N.E.Br., plants widely grown in China (http://www.theplantlist.org/). It often used to make tea in China, and the extract possesses strong antioxidant activity in vitro [11].

The exact renoprotection mechanisms of RPFC in STZ-induced diabetic rats remain unclear. We examined here whether RPFC could prevent renal injury in a rat model of DN, where progressive glomerular sclerosis and renal fibrosis was induced by a high-fat diet and a low-dose STZ.

**Methods**

**Drugs and reagents**

RP (the major component was purarine, then daidzin and daidzein; genistin and genistein were the least abundant) [12] and FC (essential components are flavonoids and organic acidic compounds) were provided by Shanghai Second Military Medical University. They were extracted separately using the following process: two times reflux extraction of one kilogram of each medicinal herb with 60% alcohol, the amounts of solvent of 5 folds, successively, and the extraction time of 90 min each [7]. The extract was reduced-pressure evaporated till the volume was 590 ml. RP and FC solutions were mixed at a volume ratio of 1:1 and used as the RPFC solution in this study.

**Animals and diabetic model**

Male 8-week-old Sprague-Dawley (SD) rats, weighing between 250 and 300 g, were purchased from the Laboratory Animal Center of the Academy of Zhejiang Medical Sciences (Zhejiang, Certificate No. 0012371). All rats were housed at a SPF-grade laboratory animal room in the Animal Laboratory Center of Ningbo University with 20–25 °C room temperature, 50–60% humidity, a 12/12 h light/dark cycle. The Standard diet was provided by Animal Laboratory Center of Ningbo University and the high fat diet was purchased from Pu Luteng Bio-Technique Co. Ltd. The high fat diet contained 16.9% fat and 10.2% casein in the standard diet. Rats were fed with the high fat diet (30 g per day for each rat) for 8 weeks to induce insulin resistance. The experimental diabetic model was produced by intraperitoneal injection of STZ (diluted to 1% in a 10 mmol/L citrate buffer at pH 4.5) at 25 mg/kg body weight after the rats were fasted for 12 h [13]. Rats in the control group were injected with an equal volume of the citrate buffer. Blood glucose was measured 72 h after STZ injection, and rats with blood glucose over 16.7 mmol/L were considered to be diabetic [14]. Rats were continued on the high fat diet except the rats of control group, in order to keep their blood glucose levels high.

**Experimental design**

The experimental procedure lasted 15 weeks, during which rats were kept on the standard or high fat diet according to their group assignments. RPFC were administered by intragastric gavage during the 7th–9th weeks, and rats not receiving RPFC were given normal saline. The day after the last administration of RPFC, STZ was injected, and rats not receiving STZ were injected with the vehicle (citrate buffer). 20 rats were randomly divided into five groups with four rats in each group: 1) Normal group, fed with the standard diet. 2) High fat diet (HF) group, fed with the high fat diet. 3) Diabetes mellitus (DM) group, fed with the high fat diet, gavaged with normal saline, and injected with STZ. 4) High fat diet plus RPFC prevention group (HP), fed with the high fat diet, gavaged with RPFC, and injected with the citrate buffer. 5) Diabetes mellitus plus RPFC prevention group (DP), fed with the high fat diet, gavaged with RPFC, and injected with STZ (Fig. 1). As increased time of high fat diet feeding, obesity and resistance to insulin would be more apparent [15]. The beta-islet cells susceptibility of STZ increase in rats feed with the high fat diet. So we add HF group to observe whether high-fat rats have metabolism disorders or renal injury, and add HP group to confirm whether RPFC has impacts on the above changes.

**Measurement of body weight, blood glucose, and urinary protein**

Blood glucose was measured once a week and body weight was measured once a month. Blood samples were collected from the tail vein to measure the blood glucose level. The fasting blood glucose (FBG) was measured after a 12 h fasting, and postprandial blood glucose (PBG) was measured at 2 h after a meal. Blood glucose...
level was measured with One Touch Ultra test strips and blood glucose meter (Johnson& Johnson Medical Ltd., Shanghai, China).

