Abstract

Aim: Patients with prostate cancer frequently develop osteoblastic bone metastases. Canine models are important because dogs are the only mammal to develop spontaneous prostate cancer with osteoblastic bone metastases similar to men. The mechanism by which prostate cancer induces bone formation is unclear; however, it depends on the complex interaction between prostate cancer cells and bone microenvironment. This study investigated the effects of three canine prostate cancer cell lines (Ace-1, LuMa, and Probasco) on bone formation and resorption in vitro.

Methods: Mouse calvaria were treated with conditioned medium (CM) from cell lines. Calvaria were evaluated by histology, fluorescent calcein uptake at sites of bone mineralization, medium calcium assay, and alkaline phosphatase activity. The expression of bone-related genes was measured using quantitative reverse transcription-PCR.

Results: A novel calcein uptake assay was developed to measure bone formation and mineralization in vitro. Ace-1 CM induced predominantly bone resorption in calvaria, while Probasco CM induced marked bone formation, mineralization, and healing of calvaria defects. The expression of osteoblast-related genes in calvaria showed that Probasco CM stimulated the maturation and differentiation of osteoblasts and inhibited osteoclastogenesis. Both bone modeling and remodeling were involved in Probasco CM-induced bone formation and mineralization by inhibiting remodeling with zoledronic acid. Inhibition of WNT activity by DKK-1 decreased the osteoblastic activity...
Conclusion: Probasco cells induced bone formation and mineralization in vitro that depended on the WNT signaling pathway. Probasco cells will serve as a valuable model for studying the mechanisms of osteoblastic bone metastasis in prostate cancer.

Keywords: Bone, canine, dog, metastasis, osteoblast, prostate cancer, zoledronic acid, DKK-1

INTRODUCTION

Prostate cancer (PCa) metastasizes to bone and induces “osteoblastic” metastases that are characterized by increased woven bone formation initially on endosteal bone surfaces and can eventually extend beyond the normal limits of the bone[1]. This results in a metastasis composed of prostate cancer tissue and dense bone that can be visualized by radiography. Osteoblastic metastases are often associated with bone pain, impaired mobility, and spinal cord compression that severely affect the patients’ quality of life[2].

Dogs have a lobular prostate gland that is similar to men’s in size and intact male dogs develop benign prostatic hyperplasia as they age similar to men[3]. Dogs are the only animal that develops spontaneous prostate cancer with metastasis to bone with the formation of osteoblastic metastases[4]. Most metastatic prostate cancers in dogs are androgen-independent and represent a valuable animal model for late-stage androgen-resistant prostate cancer as it occurs in men[5].

Our laboratory has developed multiple prostate cancer cell lines from spontaneous prostate cancer in dogs. This investigation utilized three different cell lines (Ace[6], Probasco[7], and LuMa[8]) that vary in their in vivo phenotype in athymic mice. All metastasize to bone and induce different degrees of osteoblastic bone formation and osteoclastic bone resorption.

Several factors have been implicated as contributing to osteoblastic metastasis including WNT proteins[9]. WNT proteins are soluble glycoproteins that bind to receptor complexes composed of LRP5/6 and Frizzled proteins[10]. Although the function of WNTs in bone development and homeostasis is well elucidated, the role of WNT signaling in the interaction between PCa cells and bone microenvironment is complicated and remains poorly understood. Previously we have shown that inhibition of bone formation by prostate cancer by blocking WNT action with DKK-1 resulted in increased bone metastases and more rapid tumor growth[11]. This suggests that bone formation in osteoblastic metastases plays an important role in the regulation of cancer progression.

Since the pathogenesis of bone formation in prostate cancer bone metastases is unknown, we developed a novel in vitro model system for measurement of bone formation. In vitro cultures of mouse calvaria is a time-honored system for culture of bone tissue and is especially useful for investigating in vitro osteoclastic bone resorption[12]. Measurement of in vitro bone formation using calvaria has been less utilized because it is more technically challenging. There are few reports of bone formation induction in vitro in calvaria. We have optimized the use of calcine, a fluorescent biomarker of bone mineralization, to permit measurement of bone formation in vitro[13].

