Introduction

Diabetes mellitus is increasingly prevalent worldwide. Diabetic nephropathy (DN) is one of the most common complications in diabetes mellitus, which is the leading cause of end-stage renal disease (1). The characteristic features of this disease are persistent albuminuria and structural alterations, such as thickened glomerular basement membrane and progressive accumulation of extracellular matrix proteins in the glomerular mesangium (2). Progressive loss of renal function, which ultimately results in end-stage renal disease requiring dialysis or transplantation, is an increasing health problem around the world (3).

The early stage of diabetic mellitus is characterized by renal hyperfiltration, which promotes the eventual development of DN. Although preglomerular afferent arteriolar dilation and diminished responsiveness of this vascular segment to a variety of vasoconstrictors have been proposed to be a contributing factor (4), increasing evidence suggests that dysfunction of glomerular mesangial cells (MCs) is also involved in the diabetic hyperfiltration (5 – 7). MCs are important targets of metabolic abnormalities in diabetes and contribute to the functional and structural abnormalities of DN. MCs are located within glomerular capillary networks, regulating effective filtration surface area. MC contractility is impaired in diabetes and this impairment disrupts normal glomerular hemodynamics (8). Decreased Ca2+ influx contributes
to the impaired contractile function of MCs, presumably due to altered calcium channel number and/or activity (9).

Recently, canonical transient receptor potential (TRPC) proteins have been proposed as Ca\(^{2+}\)-permeable cation channels that are activated in response to stimulation of G-protein-coupled receptors (10 – 12). The TRPC family includes seven related members, designated as TRPC 1 – 7 (13). TRPC6 in this family is expressed in MCs and participates in angiotensin (Ang) II-induced MC contraction (14 – 16). Our previous study showed that in glomeruli isolated from streptozotocin (STZ)-induced diabetic rats, TRPC6 was markedly reduced compared with the glomeruli of control rats (15, 17). Furthermore, TRPC6 mRNA in MCs was also significantly decreased by high glucose (17). Accumulating evidence implies that downregulation of TRPC6 leads to significantly decreased Ca\(^{2+}\) signaling and hypocontractility of diabetic MCs. Thus, the reduced mesangial responsiveness to vasoactive agents seen in diabetes may be partially due to reduced TRPC6 expression (17). Multiple pathogenic mechanisms are now believed to contribute to DN, including inflammatory cytokines, autacoids, and oxidative stress. Among them, oxidative stress has been suggested to play an important role in the pathogenesis of DN (18 – 20). An increased production of reactive oxygen species (ROS), including superoxide anion, contributes to the pathogenesis of renal injury in diabetes mellitus (21). Superoxide dismutase (SOD) is the major antioxidant enzyme for superoxide removal, which converts superoxide into hydrogen peroxide and molecular oxygen. The hydrogen peroxide is further detoxified to water by catalase or glutathione peroxidase (22). As SOD and catalase activities are decreased in diabetes, the improvement of the antioxidant system including SOD could be a potential therapeutic target in diabetic nephropathy (23).

Tempol (4-hydroxy-2,2,6,6-tetramethyl-piperidine-1-oxyl) is a stable, metal-independent low-molecular weight SOD mimetic with an excellent cell permeability. The reaction of tempol with superoxide anion to form hydrogen peroxide accounts for its “SOD mimetic” action. As reported, tempol normalized blood pressure in hypertensive rats and dilated afferent arterioles of diabetic rabbits (24, 25). Also, tempol protected the kidney from ischemic damage and inhibited hypertensive renal damage in the Dahl salt-sensitive model (26, 27). However, few studies have investigated the preventative effect of tempol on DN and its possible mechanisms of action. Thus, the present study was carried out to examine the effect of tempol on progression of DN and the expressions of several diabetes-associated proteins in a streptozotocin rat model of diabetes. The findings in this study may provide a theoretical basis for clinical application of tempol.

