Circular RNA DHX33 promotes malignant behavior in ccRCC by targeting miR-489-3p/MEK1 axis

Jie Wang1,2,*, Jian-Qiu Zhang1,2,*, Xiao-Lei Zhao3,*, Jing-Yu Lu4, Ze-Ming Weng1,2, Zhen-Min Ding4,*, Feng-Qiang Yang1,2,5

1Department of Urology, Ninghai First Hospital, Zhejiang 315600, China
2Department of Urology, Ninghai Hospital, Branch of Shanghai Tenth People’s Hospital, Zhejiang 315600, China
3Department of Urology, Huaihe Hospital of Henan University, Kaifeng 475000, China
4Department of Anesthesia, Shanghai East Hospital, Tongji University School of Medicine, Shanghai 200120, China
5Department of Urology, Shanghai Tenth People’s Hospital, Tongji University, Shanghai 200072, China

*Equal contribution

Correspondence to: Zhen-Min Ding, Ze-Ming Weng, Feng-Qiang Yang; email: zhenminding81@163.com, wenzem@sina.com, 05fengqiangyang@gmail.tongji.edu.cn

Keywords: clear cell renal cell carcinoma, circDHX33, miR-489-3p, MEK1

ABSTRACT

Mounting evidence indicates that circular RNAs modulate the initiation of clear cell renal cell carcinoma (ccRCC). However, their specific roles in the malignancy of ccRCC is understudied. Here, we present a novel circular RNA, circDHX33, that is up-regulated in ccRCC cell lines and tissues. Upregulated circDHX33 in ccRCC patients significantly correlates with advanced TNM stage and metastasis. Suppressing circDHX33 expression inhibits the proliferation and invasion of cultured cells, and suppresses tumor growth in vivo. Mechanistically, we show that circDHX33 promotes ccRCC progression by sponging miR-489-3p and modulating MEK1 expression. In conclusion, our findings suggest that circDHX33 plays a role in promoting ccRCC via the miR-489-3p/MEK1 axis and may serve as a novel therapeutic target for the treatment of ccRCC patients.

INTRODUCTION

Renal cell carcinoma (RCC) is a type of cancer affecting renal tubular epithelial cells, and has a high mortality rate [1]. It has many subtypes with clear cell renal cell carcinoma (ccRCC) being the most common type [2]. The mainstay treatment for ccRCC is surgery [3]. Although recent advances in its management have improved the survival rate of ccRCC, its prognosis following metastasis or recurrence remains poor [4, 5]. It is therefore necessary to improve our understanding on the molecular basis of this cancer.

Circular RNA (circRNA) is a novel class of endogenous RNA formed via covalent joining of the 3’ end to 5’ ends of RNA molecules [6, 7]. Mounting evidence suggests that circRNAs expression is dysregulated in various diseases, including cancers. Ge et al reported that low circMTO1 expression leads to low proliferation and invasion in colorectal cancer through a mechanism that involves the Wnt/beta-catenin signaling pathway [8]. Shao et al have reported that hsa_circ_0075341 promotes cervical cancer by sponging miR-149-5p [9]. Elsewhere, it was found the hsa_circ_002577/miR-197/CTNND1 axis promotes Wnt/β-catenin signaling to promote endometrial carcinoma progression [10]. However, the mechanisms through which most circRNAs influence tumor progression remain unknown.

In this study, a novel 399 bp circRNA (hsa_circ_0000740), named circDHX33 is reported. This circRNA is formed by the splicing of DHX33 mRNA. We find that circDHX33 is elevated in ccRCC patients and correlates with advanced metastasis and
clinical stage. Mechanistically, circDHX33 promotes ccRCC by sponging miR-489-3p thereby upregulating MEK1 expression, accelerating cancer growth and metastasis. Collectively, these results imply that circDHX33/miR-489-3p/MEK1 axis could be therapeutically targeted to improve ccRCC outcomes.