The overall goal of this investigation was to collect conditioned medium (CM) from cultured canine prostate cancer cell lines and measure their effect on bone formation and resorption and bone healing in vitro. In addition, the mRNA expression for bone-specific proteins and candidates for induction of bone
formation were measured. DKK-1 was used as a broad inhibitor of WNT signaling to determine the role of the WNT pathway on bone formation *in vitro*.

**METHODS**

**Cell lines, conditioned medium, and chemicals**
Four canine prostatic cancer cell lines were used in this study including Ace-1[^6], Probasco[^7], Probasco+DKK-1 (Probasco cells transduced with human DKK-1), and LuMa[^8] (all cells were established by our laboratory). Cells were cultured in DMEM/F12 (Gibco, Carlsbad, CA) containing 10% fetal bovine serum (FBS), 1% penicillin/streptomycin and 100 μg/mL Normocin (Invivogen, San Diego, CA). All cells were serially passage using trypsinization and maintained at 37 °C and 5% CO₂ in a humidified atmosphere.

CM was prepared by incubating near confluent cells in T75 flask for 24 h with serum-free BGJb culture medium (Gibco) with 0.1% bovine serum albumin (BSA) fraction V (Sigma-Aldrich Corp, St Louis, Missouri) and 100 μg/mL Normocin. CM was stored in aliquots at -80 °C until used. ZOL (zoledronic acid, Zomet™, Novartis) was purchased from the James Cancer Center at Ohio State University.

**In vitro cultures of mouse calvaria**
Calvaria of 4- to 7-day-old CD1 pups were aseptically removed and dissected from surrounding soft tissues, cut in half, and incubated in 1 mL serum-free BGJb medium or conditioned medium for 2-6 days at 37 °C and 5% CO₂ in a humidified atmosphere. Medium was changed every other day in both control (BGJb + calvaria only) and treatment groups (50% BGJb + 50% CM + calvaria) until the end of the experiment.

**Wound healing of calvaria defect**
An *in vitro* bone healing model was used to evaluate the induction of new bone formation by Probasco conditioned media. A 1.5 mm bone disk was punched and removed from the parietal bone on both sides in each calvaria using biopsy punches (Braintree Scientific, Braintree, MA) to create artificial defects. Calvaria with defects were treated with CM or control (plain media) for 6 days. The calvarial defects were imaged at day 6 using a dissecting microscope (Nikon SMZ-U) and the repaired areas were quantified using imageJ (NIH, version 2.0).

**Calcein uptake and bone formation with mineralization**
A novel *in vitro* bone formation and mineralization assay was developed. Each half calvarium was cultured in BGJb medium or 50% conditioned medium for 6 days. Calcein (Sigma-Aldich), a green, fluorescent dye with excitation and emission wavelengths of 495/515 nm that binds to calcium at sites of bone formation and mineralization, was added to the cultures (0.4 μg/mL) at day 4. At the end of the experiment, calvaria were fixed in 10% neutral-buffered formalin at room temperature for 1.5 h. A bone disk (1.5 mm) was removed from each side of the fixed mouse calvaria using dermatology biopsy punches and then mounted on glass slides. Newly mineralized bone labelled by calcein was detected and imaged using a Nikon E600 fluorescent microscope with a digital camera. Total fluorescence intensity was quantified using imageJ (NIH, version 2.0) to measure the amount of new bone formation.

**Histopathology**
Fixed calvaria were decalcified (10% EDTA, pH 7.4, for 24 h at 4 °C), paraffin-embedded, 4 μm sections cut, mounted on glass slides, and stained with hematoxylin and eosin (H&E) for examining the changes of calvaria thickness. Undecalcified calcein-treated calvaria were embedded in OCT, 10 mm sections cut on a cryomicrotome, and mounted for viewing calcein uptake.
Alkaline phosphatase activity assay
3.5 mm calvaria discs (after 6 days of incubation with BGJb or CM) were extracted in 500 μL 0.01% triton X-100 (containing 0.02% azide) for 72 h at 4 °C, and extracted protein was collected. Alkaline phosphatase (ALP) activity was measured by the time-dependent formation of p-nitrophenolate (pNP) from ALP substrate solution (pNPP) at room temperature for 1 h [14]. Typically, 10 μL of extract was incubated in a total volume of 100 μL containing 4.7 mM pNPP, 1 mM MgCl₂ and 1.02 M diethanolamine buffer, pH 9.8, in a 96-place microtiter plate. Each sample was measured in duplicate. The reactions were initiated by addition of substrate, and the time-dependent increase in absorbance at 405 nm was measured every 5 min (Synergy™ HTX Multi-Mode Microplate Reader). ALP activity was calculated as pNP produced/min using a standard curve.

