Abstract. The aim of the present study was to investigate the effects of the heat shock protein (HSP)90 inhibitor AUY922 on the proliferation and migratory ability of renal cell carcinoma (RCC) cells. The expression of HSP90 was measured in vitro using western blotting and quantitative polymerase chain reaction in the ACHN and 786-O cell RCC lines, and also in the immortalized normal human proximal tubule epithelium HK-2 cell line. The effects of the time and concentration of AUY922 administration were investigated in the ACHN and 786-O cells, and the cell proliferation was measured using an MTT assay. A Transwell assay was performed to evaluate the migratory ability of ACHN cells following treatment with AUY922 at concentrations of 10, 50 and 100 nM. Western blot analysis and reverse transcription polymerase chain reaction revealed that HSP90 mRNA and protein were overexpressed in the two RCC cell lines compared with the HK-2 cell line. AUY922 inhibited the proliferation of the ACHN and 786-O cells in a time- and concentration-dependent manner, and the migratory ability of the ACHN cells was markedly suppressed subsequent to treatment with AUY922. The present data suggest that the pathogenesis of human RCC may be mediated by HSP90. It was also indicated that the specific HSP90 inhibitor AUY922 plays a therapeutic role in the treatment of RCC, and therefore, HSP90 may be a selective target for molecular-targeted treatments of RCC.

Introduction

Renal cell carcinoma (RCC) accounts for ~90% of all kidney cancer cases, which in turn account for 2-3% of all cancers worldwide (1). RCC is the most lethal urological tumor and is the sixth leading cause of cancer-associated mortality in Western countries (1). Although the prevalence of RCC varies between populations and geographic regions, the incidence of RCC has increased steadily over previous years (2,3). With the increased use of various imaging modalities, an increased number of RCC cases have been diagnosed at an early stage of disease. However, 20-30% of patients with RCC demonstrate evidence of metastases at the time of the initial presentation (4). Although radical nephrectomy is an effective treatment for early-stage and local RCC, 30% of patients develop metastatic disease subsequent to surgery (5). Patients with metastatic RCC possess a poor prognosis, and therapeutic options are limited (6). The median survival of the cohort of patients with RCC enrolled in a previous study was only 1.5 years, with <10% of patients surviving for five years (7). Therefore, the overall prognosis of patients with RCC is limited, indicating the requirement for improved therapies directed at potential molecular targets.

Heat shock protein 90 (HSP90) is a cellular molecular chaperone that is important for protein assembly, folding and stability (8). The client proteins of HSP90, including human epidermal growth factor receptor 2, Raf, mutant p53 and vascular endothelial growth factor (VEGF) receptor, play critical roles in oncogenesis (9). Numerous studies have demonstrated that HSP90 is overexpressed in solid tumors, and have reported that the inhibition of HSP90 exerts a promising therapeutic effect on certain cancers (10,11). Previously, inhibition of HSP90 has been performed using the low molecular weight adenosine triphosphate (ATP)-competitive non-geldanamycin HSP90 inhibitor AUY922. This inhibitor has been speculated to offer advantages over ansamycin benzoquinone HSP90 inhibitors, including 17-allylamino-17-demethoxygeldanamycin (17-AAG), based on the independence of AUY992 from NAD(P)H:quinone oxidoreductase 1 metabolism, P-glycoprotein expression and favorable aqueous solubility (12). In the present study, the anti-tumor efficacy of the HSP90 inhibitor AUY922 was evaluated in the RCC ACHN and 786-O cell lines in vitro.

Materials and methods

Cell culture and reagents. The immortalized normal human proximal tubule epithelial HK-2 cell line was purchased from...
the American Type Culture Collection (Manassas, VA, USA). The human RCC 786-O and ACHN cell lines were obtained from the Cell Bank of Type Culture Collection of the Chinese Academy of Sciences (Beijing, China). The HK-2 cells were cultured in keratinocyte serum-free medium (Gibco Life Technologies, Carlsbad, CA, USA), and the 786-O and ACHN cells were cultured in RPMI-1640 medium (HyClone, Logan, UT, USA), supplemented with 10% fetal bovine serum (FBS) and 50 μg/ml streptomycin. All cells were cultured in a sterile incubator that was maintained at 37°C with an atmosphere of 5% CO₂. The HSPP90 inhibitor AUY922 was purchased from Selleck Chemicals (Houston, TX, USA). The compounds were dissolved in dimethylsulfoxide (DMSO) to generate a 5% DMSO stock solution and then stored at -20°C. The mouse anti-human HSP90 monoclonal antibody was obtained from Abcam (Cambridge, MA, USA; catalog no., ab13492).

