Bone Marrow Stromal Cells Create a Permissive Microenvironment for Myeloma Development: A New Stromal Role for Wnt Inhibitor Dkk1

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Abstract
The rapid progression of multiple myeloma is dependent upon cellular interactions within the bone marrow microenvironment. In vitro studies suggest that bone marrow stromal cells (BMSC) can promote myeloma growth and survival and osteolytic bone disease. However, it is not possible to recreate all cellular aspects of the bone marrow microenvironment in an in vitro system, and the contributions of BMSCs to myeloma pathogenesis in an intact, immune competent, in vivo system are unknown. To investigate this, we used a murine myeloma model that replicates many features of the human disease. Coinoculation of myeloma cells and a BMSC line, isolated from myeloma-permissive mice, into otherwise nonpermissive mice resulted in myeloma development, associated with tumor growth within bone marrow and osteolytic bone disease. In contrast, inoculation of myeloma cells alone did not result in myeloma. BMSCs inoculated alone induced osteoblast suppression, associated with an increase in serum concentrations of the Wnt signaling inhibitor, Dkk1. Dkk1 was highly expressed in BMSCs and in myeloma-permissive bone marrow. Knockdown of Dkk1 expression in BMSCs decreased their ability to promote myeloma and the associated bone disease in mice. Collectively, our results show novel roles of BMSCs and BMSC-derived Dkk1 in the pathogenesis of multiple myeloma in vivo. Cancer Res; 72(9); 2183–9. ©2012 AACR.

Introduction
The bone marrow is well known to support tumor growth and osteolytic bone disease in multiple myeloma. However, the contributions of this specialized microenvironment during early stages of myeloma development are poorly understood, largely because of the need for in vivo experiments that accurately replicate the complexity of tumor–host cell interactions. To address this, we have exploited the unique opportunities provided by the well-characterized 5T Radl myeloma model, in which transplantation of 5T myeloma cells into recipient mice of the specific C57Bl/KaLwRij strain results in propagation of myeloma with many features of the human disease (1). One important feature of the Radl model is that 5T myeloma cells will only grow in syngeneic KaLwRij mice, and not in closely-related C57Bl6 mice. Because there is an emerging role for host fibroblasts in the establishment and progression of solid tumors (2–4), we hypothesized that fibroblasts, or bone marrow stromal cells (BMSCs), from myeloma-permissive KaLwRij mice may promote myeloma development in environments that are otherwise not permissive for myeloma. Furthermore, increasing evidence for elevated Dickkopf 1 (Dkk1) in patients with monoclonal gammopathy of undetermined significance (MGUS; refs. 5–7) suggests a potential role for host-derived Dkk1 in the early stages of myeloma. We used a BMSC line derived from KaLwRij mice to determine the function of BMSCs and BMSC-derived Dkk1 in myeloma pathogenesis in vivo.

Materials and Methods

Cell culture
5TGM1-GFP myeloma cells were cultured as previously described (8). 14M1 BMSCs were originally isolated from KaLwRij mice bearing 5T myeloma. Bone marrow was flushed from long bones, and cells cultured in Dulbecco’s Modified Eagle’s Medium (DMEM) with 10% fetal calf serum (FCS) and L-glutamine. Media were changed every 3 to 4 days, and adherent cells were maintained in culture. The cell line was generated by continuous growth in vitro, with no requirement for cellular or viral immortalizing genes. Aliquots of early passage cells were frozen in 10% dimethyl sulfoxide/90% FCS and stored in liquid nitrogen. ST2 BMSCs were purchased from Riken cell Bank and maintained in DMEM media with 10% FCS and L-glutamine.

Stable cell lines
14M1 BMSCs were transfected with 1 μg Dkk1shRNA or scrambled control (Santa Cruz Biotechnology) using short
hairpin RNA (shRNA) transfection reagent and standard protocols. Cells were cultured in DMEM with 10 μg/mL puromycin. Knockdown of Dkk1 expression was confirmed by ELISA, and cells were continuously cultured in puromycin to maintain knockdown. ST2 BMSCs were transfected with 250 ng full-length Dkk1 cDNA (OriGene) or empty vector control according to the manufacturer’s instructions. Cells were cultured in DMEM with 800 μg/mL neomycin. Overexpression of Dkk1 was confirmed by ELISA, and cells were continuously cultured in neomycin to maintain expression.

**Differentiation studies**

BMSCs were cultured in either control media, osteoblast differentiation media (10 mmol/L β-glycerophosphate, 50 μg/mL ascorbic acid, 10 ng/mL BMP-2), or adipocyte differentiation media (10 nmol/L dexamethasone, 5 μg/mL insulin) for 7 days (osteoblast differentiation) or 10 days (adipocyte differentiation). Osteoblasts were identified by alkaline phosphatase staining, using a BCIP/NBT substrate kit and adipocytes by Oil Red-O staining.