Materials and Methods

Animals and experimental protocols

Male Sprague-Dawley rats (up to 8 weeks of age) were purchased from the Experimental Animal Center of Anhui (Hefei, China) (Certificate No. SCXK 2005-001). The animals were housed individually in cages at constant humidity (55% ± 5%) and temperature (25°C ± 2°C) with an electrically controlled 12-h light–dark cycle. Food and water were provided ad libitum. This study was approved by the Animal Care and Use Committee of Anhui Medical University, and all studies were conducted in accordance with “Guiding for the Care and Use of Laboratory Animals” adopted by the Committee on the Care and Use of Laboratory Animals of Anhui Medical University. Diabetes was induced by a single injection of STZ (Sigma Chemical Co., St. Louis, MO, USA) into the rat tail vein at 60 mg/kg in sodium citrate buffer (0.01 M, pH 4.5) after fasting overnight. Age- and weight-matched control rats were given an equivalent volume of the vehicle (citrate buffer) alone. The fasting blood glucose (FBG) levels in tail vein blood were determined 72 h later and periodically thereafter with a glucometer (LifeScan OneTouch glucometer; Johnson & Johnson, New Brunswick, NJ, USA). STZ-injected rats with sustained elevation of blood glucose above 16.7 mM were designated as diabetic rats (28). The diabetic rats were randomly divided into 2 groups matched for body weight and blood glucose: untreated DM group (Model group, n = 10) and a diabetic group given tempol (Sigma Chemical Co.) in their drinking water (1 mM) for 6 weeks (Tempol group, n = 10), beginning 24 h after the induction of diabetes. Age- and weight-matched rats with sodium citrate buffer injection alone served as controls (Control group, n = 10).

Sample preparation

Body weight, daily food intake, daily water intake, and daily urine output of all rats were measured at pre-therapy, 4 and 6 weeks after tempol treatment. Urine was collected via a metabolic cage. After 6 weeks of tempol treatment, the rats after fasting overnight were anesthetized with 10% chloral hydrate (30 mg/kg body weight intraperitoneally), and then blood was drawn from the abdominal aorta for measuring blood chemical parameters. After blood sampling, kidneys were processed for biochemical, histologic, and immunochemical examinations as described below.
Measurement of urinary parameters

After 24-h urine collection, the samples were centrifuged (3000 rev/min, 4°C, 10 min) and the supernatant was stored at −20°C. The urine samples were used to calculate urine volume and urinary albumin excretion (UAE). Urine protein was measured by the Bradford method (Bio-Rad, Richmond, VA, USA). Meanwhile, urinary TGF-β1 excretion in daily urine was determined by Enzyme-linked Immunosorbent Assay (ELISA) (Senxiong Bio-Tech, Shanghai, China).

Measurement of blood parameters

The FBG levels in tail vein blood were determined with a glucometer at pre-therapy, 4 and 6 weeks after tempol treatment. Serum samples from rats were collected for measurement of glycosylated serum protein (GSP), fasting serum insulin (FINS), creatinine (Cr), and blood urea nitrogen (BUN). The detection was carried out using diagnostic kits (Jiancheng Institute of Bioengineering Company, Nanjing, China). GSP was determined with the fructosamine assay. BUN was reacted with diacetyl to produce a red compound and measured by spectrophotometer. Serum Cr was measured by the modified picric acid method. FINS was assayed by ELISA (Yuanye Bio-Tech, Shanghai, China). Insulin sensitivity index [ISI = 100 / (FBG × FINS)] was applied to assess the status of insulin resistance.

Measurement of MDA level and activities of SOD and GSH-Px in the kidney

After blood sampling, the kidneys were quickly removed from the animals and the surrounding fat was removed. They were washed in a sterile saline solution and weighed. The renal tissue was homogenized in a nine volumes of ice-cold normal saline, and then the homogenates were centrifuged (4000 rev/min, 4°C, 10 min). The level of malondialdehyde (MDA) and activities of superoxide dismutase (SOD) and glutathione peroxidase (GSH-Px) in the kidney were measured following the manufacturer’s instruction (Jiancheng Institute of Bioengineering Company, Nanjing, China). 