RESULTS
circDHX33 is highly expressed in ccRCC

To detect aberrantly expressed circRNAs in ccRCC tissues, we took advantage of a microarray expression profile data of circRNAs in ccRCC tissues (GSE137836). Analysis revealed that 211 circRNAs were differentially expressed between metastatic and primary groups (fold change >2.0) and p < 0.05 (Figure 1A, 1B). GO term analysis revealed that these differentially expressed circRNAs are involved in various physiological processes, molecular functions, and cell signaling pathways (Figure 1C). Next, we selected the top 5 upregulated circRNAs in dataset GSE137836 and used RT-qPCR to validate their upregulation in ccRCC tissues. A novel circRNA hsa_circ_0000740 (circDHX33) was found to be the most significantly upregulated circRNA in the ccRCC tissues relative to normal controls (Figure 1D).

We therefore decided to further characterize circDHX33. CircDHX33, which is located on chromosome 17, consists of 2 exons (Exons 2-3) that are spliced from DHX33 mRNA (Figure 2A). We then confirmed the circular characteristics of circDHX33. To this end, genomic DNA and cDNA from some ccRCC tissue were used as templates to amplify circDHX33

![Figure 1. Screening of ccRCC-related circRNAs in GSE137836 dataset. (A, B) Hierarchical clustering analysis and volcano plots of significantly differentially expressed circRNAs in ccRCC tissues from the GSE137836 dataset. (C) GO enrichment analysis for dysregulated circRNAs gene symbols. (D) 5 upregulated circRNAs in 8 paired ccRCC tissues was determined by using RT-qPCR. *p<0.05, **p<0.01.](image-url)
using divergent primers. We did not obtain amplification products for genomic DNA (Figure 2B). Moreover, our data revealed that circDHX33 resisted RNase R, revealing that circDHX33 circularized the RNA (Figure 2C). RT-qPCR showed that circDHX33 expression in ccRCC was significantly enhanced and correlated with advanced TNM stage and metastasis (Figure 2D–2F). These data indicate that circDHX33 has an important function in ccRCC progression.

Knockdown of circDHX33 inhibits the proliferation and invasion of ccRCC cells

To assess the biological functions of circDHX33 in ccRCC progression, mRNA levels of circDHX33 in RCC cells were estimated. Results indicated that circDHX33 level was markedly upregulated in the RCC cell lines 786O, Caki1, A498, Caki2, and ACHN, relative to HK2 cell lines (Figure 3A). We then transfected si-circDHX33 into 786O and A498 cell lines (Figure 3B). Cell proliferation analysis using CCK-8 and colony formation tests indicated that circDHX33 inhibition strongly decreased proliferation (Figure 3C–3F). Analysis of cell invasion showed that down-regulation of circDHX33 markedly inhibited invasion of 786O and A498 cells (Figure 3G–3H). These findings suggested that circDHX33 silencing represses the proliferation and metastasis of RCC cells in vitro.

circDHX33 targets miR-489-3p as a miRNA sponge

Multiple studies have demonstrated that circRNAs modulate cellular function by sponging miRNA and therefore suppressing their function [11–13]. The nuclear-cytoplasmic fractionation assay showed that the cytoplasmic fraction contained higher circDHX33 levels relative to the nuclear fraction (Figure 4A). We then used miRanda and RNAhybrid databases to predict potential target miRNAs for circDHX33 (Figure 4B and 4C). Pull-down assay revealed that miR-489-3p was strongly associated with the circDHX33 probe in both 786-O and A498 cells (Figure 4D–4E). Figure 4F showed a schematic representation of miR-489-3p binding sites in circDHX33. A luciferase assay revealed that the miR-489-3p mimics suppressed luciferase activity in circDHX33-WT group (Figure 4G). The RIP assay confirmed an association between circDHX33 and miR-489-3p in RCC cells (Figure 4H).

