TMEM45A is involved in renal fibrosis in rats by regulating Jagged1/Notch pathway

Jian-zí Lan¹, Qi Pan¹, Qing Tao¹, Xin Shi¹, Wei-zhun Wang¹, Bo-wen Xu¹, Jie-wei Luo²,³

¹Department of Traditional Chinese Medicine, Shanghai East Hospital, Tongji University School of Medicine, NO.150, Jimo Road, Pudong District, Shanghai 200120, China, ²Provincial Clinical Medical College, Fujian Medical University, NO. 88, Jiaotong Road, Fuzhou 350001, China, ³Department of Traditional Chinese Medicine, Fujian Provincial Hospital, NO. 134, Dongjie Road, Fuzhou 350001, China

TABLE OF CONTENTS

1. Abstract
2. Introduction
3. Materials and methods
   3.1. Chemicals and reagents
   3.2. Cell culture
   3.3. Induction of renal fibrosis
   3.4. Experimental groups
      3.4.1. Effects of different concentrations of TGF-beta 1 on mRNA and protein expression of TMEM45A and Notch1 in NRK-49F cells
      3.4.2. Effects of TMEM45A over-expression on NRK-49F cells
      3.4.3. Effects of TMEM45A siRNA on TGF-beta 1 treated NRK-49F cells
   3.5. H&E and Masson trichrome staining
   3.6. Serum creatinine, total urinary protein and BUN
   3.7. Determination of hydroxyproline, Col-I, Col-III, TNF-alpha and IL-6
   3.8. Real-time PCR analyses
   3.9. Western blotting
   3.10. Statistical analysis
4. Results
   4.1. Effects of 5/6 Nx on rats
   4.2. Increased TMEM45A and Notch1 expression by TGF-beta 1 in NRK-49F cells
   4.3. Effects of TMEM45A OE on NRK-49F cells
   4.4. Effects of TMEM45A siRNA on TGF-beta 1 treated NRK-49F cells
5. Discussion
6. Acknowledgments
7. References

1. ABSTRACT

Fibrosis, or the excess deposition of fibrous tissue, is a critical feature of chronic kidney disease. Here, using renal fibrotic rat as a model, which was established via 5/6 nephrectomy (Nx), the role of TMEM45A transmembrane protein in renal fibrosis was investigated. The results indicated that 5/6 Nx gradually led to histopathological abnormalities and loss of kidney function in rats, which correlated with upregulation of TMEM45A and Notch1. Interestingly, in NRK-49F renal cells, overexpression of TMEM45A resulted in up-regulation of extracellular matrix
TMEM45A is involved in renal fibrosis

(ERM) components as well as induction of Notch-1 and Jagged-1. These effects were weakened by DAPT, an inhibitor of the Notch pathway, suggesting an important role of Notch signaling in mediating the functions of TMEM45A in NRK-49F cells. Moreover, TMEM45A knockdown by TMEM45A siRNA in NRK-49F cells diminished TGF-β1-induced upregulation of ECM components, inflammatory cytokines, Notch-1 and Jagged-1. Correspondingly, TGF-beta 1 exhibited profibrogenic like effect in NRK-49F cells and induced TMEM45A and Jagged1/Notch expression. Collectively, these results demonstrate that TMEM45A plays an important role in renal fibrosis by regulating ECM components and Jagged1/Notch pathway.

2. INTRODUCTION

Chronic kidney disease (CKD) is a life-threatening condition characterized by irreversible and progressive loss of kidney functions due to loss of renal units and replacement by extracellular matrix (ECM), which ultimately leading to fibrogenesis and renal scarring (1, 2). Transforming growth factor-beta 1 (TGF-beta 1) is a crucial factor that drives fibrosis in various forms of human CKDs (3-5). Specifically, TGF-beta 1 has been shown profibrogenic effect in CKDs by activating downstream Smad signaling in both experimental animal models and human kidney diseases (6, 7). Additionally, TGF-beta 1 displays profibrogenic-like effects in vitro by promoting ECM synthesis and inhibiting ECM degradation, as shown in mTECs and NRK-49F cells (8, 9).

