YAP is overexpressed in clear cell renal cell carcinoma and its knockdown reduces cell proliferation and induces cell cycle arrest and apoptosis

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Abstract. Yes-associated protein (YAP) has been reported to be an oncogene in a number of malignancies. It constitutes an important regulatory mechanism for the Hippo pathway, a key regulator of cell growth and apoptosis. The present study aimed to investigate the clinical significance and the role of YAP in the development of clear cell renal cell carcinoma (ccRCC). YAP expression levels were compared between ccRCC and adjacent normal renal tissues by RT-PCR and immunohistochemistry, respectively. YAP expression levels were then detected in ccRCC cell lines 786-0 and ACHN, as well as in human embryonic kidney 293 cells (HEK-293) using western blotting. Three specific YAP-shRNA lentiviral vectors were constructed and transfected into 786-0 cells, and then the mRNA and protein levels of YAP and downstream transcription factor TEAD1 were detected. Finally, the effects of YAP silencing on proliferation and the cell cycle distribution of 786-0 cells were detected by Cell Counting Kit-8 (CCK-8) and flow cytometry (FCM), respectively. The apoptosis rate was also analyzed by FCM. It was observed that the expression levels of YAP mRNA and protein in ccRCC tissues were higher than these levels in the adjacent normal renal tissues. The expression of YAP protein in ccRCC tissues was significantly correlated with clinical stage and differentiation. The YAP protein levels in the two ccRCC cell lines 786-0 and ACHN were significantly higher than that in the HEK-293 cells. Additionally, treatment of 786-0 cells with YAP-shRNA lentiviral vectors significantly reduced the expression levels of YAP and TEAD1 mRNA and protein. Further analyses in 786-0 cells in which YAP was decreased, revealed that cell proliferation was inhibited, cell cycle was arrested at the G1 phase and apoptosis was increased. These results indicate that YAP is an underlying oncogene in ccRCC and it may be a promising biomarker and therapeutic target of ccRCC.

Introduction

As one of the most common malignant tumors in the urinary system, renal cancer accounts for more than 270,000 new cases worldwide each year (1). Clear cell renal cell carcinoma (ccRCC) is the most common histological subtype, accounting for 75-85% of all RCC cases. The incidence of RCC has steadily increased in many countries (2). However, the treatment options for RCC are limited, and the mechanisms of the pathogenesis of renal carcinoma are not completely defined. Therefore, new treatment strategies for patients with RCC are urgently needed. With continued research on the mechanisms of tumors, targeted therapy has made remarkable progress (3). In recent years, novel markers and therapeutic targets of RCC have continued to emerge. Nevertheless, to date, no biomarkers have been used in RCC in the routine clinic (4).

The Hippo signaling pathway, initially discovered in Drosophila, is a potent regulator of cell growth and apoptosis (5). A series of biochemical and genetic studies identified this pathway whose components are highly conserved between Drosophila and mammals (6,7). In mammals, the components of the Hippo pathway include MST1/2, Sav1, LATS1/2, YAP, TAZ and TEAD (8,9). MST1/2, Sav1 and LATS1/2 are regarded as upstream kinases of the mammalian Hippo pathway. It is presumed that through phosphorylation of YAP, the upstream kinases LATS1/2 inhibit YAP transcription activity (10). YAP, TAZ and TEAD were identified as downstream transcription factors of this pathway (11).

Yes-associated protein (YAP), the mammalian homolog of Drosophila Yorkie (Yki), is a 65-kDa protein which was originally identified due to its interaction with the Src family tyrosine kinase c-Yes (12). Recent studies have found that the transcriptional co-activator YAP is a key nuclear effector of the Hippo pathway which regulates organ size by governing cell proliferation and apoptosis (13,14). The TEAD family,
located in the downstream of the Hippo pathway, are major mediators of YAP transcriptional activity (15,16). The role of YAP which interacts with the TEAD family of transcription factors is essential for YAP-mediated tumor growth and metastasis (17). Recently, YAP has been shown to be a candidate oncogene in recurrent amplification at human chromosome 11q22 (18). Overexpression of YAP protein and its nuclear localization were observed in colon, lung, pancreatic, hepatocellular, ovarian and prostate carcinomas (16,19,20).