**ELISA, western blotting, and flow cytometry**

The concentration of Dkk1 in conditioned media or sera was measured by ELISA, according to the manufacturer’s instructions (R&D Systems). Sera were assayed for monoclonal mouse IgG2b paraprotein as described (8). Western blotting and flow cytometry were conducted as previously described, and outlined in Supplemental Data (9).

**In vivo studies**

Studies were conducted with weight-matched, 8- to 10-week-old female C57BL/KaLwRijHsd (Harlan Netherlands B.V.) or C57Bl6 mice (Harlan U.S.). Studies were approved by the Institution of Animal Care and Use Committee at Vandebilt University. C57Bl6 or C57Bl/KaLwRij mice were intravenously inoculated with (i) 10^6 5TGM1-GFP, 10^5 ST2 BMSCs, 5^6 or 5^6 were inoculated intravenously with (i) 10^6 5TGM1-GFP, 10^5 ST2 BMSCs, 5^6 14M1 BMSCs, 10^6 14M1-Dkk1KD BMSCs, 5^6/14M1 BMSCs had many characteristics of both the original KaLwRij model and accumulation of plasma cells within the paraprotein as described (8). Western blotting and flow cytometry were conducted as previously described, and outlined in Supplemental Data (9).

**Results and Discussion**

The importance of the host microenvironment in myeloma is exemplified by the 5T myeloma model, in which KaLwRij mice possess a unique host microenvironment that is permissive for 5T myeloma growth (10). We were able to induce myeloma development in nonpermissive C57Bl6 mice by inoculating 5TGM1 myeloma cells in the presence of BMSCs isolated from KaLwRij mice (14M1 BMSCs). Importantly, myeloma induced by coinoculation of myeloma cells and 14M1 BMSCs had many characteristics of both the original syngeneic KaLwRij model, and human myeloma. Mice showed a time-dependent increase in serum paraprotein concentrations similar to the original KaLwRij model and accumulation of myeloma cells in bone marrow and spleen (Fig. 1A and B). Osteolytic bone disease is a major feature of multiple myeloma, and histomorphometric and microCT analysis confirmed a significant increase in osteolytic lesion number and osteoclast number, meanwhile showing a significant decrease in trabecular bone volume and osteoblast number (Fig. 1C, Supplementary Fig. S1). Histologic analysis revealed that C57Bl6 mice inoculated with 5TGM1 myeloma cells in the presence of 14M1 BMSCs displayed a striking similarity to the original KaLwRij myeloma model, with replacement of bone marrow by myeloma cells, loss of trabecular bone, and erosion through cortical bone (Fig. 1D). Inoculation of 5TGM1 myeloma cells alone into C57Bl6 mice did not result in the development of myeloma, with no significant accumulation of plasma cells within the bone marrow or evidence of osteolytic bone disease. There is increasing evidence from solid tumors to support a role for fibroblasts in tumor progression (11, 12). In contrast to.
previously published studies, in which the fibroblasts were enhancing tumor growth, we have identified a role for fibroblasts or BMSCs whereby the tumor is unable to grow unless the fibroblasts are present, highlighting the critical role of these cells in myeloma pathogenesis.

Coinoculation of 14M1 BMSCs with 5TGM1 myeloma cells resulted in myeloma development in an otherwise nonpermissive microenvironment. In contrast, coinoculation of 5TGM1 myeloma cells and 14M1 BMSCs significantly increased serum IgG2b concentrations (A) and tumor burden within bone marrow and spleen (B). Coinoculation of 14M1 BMSCs and 5TGM1 myeloma cells resulted in the development of myeloma bone disease, associated with an increase in osteolytic lesions. D, histologic sections showed the similarity between C57Bl6 mice coinoculated with 14M1 BMSCs and 5TGM1 myeloma cells as compared with the original 5T model, in which KaLwRij mice are inoculated with 5TGM1 myeloma cells. Arrows indicate areas where tumor cells have eroded through cortical bone. *P < 0.05; **P < 0.01; ***P < 0.001 as compared with 5TGM1.

