Abstract. Zoledronic acid (ZA) is the current standard of care for the therapy of patients with bone metastasis or osteoporosis. ZA inhibits the prenylation of small guanine-5′-triphosphate (GTP)-binding proteins, such as Ras, and thus inhibit Ras signaling. The present study demonstrated that ZA inhibited cell proliferation and the pentose phosphate pathway (PPP) in bladder cancer cells. In addition, the expression of glucose-6-phosphate dehydrogenase (G6PD, the rate-limiting enzyme of the PPP) was found to be inhibited by ZA. Furthermore, the stability of TAp73, which activates the expression G6PD was decreased in zoledronic acid treated cells. Decreased levels of Ras-GTP and phosphorylated-extra-cellular signal-regulated kinase 1/2 were also observed following treatment with ZA. This may be due to the fact that activated Ras was reported to stabilize TAp73 inducing its accumulation. The inhibition of Ras activity by PT inhibitor II also significantly reduced the levels of TAp73 and G6PD and the PPP flux. Moreover, knockdown of TAp73, attenuated the PPP flux and eliminated the affection of ZA on the PPP flux. In conclusion, it was proposed that ZA can inhibit stability of TAp73 and attenuate the PPP via blocking Ras signaling in bladder cancer cells.

Introduction

Bladder cancer (BC) is a common malignancy of the urinary tract (1,2). Bone metastasis occurs in ~40% of patients with BC (3,4). Zoledronic acid (ZA) is effective in patients with bone metastases from bladder cancer (5-7). In addition to its bone-protective effects, ZA can prevent tumor progression (8). Due to its inhibition of the prenylation of small GTP-binding proteins, such as Ras, ZA inactivates Ras signaling and inhibits Ras-dependent cell proliferation (9-11).

P73 was identified to be a member of the p53 family and is frequently overexpressed in human tumors (12-14). Although p53 is firmly established to be a tumor suppressor, the role of p73 in human tumorigenesis is not well understood (15,16). The p73 gene contains two promoters and thus encodes the transcriptional domain-containing (TAp73) and the amino-deleted (ΔNp73) isoforms (17). TAp73 isoforms contain an amino-terminal transactivation domain and thus can activate the promoters of p53-target genes and induce apoptosis (18,19). ΔNp73 isoforms, which lack the transactivation domain of TAp73 protein and retain the DNA-binding and oligomerization domains, act as dominant-negative inhibitors for p53 family members by forming inactive hetero-oligomers or competing for p53-DNA-binding (20).

During rapid cell growth, adequate levels of intracellular nicotinamide adenine dinucleotide phosphate (NADPH), generated predominantly through the pentose phosphate pathway (PPP), are critical for cell survival. NADPH is required for DNA, protein and lipid biosynthesis, and is also required to generate sufficient material to support cancer cell proliferation (21-23). The tumor suppressor p53 was reported to bind to glucose-6-phosphate dehydrogenase (G6PD), the rate-limiting enzyme of the PPP, and inhibit its activity. Thus, p53 can suppress the glucose consumption and NADPH production of cells (24). Conversely, TAp73 was reported to activate the expression of G6PD and thus promote the PPP flux and NADPH production (25). The present study investigated the effects of ZA on the PPP flux and the proliferation of tumor cells.

Materials and methods

Reagents. ZA and cycloheximide (CHX) were purchased from Sigma-Aldrich (St. Louis, MO, USA). The mouse monoclonal anti-human Ras (05-1072; 1:1,000 dilution) and rabbit
monoclonal anti-human phospho-extracellular signal-regulated kinase (ERK) 1/2 (Thr202/Tyr204 and Thr185/Tyr187; 05-797R; 1:1,000 dilution) were obtained from Millipore (Bedford, MA, USA). The rabbit polyclonal anti-human G6PD (8866; 1:1,000 dilution) and β-actin (4967; 1:3,000 dilution) antibodies were obtained from Cell Signaling Technology Inc. (Beverly, MA, USA). Rabbit polyclonal anti-human p73 (ab37797; 1/800 dilution) was from Abcam (Cambridge, MA, USA). FPT inhibitor II was purchased from Santa Cruz Biotechnology Inc. (Dallas, TX, USA). The real-time PCR Master Mix kit (SYBR green PCR master mix) was purchased from Takara Bio Inc. (Dalian, China). To evaluate Rac activity [Ras-guanosine-5′-triphosphate (GTP) levels], affinity precipitation of active Rac was performed using a Ras activation assay according to the manufacturer's instructions (Millipore).