Despite of the growing evidence of YAP as a crucial regulator of human types of cancers, its involvement in ccRCC is still obscure. In the present study, we attempted to identify the role of YAP in ccRCC. We evaluated the expression levels and clinical significance of YAP in ccRCC tissues. In addition, we investigated the effects of YAP silencing via YAP-shRNA lentiviral vectors transfected into ccRCC cell lines. These findings suggest that YAP may serve as a potential target in the treatment of ccRCC.

Materials and methods

Tissue samples and cell culture. Tissue samples from radical nephrectomy were obtained from the First Affiliated Hospital of Chongqing Medical University from March 2010 to November 2011. The tissue samples consisted of 30 cases of ccRCC tissues and 30 cases of adjacent normal renal tissues (4 cm or more from the tumor). Samples were confirmed by a pathologist and stored at -80˚C. The present study was conducted with approval from the Ethics Committee of the First Affiliated Hospital of Chongqing Medical University. Informed consent was obtained from all patients. Human ccRCC cell lines 786-0 and ACHN and the human HEK-293 cell line were purchased from ATCC. ACHN and 786-0 and ACHN and the human HEK-293 cell line were purchased from ATCC. ACHN and the human HEK-293 cell line were purchased from ATCC. ACHN and the human HEK-293 cell line were purchased from ATCC. ACHN and the human HEK-293 cell line were purchased from ATCC. ACHN and the human HEK-293 cell line were purchased from ATCC. ACHN and the human HEK-293 cell line were purchased from ATCC. ACHN and the human HEK-293 cell line were purchased from ATCC. ACHN and the human HEK-293 cell line were purchased from ATCC.

RT-PCR. The tissues and the cultured cells were collected. Total RNA were extracted using the TRIzol kit (Takara, Osaka, Japan). The total RNA quality was detected by a UV spectrophotometer (Bio-Rad, Hercules, CA, USA). The cDNA was synthesized reversely for the cDNA and PCR reaction amplification of YAP and TEAD1 genes, and GAPDH as an internal reference. Specific primers were designed: YAP-F, 5'-GCTCATTCCTCTCCAGCTT-3', and YAP-R, 5'-TCCTAACAGTGCCACCTAT-3'. Negative and GAPDH were designed as follows: 5'-GCTCATTCCTCTCCAGCTT-3', and GAPDH-F, 5'-CCGTTTCCCAGACTACCTT-3'. Non-silencing shRNA (5'-TTCTCCGAACGTGACGCTT-3') was synthesized as the negative control (NC shRNA). The YAP lentiviral interference vector was designed and constructed by Shanghai SBO Medical Biotechnology. Briefly, for generation of shRNA-YAP or control lentiviral vector, the lentivirus expressing plasmids pLTR-G and pNL-EFGP/CMV/WPREdU3 which were carrying shRNA, along with lentivirus packaging plasmids pCD/NL-BH-D (both from Addgene, USA) were added into 293T cells for lentiviral vector packaging. Eventually, the lentivirus was extracted and lentivirus concentrations were verified. For lentivirus transfection, 786-0 cells were grown to 30-50% confluency and transfected with three pairs of YAP-shRNAs and the non-silencing control shRNA at an optimized multiplicity of infection (MOI) of 20 with Polybrene at a concentration of 5 µg/ml. The successful transfection of YAP-shRNA lentiviral vectors was observed by fluorescence microscope 96 h after transfection. The YAP-shRNA infected cells were detected as GFP-positive cells under fluorescence microscopy at a magnification of x200 (Olympus, Tokyo, Japan). To detect the transfection efficiency, the mRNA and protein levels of YAP were respectively assessed by RT-PCR and western blotting 96 h after transfection.