14M1 BMSCs were originally isolated from myeloma-bearing KaLwRij mice, and so represent host stromal cells from the permissive KaLwRij bone marrow microenvironment. 14M1 BMSCs expressed both vimentin and fibroblast-specific protein-1, indicative of fibroblasts (Fig. 2B). Immunocytochemistry for FSP-1 showed expression by all cells, confirming that 14M1 BMSCs are comprised entirely of fibroblasts. (Supplementary Fig. S2). Further characterization showed no osteogenic or adipogenic differentiation potential, commonly seen in BMSCs (Fig. 2C). In vivo analysis confirmed engraftment of 14M1 BMSCs, with flow cytometric analysis detecting approximately 13% BMSCs in the bone marrow following coinoculation of 14M1 cells labeled with DsRedII with myeloma cells (Supplementary Fig. S3). Inoculation of 14M1 BMSCs was unable to support the growth of human myeloma cells and had no effect on the proportion of B, T, or NK cells suggesting that 14M1 cells were not immunosuppressive (Supplementary Fig. S4). To further characterize the in vivo effects of 14M1 BMSCs promote myeloma development in nonpermissive C57Bl6 mice. C57Bl6 mice were inoculated with either 10⁶ 5TGM1-GFP cells alone, 5 × 10⁵ 5TGM1-GFP + 5 × 10⁵ 14M1 BMSCs or vehicle control (PBS). Coinoculation of 5TGM1 myeloma cells plus 14M1 BMSCs significantly increased serum IgG₂b concentrations (A) and tumor burden within bone marrow and spleen (B). Coinoculation of 14M1 BMSCs and 5TGM1 myeloma cells resulted in the development of myeloma bone disease, associated with an increase in osteolytic lesions. D, histologic sections showed the similarity between C57Bl6 mice coinoculated with 14M1 BMSCs and 5TGM1 myeloma cells as compared with the original 5T model, in which KaLwRij mice are inoculated with 5TGM1 myeloma cells. Arrows indicate areas where tumor cells have eroded through cortical bone. *P < 0.05; **P < 0.01; ***P < 0.001 as compared with 5TGM1.
BMSCs, 5TGM1 myeloma cells and 14M1 BMSCs were coinoculated into KalwRij mice, which are already permissive for 5T myeloma development. Although 5TGM1 myeloma cell inoculation resulted in tumor growth and osteolytic bone disease, the coinoculation of 14M1 BMSCs did not further increase tumor burden in bone marrow or spleen (Fig. 2D) or osteolysis (data not shown), suggesting that the 14M1 BMSCs were not acting directly on the myeloma cells to increase myeloma growth.

The inability of 14M1 BMSCs to promote myeloma in an already permissive environment suggested that the cells might instead act directly on the host microenvironment to render it

Figure 2. Characterization of 14M1 BMSCs in vitro and in vivo. C57Bl6 mice were inoculated with either $10^5$ 5TGM1-GFP cells alone, $5 \times 10^5$ ST2 BMSCs, or vehicle control (PBS). KaLwRij mice were inoculated with 5TGM1 myeloma cells as a positive control. A, coinoculation of ST2 BMSCs with 5TGM1 myeloma cells had no effect on tumor burden, as measured by serum IgG2bk. B, Western blot analysis showed expression of vimentin and FSP-1 by 14M1 BMSCs. C, in contrast to ST2 BMSCs, 14M1 BMSCs did not differentiate into osteoblasts or adipocytes. KalwRij mice were inoculated with either $10^5$ 5TGM1-GFP cells alone, $5 \times 10^5$ 5TGM1-GFP + $5 \times 10^5$ ST2 BMSCs, or vehicle control (PBS). D, coinoculation of 14M1 BMSCs with 5TGM1 myeloma cells had no effect on tumor burden, as measured by serum IgG2bk.

Figure 3. 14M1 BMSCs have direct effects to decrease bone formation. C57Bl6 mice were inoculated with $10^5$ 14M1 BMSCs and sacrificed after a period of 4 weeks. Inoculation of 14M1 BMSCs had no effect on osteoclast numbers, but significantly decreased osteoblast numbers (A), rates of bone formation (B), and increased serum concentrations of Dkk1 (C). D, 14M1 BMSCs secrete high concentrations of Dkk1. *, $P < 0.05$; **, $P < 0.01$, as compared with control.
permissive for myeloma. To investigate this, we inoculated C57Bl6 mice with 14M1 BMSCs alone. After 4 weeks, we detected a 23.4% decrease in trabecular bone (control, 7.38 ± 1.50 vs. 14M1 BMSCs, 5.65 ± 2.13), associated with a significant decrease in osteoblast number and bone formation rates, but no change in osteoclast number (Fig. 3A and B). In control experiments, inoculation of ST2 BMSCs was found to have no effect on trabecular bone volume, bone cell number, or bone formation rates (data not shown). In support of a decrease in trabecular bone volume contributing to myeloma development, KaLwRij mice were also found to have a significant reduction in trabecular bone volume (Supplementary Fig. S5) when compared with age- and sex-matched C57Bl6 mice.