Histological examination

The renal samples were fixed in 4% paraformaldehyde, embedded with paraffin, and cut into sections of about 4–μm thickness and stained with periodic acid–Schiff (PAS) staining. The slides were examined under light microscopy with magnification of 400 × by a pathologist kept unaware of the experimental profile. The purple color in glomeruli was defined as “positive” and the positive area of each glomerulus was measured using the JD801 morphological microscope image analysis system (JEDA Science-Technology Development Co., Ltd., Jiangsu, China). The positive score (PS) of glomeruli in one rat was calculated by averaging positive scores of 30 glomeruli from the same rat using the following formula:

$$PS = \frac{\sum_{i=1}^{30} \[\frac{AP_i}{AG} \times 100\]}{30}$$

where AP represents the positive area and AG represents the area of the glomerulus. Six rats were examined in each group.

RT-PCR

Total kidney RNA was extracted using TRIzol (Invitrogen, Darmstadt, Germany) according to the instructions from the manufacturer. The extracts were re-suspended in 20 μl of DEPC-treated water. RNA concentration was determined using a biophotometer (Shanghai Scientific China, Shanghai, China). Four micrograms of RNA underwent reverse transcription to generate cDNA using random hexamer primers and Primescript™ RTase (Takara Bio, Otsu). PCR was conducted using a RT-PCR kit (Promega, Madison, WI, USA). Primer sequences (Shanghai Sangon Bio-Tech, Shanghai, China) and annealing temperatures for TGF-β1, Coll IV, TRPC6, and β-actin PCR are listed in Table 1. PCR products and a 1-kb DNA molecular weight marker (Promega) were then electrophoresed on a 1% agarose gel, and the gel was visualized and photographed using a gel imaging system (Biosens SC810X; Shanghai Bio-Tech, Shanghai, China). The gel imaging software was used for quantitative analyses. Relative quantities of target gene expression were compared after normalization to the value of β-actin, a housekeeping gene.

<table>
<thead>
<tr>
<th>mRNA</th>
<th>Sense primer</th>
<th>Antisense primer</th>
<th>Anneal temperature (°C)</th>
<th>PCR target (bp)</th>
<th>Cycles</th>
</tr>
</thead>
<tbody>
<tr>
<td>TGF-β1</td>
<td>5'-CTG TCC AAA CTA AGG CTC GC-3'</td>
<td>5'-AGA CAG CCA CTC AGG CGT A-3'</td>
<td>55</td>
<td>432</td>
<td>38</td>
</tr>
<tr>
<td>Collagen IV</td>
<td>5'-ATT CCT TTG TGA TGC ACA CCA G-3'</td>
<td>5'-AAG CTC TAA GCA TCC GCG TAG TA-3'</td>
<td>56</td>
<td>151</td>
<td>38</td>
</tr>
<tr>
<td>TRPC6</td>
<td>5'-AAA GAT ATC TTC AAA TTC ATG GTC-3'</td>
<td>5'-CAC GTC CGC ATC ATC CTC AAT TCC-3'</td>
<td>54</td>
<td>327</td>
<td>38</td>
</tr>
<tr>
<td>β-Actin</td>
<td>5'-AGC ATT TGC GGT GCA CGA TGG AGG G-3'</td>
<td>5'-ATG CCA TCC TGC GTC TGG ACC TGG C-3'</td>
<td>53</td>
<td>606</td>
<td>38</td>
</tr>
</tbody>
</table>

Table 1. Oligonucleotide primers used for cDNA amplification
Statistical analyses

Data were expressed as means ± S.D. One-way analysis of variance (ANOVA) was used to test differences among groups and the Student-Newman-Keuls (SNK) test was adopted to assess which two groups have significant difference. Value of $P < 0.05$ was considered to be significant.

Results

Physical and metabolic parameters

Data were obtained at pre-therapy, 4 and 6 weeks after tempol treatment. It was observed that the body weight increased in the control group. The body weights in the model group were significantly lower in comparison with the control group ($P < 0.01$) at the end of the experiment. In contrast, the weights in diabetic rats after tempol treatment were higher than those in the model group.