**Cell culture.** T24 human bladder cancer cells and 293T cells were purchased from the Type Culture Collection of the Chinese Academy of Sciences (Shanghai, China). Cells were cultured in RPMI-1640 medium (Invitrogen Life Technologies, Carlsbad, CA, USA) supplemented with 10% fetal bovine serum, penicillin (100 U/ml) and streptomycin (100 µg/ml) (all from Invitrogen Life Technologies) in a humidified atmosphere of 5% CO₂ at 37°C.

mRNA extract and reverse transcription-quantitative polymerase chain reaction (RT-qPCR). To examine the expression of G6PD and TAp73 mRNA in T24 cells, RT-qPCR was performed. Total RNA was extracted from the cell lines using the RNAiso reagent (Takara, Otsu, Japan). The first-strand cDNA was generated using TransScript First-Strand cDNA Synthesis SuperMix kit (Transgen, Beijing, China). Amplification and data acquisition were run on a real-time PCR system (ABI Prism 7500; Applied Biosystems, Foster City, CA, USA) for SYBR green PCR master mix. The quantity of the cDNA sample was 1 µl. RT-PCR was performed by 3 min incubation at 95°C and 40 amplification cycles (95°C for 10 sec; 56°C for 15 sec; and 72°C for 35 sec). β-actin served as a control. The primer sequences used were as follows: Forward: 5′-GTACCTGGCATTGCTGATGACT-3′ and reverse: 5′-CCGCTCATTGCGCAATGGTGAT-3′ for TAp73. The primer sequences used were as follows: forward: 5′-GTACCTGGCATTGCTGATGACT-3′ and reverse: 5′-CCGCTCATTGCGCAATGGTGAT-3′ for G6PD; and forward: 5′-GCA CCTACTTGGACCTCCTCCC-3′ and reverse: 5′-GCACTTGCTGAGCAAATTTGAA-3′ for TAp73.

**Metabolism assays.** Glucose consumption was measured in the cell lysates with Glucose Uptake Colorimetric Assay kit according to the manufacturer's instructions (Millipore). The production of NADPH was measured in the cell lysates with NAD⁺/NADH Quantification Colorimetric kit (BioVision, San Francisco, CA, USA). Following the manufacturer's instructions for these measures.

**Vector and cell transfection.** pCMV6-TAp73 (NM_005427) was obtained from OriGene (Rockville, MD, USA). For the transfection of the T24 cells, pCMV6 empty vector (OriGene, Rockville, MD, USA) and pCMV6-TAp73 vector were transfected into cells using Lipofectamine™ 2000 (Invitrogen Life Technologies). After 6 h, the medium was refreshed and cultured for 48 h. Then the cells were treated with ZA (200 µM).

**Viral production and infection.** Expression plasmids for small hairpin (sh)RNAs of TAp73 were made in pLKO.1 puro plasmids (Sigma-Aldrich). The targeted sequences were: 5′-GATTTCCGACATGAGCAGTCTT-3′ (sh1) and 5′-CAG GGGTTACAGAGCATT-3′ (sh2). A negative control vector containing scrambled shRNA was also obtained from Sigma-Aldrich. The plasmids were prepared with a plasmid maxi kit and transfected in 293T cells (Invitrogen Life Technologies) with the Lipofectamine™ 2000 to produce lentiviral particles. Then, the T24 cells were infected with the lentiviral particles for 24 h. Cells were then selected with 2 µg/ml puromycin (Sigma-Aldrich) for 72 h.

