Tunicamycin suppresses renal cell carcinoma cells growth and metastasis through regulation of ERK/AKT signaling pathway.

Yiping Jia¹, Yuan Yang², Yan Luo*³

¹Department of Ultrasound, No. 4 West China Teaching Hospital, Sichuan University, Chengdu, PR China
²Department of Microbiology, West China School of Preclinical and Forensic Medicine, Sichuan University, Chengdu, PR China
³Department of Ultrasound, West China Hospital, Sichuan University, Chengdu, PR China

Abstract

Renal Cell Carcinoma (RCC) is a higher malignant degree tumor in urinary system. Evidences have suggested that tunicamycin can inhibit many human cancer growth and metastasis. The purpose of this study is to investigate the role of tunicamycin in the progression of RCC cells and RCC mouse model. Results showed that tunicamycin significantly suppressed ACHN and RuCa cells growth, migration and invasion in vitro. We demonstrated that tunicamycin treatment arrested cell cycle of ACHN and RuCa cells. In addition, results indicated that tunicamycin treatment induced apoptosis and down-regulated ERK and AKT expression and phosphorylation levels in ACHN and RuCa cells. Mechanism research showed that overexpression of ERK increased AKT expression and phosphorylation levels and canceled Tunicamycin-down-regulated AKT expression and phosphorylation levels in ACHN and RuCa cells. In addition, overexpression of ERK inhibited Tunicamycin-suppressed growth and aggressiveness of ACHN and RuCa cells. Further, in vivo assays showed that Tunicamycin markedly inhibited tumor growth in ACHN-bearing mouse model via increasing apoptosis. In conclusion, these results indicate that tunicamycin could inhibit growth of RCC cells both in vitro and in vivo via ERK/AKT signal pathway, suggesting tunicamycin may be an efficient anti-cancer agent for RCC therapy.

Keywords: RCC, Tunicamycin, Metastasis, Apoptosis, ERK/AKT.

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Introduction

Renal Cell Carcinoma (RCC) is originated from urinary tubular epithelium and RCC represents 80-90% in human malignant tumors [1,2]. Among urologic neoplasm, the incidence of RCC is the third highest and growing at percent of 6.5 average year perannumpa, and the majority of RCC patients eventually develop metastatic RCC [3,4]. Pathological classification of treatment and prognosis of kidney cancer has important guiding significance for patients with RCC [5,6]. The performance of the RCC is various and mainly presents hematuria, flank pain and swelling for clinical patients [7,8]. RCC is highly aggressive and generally resistant to chemotherapy, radiotherapy, and hormone therapy compared to that many other malignancies [9,10]. Therefore, new therapeutic alternatives and elaboration of molecular mechanism are needed in order to provide more effective treatments of malignancy of RCC.

Tunicamycin is nucleotide antibiotic produced by Streptomyces lyosuperficus. A large numbers of reports have suggested that Tunicamycin can be widely suppressed human cancer cells growth and metastasis [11,12]. Evidences have showed that inhibition of N-linked glycosylation by tunicamycin induces E-cadherin-mediated cell-cell adhesion and inhibits cell proliferation in undifferentiated human colon cancer cells [13]. Huong et al. have showed that proteasome inhibitor-I can enhance tunicamycin-induced chemosensitization of prostate cancer cells through up-regulation of NF-kappaB and CHOP expression [14]. Further, tunicamycin sensitizes human melanoma cells to tumor necrosis factor-related apoptosis-inducing ligand-induced apoptosis by up-regulation of TRAIL-R2 via the unfolded protein response [15]. However, no reports have investigated the therapeutic role of Tunicamycin on RCC mouse model.

In this study, we analysed the inhibitory effects of Tunicamycin on RCC cells in RCC cells and RCC mouse model. We investigated the potential signal mechanism mediated by Tunicamycin in breast cancer cells. We also identified apoptosis rate of CRC cells were induced by Tunicamycin. Results presented that Tunicamycin may be an efficient anti-cancer agent for the treatment of RCC via regulation of ERK/AKT signaling pathway.
Materials and Methods

Ethic statement

This study was performed by the recommendations in the Guide for the Care and Use of Laboratory Animals of Sichuan University. All animals’ protocols were experimented in accordance with National Institutes of Health and approved by Committee on the Ethics of Animal Experiments Defence Research.

Cells and reagents

ACHN and RuCa were purchased from American Type Culture Collection (ATCC). All tumor cells were cultured in DMEM (Sigma-Aldrich) medium (Gibco, CA, USA) supplemented with 10% Fetal Serum (FBS) (Invitrogen, CA, USA). All cells were cultured in a 37°C humidified atmosphere of 5% CO2.

