Cancer Therapy: Preclinical

Targeting of Bone-Derived Insulin-Like Growth Factor-II by a Human Neutralizing Antibody Suppresses the Growth of Prostate Cancer Cells in a Human Bone Environment

Taichi Kimura¹,³, Takeshi Kuwata¹, Satoshi Ashimine¹,³, Manabu Yamazaki¹, Chisako Yamauchi¹, Kanji Nagai², Akashi Ikehara³, Yang Feng⁴, Dimiter S. Dimitrov⁴, Seiichi Saito³, and Atsushi Ochiai¹

Abstract

Purpose: Advanced prostate cancer frequently involves the bone, where the insulin-like growth factor (IGF)-II is abundant. However, the importance of IGF-II in bone metastasis from prostate cancer is uncertain. The present study was aimed at examining the therapeutic importance of targeting IGF-II in bone metastases from prostate cancer.

Experimental Design: We investigated whether inhibiting IGF-II using a human neutralizing antibody (m610) suppresses the growth of prostate cancer cells in a human bone environment. Human MDA PCa 2b prostate cancer cells were inoculated into human adult bone implanted into mammary fat pad of nonobese diabetic/severe combined immunodeficient mice or inoculated into mammary fat pad of the mice without human bone implantation. The mice were treated with m610 or a control antibody (m102.4) once weekly for 4 weeks immediately after inoculation with MDA PCa 2b cells.

Results: Histomorphologic examination indicated that m610 treatment significantly decreased the MDA PCa 2b tumor area in the human bone compared with the control. Ki-67 immunostaining revealed that the percentage of proliferating cancer cells in the m610-treated bone tumor sections was significantly lower than that in the control. m610 had no effect on MDA PCa 2b tumor growth in the absence of implanted human bone. m610 prevented the in vitro IGF-II–induced proliferation of MDA PCa 2b cells.

Conclusions: Our results indicate that IGF-II plays an important role in the prostate cancer cell growth in human bone, suggesting that targeting it by neutralizing antibodies offers a new therapeutic strategy for bone metastasis from prostate cancer. Clin Cancer Res; 16(1); 121–9. ©2010 AACR.
functions: both IGF-I and IGF-II activate the IGF-I receptor (IGF-IR), whereas IGF-II also can activate the insulin receptor. Moreover, IGF-II is the most abundant growth factor in human bone, and the IGF-II level is nine times higher than IGF-I level. In contrast to human bone, IGF-I is three times higher than IGF-II in mouse bone (11). Interestingly, in our bone-transplanted model, human prostate cancer cells metastasized to implanted human bone but not to mouse bone (5). These findings suggest that IGF-II is more supportive of human prostate cancer than IGF-I in the human bone environment, which also supports the concept that IGF-II might be a potentially useful target for the treatment of bone metastases from prostate cancer.

We recently developed a fully human monoclonal antibody (mAb; m610), which is highly specific for IGF-II but not for either IGF-I or insulin (12). To explore the mechanism of IGF-II in the growth of prostate cancer cells in human bone and the therapeutic potential of inhibiting IGF-II by a neutralizing antibody, we used our HAB implanted model and a fully human mAb, m610, specific for IGF-II. Our results indicate that IGF-II plays an important role in the prostate cancer cell growth in human bone and that the human mAb m610 and other anti–IGF-II mAbs could be promising candidate therapeutics for treatment of prostate cancer.

Materials and Methods

Cell line and cell culture. An androgen-responsive human prostate cancer cell line, MDA PCa 2b, was purchased from the American Tissue Culture Collection. The cells were maintained in BRFF-HPC1 medium (Athena Enzyme System) containing 20% fetal bovine serum and were incubated at 37°C in a humidified atmosphere of 5% CO₂. Animal care. Male nonobese diabetic/severe combined immunodeficient mice were obtained from CLEA Japan.

The mice were maintained under specific pathogen-free and temperature-controlled air conditions according to institutional guidelines. The mice used in the experiments were all 6 to 8 weeks old.