Immunohistochemistry. The tissues were fixed with 10% formaldehyde (ZSGB-Bio, Beijing, China) embedded in paraffin, sliced into 4-µm sections and used for staining. In brief, the sections were rehydrated, and antigen retrieval was performed by a microwave vacuum histoprocessor (RHS-1; Bergamo, Italy) at 121˚C for 15 min. After being blocked with goat serum (Gibco-BRL), the primary antibody against YAP (Santa Cruz, Dallas, TX, USA) was diluted (1:200) and incubated with the tissue sections overnight in a humidity chamber at 4˚C. The sections were treated with horseradish peroxidase (HRP)-labeled secondary antibody (ZSGB-Bio) and DAB, and the results were observed with a microscope. The tissues were scored according to the percentage of the positively stained area and staining intensity. The percentage of the positive area was graded as 0 (≤5%), 1 (6-25%), 2 (26-50%), 3 (>51%), and the staining intensity was graded as 0-2 (i.e., negative, 0; weak, 1 and strong, 2). The two grades were multiplied and tissues were assigned to one of three levels: 0, negative; 1-4, weak positive; and 5-6, strong positive.

Western blotting. The YAP antibody was purchased from Santa Cruz Biotechnology (Santa Cruz, CA, USA) and the TEAD1 antibody was purchased from Proteintech Group, Inc. (Chicago, IL, USA). Total protein was extracted from the tissue homogenates and the cell lysate by RIPA buffer (50 mM Tris, pH 7.4, 150 mM NaCl, 1% Triton X-100, 1% Nonidet-P40, 0.5% sodium deoxycholate, 0.1% SDS, 1 mM EDTA and PMSF) (Beyotime, Shanghai, China). The lysate containing 50 µg protein quantified by the BCA protein assay kit for each sample was subjected to SDS-PAGE, and was then transferred to a 0.45-µm polyvinylidene fluoride (PVDF) membrane (both from Beyotime). The membrane was blocked in Tris-buffered saline-Tween 0.1% (TBST) with 5% skimmed milk powder for 2 h at 37˚C. The primary antibodies were diluted (YAP 1:200, TEAD1 1:200 and β-actin 1:1,000). The proper species and diluted HRP-labeled goat anti-rabbit, goat anti-mouse, and goat anti-human secondary antibodies were diluted (YAP 1:200, TEAD1 1:200 and β-actin 1:1,000) and incubated at room temperature for 1 h. The membrane was washed with TBST, incubated with HRP-conjugated secondary antibodies, and visualized with an enhanced chemiluminescence (ECL) kit (Thermo Scientific, Waltham, MA, USA) for 1 min at room temperature. The protein bands were detected by an ImagePro Plus imaging system (Media Cybernetics, Bethesda, MD, USA) and normalized to β-actin as an internal reference.
secondary antibodies were added, and Western blot results were analyzed by Bio-Rad software.

**Proliferation assay.** The 786-0 cells were seeded in 96-well plates for 12 h at a concentration of $5 \times 10^3$ cells/well. Cells were transfected with shYAP-3 or NC shRNA lentiviral vectors. After treatment for 48, 96 and 144 h, 10 µl of Cell Counting Kit-8 (CCK-8) reagent (Sigma, USA) in 90 µl RPMI-1640/well was added to the cell culture medium and then incubated at 37˚C for 1 h. The absorbance was measured with a microplate reader (Spectramax M2; MD, USA) at 450 nm.

**Cell cycle assay by flow cytometry (FCM).** The 786-0 cells were transfected with the shYAP-3 lentiviral vector which was the most efficient inhibition transfected group or NC shRNA lentiviral vectors. After treatment for 48, 96 and 144 h, 10 µl of Cell Counting Kit-8 (CCK-8) reagent (Sigma, USA) in 90 µl RPMI-1640/well was added to the cell culture medium and then incubated at 37˚C for 1 h. The absorbance was measured with a microplate reader (Spectramax M2; MD, USA) at 450 nm.

**Apoptosis assay.** The 786-0 cells were transfected with shYAP-3 or the NC shRNA lentiviral vector for 96 h, then trypsinized and washed with PBS, resuspended in 100 µl binding buffer, and stained with 7-aminoactinomycin D (7-AAD) and PE-conjugated Annexin V (Bioscience). The apoptosis rate of each group was measured with a flow cytometer (Beckman Coulter, USA).