Notch is a highly conserved key regulator of cell differentiation, specification, and tissue patterning. In mammals, there are 5 canonical ligands (Jagged 1, Jagged 2, Delta-like ligand 1, 3, and 4) and 4 Notch receptors (Notch1-4) in mammals. All the ligands and receptors are type I single-pass transmembrane receptor proteins (10, 11). Notch signaling is diminished in normal healthy adult kidney but induced in chronic and acute kidney injuries (12). Moreover, increased glomerular epithelial Notch signaling has been closely associated with glomerulosclerosis, albuminuria and other chronic kidney disease, and the severity of fibrosis is closely correlated with Notch1 activation in the tubulointerstitium (13-15).

TMEM45A (also known as DNAPTP4, FLJ10134 or DERP7) is a multi-pass membrane protein that belongs to a large family encoding predicted transmembrane (TMEM) proteins. Previously, it was shown that TMEM45A was highly associated with epidermal keratinization (16). In addition, TMEM45A knockdown by siRNA resulted in reduced expression of matrix metalloproteinases such as MMP2 and MMP9 (17), which were associated with ECM degradation (18, 19). These findings suggest that TMEM45A is a potential regulator of ECM. However, very little is known regarding the regulatory effects of TMEM45A in this field, and the precise molecular mechanism remains unclear. In this study, surgically-induced renal fibrotic rat was generated via 5/6 nephrectomy (5/6 Nx) and used as a model to examine the expression and role of TMEM45A in renal fibrosis (20). Moreover, the regulatory functions of TMEM45A were investigated in NRK-49F cells in vitro.

3. MATERIAL AND METHODS

3.1. Chemicals and reagents

Hematoxylin-eosin (H&E) and Masson-trichrome staining kits were obtained from Beyotime Biotech (Shanghai, China). Hydroxyproline, total urinary protein, serum creatinine and blood urea nitrogen (BUN) assay kits were purchased from Jiancheng Bioengineering Institute (Nanjing, China). Trizol reagent was obtained from Invitrogen Co. (Carlsbad, CA, USA). Primary antibodies for TMEM45A, fibronectin (FN) and connective tissue growth factor (CTGF) were obtained from Abcam Biotech (Cambridge, MA, USA). Primary antibodies for Jagged-1and Notch1 were obtained from Santa Cruz Biotech (CA, USA). Primary antibody of GAPDH was obtained from CST Inc. (Cell Signaling Technology, Beverly, MA, USA). ELISA kits to measure tumour necrosis factor-alpha (TNF-alpha), IL-6, Collagen-I (COL-I) and Collagen-III (COL-III) concentrations were purchased from CST. Recombinant human TGF-beta 1 was purchased from PeproTech (Rocky Hill,
TMEM45A is involved in renal fibrosis

Table 1. Primer sequences

<table>
<thead>
<tr>
<th>Description</th>
<th>Forward primer 5’-3’</th>
<th>Reverse primer 5’-3’</th>
</tr>
</thead>
<tbody>
<tr>
<td>siRNA (376-394)</td>
<td>GGACTTTCTTCATGAT</td>
<td>GATGAAGAAAGTCCTTTTT</td>
</tr>
<tr>
<td></td>
<td>CCGGGGACTTTTCCATGATGAGTCTCGAGATCAT</td>
<td>ATCATGGAAGAAAGTCC</td>
</tr>
<tr>
<td>TMEM45A OE CDS (345-1166)</td>
<td>GCGAATTCATGGGAGTTCTATGAGTCTCAT</td>
<td>CGGGATCTACTTTTCTTCTGATTCTTGT</td>
</tr>
<tr>
<td>Tmem45A</td>
<td>GTGCGTCTGATGACCCAATCC</td>
<td>ACATTCCACCTCCGGCTAC</td>
</tr>
<tr>
<td>Notch1</td>
<td>TGCCCAATCCACAGAGACAC</td>
<td>CATCCACAGCATTGACAC</td>
</tr>
<tr>
<td>GAPDH</td>
<td>GGAGTCTACTGCGGTCTTACC</td>
<td>ATGAGCCCTTCACGATGC</td>
</tr>
</tbody>
</table>

NJ, USA). DAPT (Notch pathway inhibitor (21)) was purchased from Sigma-Aldrich (St. Louis, MO, USA). Fetal bovine serum was obtained from Gibco Company (Cleveland, USA). Dulbecco’s modified Eagle’s medium (DMEM) was purchased from Hyclone (Logan, Utah, USA).