Calcium concentration in calvaria medium
Calvaria culture medium supernatant was collected on day 1 and day 2, and the total medium calcium was measured as described by Lorentz [15].

Dickkopf-1 enzyme-linked immunosorbent assay
DDK-1 levels in conditioned medium were measured using the DuoSet Human DKK1 enzyme-linked immunosorbent assay (ELISA) Kit, as recommended by the manufacturer (R&D Systems, Minneapolis, MN). The lowest standard of the assay was 62.5 pg/mL.

RNA extraction and quantitative reverse-transcription polymerase chain reaction
Total RNA was extracted from calvaria using RNeasy mini kit (Qiagen). Then total RNA (0.5 μg) was reverse transcribed in cDNA using the Superscript II First Strand cDNA synthesis kit (Invitrogen). Quantitative RT polymerase chain reaction (PCR) was performed for the reference gene, ubiquitin C (UBC), osteoblast-related genes including runt-related transcription factor 2 (RUNX2), Osterix (OSX), Alkaline phosphatase (ALPL), receptor activator of nuclear factor kappa-B ligand (RANKL), osteoprotegerin (OPG), osteocalcin (OCN), integrin-binding sialoprotein (IBSP), parathyroid hormone 1 receptor (PTH1R), osteomodulin (OMD), and osteonectin (ON), and osteoclast-related genes including matrix metalloproteinase 9 (MMP9) and cathepsin K (CTSK). The primers are listed in Table 1. The thermal cycle conditions were: 95 °C, 3 min; (95 °C, 10 s; 55 °C, 30 s; 75 °C, 30 s) ×39; 95 °C, 1 min; 55 °C, 1 min. The expression of targeted genes was normalized by UBC and quantified using the 2^ΔΔCt method.

Statistical analysis
All data were displayed as mean ± standard deviation. Paired student’s t-test was used for bivariate analyses. ANOVA was used to analyze multivariate comparisons. Data with P values ≤ 0.05 were considered statistically significant. Statistical calculation was performed using GraphPad Prism version 9 (La Jolla, CA).

RESULTS
Effect of canine prostate cancer cell conditioned medium on bone formation and resorption in vitro
Ace-1, LuMa, and Probasco are known to metastasize to bone and induce different phenotypes of bone metastasis in vivo [5]. Ace-1 induces mixed osteoblastic/osteolytic bone metastasis, LuMa induces osteoblastic bone metastasis, and Probasco induces osteoblastic bone metastasis with abundant new bone formation. Calvaria were treated with CM from the three canine prostate cancer cells to determine if they would induce similar phenotypes in bone in vitro as compared to bone metastases in vivo. The ability of CM to induce bone formation was measured by uptake of the fluorescent dye, calcine. The results showed that Probasco CM induced bone formation with mineralization, while Ace-1 CM induced significantly less bone formation compared with controls. No difference in bone formation was seen between LuMa CM-treated calvaria and controls (calvaria treated with medium alone) [Figure 1A and B].
Table 1. Primers used for qRT-PCR with mouse calvaria treated with control media or conditioned media collected from canine PCa cell lines