Western blot analysis. The cells were lysed in lysis buffer containing protease inhibitor. The protein concentration was determined using a bicinchoninic protein assay kit (Pierce Biotechnology, Inc., Rockford, IL, USA). In total, 30 μg of the respective tissue proteins were separated by SDS-PAGE performed on 10% gels and then transferred onto polyvinylidene fluoride membranes (Millipore, Billerica, MA, USA). The membranes were blocked with 5% non-fat milk and then incubated with mouse anti-human HSPP90 monoclonal antibody (1:100; Abcam) and mouse anti-actin monoclonal antibody (catalog no. 0869100; 1:10,000; MP Biomedicals, Santa Ana, CA, USA). The membranes were washed three times for 10 min each using Tris-buffered saline comprising 50 mM Tris (pH 7.4), 0.9% NaCl and 0.05% Tween-20 (TBS-T), and then incubated with phycoerythrin-conjugated secondary antibodies (donkey anti-mouse IgG H&L; catalog no., ab7003; Abcam). The membranes were subsequently washed again with TBS-T three times, with each wash lasting for 10 min. The target protein bands were visualized using the enhanced chemiluminescence method. All western immunoblot analyses were performed three times.

Total RNA extraction and complementary (c)DNA synthesis. Total RNA was isolated from the HK2, 786-O and ACHN cell lines using TRIzol Reagent (Invitrogen, Carlsbad, CA, USA), according to the manufacturer's instructions. The concentration and quality of the extracted total RNA was determined by measuring the optical density (OD) at 260 and 280 nm, and then calculating the OD_{260}/OD_{280} ratio. cDNA was synthesized from 2 μg of total RNA using a Reverse Transcription System kit (Promega, Madison, WI, USA), according to the following protocol obtained from the reaction kit. Briefly, the samples were pre-incubated at 70°C for 10 min, cooled on ice and then added to a reaction mixture consisting of 10 mmol/l deoxynucleotide triphosphate mixture, 25 mmol/l MgCl₂, 15 units avian myoblastosis virus reverse transcriptase, 10X reverse transcription buffer, 0.5 units RNasin™ and 0.5 μg oligo-(dT)₁₅ primer (all provided with the Reverse Transcription System kit). The reaction mixture was made up to a final volume of 20 μl and was incubated successively at 44°C for 15 min, 99°C for 5 min and 4°C for 5 min. The cDNA was maintained at 20°C prior to use.

Reverse transcription-quantitative polymerase chain reaction (RT-qPCR). RT-qPCR was performed using SYBR Master Mix (Takara Bio Inc., Otsu, Japan) on an ABI Prism 7900HT Sequence Detection System (Applied Biosystems Life Technologies, Foster City, CA, USA). The human GAPDH gene was used as an endogenous control for sample normalization. The results were expressed as the fold change in expression relative to the expression of GAPDH. The PCR primers were as follows: GAPDH forward, 5'-GGGACACAAGCTTTCGCTT-3' and reverse, 5'-GACAAGCTTCCGTGTCCTGC-3'; and HSPP90 forward, 5'-TGTTTCAGTGTCCTCGGTGGC-3' and reverse, 5'-TGTTGGTGTCTTGGGCTCGGT-3'.