Implantation of HAB and induction of bone tumors. After obtaining the informed consent of the patients, HAB was obtained from lung or esophagus cancer patients (age range, 44-82 years; mean, 66.5 years) who had undergone a pulmonary lobectomy or esophagectomy in the Division of Thoracic Oncology, National Cancer Center Hospital East. The implantation of HAB into nonobese diabetic/severe combined immunodeficient mice was done as described previously, with some modifications (5). Briefly, morselized cancellous bone (200 mm³) from HAB was implanted into mammary fat pad (MFP) of the mice through a small skin incision in the right flank. In a preliminary examination, graft survival was ascertained by histologic examination and the detection of human IgG in the mouse sera using Western blotting at 4 weeks after the implantation. At 4 weeks after bone implantation, single-cell suspensions (8 × 10⁶ cells/100 μL serum-free medium) of MDA PCa 2b cells were directly injected into the marrow spaces of the implanted HAB.

Induction of MFP tumors. The single-cell suspensions prepared as mentioned above were injected into the MFP of the age-matched nonobese diabetic/severe combined immunodeficient mice (10-12 weeks old) through a small skin incision in the right flank.

Human antibodies. An anti–IGF-II fully human mAb (IgG1 m610), which binds to IGF-II but not to IGF-I and insulin, was identified by screening of a large antibody library in the authors' laboratory (National Cancer Institute-Frederick, NIH; ref. 12). To obtain sufficient quantities of m610, m610 was produced from a permanent cell line developed by transfection of CHO-K1 cells and selection of the highest producing clone (13). A control human antibody (IgG1 m102.4), which was produced from a permanent CHO cell line under the same conditions as m610 (14), was chosen to be as closed in sequence as possible to m610 but with entirely different specificity. Compared with m610, m102.4 has identical Fc, hinge and CH1, very similar CL, and similar frameworks of the variable domains but different complementarity-determining regions to ensure specificity to Hendra and Nipah virus.

Protocol for m610 treatment. In the model involving the induction of bone tumor in HAB (HAB model), m610 (1 or 10 mg/kg) or a control antibody (10 mg/kg; n = 8 per group) was intraperitoneally administered to the mice at weekly intervals for 4 weeks starting immediately after the inoculation of the MDA PCa 2b cells into the HAB. In the model involving the induction of tumors in MFP (MFP model), m610 (10 mg/kg) or control antibody (10 mg/kg; n = 5 or 4 per group, respectively) was administered to the mice at weekly intervals for 4 weeks after the inoculation of the cells into the MFP. At 4 weeks after the start of the m610 treatment, the mice were sacrificed and
the HAB tissues and mouse organs (lung, liver, spleen, kidney, and lumbar vertebrae) in the HAB model and the MFP tissues in the MFP model were harvested. The body weight of mouse was measured at the time of HAB implantation, the start of the m610 treatment, and the time of sacrifice.

**Histomorphometric examination.** Harvested specimens were fixed in 10% neutral buffered formalin and embedded in paraffin. The bone specimens were decalcified with EDTA solution (Wako) before paraffin embedding. Two-μm-thick sections were prepared from each specimen. Then, we analyzed the tumor burden in the HAB and the MFP by histomorphometrically examining sections stained for PSA, as described previously (6). Briefly, tissue sections of the HAB and MFP specimens were cut at three levels far enough apart (>200 μm) to avoid replicating the sampling of a single surface event; immunostaining for PSA was done using the EnVision Plus System HRP kit (Dako) according to the manufacturer’s protocol. An anti-PSA polyclonal antibody (1:800; Dako) was used as a primary antibody. Antigen retrieval was done. The total tissue area and the total tumor area immunostained by the anti-PSA antibody were determined using image analysis software (Image J version 1.38v; NIH). The results were expressed as the absolute tumor area (mm²) and the tumor area as a percentage of the tissue section area (%).

**Immunohistologic examination.** The immunostaining procedure used for IGF-IR, insulin receptor, Ki-67, and cleaved caspase-3 was the same as described above. However, microwaving (95°C, 20 min) in citrate buffer (pH 6.0) or in Target Retrieval Solution High pH Buffer (Dako) was required for Ki-67 antigen retrieval or cleaved caspase-3 antigen retrieval, respectively. An anti-IGF-IR mAb (1:100; Chemicon), an anti–insulin receptor mAb (1:100; Chemicon), an anti-Ki-67 mAb (1:50; Dako), or an anti–cleaved caspase-3 polyclonal antibody (1:200; Cell Signaling Technology) was used as a primary antibody.