<table>
<thead>
<tr>
<th>Gene</th>
<th>Forward primer (5’-3’)</th>
<th>Reverse primer (5’-3’)</th>
</tr>
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<tbody>
<tr>
<td>UBC</td>
<td>CGTCGAGCCCGAGTTACCAAAAGG</td>
<td>CCCCATCACACCAAGAAAGCAACAG</td>
</tr>
<tr>
<td>RUNX2</td>
<td>CGACAGTCGACACTGCTTG</td>
<td>TACCTTCGAGGGCTCACAA</td>
</tr>
<tr>
<td>OSX</td>
<td>ATGGCGTCTCTCTGGTGG</td>
<td>ATGGCGTCTCTCTGGTGG</td>
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<td>ALPL</td>
<td>GGCGATATGAGGTACATCCA</td>
<td>GTGATCGACAGTGCTGAGT</td>
</tr>
<tr>
<td>RANKL</td>
<td>ACACCTCACATCATGGCCT</td>
<td>CTTAAGCGCTATGATGATCTGG</td>
</tr>
<tr>
<td>OPG</td>
<td>AGCGTCTGAAGGTGAGG</td>
<td>TGAGTGCCCGGAGAT</td>
</tr>
<tr>
<td>OCN</td>
<td>CTACAGATGCGCAGGGCCC</td>
<td>CCAAGGTAGACCCGGAGTCT</td>
</tr>
<tr>
<td>IBSP</td>
<td>CGGCGATGATGGGCAAAGAGGG</td>
<td>GCTAGCGTACACACCGGAGAG</td>
</tr>
<tr>
<td>PTH1R</td>
<td>GGCGCTGGGAGGTGTT</td>
<td>CGACTCTTCATGGTCATGAACCTG</td>
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<tr>
<td>OMD</td>
<td>GTGGAGTGGAGTTGCTGGA</td>
<td>TGAAGGGTGGAGTTGCTGAGT</td>
</tr>
<tr>
<td>ON</td>
<td>GCTCAGAATGCTGGGAGAGAG</td>
<td>CTCACGACGTCATGGGAGT</td>
</tr>
<tr>
<td>MMP9</td>
<td>CATTGGGATGGATAGGGAT</td>
<td>ATTTGGAAACCTACACGGCC</td>
</tr>
<tr>
<td>CTSK</td>
<td>CTCTCAATAGCTGCAACAGA</td>
<td>CCTCTGATTTAGCTGCTT</td>
</tr>
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PCa: Prostate cancer.

ALP activity was detected in protein extracted from PCa CM-treated calvaria. Consistent with bone mineralization, calvaria treated with Ace-1 CM showed the lowest ALP activity, there was no difference between control and LuMa CM-treated calvaria, and there was a marked increase in ALP activity in Probasco CM-treated calvaria [Figure 1C].

The effect of CM on bone resorption by osteoclasts was measured. Calvaria were treated with CM or control for 2 days. Calcium concentration in culture media was measured on day 1 and day 2. Ace-1 and LuMa CM increased bone resorption (greatest for Ace-1 CM) resulting in more calcium released into the medium compared to control, while there was no difference in bone resorption between Probasco CM-treated calvaria and control [Figure 1D]. These data demonstrated that CM from three canine prostate cancer cells induces different degrees of bone formation and resorption in vitro similar to the in vivo phenotype of bone metastases.