Cell viability measurement. Cell viability was analyzed using an MTT assay (Sigma-Aldrich, St. Louis, MO, USA). The cells were divided into three groups: the negative control group, which comprised cells treated with solvent DMSO; the blank control group, comprising cells treated with RPMI-1640 medium; and the experimental group. The cells of the experimental group were grown to 70-80% confluency in 96 well plates. AUY922 was then administered at a final concentration of 0.1, 0.5, 1, 5, 10, 50 or 100 nmol/l for treatment times of 24, 48 or 72 h. The blank group was treated with the addition of culture fluid. Subsequent to the reaction with the drugs for 24, 48 and 72 h, the cells were then treated with 10 ml/well MTT for 4 h at 37°C. The cells underwent absorbance readings at 570 nm using a 96-well microplate reader to obtain the OD values for the experimental (OD_{experimental}), blank control (OD_{blank}) and negative control (OD_{negative}) groups. The OD values were normalized to cells treated with 0 nmol/l of AUY922. The percentage of residual cell viability was calculated as follows:

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\text{Residual cell viability} (\%) = \frac{(\text{OD}_{\text{experimental}} - \text{OD}_{\text{blank}})}{(\text{OD}_{\text{negative}} - \text{OD}_{\text{blank}})} \times 100
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The assays were performed three times.

Cell migration assay. The motility capability of the cells was measured in vitro using Transwell chambers (Corning Inc., Corning, NY, USA). Four groups of ACHN cells (10⁵ per group, treated with 10, 50 or 100 nmol/l AUY922, or blank control medium) were seeded onto the upper wells with serum-free medium. Medium supplemented with 20% FBS was plated into the bottom wells to act as a chemoattractant. Subsequent to a 48-h period of incubation, the cells were fixed with methanol and stained with 1% crystal violet for 30 min at 37°C. The cells that remained on the upper surface of the membranes were removed, while those on the lower surface were counted. Images of the migrated cells were captured under a microscope (CKX41SF; Olympus Corporation, Tokyo, Japan).

Statistical analysis. Data analyses were performed using SPSS software, version 15.0 (SPSS, Inc., Chicago, IL, USA). The results were expressed as the mean ± standard deviation for continuous variables, and as the number and percentage for discrete variables. Differences in cell viability among groups was assessed using the t-test or two-way analysis of variance.
The Matrigel invasion assay revealed that the average number of cells migrating across the Matrigel-coated membrane in the negative control group was increased compared with the number in the AUY922-treated groups. The group of ACHN cells treated with 100 nM AUY992 yielded a decreased number of cells that migrated through the polycarbonate membrane compared with the group treated with 10 nM AUY992 (P<0.05) (Fig. 4). These results reveal that AUY992 performs a critical role in the inhibition of RCC cell migration.

Discussion

As the third most common genitourinary malignancy, RCC is not only a single oncological concept, but also a complex entity with respect to the etiology, histology and prognosis of the disease (13). Radical surgery remains the only potentially curative therapeutic option for RCC (14). Although an increased number of small renal masses are being detected, approximately one-third of patients present with metastatic disease (15). Metastatic RCC has demonstrated a poor response to chemotherapy and hormonal therapy. For patients with metastasized RCC that do not undergo complete resection, the five-year survival rate is <10% (7). Significant advances in the understanding of the biology of RCC tumors have led to the development of molecular therapies targeting the VEGF and mammalian target of rapamycin (mTOR) pathways, resulting in a significant improvement in the overall survival rate and quality of life experienced by patients (16).

Although its molecular pathophysiology is not completely understood, an improved understanding of the molecular pathogenesis of RCC, particularly in genomics, proteomics and metabolomics, has attracted increasing attention during the last decade (17). These previous studies have allowed for the development of novel therapeutics that target the VEGF and mTOR pathways (18). HSP90 is the most abundant cytosolic HSP and regulates the maturation and stability of various proteins that are crucial for multiple cell signaling processes (19). In a variety of cancers, including hepatocellular carcinoma, breast cancer, non-small-cell lung cancer and prostate cancer, HSP90 is overexpressed and may contribute to tumor cell survival by mediating the maturation and stability of a variety of client proteins, such as VEGF and hypoxia inducible factor-1α (HIF-1α) (20). In the present study, RT-qPCR and western blot analysis were performed to detect the expression of the mRNA and protein of HSP90 in the RCC ACHN and 786-O cell lines, and also in the immortalized normal human proximal tubule epithelial HK-2 cell line. Western blot analysis and RT-qPCR revealed that the mRNA and protein of HSP90 were each overexpressed in the two RCC cell lines compared with the HK-2 cell line (Fig. 2).