To evaluate the proliferative and apoptotic statuses of the tumor cells, we counted the number of Ki-67–positive and cleaved caspase-3–positive cancer cells and the total cancer cells in three high-power fields of the most strongly stained tumor areas and then calculated the percentages of these immunostaining positive cancer cells to the total cancer cells. The necrotic area was not included in the evaluation of apoptosis. The data from samples with the presence of cancer cells were presented.

**Determination of serum PSA levels.** Blood samples were obtained from the mice before sacrifice. The serum PSA levels were determined using a chemiluminescent enzyme immunoassay kit (Hybritech-PSA; Beckman Coulter) according to the manufacturer’s protocol.

**Proliferation assay.** MDA PCa 2b cells (5 × 10⁶) seeded on a 6-cm dish were cultured in complete growth medium for 48 h. The cells were rinsed and cultured in serum-free F12K for 6 h. Then, the cells were treated with the serum-free medium alone, with IGF-I (100 ng/mL; R&D systems), or with IGF-II (100 ng/mL) for 15 min. Immunoprecipitation and Western blot analysis procedures were done as described previously (15). Two hundred micrograms of protein from each cell lysate were, respectively, immunoprecipitated with an anti–IGF-IR polyclonal antibody (2 μg; clone C-20; Santa Cruz Biotechnology) or with an anti–insulin receptor polyclonal antibody (2 μg; clone C-19; Santa Cruz Biotechnology) for 2 h at 4°C, and each immunoprecipitate was collected using protein G Sepharose. Each sample (derived from 20 μg total cell lysate) was fractionated by 7.5% SDS-PAGE and transferred to polyvinylidene fluoride membrane (Millipore). The membranes were probed with an anti-phosphotyrosine mAb (1:2,000; clone 4G10; Upstate Biotechnology) and visualized by an enhanced chemiluminescence (Amersham Biosciences). After stripping the membrane, total IGF-IR or insulin receptor in the immunoprecipitates was detected with C-20 or C-19 antibody, respectively.

In an assay for the effect of IGF-II on phosphorylation of Akt and mitogen-activated protein kinase (MAPK), the cells were treated with the serum-free medium or with IGF-II (1, 10, or 100 ng/mL) for 15 min. In an assay for the inhibitory effect of m610 on the IGF-II–induced phosphorylation of IGF-IR, insulin receptor, and Akt, the cells were preincubated for 30 min with m610 (0.1, 1, or 10 μg/mL) before IGF-II treatment (100 ng/mL). Twenty-microgram proteins from the lysates were subjected to a Western blot analysis. To detect phosphorylated IGF-IR and insulin receptor, a polyclonal antibody specific for both phosphorylated Tyr1135/1136 of IGF-IR and phosphorylated Tyr1129/1136 of insulin receptor β was used (1:1,000; Biosource International). Phospho-Akt, total-Akt, phospho-MAPK, or total MAPK was, respectively, detected with an anti–phospho-Akt (Ser⁴⁷³) polyclonal antibody, an anti-Akt polyclonal antibody, an anti–phospho-MAPK (Tyr²⁰²/²⁰⁴) mAb, or an anti-MAPK polyclonal antibody (1:1,000 each; Cell Signaling Technology).

**Statistical analysis.** Data are expressed as mean ± SE. Variables were compared between experimental and control groups using Student’s t test. The relation between the percentage of the tumor area and the serum PSA level was assessed using a linear regression analysis. Statistical calculations were done on a Windows personal computer with the GraphPad PRISM software version 4.03 (GraphPad Software). P < 0.05 was considered statistically significant.
Results