Clinically, microalbuminuria was defined as urine albumin excretion rate (UAER) in the range of 20–200 μg/min or urine albumin at 30–300 mg/g creatinine [16]. Rats were considered to have nephropathy if they displayed microalbuminuria. Rats were housed individually in metabolic cages for 24 h to collect urinary samples analyzed with a MODULAR P800 Automation Biochemist Analyzer (Roche, Basel, SWIT).

Oral glucose tolerance test (OGTT), and measurement of insulin and other biochemical parameters
The oral glucose tolerance test was performed on overnight-fasted rats at the end of the study. Rats were fasted for 12 h and then given a glucose solution (2 g/kg body weight), and blood samples were collected before and at 30, 60, and 120 min after the glucose solution was given [17]. Insulin and other biochemical parameters were measured at the end of study. Blood samples were collected from the femoral artery into EDTA-anticoagulant tubes, collected the serum for the detection of serum insulin, total protein (TP), high-density lipoprotein cholesterol (HDL-C), low-density lipoproteins (LDL-C), total cholesterol (TC), triglycerides (TG), urea nitrogen (BUN), creatinine (CREA) and uric acid (UA). The levels of serum insulin were measured with a Rat Insulin ELISA kit (Qiaodu, Shanghai, China). Other biochemical indicators were measured with a MODULAR P800 Automation Biochemist Analyzer (Roche, Basel, SWIT). At last, rats were sacrificed and kidney tissues were saved standby.

Histological examination of the kidney
Renal tissues were fixed for 48 h, dehydrated through a graded series of ethanol, embedded in paraffin wax, and cut into 3 μm sections. The sections were subjected to hematoxylin and eosin (H&E), periodic acid schiff (PAS), and Masson trichrome staining and observed under a microscope.

Real time reverse transcriptase-polymerase chain reaction (RT-qPCR)
Total RNA was extracted from renal tissues using TRIzol reagent (Invitrogen, USA). 1 microgram of total RNA was for reverse transcription. Polymerase chain reactions were performed in a Lightcycler 480 II Authorized Thermal Cycler (Roche, Basel, SWIT) using the following protocol: denaturation at 95 °C for 5 min, followed by 45 cycles of 95 °C 10 s, annealing 20 s, and 72 °C 30 s [18]. The primers used were as follows: α-SMA, F: 5′-CATTTGCTGACGAGGATGCAGAA-3′, and R: 5′-GAGCATTGGCGTGACAA-3′; collagen IV, F: 5′-GTTGGTCTACCGGACCTC-3′, and R: 5′-GTTGGTCTACCGGACCTC-3′; β-actin, F: 5′-CTGAACTGCTAAGGCCACG-3′, and R: 5′-GAC-CAGAGGCTACACGGACAA-3′. Relative mRNA levels were determined with the 2−ΔΔCt method using the gene β-actin as the internal reference.

Western blot analysis
Renal tissues of 40 mg were lysed for protein extraction. Samples (50 μg) were separated by 8% sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE) gels and transferred to polyvinylidene fluoride membranes. The membranes after blocked were incubated with primary antibodies to Phosphoinositide 3-kinase (PI3K) (p85) (Abcam, Cambridge, MA, 1:1000), protein kinase B (AKT) (Signalway Antibody LLC, Maryland, USA, 1:1000), and α-SMA (Abcam, Cambridge, MA, 1:200) at 4 °C overnight. Then incubated with a HRP-labeled secondary antibody (Beyotime, China). The immunoreactions were detected with a gel imaging and analysis system (Tanon, Shanghai, China). The density values of bands were quantified using the software Image J (NIH, Maryland, USA).