As shown in Table 2, daily food and water intake (normalized to rat body weight) of the control group were unchanged throughout the experiment. However, the daily food and water intake in the model group were significantly higher than those in the control rats. Also, daily food and water intake significantly decreased ($P < 0.01$) in the tempol group. Similarly, urine output (UO, normalized to rat body weight) in the control group did not change throughout the experiment. In contrast, UO was significantly higher in model group than in the control group ($P < 0.01$). After tempol treatment, UO was lower than that in the model group ($P < 0.01$). An increase in UAE amount is the earliest manifestation of renal disorder in diabetes. The time course of daily UAE was shown in Table 2. UAE in the model group increased significantly compared with that in the control group ($P < 0.01$). Tempol treatment decreased UAE of diabetic rats significantly ($P < 0.01$) in comparison with the model group at the end of the experiment. Likewise, urinary TGF-β1 excretion in daily urine in the model group increased significantly compared with that in the control group ($P < 0.01$). Tempol administration decreased this parameter of diabetic rats significantly ($P < 0.01$). These results indicate that hypermetabolism in diabetic rats was persistent.

As shown in Table 2, FBG and GSP were significantly higher in the model group than those in the control group ($P < 0.01$). Tempol treatment showed a decreased tendency in FBG and GSP and the difference was significant ($P < 0.01$). However, blood glucose did not return to normal level at the end of the experiment.

At the 6th week after tempol treatment, FINS levels in the model group increased markedly compared with those in the control group, but it was decreased by tempol treatment. Meanwhile, ISI of the model group was lower than that in the control group ($P < 0.01$). Increased ISI of

<p>| Table 2. Effect of tempol on the physiological parameters in diabetic rats induced by STZ |</p>
<table>
<thead>
<tr>
<th>Parameter</th>
<th>Pre-therapy</th>
<th>4 weeks</th>
<th>6 weeks</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Control</td>
<td>Model</td>
<td>Tempol</td>
</tr>
<tr>
<td>BW, g</td>
<td>250 ± 18</td>
<td>228 ± 19*</td>
<td>234 ± 25</td>
</tr>
<tr>
<td>Daily food intake, g/100 g BW</td>
<td>9 ± 2</td>
<td>12 ± 3*</td>
<td>14 ± 3*</td>
</tr>
<tr>
<td>Daily water intake, g/100 g BW</td>
<td>15 ± 3</td>
<td>59 ± 8*</td>
<td>62 ± 18**</td>
</tr>
<tr>
<td>UO, ml/100 g BW</td>
<td>8 ± 3</td>
<td>49 ± 10**</td>
<td>52 ± 13**</td>
</tr>
<tr>
<td>Urinary albumin excretion, mg/day</td>
<td>6.47 ± 0.89</td>
<td>24.88 ± 4.97**</td>
<td>23.35 ± 5.02**</td>
</tr>
<tr>
<td>Urinary TGF-β1 excretion, ng/day</td>
<td>3.55 ± 0.71</td>
<td>18.80 ± 4.27**</td>
<td>20.03 ± 4.56**</td>
</tr>
<tr>
<td>GSP, mM</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
</tr>
<tr>
<td>FBG, mM</td>
<td>4.9 ± 0.5</td>
<td>20.0 ± 2.3**</td>
<td>20.5 ± 2.0**</td>
</tr>
<tr>
<td>FINS, mU/L</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
</tr>
<tr>
<td>ISI</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
</tr>
<tr>
<td>KW/BW, mg/g</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
</tr>
</tbody>
</table>

BW, body weight; UO, urine output (was normalized to rat body weight); GSP, glycosylated serum protein; FBG, fasting blood glucose; FINS, fasting serum insulin; ISI, insulin sensitivity index = 100 / (FBG x FINS); KW, kidney weight (both kidneys); KW/BW, ratio of kidney weight to body weight; Control, nondiabetic rats; Model, diabetic rats; Tempol, diabetic rats treated with tempol (was supplemented to the rat’s drinking water at a concentration of 1 mM); ND, not determined. Data are presented as means ± S.D. n = 10 rats in each group. *P < 0.05, **P < 0.01 vs. control rats; *P < 0.05, **P < 0.01 vs. model rats.
Renal Protection by Tempol in DM rats

Diabetic rats was observed in the tempol group compared with the model group. Therefore, tempol treatment could elevate ISI of diabetic rats. Furthermore, the kidney weight / body weight ratio (KW/BW) was significantly increased in the model group in comparison with the control group ($P < 0.01$). After tempol administration, KW/BW was decreased significantly compared with that in the model group ($P < 0.01$). In addition, the serum levels of BUN and Cr in the model group were markedly higher than those of the control group ($P < 0.01$). The creatinine clearance rate of the model group decreased obviously compared with that in the control group. Tempol significantly affected the elevated BUN and Cr concentration of diabetic rats ($P < 0.01$). Meanwhile, tempol treatment significantly improved the creatinine clearance rate of the diabetic group ($P < 0.01$) (Table 3).