Effect of m610 on the growth of bone tumors from MDA PCa 2b cells in HAB. To investigate the effect of m610 on the growth of bone tumors from prostate cancer cells in human bone tissue, m610 (low dose, 1 mg/kg; high dose, 10 mg/kg) or control antibody (10 mg/kg) was administered to the mice with HAB implantation according to the protocol (Fig. 1). At 4 weeks after the treatments, a histomorphologic analysis of the HAB specimens was done. All 8 of the control antibody-treated mice, 7 of the 8 mice in the low-dose group, and 6 of the 8 mice in the high-dose group developed bone tumors. The mean total tumor areas in the low-dose and high-dose groups were 0.8 ± 1.5 and 0.8 ± 1.2 mm², respectively; these values were significantly smaller than that in the control group (2.2 ± 0.9 mm²; P = 0.0397 and 0.0152, respectively; Fig. 2A and B). The mean percentages of the tumor area to the total tissue section area in the low-dose and high-dose groups were 0.7 ± 1.4% and 0.6 ± 0.9%; these values were also significantly lower than that in the control (2 ± 0.8%; P = 0.0438 and 0.0075, respectively; Fig. 2C).

To further assess the effect of m610 on tumor growth, serum PSA level, which is widely used for the diagnosis and therapeutic evaluation of prostate cancer, was measured. The serum PSA levels of the m610-treated groups were significantly lower than that of the control group, in parallel with the reduction in the tumor area (low-dose group, P = 0.0384; high-dose group, P = 0.0072; Fig. 2D). Additionally, a strong correlation was observed between the percentage of the tumor area and the serum PSA values (r² = 0.7656; P < 0.0001, linear regression analysis).

Together, these results of the histomorphologic examination and the PSA measurement show that m610 significantly suppressed the growth of bone tumors induced by MDA PCa 2b cells in implanted HAB, although a dose escalation effect of m610 on the response was not observed.

Immunohistologic examination of m610-treated bone tumor sections in the HAB model. To understand the processes by which the bone tumor growth was suppressed by m610, immunohistologic examinations were done. First, we examined the expression levels of IGF-IR and insulin receptor, which are functional receptors of IGF-II, on the bone tumors by MDA PCa 2b cells in HAB. IGF-IR and insulin receptor were diffusely expressed in both control and m610-treated bone tumors (Fig. 3A and B), and no differences in the expression levels of these receptors were

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**Fig. 1.** Summary of treatment protocol. On day 0, nonobese diabetic/severe combined immunodeficient mice were either implanted with HAB into MFP tissues or not. After 4 wk, MDA PCa 2b cells were inoculated into the implanted HAB or into MFP tissues of the mice without implantation. Weekly administration of m610 was started immediately after the cell inoculation. At 4 wk after the treatment, the mice were sacrificed for histologic analysis and measurement of serum PSA.

**Fig. 2.** m610 suppresses the growth of bone tumors from MDA PCa 2b cells in HAB. A, representative macroscopic images of the PSA-stained HAB sections from control and m610-treated groups. Localized PSA-positive MDA PCa 2b tumor foci were observed. Bar, 2 mm. Sections were histomorphologically analyzed for (B) total tumor area and (C) percentage of tumor area to total HAB tissue area. D, serum PSA values in blood samples of the groups. Control, control antibody group; m610 (1), low-dose (1 mg/kg) group; m610 (10), high-dose (10 mg/kg) group. Mean ± SE for 8 mice in each group. *P < 0.05; **P < 0.01, compared with the control.
microscopically observed between the bone tumors. These findings indicate that the suppressive effect of m610 on the bone tumor growth was not caused by alterations in the expression levels of these receptors. Next, immunohistologic staining for Ki-67 or cleaved caspase-3 was done to investigate the proliferative and apoptotic statuses of the bone tumors. Compared with the control, an apparent decrease in the number of cancer cells with Ki-67–positive nuclei was observed in the m610-treated bone tumor sections (Fig. 4A), whereas a slightly increased number of cleaved caspase-3–positively stained cancer cells was observed (Fig. 4B). Then, the percentages of positively immunostained cancer cells out of the total cell numbers were calculated in three high-power fields with the most highly stained tumor areas. As shown in Fig. 4C, the percentages of Ki-67–positive cancer cells in the m610-treated groups were significantly lower than that in the control (low-dose group, \( P = 0.0007 \); high-dose group, \( P = 0.0002 \)). Meanwhile, the mean percentages of cleaved caspase-3–positive cancer cells in the m610-treated groups were slightly higher than that in the control but not significantly (Fig. 4D; low-dose group, \( P = 0.1937 \); high-dose group, \( P = 0.0601 \)). These findings show that m610 suppressed the proliferation of MDA PCa 2b cells in HAB but did not lead to a significant increase of apoptosis, although a trend to an increase was observed for the high-dose group.