**Western blot analysis.** Cells were lysed in a radioimmunoprecipitation assay buffer containing and complete Protease/Phosphatase Inhibitor Cocktail (Cell Signaling Technology, Inc.). The lysate was centrifuged at 11,500 x g for 15 min at 4°C and the supernatant was collected. Protein concentration was determined using a Bradford protein assay kit (Bio-Rad, Hercules, CA, USA). The proteins were separated by 10% SDS-PAGE gel (RSBM, Taiyuan, China) and then transferred onto a polyvinylidene difluoride membrane (Millipore). The membranes were blocked with 10% milk in Tris-buffered saline with Tween-20 (TBST) for 1 h and then were incubated with antibodies overnight at 4°C. After washing with TBST three times, the membranes were probed with horseradish peroxidase-conjugated secondary antibodies in TBST for 1 h at room temperature, then washed with PBST three times. The immunobinding signals were detected by a chemiluminescence kit (Millipore).

**Cell proliferation analysis.** Cell proliferation was examined by using a Cell Counting Kit-8 (CCK-8; Dojindo, Kumamoto, Japan). Cells were plated and treated with ZA (200 µM) in 96-well plates at 2,000 cells per well and cultured in growth medium for 20 h. After 20 h, CCK-8 (10 µl) was added to each well containing 100 µl RPMI-1640 medium. Then the plate was incubated for 2 h at 37°C. Absorbance was measured at 450 nm using a microplate reader (SpectraMax M5 Microplate Reader; BioTek, Winooski, VT, USA).

**Statistical analysis.** Data were analyzed using Student's t-test. Graphpad 6.01 Prism software (GraphPad, Inc., La Jolla, CA, USA) was used for statistical analyses. P<0.05 was considered to indicate a statistically significant difference.

**Results.**

ZA inhibits the proliferation of bladder cancer cells and attenuates the PPP. To confirm the effects of ZA on the proliferation of bladder cancer cells, T24 human bladder cancer cell lines were treated with ZA (200 µM) for 20 h. As shown in Fig. 1A, ZA significantly reduced the proliferative activity of T24 cells. The PPP and NADPH produced in the PPP are required for rapid cancer cell growth. Thus, it was examined whether ZA affects the PPP flux in bladder cancer cell lines. It was demonstrated that glucose consumption and NADPH were inhibited.
by treatment with ZA in T24 cells (Fig. 1B). Furthermore, G6PD, the rate-limiting enzyme of the PPP, was found to be downregulated at the mRNA and protein level (Fig. 1C).

**ZA decreases the stability of TAp73.** TAp73 was reported to activate the expression of G6PD and thus promote the PPP flux and NADPH production (25). As shown in Fig. 2A, it was demonstrated that the levels of TAp73 decreased following treatment with ZA in T24 cells (Fig. 2A). The expression of TAp73 mRNA was then examined by RT-qPCR. The data show that there were no significant changes in TAp73 mRNA levels following treatment with ZA (Fig. 2B). It was hypothesized that the regulation of TAp73 levels by ZA may be the result of protein stability regulation. The TAp73 stability in T24 cells was then determined by treatment with CHX. As shown in Fig. 2C, following pre-treatment with ZA, the stability of TAp73 decreased.

Stability of TAp73 regulated by ZA may depend on the inhibition of the activity of Ras. Activated Ras was reported
to stabilize TAp73 inducing its accumulation (26). The activity of Ras in T24 cells treated with ZA was then examined. GTP-bound Ras (Ras-GTP), a marker of Ras activation, as well as the quantity of ERK1/2 phosphorylation were decreased in T24 cells following treatment with ZA for 20 h (Fig. 3A). FPT inhibitor II, exhibited inhibitory effects of Ras farnesylation and activity, was substituted for ZA to treat T24 cells. As shown in Fig. 3B, FPT inhibitor II significantly reduced the levels of Ras-GTP and TAp73. In addition, FPT inhibitor II also inhibited the expression of G6PD (Fig. 3C), glucose consumption and NADPH production (Fig. 3D). These results support the role of ZA in the regulation of the stability of TAp73 via inhibiting the activity of Ras.
ZA regulates PPP through the regulation of TAp73. In order to further investigate the function of ZA in the regulation of the PPP, following transfection with TAp73 for 48 h (Fig. 4A), the inhibitory effect on the PPP by ZA, as well as the glucose consumption and NADPH production was attenuated by TAp73 overexpression (Fig. 4B). Furthermore, the expression of G6PD was also upregulated (Fig. 4A). These results further support that ZA inhibits the expression of G6PD and then attenuates the PPP through the downregulation of TAp73.