Immunohistochemical staining
Renal slides (3 μm) were used to perform immunohistochemical staining of α-SMA and collagen IV on renal tissue sections [19] with the following primary antibodies: monoclonal mouse anti-rat α-SMA (Abcam, Cambridge, MA, 1:50) and polyclonal rabbit anti-rat collagen IV (Abcam, Cambridge, MA, 1:200). Quantitative analysis of the brown positive staining in glomerulus and tubules was performed using Image-Pro Plus 6.0 (Media Cybernetics, USA).

Statistical analysis
All the data are presented as mean ± standard error of the mean (S.E.M.). The differences among groups were analyzed by randomized block design analysis of variance by using SPSS (version 13.0) software. P values below 0.05 were considered statistically significant.

Results
Blood glucose, body weight, and urinary protein measurement and OGTT
All the rats were alive at the end of study. FBG levels were measured from the 4th week till the end of the study (Fig. 2a). The rats in HF, DM, and DP groups showed higher FBG levels from the 5th to the 15th week than the normal group. After the injection of STZ, both FBG and PBG levels in both the DM and DP groups rose rapidly. However, FBG levels in the DP group declined slowly, and were even lower than in HF group from the 10th week to the 15th week. However, FBG levels as well as PBG levels in the HP group started to decrease from the 7th week and became lower than those in the HF
group and comparable to those in the normal group. PBG levels 2 h after a meal were also measured starting from the 4th week (Fig. 2b). The rats in the HF, DM and DP groups showed higher PBG levels from the 8th to the 15th week than the normal group. However, from the 10th week, PBG levels in the DP group declined slowly, and were significantly higher than those in the HF group but lower than those in the DM group till the end of the study. This could be due to RPFC prevented effectively the increase in serum glucose in the model.

As shown in Fig. 2c, body weights increased significantly in the HF, DM, DP and HP groups as compared with the normal group. Compared with the normal group, urinary protein was significantly higher in all other groups. Moreover, urinary protein in the HP group was lower than in the HF group till the end of the study. This could be due to RPFC prevented effectively the increase in serum glucose in the model.

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Biochemical parameters
Table 1 shows the measurement results of biochemical parameters in all rats. Compared with the normal group, LDL-C and TC levels were significantly different in the DM group ($p < 0.05$), and TC level was significantly increased in the other four groups, and HDL-C level was significantly increased in the HF and DP groups. These data indicate that oral administration of RPFC may improve lipid metabolism.

Renal histology
Glomerular hypertrophy was observed in the HF group, as glomerular extracellular matrix accumulation and renal capsule constriction, indicators of early DN, were evident in the DM group. So, renal injury was more severe in the DM group than those in the HF group. Mesangial matrix
expansion and renal capsule constriction were visible in HP and DP groups, but were much less severe than in the corresponding RPFC untreated groups (Fig. 3a).

The hyperplastic state of collagen was observed by masson trichrome staining and glycogen was observed by PAS staining (Fig. 3b and c). The sedimentary hepitin granules (arrow) were observed in HF and DM groups (Fig. 3b), and collagen deposition (arrow) was particularly evident in DM groups (Fig. 3c). In addition, the renal tubular appeared foamy in the HF and DM groups because of the accumulation of fats and glycogen, while RPFC treatment reduced the accumulation of fat and glycogen in the kidney.

**Expression of PI3K and AKT in renal tissues in rats**

PI3K and AKT play pivotal roles in regulating cell proliferation and apoptosis in the kidney [20]. We quantified the

|                | N       | HF      | DM      | HP      | DP      |
|----------------|---------|---------|---------|---------|---------|
| TP             | 66.60±4.24| 80.77±7.45| 64.47±7.06| 74.50±1.92| 78.17±6.09|
| HDL-C          | 0.86±0.20| 1.39±0.31*| 1.13±0.11| 1.05±0.14| 1.36±0.28*|
| LDL-C          | 0.56±0.02| 0.59±0.04 | 0.71±0.07*| 0.62±0.04| 0.59±0.04  |
| TC             | 1.62±0.01| 2.29±0.33*| 2.81±0.04*| 1.91±0.25*| 2.55±0.03*|
| TG             | 2.43±1.73| 2.77±0.68 | 2.47±0.21| 2.75±0.32| 2.53±1.08  |
| BUN            | 5.40±0.86| 5.73±0.51 | 5.25±0.78| 5.66±0.28| 5.95±0.64  |
| CREA           | 23.00±4.24| 21.00±2.65| 23.00±2.00| 20.00±1.00| 21.50±0.71|
| UA             | 76.50±17.68| 129.33±37.31| 103.3±14.14| 127.67±47.26| 97.50±13.44|

