Involvement of the mevalonate pathway in the antiproliferative effect of zoledronate on ACHN renal cell carcinoma cells

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\textbf{Abstract}. Renal cell carcinoma (RCC) has been shown to be resistant to chemotherapy and radiotherapy. In order to examine the potential of zoledronate (ZOL), a bisphosphonate, as an anticancer agent, we investigated the effects of ZOL on RCC cells and the involvement of the mevalonate pathway in antiproliferative effects, as well as the effects of ZOL administration on mice inoculated with RCC. ACHN cells were used and cell viability was measured via intracellular reductase activity. Chromatin condensation was detected by Hoechst 33342 staining. Proteins were detected by western blot analysis. Tumor volume was measured bidimensionally in mice inoculated with ACHN cells after vehicle or ZOL subcutaneous administration. ZOL exhibited antiproliferative effects with an IC\textsubscript{50} value of 2.29±0.53 \textmu M in ACHN cells and chromatin condensation was observed when treated with ZOL. Farnesol (FOH) and geranylgeraniol (GGOH), precursors of farnesyl pyrophosphate and geranylgeranyl pyrophosphate, exhibited potency to rescue cells treated with ZOL. Additionally, Ras and RhoA proteins located in the membrane fraction decreased when treated with ZOL and recovered by FOH or GGOH treatment, suggesting that ZOL inhibited the mevalonate pathway, thereby suppressing the translocation of prenylated Ras and RhoA proteins to membrane fractions. An in vivo study showed the inhibitory potential of ZOL on tumor growth in mice without changes in body weight. Our study showed that ZOL could be useful as an anticancer agent for the treatment of RCC, and the mevalonate pathway could be an efficient target for novel therapeutic agents against RCC.

\textbf{Introduction}

Renal cell carcinoma (RCC) can be resistant to chemotherapy and radiotherapy, making nephrectomy the most effective treatment (1). Although nephrectomy can be expected to cure patients without metastatic involvement, the 5-year survival rate of those with metastasis is <10%. In addition, recent advances in drug development have allowed us to use novel agents for RCC targeting the cancer-specific pathway, such as bevacizumab, sorafenib and sunitinib (2-4). Although they prolonged overall survival in patients with advanced RCC, the effect was still insufficient; thus, it is necessary to discover novel chemotherapy for RCC. There have been studies by us and other groups, showing the possibility of other targets to treat RCC (5-7).

Bisphosphonates have been widely used for the treatment of bone disease with potent antiresorptive activity, especially nitrogen-containing bisphosphonates, including alendronate (ALN), risedronate (RISE), and zoledronate (ZOL), which have been characterized to have strong inhibitory effects on the mevalonate pathway, thereby inhibiting the prenylation of Ras and Rho inducing cell apoptosis (8,9). ZOL is the most potent nitrogen-containing bisphosphonate and it is widely used for bone diseases caused by metastasis of breast, prostate, bladder, and renal cancer (10,11). In addition, ZOL has been shown to have cytotoxic effects on cancer cells, including multiple myeloma and pancreatic, colon, prostate, and breast cancer (12-18), which anticipate direct effects on cancer cells (19). In addition, ZOL has been suggested to have cytotoxic effects mediated by caspase-dependent apoptotic cell death in RCC (20,21); however, limited information is available in regard to the mevalonate pathway. ZOL potency in vivo is also unknown in RCC. The aims of this study were to investigate the cytotoxic effects of ZOL in ACHN cells, derived from RCC, and to clarify the importance of the mevalonate pathway and the potency of ZOL in mice inoculated with ACHN cells.

\textbf{Materials and methods}

\textbf{Materials}. Alendronate (ALN) and Risedronate (RISE) were purchased from LKT Laboratories (St. Paul, MN), and zoledronate (ZOL) was purchased from Toronto Research
Chemicals (Ontario, Canada). Trans, trans-farnesol (FOH), geranylgeraniol (GGOH) and cholesterol (CHO) were obtained from Sigma-Aldrich (St. Louis, MO). All other agents were of reagent grade.