**Statistical analysis.** Statistical analysis was performed with SPSS 19.0 software. Data are presented as the means ± standard deviation. The Chi-square trend test was used to assess the correlation between YAP protein expression and clinicopathologic factors of the renal cancer cases. A Student's two-tailed t-test was used to compare two groups. One-way ANOVA was used in comparison of means among multiple groups. p<0.05 was considered to indicate a statistically significant result.

**Results**

**Expression of YAP in ccRCC tissues and correlation with clinicopathological factors.** As determined by RT-PCR, the expression level of YAP mRNA in the ccRCC tissues was $0.569±0.066$, while that in adjacent normal renal tissues was $0.515±0.068$. The expression level of YAP mRNA in ccRCC tissues was significantly higher than that in adjacent normal renal tissues (p<0.05) (Fig. 1).

The expression of YAP protein in 30 cases of ccRCC tissues and 30 cases of adjacent normal renal tissues was detected by immunohistochemistry. In the ccRCC tissues, 63.3% (19/30) of the samples exhibited positive expression, and the positive areas were primarily located in the nucleus and cytoplasm (Fig. 2A). In contrast, the positive rate of YAP protein expression in the normal tissues was 33.3% (10/30); most cases showed weak
positive expression and the YAP protein was confined to the renal collecting duct system (Fig. 2B). The positive rate of YAP protein in the ccRCC tissues was obviously higher than that in the adjacent normal renal tissues (p<0.05).

We further analyzed the correlation of YAP protein expression with clinicopathological factors of ccRCC. As shown in Table I, there was no obvious relationship between YAP protein expression and gender, age, renal vein metastasis or tumor size. However, the positive rate of YAP protein expression was 33.3% (3/9) in well differentiated, 61.5% (8/13) in moderately differentiated and 88.9% (8/9) in low differentiated tumors (p=0.018). In addition, the positive rate of YAP protein expression was 47.1% (8/17) in stage I-II and 84.6% (11/13) in stage III-IV cases (p=0.034). These results showed that the expression level of YAP protein was closely correlated to clinical stage and differentiation in the ccRCC tissues.

Expression of YAP in 786-0 cells is suppressed by shRNA lentiviral vectors. We examined the expression level of YAP protein in ccRCC cell lines 786-0 and ACHN and in the human embryonic kidney HEK-293 cells using western blotting. As shown in Fig. 3A, the YAP protein levels in the two ccRCC cell lines 786-0 and ACHN were significantly higher than that in the HEK-293 cells (p<0.05). To determine whether YAP plays a role in tumorigenesis of renal cancer, three specific YAP-shRNA lentiviral vectors,
shYAP-1, shYAP-2 and shYAP-3, were successfully constructed and transfected into 786-0 cells. After 96 h of transfection, as shown in Fig. 3B, 786-0 cells were successfully transfected with the YAP-shRNA lentiviral vectors. Three YAP interference groups, as well as a blank control group (BC group, cells only) and negative control group (NC group) were established from the 786-0 cells. Furthermore, the efficient knockdown of YAP expression was detected by RT-PCR and Western blot analysis. RT-PCR analysis showed that the relative mRNA expression level of YAP in the YAP-shRNA transfection cells was markedly lower than levels in the BC and NC groups (p<0.05) (Fig. 3C). Similarly, the expression of YAP protein was also significantly inhibited as confirmed by western blotting (p<0.05) (Fig. 3D). The lowest level of YAP expression was observed in the cells transfected with the shYAP-3 lentiviral vectors. These results indicated that the shYAP-3 lentiviral vector had the highest inhibition rate of the three specific YAP-shRNA lentiviral vectors. Therefore, the shYAP-3 lentiviral vector was chosen for subsequent experiments.

Knockdown of YAP inhibits expression of TEAD1. RT-PCR analysis showed that the relative mRNA expression level of TEAD1 in the shYAP-3-transfected cells was markedly lower than levels in the BC and NC group cells (p<0.05) (Fig. 4A). As shown in the western blot analysis (Fig. 4B), we also observed that the protein expression level of TEAD1 was significantly decreased compared with the levels in the BC and NC groups (p<0.05), indicating that silencing of YAP inhibited the expression level of TEAD1, which is a downstream gene in the Hippo pathway.