We next explored miR-489-3p expression in ccRCC using RT-qPCR analysis and discovered that it was downregulated in ccRCC tissues and cell lines

Figure 2. circDHX33 was highly expressed in ccRCC. (A) Schematic diagram of circDHX33. (B) Random hexamer or oligo (dT)18 primers were used in the reverse transcription experiments. (C) The relative RNA levels were determined by qRT-PCR. (D) The expression of circDHX33 in 47 paired ccRCC tissues was detected by qRT-PCR. (E, F) High circDHX33 expression was linked to advanced TNM stage and metastasis in ccRCC patients. *p<0.05, **p<0.01.
Kaplan-Meier analysis indicated that low miR-489-3p levels correlated with poor RCC survival (Figure 5C, 5D). Further analysis showed that circDHX33 expression was negatively correlated with miR-489-3p level in ccRCC tissues (Figure 5E). CircDHX33 suppression elevated miR-489-3p expression in 786O cells and A498 cells (Figure 5F). Moreover, rescue experiments indicated that miR-489-3p inhibitors abolished the effects of circDHX33 silencing on 786O cells proliferation and invasion abilities (Figure 5G, 5H). Together, these findings suggest that circDHX33 sponges miR-489-3p in ccRCC.

**MEK1 is a downstream target of miR-489-3p in ccRCC**

To explore the mechanism by which miR-489-3p functions in RCC, we searched TargetScan, Starbase, PITA, and RNA22 with the aim of identifying its target genes. This analysis revealed MEK1 as a target of miR-489-3p with a specific binding site at the 3'UTR (Figure 6A, 6B). GEPIA analysis showed that MEK1 expression was upregulated in ccRCC tissues (Figure 6C), and this was validated by RT-qPCR and IHC analysis (Figure 6D, 6E). A Kaplan-Meier analysis indicated that elevated MEK1 expression significantly correlated with poor prognosis (Figure 6F, 6G). MiR-489-3p mimics significantly suppressed MEK1 expression in 786O and A498 cells (Figure 6H, 6I). Additionally, a luciferase reporter assay revealed that miR-489-3p mimics significantly suppressed luciferase activity in the MEK1-Wt group but not in the MEK1-Mut group (Figure 6J). Moreover, correlation analysis showed that miR-489-3p expression negatively correlated with MEK1 expression in ccRCC tissues (Figure 6K). Taken together, these findings indicate that MEK1 is downstream target of miR-489-3p in ccRCC.

---

**Figure 3.** circDHX33 promoted the proliferation and invasion in ccRCC cells. (A) The expression of circDHX33 in RCC cell lines. (B) The efficiency of circDHX33 knockdown in 786O and A498 cells. (C-F) Cell viabilities were assessed by CCK-8 and colony formation and tests. (G, H) Cell invasion was assessed using Transwell assay. *p<0.05, **p<0.01.
circDHX33 promotes ccRCC progression through the miR-489-3p/MEK1 axis

Next, we investigated whether circDHX33 influences ccRCC progression via miR-489-3p/MEK1 axis. RT-qPCR showed that silencing circDHX33 decreased MEK1 expression in 786O cells and A498 cells, and that this effect was reversed by inhibitors of miR-489-3p (Figure 7A). Correlation analysis revealed a positive association between circDHX33 expression and MEK1 expression in ccRCC tissues (Figure 7B).

Finally, EdU and transwell assays showed that MEK1 overexpression abolished the effects of circDHX33 inhibition on the proliferation and invasion in 786O cells (Figure 7C, 7D).

circDHX33 promotes ccRCC cell growth in vivo

Next, we determined the functions of circDHX33 in tumor growth in vivo. A significant inhibition of tumor growth was observed following circDHX33 knockdown (Figure 8A). Measurement of tumor volume and weight indicated...
they were markedly lower in the circDHX33 knockdown group relative to control group (Figure 8B, 8C). IHC analysis of Ki67 staining revealed that circDHX33 knockdown suppressed tumor cell proliferation in vivo (Figure 8D). RT-qPCR assay revealed that miR-489-3p was upregulated in sh-circDHX33 group and MEK1 was downregulated in the sh-circDHX33 group (Figure 8E). Taken together, these observations indicate that circDHX33 may enhance MEK1 expression by acting as a ceRNA for miR-489-3p, thereby promoting ccRCC progression.