Quantitative reverse-transcription polymerase chain reaction of CM-treated calvaria in vitro
The expression of 12 bone-related genes including, an osteoblastogenesis maker (RUNX2), osteoblast differentiation-to-maturation makers (OSX, ALPL, RANKL, OPG, OCN, IBSP), osteoblast mineralization makers (PTH1R, OMD, and ON), and osteoclast-related genes (MMP9 and CTSK) was measured by quantitative reverse-transcription polymerase chain reaction in calvaria treated with CM or control for 3 days in order to investigate the potential mechanisms by which CM induced bone formation or resorption. Ace-1 CM upregulated the mRNA expression of RANKL over 60-fold, the expression of OPG 1.2-fold, and the expression of MMP9 [Figure 2A]. This increased the ratio of RANKL/OPG mRNA over 50-fold in treated calvaria, which is consistent with increased in vitro bone resorption and the observed increase of calcium in the media released from Ace-1 CM-treated calvaria. In addition, the mRNA expression level of RUNX2, OSX, ALPL, OCN, IBSP, PTH1R, OMD, and ON was decreased markedly in Ace-1 CM-treated calvaria compared to controls, which was consistent with the result that Ace-1 CM had the least ability to stimulate bone mineralization in calvaria in vitro. In LuMa CM-treated calvaria, the relative mRNA expression levels of OSX, ALPL, OCN, PTH1R, and ON were downregulated compared to controls [Figure 2B]. In Probasco CM-treated calvaria, the expression of RANKL, MMP9, and CTSK was significantly increased, which may be related to the increased bone turnover induced by Probasco CM [Figure 2C].
In addition, the mRNA expression of the genes was measured in LuMa or Probasco CM-treated calvaria and directly compared to Ace-1 CM-treated calvaria (as control) in order to investigate the different effect of the prostate cancer cell lines on calvaria. The relative mRNA expression level of RUNX2, OSX, ALPL, OCN, PTH1R, and OMD was significantly increased, RANKL was markedly decreased, and there was no change in the expression of OPG in both Probasco and LuMa CM-treated calvaria compared to Ace-1 CM. The expression of IBSP was significantly increased in Probasco CM-treated calvaria [Figure 2D and E]. The results demonstrated that osteoblastic cell lines, LuMa and Probasco, induced bone formation by the stimulation of maturation of osteoblasts to form mineralized bone, and the inhibited osteoclastogenesis in vitro.
Figure 2. Expression of bone-related genes in calvaria treated with CM or control. Graphs represent the fold changes of mRNA expression of RUNX2, OSX, ALPL, RANKL, OPG, OCN, IBSP, PTH1R, MMP9, and CTSK in (A) Ace-1 CM-treated calvaria relative to control; (B) LuMa CM-treated calvaria relative to control; (C) Probasco CM-treated calvaria relative to control; (D) Probasco CM-treated calvaria relative to Ace-1 CM-treated calvaria; (E) LuMa CM-treated calvaria relative to Ace-1 CM-treated calvaria; and (F) Probasco CM-treated calvaria relative to LuMa CM-treated calvaria. Data were expressed as mean ± SD. Significant differences were marked with asterisks (*$P < 0.05$, **$P < 0.01$, ***$P < 0.001$, ****$P < 0.0001$). Note differences in the scales of the Y axes. CM: Conditioned medium.

Furthermore, the comparison of the mRNA expression between Probasco CM-treated calvaria and LuMa CM-treated calvaria revealed that Probasco had greater osteoblastic ability compared to LuMa, which is consistent with the observed bone mineralization on calvaria induced by Probasco CM [Figure 2F].

Effect of Probasco CM on bone formation in mouse calvaria defects
Probasco CM improved the healing response of calvarial defects when compared to control calvaria [Figure 3A]. The repaired area in the defect of Probasco CM-treated calvaria was increased 2-fold compared to control [Figure 3B], which demonstrated that Probasco CM stimulated bone formation in bone defects in vitro.

Probasco CM increased bone formation through bone remodeling and modeling
Bone shape and structure are a result of bone modeling (bone formation by osteoblasts independent of osteoclasts) and bone remodeling (the removal of mineralized bone by osteoclasts followed by the formation of new bone matrix by osteoblasts that subsequently becomes mineralized). To determine if bone modeling was involved in Probasco CM-induced bone formation, bone remodeling was inhibited by blocking osteoclast formation and function with the bisphosphonate[12], zoledronic acid (ZOL; Zometa™).

Inhibition of bone remodeling greatly decreased bone formation in the control group; however, Probasco CM was still able to induce bone formation [Figure 4A and B], which revealed that both remodeling and modeling were involved in Probasco CM-induced bone formation. Additionally, calvaria treated with Probasco CM were evaluated by histology. Frozen sections and H&E staining showed that calcein was deposited in the bone matrix and marked periosteal thickening was induced by Probasco CM [Figure 4C and D].
Figure 3. Effect of Probasco CM on mouse calvarial defects. (A) Representative microscopic images. Calvaria with defects (red circle showing the initial 1.5 mm round artificial defect before treatment) incubated with only BGJb had less repaired area (white arrow) compared to calvaria treated with Probasco CM. (B) Repaired area was quantified. Data were expressed as mean ± SD. Significant differences were marked with asterisks (***P < 0.001). CM: Conditioned medium.