**Effect of m610 on the growth of tumors from MDA PCa 2b cells in MFP.** To examine whether m610 exerts a direct growth-inhibitory effect on MDA PCa 2b cells, m610 (10 mg/kg) or control antibody (10 mg/kg) was administered to the mice without HAB implantation according to the protocol (Fig. 1). Three of the four control antibody-treated mice and four of the five m610-treated mice developed tumors in the MFP tissue. The mean total tumor areas of the control and m610-treated tumors in the MFP model were 1.1 ± 1.5 and 1.3 ± 1.4 mm², respectively (\( P = 0.8787 \); Fig. 5A). The mean percentages of the tumor area to the total tissue section area in the control and m610-treated group were 1.3 ± 1.8% and 2 ± 2.1%, respectively (\( P = 0.6332 \)). The serum PSA levels of the control and m610-treated group were 0.6 ± 0.9 and 2.7 ± 1.2 ng/mL, respectively (\( P = 0.2912 \)). An immunohistologic examination revealed that IGF-IR and insulin receptor were focally expressed on both control and m610-treated tumors in the MFP tissue and that the expression levels of the receptors were not different between the tumors (Supplementary Fig. S1A and B). Ki-67 immunostaining revealed that the proliferative statuses were similar between control and m610-treated tumors, consistent with the results for tumor size (Fig. 5B; Supplementary Fig. S1C; \( P = 0.7462 \)). Together, these findings indicate that m610 had no effect on the growth of MDA PCa 2b cells in MFP, unlike in HAB, showing that the inhibitory effect of m610 on tumor cell proliferation is restricted to within the bone.

**Inhibition of IGF-II–induced cell proliferation and signaling in MDA PCa 2b cells by m610.** Using a cell proliferation assay, we confirmed that m610 inhibits the IGF-II–induced proliferation of MDA PCa 2b cells. Exogenous IGF-II dose-dependently induced the proliferation of MDA PCa 2b cells (Fig. 6A, lanes 1-4). The control antibody
had no inhibitory effect on the IGF-II-induced cell proliferation (Fig. 6A, lane 5). m610 at doses of 1 and 10 μg/mL significantly inhibited the IGF-II-induced cell proliferation, compared with the control antibody group (Fig. 6A, lanes 7 and 8; \( P = 0.0049 \) and \( 0.0039 \), respectively), but m610 had no effect on the proliferation of MDA PCa 2b cells without exogenous IGF-II stimulation, indicating that m610 did not have a direct effect on the cells (Fig. 6A, lane 9).

To confirm that m610 prevents signal transduction mediated by receptor interaction with IGF-II in MDA PCa 2b cells, we firstly investigated whether IGF-II activates IGF-IR and insulin receptor in the cells by immunoprecipitation-Western blot analysis. On ligand binding, IGF-IR and insulin receptor undergoes autophosphorylation of tyrosine residues in their β-subunits. Western blot analysis showed that IGF-II induces phosphorylation of tyrosines in each receptor in MDA PCa 2b cells (Fig. 6B). Note that IGF-II induces significantly higher level of insulin receptor phosphorylation than of the IGF-IR phosphorylation. In contrast, IGF-I induces much higher levels of phosphorylation of the IGF-IR than of the insulin receptor. Tyrosine phosphorylation of IGF-IR or insulin receptor leads to phosphorylation of the downstream kinases Akt and MAPK. We next examined whether IGF-II induces the downstream kinases in MDA PCa 2b cells by a Western blot analysis. IGF-II induced Akt phosphorylation in the cells but not MAPK phosphorylation (Supplementary Fig. S2). We finally investigated whether m610 prevents the IGF-II-induced IGF-IR and insulin receptor signaling in MDA PCa 2b cells by a Western blot analysis. To detect the total amount of activated statuses (IGF-IR plus insulin