3.2. Cell culture

NRK-49F cells were obtained from the SIBCB (Shanghai Institute of Biochemistry and Cell Biology, Chinese Academy of Sciences, Shanghai, China). NRK-49F cells were incubated in DMEM with 10% FBS and penicillin (100 U/ml) (37 °C with 5% CO₂) to exponential phase. Cells were then seeded into 6 well culture-plates (5×10⁵ per well) and used for later experiments when the cell density reached over 50%.

3.3. Induction of renal fibrosis

Six-week-old male Wistar rats (170 ± 20 g) were provided by the Shanghai Animal Lab Center (Shanghai, China). All animals were kept in plastic cages with free access to standard commercial rodent diet and water, in a room with controlled temperature (24 – 26 °C) and on a 12-h light/12-h dark cycle.

We used the experimental model of 5/6 nephrectomy or the remnant kidney model for generation of fibrosis in the kidney. The reduction of renal mass was achieved by surgical excision of both poles, with removal of the contralateral kidney (20). After one week of the adaptive feeding, a total of 24 rats were randomly divided into 2 groups (n = 12): control group and 5/6 (model) group. For the first step, rats were anaesthetized and two poles of the left kidney were cut out. One week later, the right kidney was removed. In addition, sham operations were carried out in the control group.

Rats were sacrificed at 4 and 8 week after operation (6 rats each group at each time point) and samples were collected. Serum creatinine, total urinary protein and BUN concentration were detected to evaluating the renal function. Relative mRNA expression and protein levels of TMEM45A and Notch1 were also examined.

3.4. Experimental groups

3.4.1. Effects of different concentrations of TGF-beta 1 on mRNA and protein expression of TMEM45A and Notch1 in NRK-49F cells

NRK-49F cells were divided into 6 groups: Control group (TGF-beta 1 at 0 ng/mL) and TGF-beta 1 treatment groups (1, 2, 5, 10 and 20 ng/mL). In the cultured NRK-49F cells, TGF-beta 1 was dissolved in PBS and respectively added into cultured NRK-49F cells. Cells were harvested after 24 h incubation. Relative mRNA expression and proteins levels of TMEM45A and Notch1 were measured via qRT-PCR or western blotting respectively.

3.4.2. Effects of TMEM45A over-expression on NRK-49F cells

TMEM45A OE/siRNA lentoviral vectors were constructed by JRDun Biotech (Shanghai, China) (shown in Table 1). Viral supernatants were...
diluted in culture medium to the required concentration and added to exponential phase monolayer cells.

NRK-49F cells were divided into 4 groups: EPC group, TMEM45A OE group, EPC + DAPT group and TMEM45A OE + DAPT group. Viral supernatants and DAPT (20 μM) were added into cells correspondingly and cells were cultured for 48 h in DMEM with 10% FBS at 37 °C with 5% CO2. Hydroxyproline concentration and CTGF, FN, Notch-I and Jagged-I protein levels were measured at the end of experiments.

3.4.3. Effects of TMEM45A siRNA on TGF-beta 1 treated NRK-49F cells

NRK-49F cells were divided into 4 groups: normal NRK-49F cells (NC), TGF-beta 1 (5 ng/mL) treatment group, TGF-beta 1 (5 ng/mL) + empty plasmid control (EPC) group and TGF-beta 1 (5 ng/mL) + TMEM45A siRNA group. Cells in the last 3 groups were treated with TGF-beta 1 (5 ng/mL) and cells in the NC group were substituted with equal amount of DMSO. After incubation for 24 h, viral supernatants were added and cells were cultured for 48 h (DMEM with 10% FBS at 37 °C with 5% CO2). The concentrations of hydroxyproline, Col-I, Col-III, TNF-alpha and IL-6 as well as was protein levels of CTGF, FN, Notch-I and Jagged-I were determined.