*Indicates p < 0.05 between other groups versus the normal group.
expression levels of PI3K and AKT by Western blot analysis. As shown in Fig. 4, the protein levels of PI3K and AKT were significantly higher \( (P < 0.05) \) in rats of the HF and DM groups than in the control rats. However, the protein levels of PI3K and AKT were lower in rats of the HP and DP groups than in the corresponding RPFC untreated groups \( (P < 0.05, \text{Fig. 4}) \). These results suggest that RPFC might protect against renal injury by decreasing the expression levels of PI3K and AKT proteins in diabetic rats.

**Expression of α-SMA and collagen IV**

As shown by real-time PCR, prevention with RPFC in rats of the DP and HP groups significantly reduced the expression of α-SMA and collagen IV as compared with the DM group. Moreover, there was a significant difference between the HF group and HP group \( (*P < 0.05) \), \( \text{Fig. 5a and b} \). The protein levels of α-SMA was significantly higher \( (P < 0.05) \) in the DM group than in the normal group (Fig. 5c and d). The expressions of both α-SMA and collagen IV proteins in the DM group were increased (Fig. 5e). These data suggest that RPFC might decrease the protein levels of α-SMA and collagen IV in diabetic rats thus alleviating renal injury.

**Discussion**

Human beings are heading for a health disaster with rising obesity levels and the increasing incidence of type 2 diabetes. However, some studies suggest diabetic nephropathy occupies for approximately 20 to 30% of diabetic. The incidence of diabetes and DN have seen a rapid increase in China over the last decade [9]. The diabetic nephropathy patients have a higher mortality, thus, effective preventive and adequate treatment are of uttermost importance. Despite the lifestyle interventions (such as smoking cessation, moderation of alcohol intake, healthy diet, weight control

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**Fig. 4** Supplement of RPFC decreased PI3K and AKT in high fat fed rats and DM rats. Protein expression of PI3K a and b, and AKT c and d. Renal PI3K and AKT increased in the kidneys of the HP and DM group rats in comparison to that of the normal group \( (*P < 0.05) \). Renal AKT decreased in the kidneys of the HP and DP group rats in comparison to that of the HF and DM group \( (**p < 0.05) \). Bars represent mean ± SD from three experiments.
and physical activity) which can help to prevent progression or occurrence of diabetic nephropathy impairment. However, the current treatment of using ACEI and ARBs are not effective against DN [21]. In recent years, several studies have shown that TCM can ameliorate renal injury in diabetic rats [14]. In our present study, RP and FC with high specific curatorial value are both natural medicine in plants. Puerarin has exhibited classic estrogen-like biological activities and shown remarkable cardiovascular protective effects in in vivo and in vitro experiments. It improves insulin resistance and lowers blood sugar and lipid levels [22]. Moreover, puerarin also has demonstrated profound anti-oxidative effects. FC is edible and is often made into nutritious haw wine, tea, and sweetened hawthorn fruit rolls. The possible mechanism of these effects was thought to be that it can induce GLUT4 protein expression [23], inhibit oxidative stress [22]. In the present study, we find that RPFC reduces microalbuminuria effectively in early DN rats. Albuminuria is a prognostic marker of DN because the presence of proteins in the urine aggravates kidney damage [24]. To date, the research on FC has focused on its role in reducing blood lipids and cholesterol levels, lowering blood pressure, and anti-lipid peroxidation, such as treating cardiovascular diseases [25], but no study on FC and renal diseases has been conducted.