**Cell culture.** ACHN, 786-O and Caki-2 cells were obtained from the Cell Resource Center for Biomedical Research Institute of Development, Aging and Cancer Tohoku University (Sendai, Japan), Summit Pharmaceuticals International (Tokyo, Japan), and DS Pharma Biomedical (Osaka, Japan), respectively, and were used as human renal carcinoma cell models. ACHN was cultured in DMEM (Wako Pure Chemical Industries, Osaka, Japan), and 786-O and Caki-2 were maintained in a culture medium consisting of RPMI-1640 (Nacalai Tesque, Kyoto, Japan). Media were supplemented with 10% heat-inactivated FBS (Invitrogen Life Technologies, Carlsbad, CA) and penicillin (50 U/ml) - streptomycin (50 µg/ml) (Nacalai Tesque) in an atmosphere of 5% CO2-95% air at 37˚C, and subcultured every 3 or 4 days with 0.05% trypsin / 0.2 mM EDTA (Nacalai Tesque). The number of passages for ACHN, 786-O, and Caki-2 cells used in this study was 37-50, 15-21 and 63-69, respectively.

**Cell viability.** Cell viability was measured with Cell QuantiBlue™ (BioAssay Systems, Hayward, CA). Briefly, cells were seeded onto 96-well plates (Asahi Glass, Tokyo, Japan) at a density of 2x10^3 (786-O) or 3x10^3 (Caki-2 and ACHN) cells/well and incubated for 24 h, and then treated with various concentrations of ALN, RISE or ZOL for 72 h. The assay utilizes the conversion of Alamar Blue reagent to resorufin fluorescence by metabolically active cells and resorufin was measured with a CytoFluor™ Series 4000 Fluorescence Multi-Well Plate Reader (PerSeptive Biosystems, Framingham, MA) at an excitation wavelength of 530 nm/emission 580 nm. IC50 were calculated according to the sigmoid inhibitory effect model, E = IC50γ / (IC50 + C). E represents the surviving fraction (% of control), and C and γ represent the drug concentration in the medium and the Hill coefficient, respectively. In addition, effects of FOH, GGOH and CHO on antiproliferation induced by 3 µM ZOL were examined by treating together with 5 µM FOH, 5 µM GGOH or 250 µM CHO.

**Detection of chromatin condensation.** For nuclei staining, cells were incubated for 24 h after seeding, and then treated with 30 µM ZOL for 24, 48 or 72 h. The nuclear chromatin of trypsinized cells was then stained with 74 µg/ml Hoechst 33342 (Nacalai Tesque) for 15 min at room temperature, and they were observed with a bright field fluorescent microscope (Vanox; Olympus, Tokyo, Japan) under UV excitation. The percentage of chromatin-condensed cells was determined by counting >450 cells in each experiment.

**Western blotting.** Cells were seeded at a density of 5x10^3 cells/dish and incubated for 24 h. They were then treated with ZOL alone or together with FOH or GGOH for 72 h, harvested by scraping and rinsed twice with ice-cold phosphate-buffered saline (PBS). The cell pellets were suspended in lysis buffer 1 (50 mM HEPES (pH 7.4), 50 mM NaCl, 1 mM MgCl2, 2 mM EDTA, 1 mM phenylmethylsulfonyl fluoride, 10 µg/ml leupeptin, 1 mM Na3VO4, 5 mM NaF, 1 mM dithiothreitol, 0.1% (w/v) Triton X-100) with a sonicator, then left to stand on ice for 10 min, followed by ultracentrifugation at 24,000 x g for 15 min with a Beckman Coulter Optima XL-100K ultracentrifuge (Beckman Coulter, Tokyo, Japan). The supernatants were subjected to western blot analysis as cytosol samples. Lysis buffer 2 (50 mM HEPES (pH 7.4), 50 mM NaCl, 1 mM MgCl2, 2 mM EDTA, 1 mM phenylmethylsulfonyl fluoride, 10 µg/ml leupeptin, 1 mM Na3VO4, 5 mM NaF, 1 mM dithiothreitol, 1% (w/v) Triton X-100) were added to the pellets followed by sonication, then left to stand on ice for 10 min, followed by ultracentrifugation at 24,000 x g for 15 min. The supernatants were subjected to Western blot analysis as membrane samples.

The samples were mixed with the same volume of Laemmli sample buffer (Bio-Rad Laboratories, Hercules, CA) including 5% β-mercaptoethanol, and then boiled for 5 min. Protein (7 µg) was fractionated by 12.5% SDS-PAGE, transferred to PVDF membrane (ATTO, Tokyo, Japan) and blocked with Tris-buffered saline, including ECL Advance™ blocking reagent (GE Healthcare, Buckinghamshire, UK). Seven micrograms of protein were analyzed using mouse anti-Ras (BD Biosciences, San Jose, CA), mouse anti-RhoA, and mouse anti-β-actin antibodies (Santa Cruz Biotechnology, Santa Cruz, CA). Horseradish peroxidase-conjugated secondary antibody against mouse IgG (GE Healthcare) was used at 1:25000 dilution with the blocking buffer, and the signals were visualized using ECL Advance detection reagents (GE Healthcare) according to the manufacturer’s procedure. Protein concentrations were determined with the BCA Protein Assay Reagent kit (ThermoFisher Scientific, Waltham, MA), and bovine serum albumin was used for the standard curve.