DISCUSSION

The initiation, development and progression of ccRCC are complex processes that involve a wide range of internal and external factors [14, 15]. CircRNAs have been reported to play important roles in the pathogenesis of ccRCC [16]. Huang et al showed that elevated levels of circRNA ABCB10 promoted the progression of cancer and predicted poor prognosis [17]. Xue et al found that circ-AKT3 reduced ccRCC metastasis by modulating the miR-296-3p/E-cadherin axis [18]. Another study reported that circPCNL2 promoted the invasiveness and proliferation of ccRCC cells by regulating the miR-153/ZEB2 axis [19]. However, the mechanisms by which circRNAs influence ccRCC are largely unknown.

In this study, we uncovered a novel circRNA (hsa_circ_0000740/circDHX33). circDHX33 is formed by DHX33 RNA splicing and comprises a circular

![Figure 5. circDHX33/miR-489-3p axis in ccRCC. (A, B) The level of miR-489-3p in ccRCC tissues and cell lines was estimated with RT-qPCR. (C, D) Low miR-489-3p levels correlated with poor overall survival rate in patients. (E) circDHX33 expression was negatively correlated with miR-489-3p expression in ccRCC tissues. (F) circDHX33 negatively regulated expression of miR-489-3p in 786O and A498 cells. (G, H) miR-489-3p suppression abolished the effects of circDHX33 inhibition on 786O cells proliferation and invasion abilities. *p<0.05, **p<0.01.](image-url)
fragment of exon 2 and 3. DHX33 is a member of the DEAD/DEAH box family of RNA helicases involved in cancer progression [20]. Here, we find that circDHX33 is markedly upregulated in ccRCC cell lines and tissues. CircDHX33 suppression decreased the proliferative capacity and invasiveness of ccRCC cells in vitro and suppressed the growth of tumors in vivo suggesting circDHX33 is a novel therapeutic target for ccRCC.

---

Figure 6. MEK1 served as the target of miR-489-3p. (A, B) MEK1 was a potential putative target gene of miR-489-3p. (C) MEK1 expression was significantly upregulated in KIRC tissues. (D, E) MEK1 expression in ccRCC tissues was determined by RT-qPCR and IHC. (F, G) High MEK1 expression was significantly associated with poor prognosis in patients. (H, I) MiR-489-3p mimics reduced MEK1 expression in 786O and A498 cells. (J) MiR-489-3p mimics reduced luciferase activity of the MEK1-Wt group. (K) miR-489-3p expression was negatively correlated with MEK1 expression in ccRCC tissues. KIRC: Kidney renal clear cell carcinoma. *p<0.05, **p<0.01.
Figure 7. circDHX33/miR-489-3p/MEK1 axis in ccRCC. (A) MiR-489-3p suppression in 786O and A498 cells abolished the effects of circDHX33 inhibition on MEK1 expression. (B) circDHX33 expression was positively correlated with MEK1 expression in ccRCC tissues. (C, D) MEK1 overexpression abolished the effects of circDHX33 inhibition on 786O cells proliferation and invasion abilities. *p<0.05, **p<0.01.

Figure 8. circDHX33 promoted tumor growth in vivo. (A–C) circDHX33 knockdown reduced tumor growth in vivo. (D) circDHX33 knockdown reduced Ki67 expression in mice tissues. (E) circDHX33 knockdown reduced MEK1 levels and increased miR-489-3p levels in mice tissues. *p<0.05, **p<0.01.
There is evidence that circRNAs modulates miRNA function by acting as a miRNA sponge [12]. MiR-489-3p is a tumor suppressor in many types of cancers. Inhibition of miR-489-3p enhanced metastasis of osteosarcoma by stimulating PAX3-MET axis [21]. It was established that miR-489 inhibited the proliferative capacity and invasiveness of bladder cancer cells [22]. LINC01446 promoted the progression of glioblastoma cells by modulating miR-489-3p/TPT1 axis [23]. Herein, we uncover that circDHX33 binds to miR-489-3p in ccRCC. We also show that MiR-489-3p is reduced in ccRCC. Moreover, we find that circDHX33-mediated suppression of cell proliferation and invasion are mitigated by miR-489-3p inhibitors. Together, these data suggest that circDHX33 exerts oncogenic effects by sponging miR-489-3p in ccRCC.