The α-SMA is involved in progressive renal changes [26] in both human and experimental rats [27]. For α-SMA is normally expressed at low levels in mesangial cells but not in podocytes. However, it becomes widely

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**Fig. 5** The expression of α-SMA and collagen IV in renal. The mRNA expression of α-SMA a and collagen IV b, the protein expression of α-SMA determined by Western blot c and d and immunohistochemical e, the protein expression of collagen IV determined by immunohistochemical f. The data are presented as mean ± SEM, n ≥ 3, *p < 0.05.
expressed when mesangial cells and podocytes are damaged and show phenotypic changes. Moreover, increased level of renal α-SMA in diabetic rats is associated with tubulointerstitial fibrosis involved in DN [28]. Glomerular mesangial cells with higher expression of α-SMA show enhanced contractility, resulting in the changes of blood flow dynamics in glomerulus, which eventually lead to glomerular sclerosis [29]. Moreover, the presence of α-SMA will induce synthesis and secretion of extracellular matrix (ECM), which also leading to glomerular sclerosis. Collagen IV can be synthesized and secreted by glomerular mesangial cells, endothelial cells, epithelial cells and tubular epithelial cells, it plays a major role in renal fibrosis in progression of DN [30].

Our findings showed that the expression of α-SMA and collagen IV were significantly increased in diabetic rats, while decreased with prevention of RPFC. The preventive activity of RPFC in diabetic rats was partly mediated by inhibition of α-SMA expression, which was consistent with previous studies showing that the high glucose milieu of diabetes can increase the synthesis of α-SMA [27]. Moreover, it was reported that rats with diabetes exhibited markedly increased renal Smad1 and collagen IV expression and extracellular matrix deposition [31]. These data suggest that RPFC might play a role in inhibiting PI3K/AKT signaling pathway, which indirectly leads to reduction of α-SMA and collagen IV.

In our previous study, RPFC did not affect normal rats. We also found that there is no difference in serum insulin levels among the five groups (data not shown). Moreover, we found that RPFC decreased blood glucose level not only under hyperglycemic but also under hyperlipidemia conditions. A recent research [32] demonstrated that puerarin had more pronounced antioxidative effect than losartan. In addition, it was demonstrated that puerarin could attenuated proteinuria and podocyte injury in diabetic rats probably through reduction of oxidative stress in the diabetic kidney [32]. Similarly, our current study suggests that RPFC might target PI3K/AKT pathway to protect renal tissues and delay the progression of DN.

Here we provide potential mechanisms to explain how herbal mixture of RP and FC affects rats with early DN. It would also be necessary to determine whether RPFC could help ameliorate renal injury in patients with more advanced DN and with other nephropathies in future studies. Based on the current study, RPFC is safe and no side effects were observed. One traditional Chinese medicine research in mice and rats has observed acute and long-term toxicity of crude flavone of pueraria gavaged for 90 days, points out that crude flavone of pueraria has no accumulated toxicity. However, it is not known whether long-term treatment with RPFC would cause any severe side effects. It has been known that RP is not suitable for people have gastritis diseases, and RP given intravenously could cause itching and nausea, while FC is not good for anyone who has digestive diseases.

**Conclusion**

In summary, our study demonstrated for the first time that RPFC was effective in preventing early DN in rats, possibly by suppressing PI3K/AKT signaling and repressing the expression of α-SMA and collagen IV in glomerulus.

**Abbreviations**

FC: Fructus crataegi; RP: Radix puerariae

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**Availability of data and materials**

The datasets analyzed during the current study available from the corresponding author on reasonable request.

**Authors’ contributions**

BB, SB, XZ, ZC conducted the actual study and the statistical analysis. MH, MZ and ZC were involved in developing the idea, designing of the study and the write up of the manuscript. All authors approved the submitted version of the manuscript.

**Consent for publication**

“Not applicable” in this section.

**Competing interests**

The authors declare that they have no competing interest.

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