Because a major mediator of osteoblast suppression in myeloma bone disease is Dkk1 (13–15), we measured Dkk1 serum concentrations, and observed a significant increase in C57Bl6 mice inoculated with 14M1 BMSCs, as compared with either C57Bl6 or C57Bl6 mice inoculated with ST2 BMSCs (Fig. 3C). 14M1 BMSCs were found to express high concentrations of Dkk1 in vitro (Fig. 3D), and investigation of Dkk1 expression in myeloma-permissive KaLwRij mice showed a significant 160.8% increase in Dkk1 serum concentrations (P < 0.05) and increased expression of Dkk1 in Kalwrij BMSCs (Supplementary Fig. S6), suggesting a possible role for Dkk1 in myeloma-permissive microenvironments. In support of this, immunohistochemistry revealed an increase in Dkk1 expression within the bone marrow of mice inoculated with 14M1 BMSCs alone, as compared with nontumor-bearing mice (Supplementary Fig. S7). Therefore, we determined the contribution of BMSC-derived Dkk1 in myeloma pathogenesis in vivo. 14M1 BMSCs were stably transfected with Dkk1 shRNA or scrambled control, and knockdown of Dkk1 expression was confirmed by ELISA (Fig. 4A). Knockdown of Dkk1 expression had no effect on the growth rate of 14M1 BMSCs in vitro (data not shown). C57Bl6 mice were inoculated with 5TGM1 myeloma cells in the presence of either 14M1 scrambled control BMSCs or 14M1 DKK1 knockdown BMSCs. Those mice inoculated with 5TGM1 cells in the presence of 14M1 DKK1 knockdown BMSCs had a significant reduction in tumor burden, as measured by serum IgG2b titers, when compared with mice coinoculated with 5TGM1 myeloma cells and 14M1 scrambled control BMSCs (Fig. 4B). Knockdown of Dkk1 expression in the 14M1 BMSCs prevented the reduction in trabecular bone volume observed in those mice coinoculated with 5TGM1 myeloma cells and 14M1 BMSCs (Fig. 4C).
and this was associated with an increase in rates of bone formation (Fig. 4D). ST2 BMSCs were stably transfected to overexpress Dkk1, however coinoculation of these BMSCs with 5TGM1 myeloma cells in C57Bl6 mice did not result in myeloma (Supplementary Fig. S8). Dkk1 plays a key role in the osteoblast suppression of myeloma bone disease, but the majority of studies to date have focused upon the role of myeloma-derived Dkk1 (13, 15). Stromal cells and osteoblasts are known sources of Dkk1 (16, 17), and this study provides compelling evidence for a key role of stromal-derived Dkk1 to promote myeloma development. There is increasing evidence to suggest that cells of the bone marrow microenvironment are altered in MGUS, and that Dkk1 is increased in patients with MGUS, associated with a loss of trabecular bone (5–7). In support of this, our studies show increased host-derived Dkk1 in myeloma-permissive KalwRij mice that is associated with a significant reduction in trabecular bone volume. Furthermore, we have discovered a novel role of BMSCs to promote myeloma establishment and progression in an otherwise nonpermissive environment. This is mediated, in part, through stromal-derived Dkk1, however the inability of overexpression of Dkk1 in normal BMSCs to support myeloma development reveals that Dkk1 alone is not permissive for myeloma development and suggests a requirement for additional factors. Furthermore, although Dkk1 expression is undetectable in vitro, STGM1 myeloma cells do express Dkk1 in vivo, raising the possibility that tumor-derived Dkk1 may compensate for a loss in host-derived Dkk1 (Supplementary Fig. S7). It is intriguing to speculate whether the effect of 14M1 BMSCs is associated with a generalized osteoblast suppression that renders the bone marrow microenvironment favorable for myeloma. In support of this, recent studies have implicated changes in the bone marrow, associated with bone loss, that contribute to the initiation of hematopoietic cell diseases (18–20). An anti-human Dkk1 neutralizing antibody (BHQ880) is currently in phase I/II clinical trials in multiple myeloma, for the treatment of the osteolytic bone disease. This study identifies a novel role for stromal-derived Dkk1 in myeloma pathogenesis, by rendering the host microenvironment favorable for myeloma growth and development. Therefore, targeting Dkk1, such as with a neutralizing antibody, may have additional applications, including preventing myeloma progression during early stages of disease development. Collectively, our studies reveal the importance of host-derived Dkk1 and BMSCs in the pathogenesis of myeloma in vivo, highlighting the importance of changes in the local bone marrow microenvironment in the early stages of myeloma development.

Disclosure of Potential Conflicts of Interest

No potential conflicts of interest were disclosed.

Authors’ Contributions

Conception and design: J.A. Fowler, G.R. Mundy, C.M. Edwards
Development of methodology: J.A. Fowler, C.M. Edwards
Acquisition of data: J.A. Fowler, S.T. Lwin
Analysis and interpretation of data: J.A. Fowler, C.M. Edwards
Writing, review, and/or revision of the manuscript: J.A. Fowler, C.M. Edwards
Administrative, technical, or material support: J.A. Fowler
Study supervision: J.A. Fowler, C.M. Edwards

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