**Animal model.** Animal experiments were performed in accordance with the guidelines of the Institutional Animal Use Committee of Mukogawa Women’s University, and the procedure was approved by the Committee. Balb/c male mice (4 weeks old) were purchased from Japan SLC (Shizuoka, Japan), and subcutaneously inoculated in the back with ACHN cells (1x106 cells/100 µl in PBS). Ten days after inoculation, ZOL was subcutaneously administered in the abdomen at a dose of 0.5 mg/kg/100 µl in PBS once a week for five weeks. Tumor size was measured bidimensionally and volume was calculated by the formula (length x width^2)/2. Mouse weight was also monitored.

**Statistics.** The data are expressed as the mean ± SD. Statistical analysis was performed with Student’s t-test for comparison of two groups and non-repeated one-way ANOVA followed by Dunnett’s post hoc test for multiple comparison. P<0.05 was considered significant.

**Results**

**Effects of bisphosphonates (ALN, RISE and ZOL) on RCC cells.** Fig. 1 shows the effects of ALN, RISE or ZOL on the proliferation in three RCC cell lines, and Table 1 shows the IC50 values. All the bisphosphonates inhibited cell growth in a concentration-dependent manner, with IC50 values of 19-156 µM for ALN, 99-329 µM for RISE, and 2-18 µM for
ZOL in three RCC cell lines. ZOL exhibited the strongest effect on the proliferation in three RCC, in ACHN cells with IC_{50} value of 2.29±0.53 µM. When cells were exposed to 30 µM ZOL, chromatin condensation, a typical morphological change in apoptosis, was found after staining with Hoechst 33342 (Fig. 2A). The percentages of cells with chromatin condensation significantly increased by ZOL treatment time-dependently (Fig. 2B).

Table I. IC_{50} values for bisphosphonates in ACHN, 786-O and Caki-2 cells.

<table>
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<tr>
<th></th>
<th>ACHN</th>
<th>786-O</th>
<th>Caki-2</th>
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<tbody>
<tr>
<td>ALN (µM)</td>
<td>18.8±5.95</td>
<td>35.2±6.14</td>
<td>156±27.3</td>
</tr>
<tr>
<td>RISE (µM)</td>
<td>99.1±7.68</td>
<td>130±22.9</td>
<td>329±8.93</td>
</tr>
<tr>
<td>ZOL (µM)</td>
<td>2.29±0.53</td>
<td>8.43±2.29</td>
<td>18.3±3.85</td>
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Cells were precultured for 24 h at a density of 2x10^3 (786-O) or 3x10^3 (Caki-2 and ACHN) cells/well in 96-well plates and treated with ALN, RISE or ZOL for 72 h. Cell viability was determined by fluorescent assay and IC_{50} values were calculated. Data are the mean ± SD of three to five experiments.

Effects of FOH, GGOH and CHO on antiproliferation induced by zoledronate. To clarify the role of the mevalonate pathway in the antiproliferative effects of ZOL, we investigated the effects of FOH and GGOH, precursors of farnesyl pyrophosphate and geranylgeranyl pyrophosphate, on cell survival. As a result, FOH and GGOH attenuated the antiproliferative effect of ZOL in RCC cells.
Next, we investigated the localization of Ras and RhoA proteins when cells were treated with ZOL. Their inactive forms are primarily localized in the cytosol, which can be translocated to the membrane as active forms after prenylation with isoprenoid provided by farnesyl pyrophosphate or geranylgeranyl pyrophosphate. Fig. 4A shows that Ras and RhoA proteins located at the membrane decreased with ZOL treatment in a concentration-dependent manner while they increased in cytosol. In addition, treatment with FOH and GGOH recovered proteins in the membrane (Fig. 4B).

Effects of zoledronate on proliferation of ACHN inoculated in mice. To clarify the effects of ZOL in controlling tumor growth, we investigated the effects of ZOL administration on mice inoculated with ACHN cells. ZOL exhibited inhibitory effects on tumor growth in the ACHN xenograft model (Fig. 5A), while the body weights of mice were not changed compared with the vehicle administration group (Fig. 5B).