Figure 4. Probasco CM increased bone formation through bone remodeling and modeling. Calvaria were divided into two groups. One group was pre-treated with zoledronic acid (ZOL, 10⁻⁵M) and the other group was treated with vehicle overnight. In each pair of calvaria, a half calvarium was treated with Probasco CM or control medium. (A) Representative images of 1.5 mm calvarial disks taken from the parietal bone. Probasco CM increased bone formation and mineralization (calcein uptake) compared to control. (B) The total fluorescence intensity was quantified. Data were expressed as mean ± SD and analyzed by paired t-test. Significant differences were marked with asterisks (*P < 0.05, **P < 0.01). (C) Frozen section of calvaria treated with control media or Probasco CM. Calcein was deposited in mineralized bone matrix on both surfaces of the calvaria. (D) H&E-stained section of calvaria. There was marked periosteal thickening and new bone formation caused by Probasco CM. CM: Conditioned medium.

**Stimulation of bone formation by Probasco CM was dependent on the WNT signaling pathway**

Our laboratory has shown that blocking WNT action with DKK-1 inhibited the osteoblastic phenotype
induced by Ace-1 in nude mice\textsuperscript{[15]}. To test whether the stimulation of bone formation by Probasco CM also depended on WNT signaling pathway, calvaria were treated with CM from either Probasco+DKK-1 or Probasco cells. Secretion of human DKK-1 by Probasco+DKK-1 cells was confirmed by ELISA. CM collected from Probasco cells had no detectable DKK-1 secretion while Probasco+DKK-1 CM contained approximately 8 ng/mL after 24 h of culture [Figure 5A]. Probasco+DKK-1 CM strongly inhibited bone mineralization [Figure 5B and C]. Calvaria pre-treated with ZOL, in which osteoclast activity and bone remodeling were blocked by ZOL, indicated that Probasco+DKK-1 inhibited bone modeling induced by Probasco CM. Frozen sections confirmed that less calcein was deposited in Probasco+DKK-1 CM-treated calvaria matrix compared with the calvaria treated with Probasco CM [Figure 5D]. In addition, there was a marked decrease in ALP activity in Probasco+DKK-1-treated calvaria [Figure 5E]. The mRNA expression of osteoblast-related genes was measured in calvaria. The level of OCN was significantly decreased in Probasco+DKK-1 CM-treated calvaria compared to calvaria treated with Probasco CM [Figure 5F]. OCN is a marker of mature osteoblasts. Osteocalcin is secreted by osteoblasts into the bone micro-environment and regulates extracellular matrix mineralization\textsuperscript{[16]}. The reduced expression of OCN supports the observed decrease in mineralization caused by Probasco+DKK-1 CM. In addition, the downregulation of OCN in Probasco+DKK-1 CM-treated calvaria reveals that blocking WNT signaling pathway by DKK-1 may inhibit osteoblast differentiation \textit{in vitro}. These results showed that inhibition of the WNT signaling pathway by DKK-1 significantly decreased bone formation induced by Probasco CM.

**DISCUSSION**

We successfully developed a novel method to measure bone formation and mineralization \textit{in vitro}. \textit{In vitro} bone resorption assays have been used for decades to measure osteoclastic bone resorption; however, it has been particularly challenging to measure bone formation \textit{in vitro}. The use of the fluorescent dye, calcein, enabled increased sensitivity for the measurement and quantitation of bone mineralization \textit{in vitro}. This newly described assay will be helpful for mechanistically investigating the pathogenesis of bone formation induced by prostate cancer.

We showed that conditioned media from canine prostate cancer cells induced bone resorption and formation \textit{in vitro} that correlated with the \textit{in vivo} phenotype of bone metastases associated with each cell line. Ace-1 CM mainly induced bone resorption, while Probasco CM significantly stimulated bone formation by both bone modeling and remodeling in mouse calvaria. Additionally, we showed that inhibition of WNT activity by DKK-1 blocked the osteoblastic activity in Probasco cells.