**Fig. 4.** m610 suppresses the proliferative status of MDA PCa 2b tumors in HAB. A, Ki-67 immunostained images of HAB sections from control and high-dose group. B, cleaved caspase-3 immunostained images. Bar, 100 μm. The percentages of positively immunostained cancer cells to the total cells were calculated. C and D, percentages of Ki-67–positive and cleaved caspase-3–positive cancer cells, respectively. CC-3, cleaved caspase-3. Mean ± SE. ***, \( P < 0.001 \), compared with the control.
receptor), which is most relevant for the degree of induced cell proliferation, an antibody specific for phosphorylated tyrosines of IGF-IR and insulin receptor was used in the analysis. As shown in Fig. 6C, the m610 dose-dependently prevented the IGF-II-induced tyrosine phosphorylation of IGF-IR/insulin receptor as well as serine phosphorylation of Akt in MDA PCa 2b cells.

**Discussion**

IGF-II is the most abundant growth factor in human bone (8) and, unlike in mouse bone, is nine times more abundant than IGF-I (11). Our previous study revealed the role of IGFs in prostate cancer growth in human bone, as shown by the suppressive effect of a neutralizing antibody (KM1468), which cross-reacts with human IGF-I and IGF-II and mouse IGF-II, on the growth of MDA PCa 2b cells in implanted HAB (6). Moreover, another study of ours indicated that tail vein–injected prostate cancer cells preferentially metastasized to implanted HAB rather than to implanted mouse bone or native bone (5). These findings suggested that IGF-II may support human prostate cancer more efficiently than IGF-I in human bone environment, in accordance with the “seed and soil” theory by Paget (16) for metastasis. Therefore, we became interested in specifically targeting IGF-II as a potential therapy for bone metastasis from prostate cancer.

Although clinical studies have suggested that IGF-II contributes to the development and progression of several cancers including prostate cancer (17–19), only a few studies by others and us have reported an in vivo antitumor effect of targeting IGF-II. O’Gorman et al. reported that the overexpression of the IGF-II receptor, which is a clearance receptor for IGF-II, on choriocarcinoma cells reduced the cell growth in vitro and in vivo, indicating that inhibiting IGF-II is a potential target for cancer therapy (20). In another strategy using neutralizing antibodies, Miyamoto et al. reported that an IGF-II neutralizing antibody partly suppressed the development of liver metastasis induced by the intrasplenic injection of colon cancer cells, whereas

**Fig. 5.** m610 does not suppress the growth of tumors from MDA PCa 2b cells in MFP. Histologic analyses in MFP model. A, total tumor area. Mean ± SE for four mice in the control group and five mice in the m610-treated group. B, percentages of Ki-67–positive cancer cells.

**Fig. 6.** m610 inhibits IGF-II–induced cell proliferation and phosphorylations of IGF-IR/insulin receptor and Akt in MDA PCa 2b cells. A, MDA PCa 2b cells were treated for 48 h with serum-free medium alone, with various concentrations of IGF-II, with the control antibody plus 100 ng/mL IGF-II, or with various concentrations of m610 in the presence or absence of 100 ng/mL IGF-II. Cells were counted using trypan blue dye exclusion. Mean ± SE of triplicate determinations. Representative of three independent experiments. #, P < 0.05, compared with serum-free medium; **, P < 0.01, compared with control antibody group. B, cells were treated with 100 ng/mL IGF-I or IGF-II. Tyrosine phosphorylation of immunoprecipitated IGF-IR or insulin receptor was examined by a Western blot analysis. WB, Western blot; IP, immunoprecipitation; p-Tyr, phosphorylated tyrosine. C, cells were preincubated with the indicated concentrations of m610, and 100 ng/mL IGF-II was added. IGF-IR/insulin receptor and Akt phosphorylation levels and the total protein levels in the lysates were examined by Western blot analysis. p-, phosphorylated; t-, total.
IGF-I neutralizing antibody significantly suppressed the development of liver metastasis (21). Because IGF-II production in rodents becomes attenuated in most tissues soon after birth unlike IGF-I production (22), the effect of IGF-II as an endocrine and/or paracrine on tumor growth may be underestimated in models using mouse-native organs.