Flow cytometry

Apoptosis of ACHN and RuCa cells were evaluated using Annexin V-fluorescein isothiocyanate (Annexin V-FITC) and Propidium Iodide (PI) apoptosis detection kit (Becton-Dickinson, CA, USA). ACHN and RuCa cells were isolated from Simvastatin- or PBS-treated mice and suspended with Annexin V-FITC and PI according to the manufacturer instructions. Fluorescence was measured with a FACS scan flow cytometry (Becton-Dickinson, San Jose, CA, USA).

MTT assays

ACHN and RuCa cells were transfected by pvector or pAKT and culture in 96-well plates for 48 h. Growing ACHN and RuCa cells were treated by 10 mg/ml Tunicamycin for 48 h. After incubation, 20 μl of MTT (10 mg/ml) in PBS solution was added to each well, the plate was further incubated for 4 h. Most of the medium was removed and 100 μl of DMSO was added into the wells to solubilize the crystals. The OD was measured by a BIO-RAD (ELISA) reader at wavelength of 450 nm.

Endogenous expression of ERK

ACHN and RuCa cells were cultured until 90% confluence and the media was then removed from culture plate. ACHN and RuCa cells were transfected by lentivirus-ERK (pERK) or lentivirus-vector (Control) by using lipofectamine 2000 (Sigma-Aldrich). Stable ERK-overexpressed ACHN and RuCa cells were treated by Tunicamycin (10 mg/ml) for 24 h, followed homogenized at 4°C for 10 min. Protein were incubated with rabbit anti-human ERK (1: 500, ab17942, Abcam), pERK (1: 500, ab64148, Abcam), pAKT (1: 500, ab81283, Abcam) or β-actin (1: 500, ab32572, Abcam), AKT (1: 500, ab27942, Abcam) and Bcl-2 (1: 500, ab79483, Abcam) for 12 hours at 4°C. Protein were incubated with rabbit anti-human ERK (1: 500, ab17942, Abcam), pERK (1: 500, ab79483, Abcam), AKT (1: 500, ab64148, Abcam), pAKT (1: 500, ab81283, Abcam) or β-actin (1: 500, ab32572, Abcam) for 12 hours at 4°C. Horseradish peroxidase (HRP)-conjugated secondary antibody (Bio-Rad, Hercules, CA, USA) was used at a 1:5,000 dilution and detected using a Western Blotting Luminol Reagent. The results were visualized by using chemi-luminescence detection system (Amersham Biosciences, Piscataway, NJ).

Cell invasion and migration assays

Sable AKT-overexpressed ACHN and RuCa cells were cultured with Tunicamycin (5 mg/ml) or PBS for 48 hours. Migration and invasion of ACHN and RuCa cells was performed in a 6 well culture plate with chamber inserts (BD Biosciences). For migration assays, 1 × 10⁴/well concentration of the ACHN and RuCa cells were placed into the upper chamber. For invasion assays, ACHN and RuCa cells (1 × 10⁵/well) were placed into the upper chamber with the Matrigel-coated membrane. Migration and invasion of ACHN and RuCa cells were counted in at least three randomly stain-field microscope every membrane.

Real-time quantitative PCR (RT-qPCR)

Total mRNA was isolated from ACHN and RuCa cells by using RNA Easy Mini Extract Kit (Sigma-Aldrich, Gaithersburg, USA). Expressions of Bcl-2, P53, Caspase-3 and Bad in ACHN and RuCa cells were calculated by using RT-qPCR kit with β-actin expression as an endogenous control. All procedures were performed according to the manufacturer’s instructions. All the primers were synthesized by Invitrogen (Invitrogen, Shanghai, China). Relative mRNA expression levels were determined by 2-ΔΔCt. The final results were presented as the n-fold manner compared to β-actin.

Analysis of cells cycle

To analyse the effects of Tunicamycin (10 mg/ml) on the cell cycle stage of ACHN and RuCa cells, flow cytometry was performed. Exponentially, culturing ACHN and RuCa cells or ERK-overexpressed ACHN and RuCa cells were treated with Tunicamycin (10 mg/ml) for 48 h. Cells were washed and trypsinized and rinsed with Phosphate-Buffered Saline (PBS). All cells were fixed in 75% ice-cold ethanol for 5 minutes and then washed with PBS three times. The fixed cells were washed with RNase A (20 μg/ml, Fermentas) and stained with propidium iodide (20 µg ml/l, Sigma-Aldrich) for 10 min at 37°C. The percentages of cells in G0/G1 and S phase were analysed using BD FACS Calibur (Becton Dickinson, NJ, USA).