3.5. H&E and Masson trichrome staining

To evaluate histological changes and collagen-like matrix deposition, renal tissue sections (4 ± 1 μm thickness) were stained with H&E staining kit and Masson trichrome staining kit according to manufacturers’ instructions. Briefly, renal tissues were fixed in 10% neutral buffered formalin followed by the standard methods of dehydration, clearing in xylene, and paraffin embedding. After de-paraffinizing, renal head tissue sections were stained by H&E and Masson trichrome. Finally, histopathological changes were examined by optical microscope (Olympus, Japan).

3.6. Serum creatinine, total urinary protein and BUN

Relative contents of serum creatinine, total urinary protein and BUN were determined using colorimetric method according to the manufacturer’s protocol.

3.7. Determination of hydroxyproline, Col-I, Col-III, TNF-alpha and IL-6

Cell samples were centrifuged at 12,000 × g for 5 min at 4 °C to obtain the cells. The total protein of each sample was determined by BCA method. Equal volumes of hydrolytic reagent was added and boiled for 20 min. After cooling to room temperature, samples were adjusted to pH 6.6 - 6.8 with phosphoric acid (or NaOH) and treated with activated carbons to decolorize for 30 min. Samples were then centrifuged at 12,000 × g for 15 min. The absorbance of supernatant fraction (1 mL for each sample) was determined at 550 nm using an automatic biochemistry analyzer (HITACHI 7170S, Japan). Finally, using standard curve, the concentration of hydroxyproline was calculated. IL-6, TNF-alpha, Col-I and Col-III contentsin NRK-49F cells were determined using ELISA assay kits according to the manufacturer’s protocol.

3.8. Real-time PCR analyses

Relative transcript levels of genes were investigated by quantitative real time (RT)-PCR. Total RNA was isolated using Trizol reagents, and 2 μg of total RNA was reverse transcribed into cDNA by SuperScript II kit (Life technologies, Carlsbad, California, USA). PCR was performed on qRT PCR machine (ABI-7500, USA) using SYBR Green reagents (Thermo, USA).

Primers used in Real-time Fluorogenic PCR Assays were designed by Primer 5.0 and synthesized by JRDun Biotech (Shanghai, China) (Table 1). Relative mRNA expression was normalized to internal control GAPDH by the delta-delta Ct method.

3.9. Western blotting

Protein lysates (35 μg) from the renal tissues and cultured cells was extracted. After the standent operation of loading to SDS-PAGE and transferred onto nitrocellulose membranes, which were subsequently blocked in 5% non-fat milk for 1 h at room temperature.
followed by incubation with primary antibodies overnight at 4 °C. After washing, membranes were incubation with HRP-conjugated secondary antibodies. Relative protein ratios were determined and normalized to GAPDH.

3.10. Statistical analysis

Data obtained from this study were expressed as mean ± SD. Statistical data analyses were carried out by Student's t-test and ANOVA on SPSS 20.0 software (SPSS Inc., USA).

4. RESULTS

4.1. Effects of 5/6 Nx on rats

According to a previous report, glomerulosclerosis occurred at 8 weeks after operation and the untreated control group showed progression with more severe sclerosis 12 weeks after 5/6 Nx (22). In the present study, histopathological changes of rat renal tissues were examined at 8 weeks after operation by H&E staining and Masson's trichrome staining. As shown in Figure 1a, renal cells were arranged normally and there were no signs of histopathological abnormalities in normal rats, whereas significant fibrosis and degeneration were detected in 5/6 Nx rats (n=6, 100%). Moreover, creatinine and BUN contents in serum, as well as total urinary protein content, were significantly higher in the 5/6 Nx rats than in control rats at both 4 and 8 weeks after operation. Furthermore, these contents were increased in a time-dependent manner (Figure 1b). Relative mRNA expression and protein levels of TMEM45A and Notch1 in 5/6 Nx rats were also up-regulated in a time-dependent manner (Figure 1c).