Animal models are critical for investigating the pathogenesis of prostate cancer-induced bone metastasis. Currently, a few animal models are available that closely mimic the osteoblastic property of human prostate cancer. Human LNCap subline C4-2B and the LuCaP23.1 tumor cell lines are two most commonly used human cell lines that have an osteoblastic phenotype; however, both of them induce mixed bone tumors as they stimulate significant bone lysis as well as formation\textsuperscript{[17,18]}. As for murine models, most transgenic mouse models do not develop bone metastases, and bone xenograft models of prostate cancer often induce mixed or osteolytic bone tumors\textsuperscript{[5]}. In the present study, we used three canine cell lines (Ace-1, LuMa and Probasco). The ability of the Probasco cells to induce osteoblastic bone formation both \textit{in vivo} and \textit{in vitro}, makes it especially valuable for investigating the mechanisms of osteoblastic metastasis in prostate cancer. In addition, by comparing the genetic expression profile of osteoblastic Probasco cells with osteolytic Ace-1 cells, it may uncover critical pathways that contribute to the osteoblastic phenotype in prostate cancer.

Normal bone development consists of two processes, bone remodeling, defined by the sequential actions of bone resorption and bone formation, and bone modeling, characterized by the independent actions of
Figure 5. Stimulation of bone formation by Probasco CM was dependent on the WNT signaling pathway. (A) Concentration of secreted human DKK-1 in Probasco CM and Probasco+DKK-1 CM. (B) Representative images of 1.5 mm calvarial disks from parietal bone. Probasco+DKK-1 significantly decreased bone formation. (C) The total fluorescence intensity was quantified. (D) Frozen section of calvaria treated Probasco+DKK-1 CM or Probasco CM. More calcein was deposited in the bone matrix. (E) ALP activity in calvaria treated with Probasco CM and calvaria treated with Probasco+DKK-1 CM. (F) The mRNA expression of bone-related genes in Probasco+DKK-1 CM-treated calvaria compared to Probasco CM-treated calvaria. Data were expressed as mean ± SD and analyzed by paired t-test. Significant differences were marked with asterisks (*P < 0.05, **P < 0.01), ***P < 0.001).

osteoblasts and osteoclasts. Histologic assessment of PCa-induced bone metastases demonstrates that the radiodense bone metastases were largely composed of woven bone which has less structural strength compared to lamellar bone. The formation of woven bone is often caused by bone modeling since the bone formation occurs rapidly and is not dependent on initial osteoclastic bone resorption. Our in vitro results showed that Probasco CM induced osteoblastic bone formation in part by bone modeling since Probasco CM induced bone formation in ZOL pretreated-calvaria.

WNTs are a large family of regulatory proteins that direct embryological development and promote bone growth, acting through both canonical and non-canonical signaling pathways. Proteins secreted by prostate cancer cells exert their function via autocrine and paracrine actions, and result in the stimulation of PCa growth and osteoblast activity. It has been reported that inhibition of WNT activity switched the mixed osteoblastic/osteolytic C4-2B cell line to a highly osteolytic cell line. Also, PCa cells treated with WNT3A or WNT5A promoted the differentiation of osteoblasts through the canonical and non-canonical pathways in a BMP-dependent and -independent manner, which demonstrated crosstalk between WNTs and BMPs in PCa-induced osteoblast differentiation. Our in vitro results, in agreement with these findings, showed that increasing DKK-1 expression in Probasco blocked WNT-induced osteoblastic activity. Based on RNA-seq analysis of canine PCa cell lines (unpublished data), we found that several WNTs proteins (WNT3, WNT5A, and WNT7B) as well as BMPs (BMP-2, BMP-4, and BMP-7) are expressed in Probasco cells. It will be promising to further investigate if both WNTs and BMPs play a role as mediators of Probasco-induced osteoblastic activity.
ALP is an important enzyme in bone formation, and it is highly expressed in mineralized bone cells\cite{23}. While the measurement of ALP expression is well developed and is a reliable indication of the osteoblastic phenotype, the regulation of ALP expression and function in the mineralization of bone is less clear. The regulation of ALP expression is mainly controlled by BMP/RUNX2/Osterix system, and the WNT signaling cascade is also involved in the process\cite{24,25}. The capacity of BMP-2 to induce ALP relies on WNT expression and the WNT/LRP5 signaling cascade. BMP-2 mediates extracellular matrix mineralization by osteoblasts partly through the induction of a WNT autocrine/paracrine loop\cite{26}. BMP-2 is highly expressed in Probasco cells compared to Ace-1 cells\cite{7}. Secreted BMP-2 in Probasco CM could be an important factor stimulating bone mineralization and the markedly increased ALP activity in CM treated-calvaria. In addition, decreased ALP activity in Probasco+DKK-1 CM treated calvaria compared to calvaria treated with Probasco CM suggests that the regulation of BMP-2 on ALP is dependent on WNT signaling. qPT-PCR analysis revealed no difference in the mRNA expression of \textit{ALPL} in Probasco CM-treated calvaria compared to control, which indicates that the induction of ALP activity may be regulated at the post-translational level.