Discussion

Nephrectomy is still a common treatment for RCC, known to be resistant to chemotherapy and radiotherapy, although novel agents, including bevacizumab, sorafenib and sunitinib, prolonged the survival rate (1-4). ZOL has been shown to have cytotoxic effects on cancer cells, including multiple myeloma and pancreatic, colon, prostate and breast cancer (12-18). In addition, ZOL has been suggested to have cytotoxic effects on RCC (19,20).

We clearly showed that three bisphosphonates, ALN, RISE and ZOL, showed antiproliferative effects on three RCC cell
lines (Fig. 1, Table 1). However, ZOL showed the inhibitory effects most potently of the three drugs, and ACHN cells were the most sensitive to each drug in three cell lines. Therefore, we chose the most effective combination, ZOL and ACHN, for further investigations.

ZOL showed antiproliferative effects on ACHN cells with an IC<sub>50</sub> value of 2.29±0.53 µM (Fig. 1), which was comparable or somewhat stronger than the data by Ullén et al., who reported that 69% viable cells were found at 12.5 µM ZOL for 72 h in ACHN cells (21). Skerjanec et al. reported that the maximum concentration of ZOL was 300-400 ng/ml (ca. 1-1.3 µM) after infusion of 4 mg ZOL (22). Although renal distribution was not available in humans, it is likely to reach the concentrations in the clinical setting.

Next, we examined the involvement of apoptosis in the effects of ZOL. Chromatin condensation was found in ACHN cells treated with ZOL in a time-dependent manner (Fig. 2), suggesting that ZOL induced apoptosis at least in part. Pandha et al. and Ullén et al. reported that ZOL induced apoptotic cell death in a caspase-dependent manner; however, they did not examine the involvement of the mevalonate pathway (20,21), although there have been reports suggesting that the mevalonate pathway is important for the mechanisms of cell death induced by bisphosphonates in multiple myeloma, prostate cancer and breast cancer (23-25).

The mevalonate pathway plays an important role in synthesizing farnesyl pyrophosphate, geranylgeranyl pyrophosphate and cholesterol (8,9,19,26). In particular, farnesyl pyrophosphate and geranylgeranyl pyrophosphate provide isoprenoid in the prenylation of Ras and Rho; thereby, they are translocated to the membrane as active forms, regulating cell proliferation and cell death (19). To clarify the contribution of the inhibition of the mevalonate pathway in ZOL-induced antiproliferative effects in RCC, effects of FOH and GGOH were examined in ACHN cells treated with ZOL. As a result, FOH and GGOH significantly recovered cell survival (Fig. 3). It has been also reported that the depletion of cholesterol induced cell apoptosis in prostate cancer and embryonic fibroblast cell (27,28), but we did not observe a change of cell survival with cholesterol treatment. Accordingly, it was suggested that the antiproliferative effects of ZOL were due to inhibition of the synthesis of farnesyl pyrophosphate, while cholesterol depletion was not involved in antiproliferation.

Also, we found a decrease of Ras and RhoA proteins localized in the membrane fraction in ACHN cells treated with ZOL in a concentration-dependent manner, and FOH and GGOH recovered the amount of Ras and RhoA proteins at membranes (Fig. 4). Nogawa et al. reported that the ZOL-induced growth inhibitory effect was circumvented by the addition of GGOH (Fig. 4). Nogawa et al. reported that the ZOL-induced growth inhibition, suggesting that geranylgeranylation was more important than farnesylation (18). We observed a stronger effect of the addition of GGOH on the antiproliferative effects of ZOL; however, our data showed that the ZOL-induced antiproliferative effect was dependent on the amount of Ras protein localized at membranes. The reason for this discrepancy was unclear but at least the Ras signal was important for the proliferation of RCC, although it was not in prostate cancer cells. Collectively, inhibition of the mevalonate pathway by ZOL induced the decreased prenylation of Ras and Rho proteins, which led to decreased active forms of Ras and Rho located at the membrane.

Next, we evaluated the antitumor effects of ZOL administration in mice inoculated with ACHN cells. Tumor growth was significantly inhibited by the administration of ZOL, while the body weights of mice did not change (Fig. 5), suggesting that marked adverse effects did not occur in this condition. Thus, efficient therapeutic effects of ZOL could be expected for the treatment of RCC.

In conclusion, it was demonstrated that ZOL inhibited the proliferation of ACHN cells and suggested that ZOL inhibited the mevalonate pathway, thereby suppressing the translocation of prenylated Ras and Rho proteins to membrane fractions. ZOL was shown to inhibit tumor growth in mice inoculated with ACHN cells in vivo; therefore, our study showed that ZOL could be useful as an anticancer agent for the treatment of RCC, and the mevalonate pathway could be an efficient target for novel therapeutic agents against RCC.

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References