**MDA level and activities of SOD and GSH-Px in renal tissue**

The MDA level and activities of SOD and GSH-Px in renal tissue are shown in Table 4. The MDA level was increased in the model group compared with that in the control group ($P < 0.01$). Tempol treatment reduced MDA level compared to that in the model group ($P < 0.01$). The activities of SOD and GSH-Px in the control group were significantly higher than those in the other two groups ($P < 0.01$). The activities of SOD and GSH-Px were markedly reduced in the model group compared with those of the control group. The tempol treatment relieved these alterations ($P < 0.01$) (Table 4).

**Renal histology and morphometric analysis**

Glomeruli were easily distinguished by their characteristic circular morphological aspect bordered by the peripheral lumen. Figure 1 shows representative histopathological changes in the kidney with PAS staining and morphometric analysis in each group at the 6th week after tempol treatment. The model group showed extensive mesangial matrix expansion (Fig. 1B), whereas no change was observed in the control rats (Fig. 1A). PAS-positive areas increased significantly in the model group compared with those in the control group ($50.51 \pm 6.67$ vs. $17.12 \pm 3.67$, $P < 0.01$) (Fig. 1D). The positive score of glomeruli PAS staining increased markedly in the model group compared with the control group. However, the positive score of glomeruli PAS staining was reduced significantly ($22.79 \pm 6.88$ vs. $50.51 \pm 6.67$, $P < 0.01$) by tempol treatment (Fig. 1D).

**The mRNA expression of TGF-β1, Col IV, and TRPC6 in the kidney**

Consistent with the results of immunohistochemistry, expression of TRPC6 mRNA in diabetic kidneys was decreased dramatically compared with the control group ($P < 0.01$). However, expression of TGF-β1 mRNA and

<table>
<thead>
<tr>
<th>Group</th>
<th>Creatinine ($\mu$M)</th>
<th>Blood urea nitrogen ($\mu$M)</th>
<th>Creatinine clearance rate (ml/min·kg)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>49.80 ± 3.65</td>
<td>5.78 ± 0.22</td>
<td>3.14 ± 0.52</td>
</tr>
<tr>
<td>Model</td>
<td>123.62 ± 18.33$^{**}$</td>
<td>15.95 ± 2.61$^{**}$</td>
<td>1.88 ± 0.17$^{**}$</td>
</tr>
<tr>
<td>Tempol</td>
<td>79.03 ± 7.68$^{**}$</td>
<td>8.98 ± 0.89$^{**}$</td>
<td>3.00 ± 0.53$^{**}$</td>
</tr>
</tbody>
</table>

The abbreviations are the same as in the legend to Table 2. Data are presented as the means ± S.D. n = 10 rats in each group. $^{**}P < 0.01$ vs. control rats; $^{*}P < 0.01$ vs. model rats.

<table>
<thead>
<tr>
<th>Group</th>
<th>T-SOD (U/mg protein)</th>
<th>MDA (nmol/mg protein)</th>
<th>GSH-Px (U/mg protein)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>153.88 ± 29.82</td>
<td>3.80 ± 1.25</td>
<td>159.30 ± 33.34</td>
</tr>
<tr>
<td>Model</td>
<td>65.68 ± 20.15$^{**}$</td>
<td>7.89 ± 1.72$^{**}$</td>
<td>82.58 ± 20.18$^{**}$</td>
</tr>
<tr>
<td>Tempol</td>
<td>95.86 ± 32.09$^{*}$</td>
<td>4.56 ± 1.51$^{**}$</td>
<td>124.51 ± 35.07$^{**}$</td>
</tr>
</tbody>
</table>