Mitogen-activated protein kinase (MEK) plays critical roles in tumor progression [24]. Huang et al reported that miR-101 regulates proliferative capacity and apoptosis in diffuse large B-cell lymphoma through MEK1 and ERK/MAPK pathway [25]. Zhang et al demonstrated that circDLST promotes gastric cancer progression by modulating the miR-502-5p/NRAS/MEK1/ERK1/2 axis [26]. In this study, we report that MEK1 is upregulated and positively correlates with circDHX33 expression. We find that MEK1 is directly targeted by miR-489-3p and circDHX33 indirectly regulates MEK1 expression. Moreover, the effects of circDHX33 inhibition on cell proliferation and invasion were reversed by MEK1 overexpression, suggesting MEK1 might be important in ccRCC development. These findings indicate that circDHX33 might regulate ccRCC growth and metastasis by regulating the miR-489-3p/MEK1 axis.

CONCLUSION

This is the pioneer study showing that circDHX33 has an oncogenic role ccRCC both in vivo and in vitro. Mechanistically, we show that the effects of circDHX33 are mediated via the circDHX33/miR-489-3p/MEK1 axis. circDHX33 might therefore be an ideal therapeutic target for ccRCC treatment.

MATERIALS AND METHODS

Clinical samples

We collected 47 paired ccRCC tissues and adjacent normal tissues from patients at the department of urology, Ninghai First Hospital and Huaihe Hospital of Henan University. The diagnosis of ccRCC was confirmed by 2 independent pathologists. Ethical approval for this study was granted by the ethics committee of the Ninghai First Hospital and Huaihe Hospital of Henan University. All participants provided written informed consents for sample collection.

Cell culture and treatment

Renal cancer cell lines, ACHN, Caki2, A498, Caki1, and 786O, and normal kidney epithelial cells line (HK2) were obtained from the American type culture collection (ATCC). These cells were grown in Dulbecco’s modified Eagle’s medium (DMEM; Gibco, Carlsbad, CA, USA) containing 1% penicillin/streptomycin (pen/strep) (Invitrogen, Carlsbad, CA, USA) and with 10% FBS (Gibco) in a humidified incubator, at 37°C, 5% CO2.

circDHX33 silencing-specific siRNAs (si-circDHX33#1 sequence is 5’- AGATCCACAGGTTGGCTATA-3’; si-circDHX33#2 sequence is 5’-ATCCACAGGTGGCTATACA-3’; si-circDHX33#3 sequence is 5’-CAGATCCACAGTGGCTAT-3’), were purchased from GenePharma (Shanghai, China). MiR-489-3p inhibitors and mimics, pcDNA.3.1 MEK1 recombinant plasmid and the respective control RNAs were purchased from Ribobio (Guangzhou, China) siRNA transfections were done with lipofectamine 2000 (Invitrogen) following the manufacturer’s directions.

Cell proliferation assay

Cell Counting Kit-8 (CCK-8; Roche, Basel, Switzerland) was used to assess the proliferation of RCC cells. After seeding 1×104 transfected cells in 96-well plates, they were cultured for 24, 48 or 72 hours. 10μl of CCK-8 was then added to the cells and cultured for 2 hours. The absorbance of the plate was read at 450nm using a microplate reader.

Colony formation assay

To assess the ability of cells to form colonies, 1×103 cells were cultured in 6-well plates with DMEM medium for 2 weeks. The culture medium was replaced at intervals of 3 days. Next, 1X PBS was used to wash colonies, which were then fixed with methanol for staining with crystal violet after 2 weeks in culture. The number of colonies were counted manually.

Transwell invasion assay

We evaluated cell invasion using transwell chambers. Briefly, tranwell chambers were coated with 80μl of Matrigel and 3×105 cells in 400μl of serum-free DMEM seeded onto the upper chamber. The lower chamber was filled with 400μl DMEM supplemented with 10% FBS. 24 hours later, un-migrated cells were scrapped off with
a cotton swab while invasive cells were stained with 0.1% crystal violet before counting under a microscope.