4.2. Increased TMEM45A and Notch1 expression by TGF-beta 1 in NRK-49F cells

Effects of different concentrations of TGF-beta 1 on mRNA and protein expression of TMEM45A and Notch1 in NRK-49F cells were determined at the end of experiment. The results indicate that both mRNA expression and protein levels of TMEM45A and Notch1 in NRK-49F cells were dose-dependently increased by TGF-beta 1 (Figure 1a-b). The optimum concentration for treatment with TGF-beta 1 was 5 ng/mL, which was used in subsequent experiments. If the treatment concentration was higher than 5 ng/mL, an excessive dose (4 times the optimum concentration) of exogenous TGF-β1 might lead to cytokine abnormalities and trigger a feedback system to suppress the expression of TMEM45A and Notch1.

4.3. Effects of TMEM45A OE on NRK-49F cells

The effect of TMEM45A overexpression on hydroxyproline concentration in NRK-49F cells was measured by UV spectrophotometry. The protein levels of CTGF, FN, Notch-1 and Jagged-1 were examined by Western blot. The results showed that hydroxyproline concentration in NRK-49F was increased by TMEM45A OE and this effect was weakened by DAPT (Figure 3a). Moreover, the protein expression of CTGF, FN, Notch-1 and Jagged-1 displayed similar trend as that of hydroxyproline (Figure 3b).

4.4. Effects of TMEM45A siRNA on TGF-beta 1 treated NRK-49F cells

The effect of TMEM45A siRNA on hydroxyproline, Col-I, Col-III, TNF-alpha and IL-6 contents as well as was of CTGF, FN, Notch-1 and Jagged-1 protein levels were determined at the end of experiment. As shown in Figure 4, the concentrations of hydroxyproline, Col-I, Col-III, TNF-alpha and IL-6 in NRK-49F cells were elevated by TGF-beta 1 treatment, and these effects were weakened upon TMEM45A knockdown by siRNA (Figure 4a). Relative protein levels showed the same trend (Figure 4b).

5. DISCUSSION

Patients with CKD show a steady decline in nephric function over time, and fibrosis is the primary cause of the pathophysiology (23). Fibrosis is involved in the accumulation of excess ECM which usually leads to loss of function when normal tissues are replaced by scarring (24). In this study, histopathological abnormalities and gradually loss of kidney function were observed in 5/6 Nx rats, which correlated with increased serum creatinine, BUN and
TMEM45A is involved in renal fibrosis

Total urinary protein. Moreover, relative mRNA and protein levels of TMEM45A and Notch1 were up-regulated in 5/6 Nx rats. The increase in TMEM45A and Notch1 expression was associated with renal dysfunction and fibrogenesis in 5/6 Nx rats at 8 weeks after operation (Figure 1).

TGF-beta 1 is a well-known pro-fibrogenic cytokine that can induce renal fibrosis via activation of both canonical (Smad-based) and non-canonical (non-Smad-based) signalling pathways (25), both of which lead to activation of myofibroblasts, overproduction of ECM components such as connective tissue growth factor, fibronectin, collagen and alpha-smooth muscle actin, and inhibition of ECM degradation (23, 26). Previous reports showed that TGF-beta 1 signaling played a role in regulating mouse hepatic stellate cell differentiation into myofibroblast via the Jagged1/Notch pathway (27), and that TGF-beta -

Figure 1. Effects of 5/6 Nx on rats. (a): H&E staining and Masson trichrome staining in control rats and 5/6 Nx rats at 8 weeks after operation. Scale bars: 5 μm. (b): Effects of 5/6 Nx on serum creatinine, BUN and total urinary protein contents in rats at 4 and 8 weeks after operation. (c): Effects of 5/6 Nx on mRNA expressions and protein levels of TMEM45A and Notc1h in renal tissues at 4 and 8 weeks after operation. Statistical data analyses were carried out by Student's t-test, **p < 0.01, compared with control group (n=6).
induced epithelial mesenchymal transition (EMT) was blocked by RNA silencing of Jagged1 and chemical inactivation of Notch (28). These results suggest that Jagged1/Notch signaling is positively
TMEM45A is involved in renal fibrosis

regulated by TGF-beta 1, a finding that was confirmed in the present study (Figure 2 and Figure 4).