Runx2 and osterix regulate the differentiation of osteoblasts. Runx2 or osterix-deficient mice have a complete absence of bone formation and osteoblasts, indicating that these two transcription factors are essential for bone development and osteoblastogenesis\cite{27}. BMP-2 can markedly upregulate the expression of \textit{RUNX2} through the activation of Smad signaling\cite{28,29}, and the expression of \textit{OSX} is also under the control of BMP-2\cite{30}. By comparing calvaria treated with CM from osteoblastic cell lines (LuMa and Probasco) to calvaria treated with CM from osteolytic cell line (Ace-1), the mRNA expression of \textit{RUNX2} and \textit{OSX} was significantly upregulated in both LuMa and Probasco CM-treated calvaria. Since there is high expression of BMP-2 in LuMa and Probasco cell lines compared to Ace-1 cells, and both \textit{RUNX2} and \textit{OSX} are target genes of BMP-2, it’s speculated that BMP-2 plays a role in canine PCa-induced bone formation.

Bone integrin-binding sialoprotein (IBSP) is the second major sialoprotein in bone. IBSP is synthesized by osteoblasts and its expression is tightly associated to mineralization. IBSP expression marks a late stage of osteoblastic differentiation and an early stage of matrix mineralization\cite{31}. The significant upregulation of IBSP was observed in Probasco CM-treated calvaria, but not in LuMa CM-treated calvaria compared to Ace-1 CM-treated calvaria, which may reflect that Probasco was more potent on the induction of mineralization compared to LuMa. Current studies on IBSP often focus on the role of IBSP in osteotropic malignancies, such as multiple myeloma, breast, prostate and lung cancer, and tumor progression and bone metastasis\cite{32,33}. Little is known about factors secreted by these malignancies that induce the expression of IBSP in bone cells.

In conclusion, Probasco, an osteoblastic canine prostate cancer cell line, induces bone formation and mineralization in mouse calvaria \textit{in vitro} by the stimulation of maturation and mineralization of osteoblasts, and the inhibition of osteoclastogenesis. Both bone modeling and remodeling are involved in Probasco-induced bone formation, and the induction depends on the WNT signaling pathway. The Probasco cell line provides a valuable and unique model for understanding the molecular mechanisms of prostate cancer-induced osteoblastic bone metastases.

\textbf{DECLARATIONS}

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Authors’ contributions
Designed overall projects; manuscript preparation and edit: Rosol TJ, Yuan S
Performed experiments: Yuan S, Hellmann DE
Analyzed and interpreted data: Yuan S, Rosol TJ, Kantake N
Technical and material support: Kantake N, Elshafae SM

Availability of data and materials
The material and data that support the findings of this study are available from the corresponding author upon request.

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Conflicts of interest
All authors declared that there are no conflicts of interest.

Ethical approval and consent to participate
All mice used were housed in the animal facilities at Ohio University. Animal experiments were performed with the approval of the Institutional Animal Care and Use Committee (IACUC) (protocol number: 18-H-030) and in accordance with the Ethical Guidelines for Animal Experimentation of Ohio University.

Consent for publication
Not applicable.

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