**RNA extraction and quantitative real-time PCR (RT-qPCR)**

Cells and tissues were lysed with Trizol reagent (Invitrogen) to isolate total RNA. RT-qPCR was performed using First Strand cDNA Synthesis Kit (TakaRa, China) as per the manufacturer’s protocol. RT-qPCR analysis was done using the ChamQ SYBR qPCR Master Mix (Vazyme Biotech, Nanjing, China) on a Roche LC 96 qPCR system (Roche, Germany). Relative mRNA levels were estimated using the 2ΔΔCT method. Primer sequences were as follows: circDHX33, 5′-GCTTGTCTCCGTCTTCCAGA-3′ (forward) and 5′-CTGAATTGCTTCACGCAGA-3′ (reverse); miR-489-3p, 5′-GGGTTGACATCATCATATAC-3′ (forward) and 5′-CAGTGGCTGTGGAGTAG-3′ (reverse); MEK1, 5′-CAAGAAGGAAGCAGGCCCCAT-3′ (forward) and 5′-GACGCCAGCACATGGGTTG-3′; U6, 5′-CCGCGCTCGGCACACATATAC-3′ (forward) and U6, 5′- ACGTTCAGAAATTTGCGTGTC-3′ (reverse); GAPDH, 5′-GAGGCCACGACATGGGTTG-3′ (forward) and 5′-CTGAAATTGCTTGGCTGTGGAGTAG-3′ (reverse).

**Dual-luciferase reporter assay**

circDHX33 segment or MEK1 3′UTR was inserted into pGL3-control plasmid. Mutations on circDHX33 and the MEK1 3′UTR were induced by changing the conserved binding sites of miR-489-3p using a gene mutation kit (Takara). The mutated sequence or wild-type seed region were co-transfected with miR-489-3p mimics or miR-NC into RCC cells using Lipofectamine 2000. Luciferase activity was measured using the Dual-Luciferase® reporter assay system (Promega, Madison, WI, USA) according to the manufacturer’s protocol.

**RNA immunoprecipitation (RIP)**

RIP was done using the Magna RIP Kit (Millipore, Billerica, MA, USA) as per the manufacturer’s instructions. Antibodies against argonaute2 (anti-AGO2) and immunoglobulin G (IgG) were used for the RIP assays. Purified RNAs were extracted and the enrichment of circDHX33 and miR-489-3p analyzed by RT-qPCR, as described previously [11].

**Tumorigenesis assay**

Twelve (12) female nude mice (athymic BALB/c) aged 4-6 weeks and weighing 16-20g were housed in pathogen-free conditions. The animals were randomly split into two groups (the sh-NC and sh-circDHX33 groups). 200μL of 1×10⁷ cells were subcutaneously injected into the back flanks of the animals. Tumors were measured each week after the tumor became visible and tumor volume calculated following with the formula: volume = (0.5 × length × width²). After 6 weeks following implantation, the mice were sacrificed and tumors were excised, weighed and photographed.

**Bioinformatics analysis**

Survival analysis for RCC patients with miR-489-3p expression was performed using the KM plot database (http://kmplot.com) and PROGgeneV2 (http://genomics.jefferson.edu/proggene/results.php). The expression of MEK1 in RCC was determined by GEPIA (gepia.cancer-pku.cn). PROTEIN ATLAS (https://www.proteinatlas.org/) was used to investigate the prognosis of MEK1 in RCC. The circDHX33 specific binding with miRNAs was identified by using miRanda and RNAhybrid.

**Statistical analysis**

All data are presented as mean ± standard deviation (SD). Statistical analysis was done using SPSS 20.0 (SPSS, Chicago, IL, USA). Student’s t-test and one-way ANOVA tests were utilized to compare differences between groups, respectively. p < 0.05 was considered statistically significant.

**CONFLICTS OF INTEREST**

The authors declared no conflicts of interest.