TMEM45A is a multi-pass membrane protein whose elevated expression has been previously associated with epidermal keratinization (29). Moreover, knockdown of TMEM45A resulted in downregulation of MMP2 and MMP9 (30), which were involved in collagen degradation (18, 19). In the present study, TMEM45A was found to be upregulated in Nx rats. In addition, TGF-beta 1 induced

Figure 3. Effects of TMEM45A OE on NRK-49F cells and the involvement of Notch-l/Jagged-l signaling. Effects of TMEM45A OE on hydroxyproline content (a) and the protein levels of CTGF, FN, Notch-l and Jagged-l (b) in NRK-49F cells. (A): EPC group, (B): TMEM45A OE group, (C): EPC + DAPT group and (D): TMEM45A OE + DAPT group. Statistical data analyses were carried out by ANOVA, *p < 0.05, **p < 0.01, compared with EPC group (n=6). Statistical data analyses were carried out by Student’s t-test, $$$p < 0.01$ compared with TMEM45A OE group (n=6). $^\#p < 0.01$, compared with EPC + DAPT group (n=3).
TMEM45A is involved in renal fibrosis

TMEM45A expression in NRK-49F cells in vitro and over-expression of TMEM45A increased hydroxyproline concentration, as well as CTGF, FN, Notch-1 and Jagged-1 protein levels in NRK-49F cells. The effects of TMEM45A overexpression on ECM components were diminished when Jagged1/Notch signaling was blocked, indicating that Jagged/Notch act downstream of TMEM45A to mediate TMEM45A functions. Moreover, TGF-beta 1-induced expression of ECM components (hydroxyproline, collagen, CTGF and FN) and inflammatory cytokines (TNF-alpha and IL-6) as well as increased protein levels of Notch-1 and Jagged-1 in NRK-49F cells. The observed effects of TGF-beta were greatly diminished by TMEM45A knockdown, suggesting a regulatory path from TGF-beta 1 to TMEM45A to Jagged/Notch1.

Taken together, the results demonstrate that TGF-beta 1 exhibits pro-fibrogenic like effects in
NRK-49F cells by upregulating Jagged1/Notch signaling and may show similar effects in 5/6 Nx rat model (Figure 1). Moreover, TGF-beta 1 induced the expression of TMEM45A, which positively regulated Jagged/Notch1 signaling. Thus, TMEM45A may be a potential biomarker in renal fibrosis. Reducing TMEM45A expression may attenuate fibrogenesis in patients with CKDs. It will be important to verify in future studies whether this regulation is observed in other renal fibrosis models as well as in clinical trials.

6. ACKNOWLEDGMENTS

This work was supported by the science and technology project of Fujian Province (No. 2018Y0012), and the construction project of Zhang Xuemei’s Academic Inheritance Studio of Famous and Aged Chinese Medicine Experts in Fujian Province.

7. REFERENCES


TMEM45A is involved in renal fibrosis

PMid:29230704


TMEM45A is involved in renal fibrosis


Abbreviations: TMEM45A: transmembrane protein 45A; FN: fibronectin; CTGF: connective tissue growth factor; ECM: extracellular matrix; transforming growth factor-beta 1: TGF-beta 1; CKD: chronic kidney disease; MMPs: matrix metalloproteinases; 5/6 Nx: 5/6 nephrectomy; COL-I: Collagen-I; COL-III: Collagen-III; BUN: blood urea nitrogen.

Key Words: 5/6 nephrectomy, TGF-beta 1, TMEM45A, Jagged1/Notch signal, NRK-49F cells

Send correspondence to: Jie-wei Luo, department of Traditional Chinese Medicine,
TMEM45A is involved in renal fibrosis

Fujian Provincial Hospital. NO. 134. Dongjie Road, Fuzhou, 350001, China, Tel: 86-0591-88217471, Fax: 86-0591-88217471, E-mail:dffjsl112@126.com