Western Blotting

ACHN and RuCa cells were treated by Tunicamycin (10 mg/ml) and harvested by scraping and lysed in RIPA buffer followed homogenized at 4°C for 10 min. Protein were analysed by SDS-PAGE assays followed transfer membrane. Protein were incubated with rabbit anti-human ERK (1: 500, ab17942, Abcam), pERK (1: 500, ab79483, Abcam), AKT (1: 500, ab64148, Abcam), pAKT (1: 500, ab81283, Abcam) or β-actin (1: 500, ab32572, Abcam) for 12 hours at 4°C. Horseradish peroxidase (HRP)-conjugated secondary antibody (Bio-Rad, Hercules, CA, USA) was used at a 1:5,000 dilution and detected using a Western Blotting Luminol Reagent. The results were visualized by using chemi-luminescence detection system (Amersham Biosciences, Piscataway, NJ).

Animal study

Male Balb/c (8 weeks old, body weight: 30-35 g) mice were purchased from Slack co., LTD (Shanghai, China). Mice were maintained at a 12 h light/dark cycle with free access to diet and water. Experimental mice were implanted ACHN cells (5 × 10⁵) into groin and were divided into two groups (n=50 in
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Tunicamycin treatment inhibited migration and invasion of ACHN and RuCa cells (Figures 1B and 1C). These results suggest that Tunicamycin treatment can inhibit growth and aggressiveness of RCC cells.

**TUNEL analysis**

For analysis the apoptosis of myocardial cells, the terminal deoxynucleotidyl transferase (TdT)-mediated dUTP nick end labeling (TUNEL) assay (Biotool) were used to detect TUNEL-positive cells. Tumor cells from experimental mice after treated with Tunicamycin or PBS were fixed with 4% paraformaldehyde solution for 60 minutes at 4°C. Subsequently, cells were incubated with equilibration buffer for 30 minutes at 4°C. The tumor sections were then incubated with 50 μl of the reaction mixture at 37°C for 60 minutes and washed 3 times with PBS. The cells nuclei were stained with 4', 6-Diamidino-2-Phenylindole (DAPI) for 60 minutes at 4°C. Finally, tumor tissues images were captured with a ZEISS LSM 510 confocal microscope at 488 nm.

**Immunohistochemistry**

Tumors from xenografted mice were fixed by using formaldehyde (10%) followed with embed in paraffin. Immunohistochemical staining was performed by an avidin-biotin-peroxidase technique. Paraffin-embedded tissue sections were prepared and epitope retrieval was performed for further analysis. The paraffin sections were subjected with hydrogen peroxide (3%) for 10~15 minutes, which subsequently were blocked by a regular blocking solution for 10~15 minutes 37°C. Finally, the sections were incubated with biotinylated goat anti-human ERK (1: 500, ab17942, Abcam) and AKT (1: 500, ab64148, Abcam), respectively, at 4°C for 12 h. Samples were washed with PBS three times and relative expression of proteins was analysed by chemi-luminescence detection system (Amersham Biosciences, Piscataway, NJ).

**Statistical analysis**

Results are calculated as mean ± SEM. Significance was established with the SPSS statistical (SPSS, USA) and GraphPad Prism 5 software (GraphPad Software, USA), Pearson's correlation coefficient or a two-tailed Student's t-test. The data are presented as the mean and SEM of three independent experiments. Differences were considered significant at *P<0.05, **P<0.01.

**Results**

**Tunicamycin treatment inhibits growth and aggressiveness of RCC cells**

We first investigated the inhibitory effects of Tunicamycin treatment on growth and aggressiveness of RCC cells. As shown in Figure 1A, tunicamycin treatment significantly inhibited ACHN and RuCa cells growth. We found that Tunicamycin treatment inhibited migration and invasion of ACHN and RuCa cells. We next analysed the efficacy of Tunicamycin for cell cycle in vitro. Our results showed that Tunicamycin treatment arrested G_{0}/G_{1} and S phase of ACHN and RuCa cells (Figures 3A and 3B). Results showed that Tunicamycin down-regulated CDK1 and CDk2 mRNA expression levels in ACHN and RuCa cells (Figures 3C and 3D). These results indicate that Tunicamycin can arrest cells cycle of RCC cell cycle through down-regulation of CDK1 and CDk2 expression levels.
investigate ERK/AKT signal pathway in RCC cells. We found that Tunicamycin treatment decreased expression and phosphorylation levels of ERK and AKT in ACHN and RuCa cells (Figures 4A and 4B). We observed that overexpression of ERK (OPERK) increased AKT expression levels and inhibited Tunicamycin-inhibited (OPERK/TC) AKT expression in ACHN and RuCa cells (Figure 4C). We found that overexpression of ERK abolished Tunicamycin-inhibited growth, migration and invasion of ACHN and RuCa cells (Figures 4D and 4F). Collectively, these results suggest that tunicamycin can regulate RCC cells growth and aggressiveness through ERK/AKT signal pathway.