**Results**

**Expression of the mRNA and protein of HSP90 in the RCC cell lines.** RT-qPCR and western blot analysis were performed to detect the expression of the mRNA and protein of HSP90 in the RCC ACHN and 786-O and immortalized normal human proximal tubule epithelial HK-2 cell lines. As shown in Fig. 1, the 786-O and ACHN cell lines possessed higher HSP90 transcript levels compared with the HK-2 cell line. Similarly, the expression of the HSP90 protein was increased in the ACHN and 786-O cell lines compared with the HK-2 cell line (Fig. 2).

**Effects of the HSP90 inhibitor AUY922 on cell growth.** In order to determine whether HSP90 may be an effective therapeutic target for the treatment of RCC, the RCC cell lines were administered with AUY992 at a final concentrations of 0.1, 0.5, 1, 5, 10, 50 or 100 nmol/l for a treatment time of 24, 48 or 72 h, and the cell survival rate was measured using an MTT assay. As shown in Fig. 3, the MTT assay revealed that AUY922 significantly inhibited the proliferation of 786-O (Fig. 3A) and ACHN (Fig. 3B) cells in a time- and dose-dependent manner (P<0.05). The half-maximal growth inhibitory concentration (IC_{50}) of AUY992 in 786-O and ACHN cells after 72 h of treatment was 10.2 nM and 7.6 nM, respectively.

**Effect of the HSP90 inhibitor AUY922 on the migration of RCC cells.** The present data revealed that the HSP90 inhibitor AUY922 inhibited the migration of the RCC ACHN cell line.
anti-tumorigenic and anti-angiogenic properties \textit{in vitro} and \textit{in vivo} in animal models, which is partly due to the ability of these agents to inhibit the function of HIF (22,23).

HSP90 inhibitors have been considerably improved. AUY922 is a member of the isoxazole HSP90-inhibitor family and inhibits ATPase activity with an IC$_{50}$ of 30 nM (24). This molecule is a non-geldanamycin analog that possesses the potential to cause prolonged target inhibition and at present, has not been associated with the same degree of hepatotoxicity as its geldanamycin counterparts. AUY922 exerts its effects by binding to the ATPase domain of the HSP90 N-terminal, preventing HSP90 from performing chaperone functions. This leads to the proteasomal degradation of numerous relevant client proteins. Administration of the single agent AUY922 has demonstrated potent preclinical anticancer activity \textit{in vitro} and \textit{in vivo} against a range of histological cell types (25-28). In the present study, RCC cells were treated with 1, 2.5, 5, 10, 25, 50 or 100 nM of AUY922 for 24, 48 or 72 h, respectively. The cell survival rate was measured using an MTT assay in order to assess the inhibitory effect of AUY922 on RCC cells. As shown in Fig. 3, the MTT assay revealed that AUY922 significantly inhibited the proliferation of 786-O (Fig. 3A) and ACHN (Fig. 3B) cells in a time- and dose-dependent manner (P<0.05). The IC$_{50}$ of AUY922 in the 786-O and ACHN cells after 72 h was 10.2 nM and 7.6 nM, respectively. AUY922 is able to inhibit the proliferation of various tumor cells according to previous studies, therefore the present results validate HSP90 as an important target in RCC treatment. The presumptive tumor suppressor function of the HSP90 inhibitor AUY922 in human RCC was further investigated in the current study using a Transwell assay. The results revealed that the migration ability was significantly suppressed by AUY922. The migration assay revealed that the ACHN cells in the AUY922-treated groups were less likely to penetrate through the polycarbonate membrane compared with the negative control group. The data revealed this inhibition to be dose-dependent. Therefore, these results reveal a critical role of AUY922 in the inhibition of renal cancer cell migration.
In conclusion, the present results validate HSP90 as an important target in RCC and treatment with the novel HSP90 inhibitor AUY922 may provide a promising strategy for antitumor therapy in RCC.

References