The abbreviations are the same as in the legend to Table 2. Data are presented as the means ± S.D. n = 10 rats in each group. $^{**}P < 0.01$ vs. control rats; $^{*}P < 0.05$, $^{*}P < 0.01$ vs. model rats.
Col IV mRNA in diabetic kidneys was increased dramatically relative to the control ($P < 0.01$). Up-regulation of TGF-$\beta_1$ mRNA and Col IV mRNA expression was obviously decreased by administration of tempol ($P < 0.05$) (Fig. 2). Compared with the model group, tempol treatment increased the expression of TRPC6 mRNA significantly ($P < 0.05$) (Fig. 2).
Renal Protection by Tempol in DM rats

Discussion

As an important complication of diabetes mellitus, DN has become the most common cause of end-stage renal failure among patients undergoing chronic hemodialysis therapy (3). DN is characterized by a series of changes, including renal hypertrophy, oxidative stress, basement membrane thickening, and extracellular matrix accumulation, which eventually results in the loss of renal function (29). Although hyperglycemia is clearly a prerequisite for the development of DN, hyperglycemia itself is insufficient for the progression of the diabetic complication. A variety of derangements associated with diabetes contribute to the development of DN. Among them, oxidative stress has been suggested to play an important role in the pathogenesis of DN (30). Enhanced oxidative stress is widely recognized as a key pathogenic factor and contributed to pathological development and progression of DN (31 – 34). Markers of oxidative stress such as ROS and reduced levels of antioxidants have been found in the blood and renal tissues in both human and experimental diabetes (35, 36).

Tempol is a stable and cell membrane–permeable SOD mimic and has been successfully used to investigate the role of ROS in renal function in intact animals (15, 37). In the present study, we used an STZ-induced diabetes rat model to demonstrate the therapeutic efficacy of tempol in early DN. Hyperglycemia, albuminuria, renal hypertrophy, and expansion of the mesangial area, which are the hallmarks of DN, were confirmed in STZ-induced diabetic rats. Tsuchida et al. (38) reported that high degree of oxidative stress in the renal tissue coincides with biochemical signs typical of DN, such as high level of UAE, which is the classical glomerular injury marker of DN. In our study, the 24-h UAE was markedly higher in diabetic rats than in control rats. Furthermore, the KW/BW ratio was significantly increased in diabetic rats in comparison with that in the control rats. These results indicate that the renal disorder in STZ-induced diabetic rats, which is early stage DN, is similar to that seen in humans (39).

Antioxidant enzyme expression in DN patients is decreased. SOD and GSH-Px constitute the principal components of antioxidants and their deficiencies can cause oxidative stress. The superoxide dismutase enzyme system is a primary determinant of superoxide removal (40). The content of MDA is a good index of intensified oxidative stress in the tissues, showing enhanced peroxidation processes (41). It has been reported that an imbalance between the production of ROS and antioxidants is believed to be involved in diabetes-induced renal failure (42). In this study, the production of MDA was enhanced in diabetic rats, indicating an oxidative stress state in experimental diabetic rats. We have demonstrated a significant diminution of SOD and GSH-Px activities in the kidney of rats after 6 weeks of STZ-induced diabetes. The findings are consistent with those previously reported in this model of diabetes (43). After tempol treatment, a significant decrease in MDA level and increase in activities of SOD and GSH-Px were observed in this study. These results suggest that tempol has a renal protective role against oxidative damage in diabetic nephropathy. Reduction of oxidative stress markers was associated with reduction in renal damage parameters. Our study demonstrated that BUN and Cr levels in serum were higher in DN rats than those in control rats. The creatinine clearance rate in diabetic rats was also obviously decreased compared with non-diabetic rats. The results of our experiment showed that continuous administration of tempol could retrieve renal dysfunction and ameliorate hyperglycemia induced by STZ. Tempol significantly affected the elevated FBG and GSP concentrations of diabetic rats. Accordingly, tempol could correct insulin resistance and the associated renal damage in STZ-induced diabetes. Meanwhile, it was also found that tempol markedly suppressed glomerular hypertrophy, the expansion of mesangial matrix and the accumulation of extracellular matrix (ECM).