**FUNDING**

Funding information National Natural Science Foundation of China, Grant/Award Numbers: 81502426, Zhejiang Youth Talent Project Foundation, Grant/Award Number: 2019RC272 and 2020RC112; Bureau of Science, Peiyi Program of Ninghai First Hospital

**REFERENCES**

1. Siegel RL, Miller KD, Jemal A. Cancer statistics, 2019. CA Cancer J Clin. 2019; 69:7–34. https://doi.org/10.3322/caac.21551 PMID:30620402

2. Rossi SH, Klatte T, Usher-Smith J, Stewart GD. Epidemiology and screening for renal cancer. World J Urol. 2018; 36:1341–53. https://doi.org/10.1007/s00345-018-2286-7 PMID:29610964

3. Hsieh JJ, Purdue MP, Signoretti S, Swanton C, Albiges L, Schmidinger M, Heng DY, Larkin J, Ficarra V. Renal cell carcinoma. Nat Rev Dis Primers. 2017; 3:17009.

www.aging-us.com
An emerging function of circRNA:

1. Zhang HD, Jiang LH, Sun DW, Hou JC, Ji ZL. CircRNA: a novel type of biomarker for cancer. Breast Cancer. 2018; 25:1–7. https://doi.org/10.1007/s12282-017-0793-9 PMID: 28721656

2. Bielecka ZF, Czarnecka AM, Solarek W, Kornakiewicz A, Szczyluk C. Mechanisms of acquired resistance to tyrosine kinase inhibitors in clear - cell renal cell carcinoma (ccRCC). Curr Signal Transduct Ther. 2014; 8:218–28. https://doi.org/10.2174/1574362409666140206223014 PMID: 25152704

3. Phillips R. Kidney cancer: FBP1 depletion feeds ccRCC. Nat Rev Urol. 2014; 11:482. https://doi.org/10.1038/nrurol.2014.200 PMID: 25091010

4. Wang K, Sun Y, Tao W, Fei X, Chang C. Androgen receptor (AR) promotes clear cell renal cell carcinoma (ccRCC) migration and invasion via altering the circHIAT1/miR-195-5p/29a-3p/29c-3p/CDC42 signals. Cancer Lett. 2017; 394:1–12. https://doi.org/10.1016/j.canlet.2016.12.036 PMID: 28089832

5. Huang Y, Zhang Y, Jia L, Liu C, Xu F. Circular RNA ABCB10 promotes tumor progression and correlates with pejorative prognosis in clear cell renal cell carcinoma. Int J Biol Markers. 2019; 34:176-83. https://doi.org/10.1177/1724600819842279 PMID: 31106654

6. Xue D, Wang H, Chen Y, Shen D, Lu J, Wang M, Zebibula A, Xu L, Wu H, Li G, Xia L. circ-AKT3 inhibits clear cell renal cell carcinoma metastasis via altering miR-296-3p/e-cadherin signals. Mol Cancer. 2019; 18:151. https://doi.org/10.1186/s12943-019-1072-5 PMID: 31672157

7. Zhou B, Zheng P, Li Z, Li H, Wang X, Shi Z, Han Q. CircPCNL2 sponges miR-153 to promote the proliferation and invasion of renal cancer cells through upregulating ZEB2. Cell Cycle. 2018; 17:2644–54. https://doi.org/10.1080/15384101.2018.1553354 PMID: 30488762

8. Fu J, Liu Y, Wang X, Yuan B, Zhang Y. Role of DHX33 in c-myc-induced cancers. Carcinogenesis. 2017; 38:649–60. https://doi.org/10.1093/carcin/bgx041 PMID: 28498893

9. Liu Q, Yang G, Qian Y. Loss of MicroRNA-489-3p promotes osteosarcoma metastasis by activating PAX3-MET pathway. Mol Carcinog. 2017; 56:1312–21. https://doi.org/10.1002/mc.22593 PMID: 27859625

10. Zhang HM, Yang FQ, Chen SJ, Che J, Zheng JH. Upregulation of long non-coding RNA MALAT1 correlates with tumor progression and poor prognosis in clear cell renal cell carcinoma. Tumour Biol. 2015; 36:2947–55. https://doi.org/10.1007/s13277-014-2925-6 PMID: 25480417