Figure 3. Tunicamycin arrests cells cycle at G0/G1 and S phase of RCC cell cycle (A, B) Tunicamycin treatment arrests G0/G1 (A) and S (B) phase of ACHN and RuCa cells. (C, D) Tunicamycin down-regulates CDK1 and CDk2 mRNA expression levels in ACHN (C) and RuCa (D) cells.

Figure 4. Tunicamycin regulates RCC cells growth and aggressiveness through ERK/AKT signal pathway. (A) Tunicamycin decreases expression and phosphorylation levels of ERK and AKT in ACHN and RuCa cells. (B) Overexpression of ERK (OPERK) increases AKT expression levels and inhibits Tunicamycin-inhibited (OPERK/TC) AKT expression in ACHN and RuCa cells. (C) Overexpression of ERK (OPERK) canceled Tunicamycin-inhibited growth of ACHN and RuCa cells. (D-E) Overexpression of ERK (OPERK) canceled Tunicamycin-inhibited migration (D) and invasion (E) of ACHN and RuCa cells.

Tunicamycin inhibits tumor in RCC-bearing mouse model

We further the in vivo effects of Tunicamycin on growth in ACHN-bearing mice. As shown in Figure 5A, we found that Tunicamycin treatment (10 mg/kg) significantly inhibited RCC tumor growth compared to PBS group. We observed that Tunicamycin treatment increased apoptotic body in tumor sections in Tunicamycin-treated mice (Figure 5B). Immunohistochemistry assays showed that Tunicamycin treatment decreased ERK and AKT expression levels in tumor tissues compared to PBS-treated tumors (Figure 5C). We also observed that Tunicamycin treatment prolonged survival of RCC-bearing mouse model compared to PBS-treated mice (Figure 5D). These results suggest that Tunicamycin treatment can significantly inhibited RCC growth in vivo.

Figure 5. In vivo efficacy of Tunicamycin in RCC-bearing mouse model (A) Tunicamycin treatment (10 mg/kg) significantly inhibits RCC tumor growth compared to PBS group. (B) Tunicamycin treatment increases apoptotic body in tumor sections in experimental mice. (C) Tunicamycin treatment decreases ERK and AKT expression in tumor sections in experimental mice. (D) Tunicamycin treatment prolongs survival of RCC-bearing mouse model compared to PBS-treated mice.

Discussion

The global incidence of RCC is increasing annually and the causes are multifactorial [17]. Though multiple new targets have been identified, and therapeutic agents are in varying stages of development, the targeted therapy remains a promising treatment modality in human cancer [18,19]. Hopefully, the approval of newer agents will translate into more accessible and efficacious options for patients and oncologists. Currently, previous study has reported the efficacy of Tunicamycin for the treatment of human cancer [11,20]. Research also has showed that Tunicamycin presents anti-tumor efficacy by killing tumor cells growth and enhancing apoptosis of tumor cells [21,22]. Findings have showed that Tunicamycin significantly inhibited growth and aggressiveness of RCC cells. We have found that Tunicamycin treatment induced apoptosis and arrested cell cycle of ACHN and RuCa cells, which contributed to long-term survival rate of RCC-bearing mouse model after treatment with Tunicamycin. Ferraro et al. have indicated that EZH2 is regulated by ERK/AKT and targets integrin alpha2 gene to control epithelial-mesenchymal transition and anoikis in colon cancer cells [23]. Unfolded protein response is required in nu/nu mice microvasculature for treating breast tumor with Tunicamycin,
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which strongly supported tunicamycin's potential for a powerful glycotherapeutic treatment of breast cancer in the clinic [24]. In this study, we reported the efficacy of Tunicamycin for the treatment of RCC. Findings have indicated that Tunicamycin induced apoptosis of RCC cells via increasing pro-apoptosis gene expression. In addition, blockage of ERK/Akt signaling pathway promoted human breast cancer cells apoptosis and death through reactive oxygen species-dependent down-regulation [25]. Furthermore, inhibition of prostate cancer proliferation is paralleled by Erk/Akt inactivation and reduced Bcl-2 and bad phosphorylation [26]. Our results showed that Tunicamycin treatment decreased Bcl-2 and bad expression, which led to inhibition of growth of RCC cells through promoting apoptosis.

In conclusion, observed activity warrants further evaluation of the therapeutic effects of Tunicamycin in RCC cells and mouse model. Our present study demonstrates that up-regulation of expression of apoptosis contributes to inhibition of RCC growth, down-regulation of the ERK/AKT signaling pathways, as well as inhibition of migration and invasion, arresting cell cycle in RCC cell.

References


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*Correspondence to
Yan Luo
Department of Ultrasound
West China Hospital
Sichuan University
PR China
Email: luoyanprof@163.com