The globally increasing number of patients with end-stage renal disease urges the identification of molecular pathways involved in renal pathophysiology, to serve as targets for intervention. TGF-\(\beta_1\) is the key regulator of extracellular matrix remodeling in the mesangium leading to mesangial expansion (44), and induction of TGF-\(\beta_1\) is a well-documented molecular event during the development of diabetic nephropathy both in vitro and in vivo (45 – 48). Moreover, collagen accumulation is the hallmark of glomerular sclerosis, and increase of Col IV plays a central role in this process. Excessive deposition of Col IV is an established feature associated with diabetic glomerulopathy (49). Studies have demonstrated that increase in TGF-\(\beta_1\) expression is caused by several stimuli, including hyperglycemia and oxidative stress (50). In particular, several investigators stated that ROS participates in TGF-\(\beta_1\) expression and that TGF-\(\beta_1\) promotes renal fibrous change that leads to renal damage (51, 52). Up-regulation of TGF-\(\beta_1\) by ROS will lead to excessive production of ECM, resulting in glomerular fibrosis, and ultimately loss of renal function (53). Interestingly, TGF-\(\beta_1\) can also up-regulate Col IV in all glomerular cell types (48). As we expected, the mRNA expression of Col IV and TGF-\(\beta_1\) was dramatically increased in diabetic rats in comparison with the control rats. The results of RT-PCR showed that significant up-regulation of Col IV and TGF-\(\beta_1\) expression was concomitant with deterioration of renal function and contrib-
uted to the excessive deposition of glomerular extracellular matrix in DN. The accumulation of TGF-β₁ and Col IV in glomeruli in diabetic rats was also inhibited by tempol at the 6th week after starting treatment. As reported, tempol prevented renal injury in the Dahl salt-sensitive hypertensive rat via reduction of oxidative stress and TGF-β₁ (54). High glucose induces ROS and up-regulates TGF-β₁ and Col IV expression in the glomerular mesangial cells (18, 55). Taken together, these protective effects of tempol are thought to be mediated through the down-regulation of suppression of TGF-β₁, Col IV, and reduction of excessive ROS production. It is our finding that tempol improved renal function of DN rats and this beneficial effect was associated with the inhibitory effect of tempol on kidney TGF-β₁ over-expression and Col IV over-production. Also, renal histological examination revealed that tempol significantly ameliorated diabetic-induced mesangial expansion. These findings imply potential efficacy of tempol in preventing the DN progression.

The early stage of diabetic mellitus is characterized by renal hyperfiltration, which promotes the eventual development of DN. Increasing evidence suggests that the dysfunction of glomerular MCs is involved in diabetic hyperfiltration. Impaired Ca²⁺ signaling has been inferred to be a major contributing factor to the dysfunction of diabetic MCs (7, 20). TRPC6 has been known to function as a receptor-operated Ca²⁺ channel in a variety of cell types (56). For instance, TRPC6 was detected and characterized as a component of the slit diaphragm multiprotein complex of glomerular podocytes, suggesting that it functions as a critical regulator of normal renal function (57). We have revealed that knockdown of endogenous TRPC6 in MCs reduced the Ang II-induced channel activity, implying that TRPC6 protein might be the channel or a critical component of the channel complexes mediating the Ang II-stimulated membrane response (17). Thus it is not surprising that TRPC6 participates in the contractile function of MCs and deficiency of this channel protein leads to impairment of agonist-induced mesangial contraction that subsequently results in supernormal glomerular filtration rate. In our study, the mRNA expression level of TRPC6 in the kidney was evaluated by RT-PCR and compared between nondiabetic rats and diabetic rats with and without tempol treatment. TRPC6 mRNA was strongly expressed in the healthy control rats. Comparing to the model group, there was a great improvement of expression of TRPC6 in the tempol group. In our previous article (15), we have found that expression of TRPC6 protein in glomeruli isolated from STZ-diabetic rats was dramatically decreased compared to the glomeruli from non-diabetic control rats. Also, tempol treatment significantly increased expression of TRPC6 protein, suggesting that antioxidant treatment efficiently suppressed the effect of diabetes on TRPC6. These results indicate that diabetes decreased the TRPC6 protein expression by repression of TRPC6 gene by ROS, suggesting a potential implication of TRPC6 in the development of DN.

In summary, the results from this study showed that chronic treatment of diabetic rats with tempol ameliorated the pathological changes in glomeruli, possibly by reducing expression levels of TGF-β₁, Col IV, and increasing abundance of TRPC6. Our findings provide a rationale for treating DN with antioxidants.

Acknowledgments

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