11. Rong D, Sun H, Li Z, Liu Y, Dong C, Fu K, Tang W, Cao H. An emerging function of circRNA-miRNAs-miRNA axis in human diseases. Oncotarget. 2017; 8:73271–81. https://doi.org/10.18632/oncotarget.19154 PMID: 29069868

12. Shen Q, He T, Yuan H. Hsa_circ_0020577 promotes endometrial carcinoma progression via regulating miR-197/CTNNB1 axis and activating Wnt/β-catenin pathway. Cell Cycle. 2019; 18:1229–40. https://doi.org/10.1080/15384101.2019.1617004 PMID: 31081718

13. Ge Z, Li LF, Wang CY, Wang Y, Ma WL. CircMTO1 inhibits cell proliferation and invasion by regulating Wnt/β-catenin signaling pathway in colorectal cancer. Eur Rev Med Pharmacol Sci. 2018; 22:8203–9. https://doi.org/10.26355/eurrev_201812_16513 PMID: 30568859

14. Zhao S, Wang C, Wang S, Zhang H, Zhang Y. Hsa_circ_0075341 is up-regulated and exerts oncogenic properties by sponging miR-149-5p in cervical cancer. Biomed Pharmacother. 2020; 121:109582. https://doi.org/10.1016/j.biopha.2019.109582 PMID: 31706100

15. Jia L, Liu C, Xu F. Circular RNA MALAT1 regulates cell proliferation and invasion by regulating Wnt/β-catenin signaling pathway. Tumour Biol. 2019; 40:15384101.2019.1617004 PMID: 31081718

16. McDermott D, Mier J, Stanbridge E, Youmans A, Febbo P, Upton M, Lechpammer M, Signoretti S. Carbonic anhydrase IX expression predicts outcome of interleukin 2 therapy for renal cancer. Clin Cancer Res. 2005; 11:3714–21. https://doi.org/10.1158/1078-0432.CCR-04-2019 PMID: 15897568

17. Ljungberg B, Campbell SC, Choi HY, Jacqmin D, Lee JE, Weikert S, Kiemeney LA. The epidemiology of renal cell carcinoma. Eur Urol. 2011; 60:615–25. https://doi.org/10.1016/j.eururo.2011.06.049 PMID: 21741761

18. Phillips R. Kidney cancer: FBP1 depletion feeds ccRCC. Eur Urol. 2014; 65:197–55. https://doi.org/10.1038/nrdp.2017.9 PMID: 28276433
22. Li J, Qu W, Jiang Y, Sun Y, Cheng Y, Zou T, Du S. miR-489 suppresses proliferation and invasion of human bladder cancer cells. Oncol Res. 2016; 24:391–98. https://doi.org/10.3727/096504016X14666990347518 PMID:28281959

23. Zhang L, Wang Q, Wang F, Zhang X, Zhang L, Tang Y, Wang S. LncRNA LINCO1446 promotes glioblastoma progression by modulating miR-489-3p/TPT1 axis. Biochem Biophys Res Commun. 2018; 503:1484–90. https://doi.org/10.1016/j.bbrc.2018.07.067 PMID:30029885

24. Johnson GL, Lapadat R. Mitogen-activated protein kinase pathways mediated by ERK, JNK, and p38 protein kinases. Science. 2002; 298:1911–12. https://doi.org/10.1126/science.1072682 PMID:12471242

25. Huang Y, Zou Y, Lin L, Ma X, Zheng R. miR-101 regulates the cell proliferation and apoptosis in diffuse large b-cell lymphoma by targeting MEK1 via regulation of the ERK/MAPK signaling pathway. Oncol Rep. 2019; 41:377–86. https://doi.org/10.3892/or.2018.6821 PMID:30365139

26. Zhang J, Hou L, Liang R, Chen X, Zhang R, Chen W, Zhu J. CircDLST promotes the tumorigenesis and metastasis of gastric cancer by sponging miR-502-5p and activating the NRAS/MEK1/ERK1/2 signaling. Mol Cancer. 2019; 18:80. https://doi.org/10.1186/s12943-019-1015-1 PMID:30953514