The present study showed that inhibiting IGF-II by m610 effectively suppresses the growth of prostate cancer cells in a human bone environment. The following results support this conclusion: (a) m610 significantly suppressed the growth of bone tumors from MDA PCa 2b cells in implanted HAB; it even decreased the number of mice developing tumors from 8 to 6, although larger number of mice are needed to confirm this effect; (b) Ki-67 immunostaining revealed that the proliferative status of m610-treated bone tumor was apparently suppressed; (c) in the absence of HAB, m610 had no effect on the growth of tumors from MDA PCa 2b cells, indicating that the suppressive effect of m610 on the tumor cells was restricted to those within HAB; and (d) in vitro assays confirmed that m610 prevents the exogenous IGF-II–induced proliferation of MDA PCa 2b cells. These results provide clear evidence of the important role of IGF-II for tumor growth in the HAB model and of an in vitro antitumor effect of m610 on metastatic bone tumor from prostate cancer through a mechanism involving the inhibition of IGF-II. They also underscore the notion that IGF-II levels in local tissue may be more relevant in tumor promotion than its plasma levels and that a paracrine mechanism of IGF-II may play a critical role in tumor growth.

The potency of m610 on the growth inhibition of MDA PCa 2b cells in the HAB model is 65%, whereas that of the previously published antibody, KM1468, is 97% compared with the respective controls: the antitumor effect of inhibiting IGF-II alone is lower than that of inhibiting both IGF-I and IGF-II in the HAB model. Despite the lower antitumor effect of m610 in the HAB model, targeting IGF-II by m610 might provide certain clinical benefits in cancer therapy for the following reasons. (a) Growth hormone feedback is not known for IGF-II, but IGF-I is regulated by this feedback. Lowering IGF-I concentration triggers feedback upregulation of the growth hormone; the growth hormone compensates for the reduced IGF-I levels. Thus, targeting IGF-I might require high concentrations of anti-IGF-I antibodies. It should be noted that KM1468 is not reactive with mouse IGF-I; therefore, its use in our HAB model does not trigger the growth hormone feedback on the IGF-I and the tumor growth. (b) Because m610 is a fully human antibody, its clinical use is less likely to induce immune reactions compared with murine antibodies.

Targeting IGF-II might provide additional therapeutic benefit in combination with other treatments. IGF-IR activation by IGF-I and IGF-II has been shown to stimulate the growth of a wide range of cancer cells (23, 24). Currently, potent mAbs against the IGF-IR are being tested in clinical trials against multiple tumor types including prostate, breast, and colon cancers and Ewing’s sarcoma (25). Importantly, it is becoming increasingly evident that insulin receptor activation by IGF-II enhances the growth of Ewing sarcoma and breast cancer in addition to the IGF-IR activation: cotargeting IGF-IR and insulin receptor is likely to be more effective than targeting the IGF-IR alone (26–28). Recently reported immunohisto logic examinations of primary human prostate cancer show that IGF-IR as well as insulin receptor are both commonly expressed on the tissues (29). Our results from the immunohistologic examinations (Fig. 3) and Western blot analyses (Fig. 6B and C) suggest that insulin receptor activation by IGF-II plays an important role in the prostate cancer cell growth in bone in addition to the IGF-IR activation. Based on these findings, IGF-II could be a promising candidate target in therapeutic strategies for cotargeting IGF-IR and insulin receptor.

If m610 were capable of suppressing the growth of prostate cancer in bone without any adverse reactions, m610 therapy might considerably improve the quality of life of patients with bone metastases of prostate cancer. In the present study, the administration of m610 did not affect the body weights of mice during the 4-week treatment period (Supplementary Fig. S3), and no adverse findings were observed in the histologic examination for the mouse organs.

In conclusion, the present study showed that an IGF-II–specific antibody, m610, can sufficiently suppress the growth of bone tumors from MDA PCa 2b cells in a human bone environment and that this effect is caused by the suppression of the proliferative status of the tumor cells. These results suggest that the targeting of bone-derived IGF-II using a neutralizing antibody offers a new therapeutic strategy for bone metastasis from prostate cancer.

Disclosure of Potential Conflicts of Interest

No potential conflicts of interest were disclosed.

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