YAP knockdown suppresses cell proliferation. To determine whether the YAP-shRNA lentiviral vector actually affects the proliferation of 786-0 cells, we examined the proliferation of the three 786-0 cell groups including the shYAP-3-transfected group, the blank group and negative control group. As shown in Fig. 5, after silencing of YAP in the 786-0 cells for 144 h, we found that there was a significant reduction in cell proliferation (p<0.05). In addition, the NC group presented a minimal effect on the proliferation of 786-0 cells compared with the blank control group (p>0.05). The results demonstrated that YAP plays an important role in cell survival, and the knockdown of the YAP gene leads to suppression of renal cancer cell proliferation.

YAP knockdown arrests the cell cycle and induces cell apoptosis. Ninety-six hours after shYAP-3 lentiviral vector transfection, the effect of YAP on the cell cycle progression was analyzed by FCM. In the shYAP-3-transfected group, we found that 61.4% of cells were in the G1 phase and 29.4% were in the S phase, compared with 51.4% of cells in the G1 phase and 39.4% in the S phase in the blank control group and 51.0% in the shYAP-3-transfected group.

Figure 5. Effect of YAP silencing on the proliferation of 786-0 cells. The absorbance values in the shYAP-3-transfected group, the blank control group and negative control group cells were measured at 48, 96 and 144 h, respectively. The absorbance values represent the mean from three independent experiments. YAP, Yes-associated protein.

Figure 6. Effect of YAP silencing on cell cycle progression and the apoptosis rate in 786-0 cells. (A) Flow cytometric analysis of the cell cycle of 786-0 cells transfected with shYAP-3 lentiviral vectors or the control group after 96 h. The cell cycle of the shYAP-3 transfection group was compared with the control group and the proportion of cells in the G1 phase was significantly increased, while the proportion of cells in the S phase was significantly reduced (p<0.05). (B) The apoptosis rate of the shYAP-3-transfected group cells was significantly higher than the rates in the BC and NC group cells (p<0.05). YAP, Yes-associated protein; BC, blank control group; NC, negative control group; shYAP-3, shYAP-3-transfected group.
of cells in G1 phase and 39.8% in S phase in the NC group (p<0.05) (Fig. 6A). No significant difference was observed in cell cycle progression between the blank control and the NC group (p>0.05). Furthermore, the apoptosis rate of cells with YAP knockdown was analyzed by FACS. The results showed that the apoptosis rate in the 786-0 cells with YAP knockdown was significantly increased as compared with the rates in the control groups (p<0.05) (Fig. 6B). The difference between the blank control group and the NC group was not obvious (p=0.582). This finding suggests that knockdown of YAP induced 786-0 cell apoptosis and G1 phase arrest.

Discussion

The Hippo pathway has been recognized as an important regulator of both organ size control and tumorigenesis (6). In mammals, the Hippo pathway negatively regulates cell growth through a kinase cascade and results in the inactivation of YAP. Hippo core components form protein kinase complexes acting in a cascade to phosphorylate YAP and induce its cytoplasmic translocation (21). YAP expression has been significantly associated with tumor metastasis, grade and stage (22). A study in lung cancer showed that YAP expression was significantly correlated with TNM stage (23).

In the present study, YAP mRNA and protein expression levels in 30 cases of ccRCC tissues and 30 cases of adjacent normal renal tissues were first detected. The results indicated that expression levels of YAP mRNA and protein in ccRCC tissues were significantly higher than those in adjacent normal renal tissues. ccRCC tissues had a high frequency of YAP-positive cells, whereas the adjacent normal tissues had a lower frequency (63.3% vs. 33.3%). Although a significant percentage of adjacent normal cases stained positive for YAP, the overall expression level was significantly higher in the tumor tissues. We then found that the expression level of YAP protein in ccRCC tissues was significantly correlated with differentiation and clinical stage. In our immunohistochemistry study, the staining intensity of YAP was tightly linked with progression from well-differentiated to poorly differentiated tumors. Furthermore, YAP protein expression was correlated with clinical stage which may serve as an independent predictor for renal cancer. In the lower stages of ccRCC, there were low levels of expression of YAP, and higher expression of YAP was accompanied by higher stages of malignancy. In previous studies, YAP was found to play an important role in tumorigenesis and to be as an independent prognostic marker in HCC and ESCC (24,25). Consistent with these research findings, the present study showed that the YAP gene is a tumor-specific gene, which is involved in the progression of ccRCC.