Knockdown of TAp73 inhibits the PPP of T24 cells independent of Ras activity or ZA treatment. shRNA-mediated TAp73 knockdown was then performed in T24 cells and the PPP flux was examined. It was demonstrated that the expression of G6PD (Fig. 5A) and the PPP flux (Fig. 5B) were inhibited following knockdown of TAp73 in T24 cells. Moreover, FPT inhibitor II and ZA did not affect the PPP flux after the knockdown of TAp73 (Fig. 5B). Similarly, the proliferation of T24 cells was not significantly changed by the FPT inhibitor II or ZA after the knockdown of TAp73 (Fig. 5C). These results confirm the role of TAp73 in the inhibition of the PPP flux by ZA.

Discussion

In the present study, it was demonstrated that ZA inhibited the proliferation of bladder cancer cells and the PPP. Moreover, ZA was found to decrease the stability of TAp73, which activates the expression of G6PD (the rate-limiting enzyme of PPP). Decreased levels of Ras-GTP and p-ERK1/2 were also found to be associated with the treatment with ZA. Furthermore, the inhibition of Ras activity by PT Inhibitor II significantly reduced the levels of TAp73, G6PD, glucose consumption and NADPH production.

ZA is effective in patients with bone metastases from bladder cancer. In addition to its bone-protective effects, ZA can prevent tumor progression (27). ZA was reported to inhibit the prenylation of small GTP-binding proteins, such as Ras. Thus, ZA was hypothesized to inactivate Ras signaling and inhibit Ras-dependent cell proliferation (9,28). Previous studies have shown that ZA inhibited the activity of Ras and hence inhibit the expression of hypoxia-inducible factor 1-α (HIF1A) (29). NADPH is required for DNA, protein and lipid biosynthesis during the rapid cell growth of cancer cells. The PPP is the predominant source of NADPH and it is often enhanced during cell cycle progression of cancer cells (23,30).

In this study, the PPP flux was found to be inhibited by ZA. In addition, the expression of G6PD, the rate-limiting enzyme of the PPP, was also found to be inhibited by ZA. Activated Ras was reported to stabilize TAp73 protein, ZA was shown to decrease the levels of Ras-GTP and p-ERK1/2 with treatment with ZA. FPT inhibitor II is the inhibitor of Ras farnesylation and its activity. Similar to ZA, FPT inhibitor II significantly reduced the levels of Ras-GTP, G6PD, TAp73 and the PPP flux. It was also demonstrated that knockdown of TAp73 resulted in an inhibition of the PPP flux. Moreover, ZA could not regulate the PPP flux in TAp73 knockdown cells. These results implied that ZA inhibits the PPP flux and the expression of G6PD via blocking Ras signaling and attenuating the stability of TAp73 in bladder cancer cells. The findings of the present study revealed a novel mechanism for ZA to regulate the PPP. Besides the bone-protective effects, ZA restrained the PPP flux of cancer
cells and inhibited their proliferation. The mechanistic target of ZA was TAp73, and thus, TAp73-overexpressing tumors may show enhanced sensitivity to ZA. Due to its targeting of TAp73, ZA may potentially be utilized in cancer therapies.

References
15. Tomkova K, Belkhiri A, El-Rifai W and Zaika AI: p73 isoforms may show enhanced sensitivity to ZA. Due to its targeting of TAp73, ZA may potentially be utilized in cancer therapies.