In previous studies, overexpression of YAP has been found in several human cancer cell types, and the oncopgenic activity of YAP has been clearly demonstrated (16,25,26). However, another study also reported that YAP may act as a tumor suppressor in certain breast carcinomas (27). The expression and activity of YAP in renal cancer cells are currently unknown. The present report showed that there was a significantly higher expression of YAP protein in ccRCC cell lines, 786-0 and ACHN, when compared with that in the HEK-293 cells. Specific shRNA lentiviral vectors targeting YAP were successfully constructed and transfected into 786-0 cells. We found that YAP mRNA and protein were effectively inhibited in the 786-0 cells. Furthermore, functional experiments were performed. The results indicated that silencing of YAP in 786-0 cells significantly inhibited cell proliferation, induced G1 phase cell cycle arrest and induced 786-0 cell apoptosis. Therefore, it is evident from the results that YAP plays an important role in the development of ccRCC.

In our initial experiment, we synthesized an siRNA targeting YAP and transfected it into 786-0 cells using Lipofectamine 2000, but it did not achieve the expected results; the effect of the suppression of the expression of the YAP gene was poor. Then, lentiviral vectors were engineered to knock down the YAP gene in 786-0 cells. All of the three YAP-shRNA interference groups were able to suppress the mRNA and protein expression levels of YAP. Further statistical analysis found that the lowest level of YAP mRNA expression in the 786-0 cells was in the shYAP-3 interference group, and the expression level of YAP protein in the shYAP-3 interference group was in accordance with the expression level of YAP mRNA. We chose lentiviral vectors for gene silencing due to their attractive properties, including high efficiency, persistent gene silencing and safety for humans (28,29). Lentiviral vectors are a powerful tool for gene transfer into a broad range of cell types (30). Lentiviral vectors producing shRNAs have widespread use in the knockdown of gene expression both in vitro and in vivo.

Acting as downstream gene of the Hippo pathway, YAP is a transcription co-activator which interacts with the PPFY-motif-containing transcription factors, including TEAD, ErbB4, P73 and Runx2 (31,32). It has been reported that YAP and TEAD1 genetically interact to promote tissue growth (33). TEAD as a new component in the Hippo pathway plays an essential role in mediating the biological functions of YAP. In humans, there are four TEAD family members. YAP and TEAD1 bind to a common set of promoters in MCF10A cells (33). Furthermore, TEAD1 has been shown to be linked to the regulation of YAP in Drosophila and mammalian epithelial cell lines (34). A recent study has provided proof of principle that inhibiting TEAD-YAP interactions is a pharmacologically viable strategy against the YAP oncoprotein (35). TEAD has been demonstrated to be important for the growth-promoting function of YAP (36). The present study showed that silencing of YAP led to the decreased expression of TEAD1 and resulted in the growth inhibition of 786-0 cells. Thus, we speculated that the relevant mechanism involved the suppression of TEAD1 expression by the knockdown of YAP which consequently affects the expression of downstream cell growth factors and thus plays functional roles in the regulation of 786-0 cell proliferation. Yet this may not be the sole mechanism, and further research on the molecular functions of the YAP gene in renal cancer is warranted.

In conclusion, these findings demonstrated that YAP is an oncogene overexpressed in ccRCC, and the expression level of YAP protein is closely correlated with clinical stage and differentiation of ccRCC tissues. Moreover, suppression of YAP expression by YAP-shRNA lentiviral vectors reduced 786-0 cell growth and arrested the cell cycle at the G1 phase and also led to induced apoptosis. Therefore, YAP could be a potential novel therapeutic target for the treatment of ccRCC.
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References


