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Immune infiltration and PD-L1 expression in the tumor microenvironment are prognostic in osteosarcoma

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Osteosarcoma patient survival has remained stagnant for 30 years. Novel therapeutic approaches are needed to improve outcomes. We examined the expression of Programmed Death Ligand 1 (PD-L1) and defined the tumor immune microenvironment to assess the prognostic utility in osteosarcoma. PD-L1 expression in osteosarcoma was examined in two patient cohorts using immunohistochemistry (IHC) ($n = 48$, $n = 59$) and expression was validated using quantitative real time PCR ($n = 21$) and western blotting ($n = 9$). IHC was used to determine the presence of tumor infiltrating lymphocytes and antigen-presenting cells (APCs) in the tumor. Expression of PD-L1 was correlated with immune cell infiltration and event-free-survival (EFS). The 25% of primary osteosarcoma tumors that express PD-L1 were more likely to contain cells that express PD-1 than PD-L1 negative tumors (91.7% vs 47.2%, $p = 0.002$). Expression of PD-L1 was significantly associated with the presence of T cells, dendritic cells, and natural killer cells. Although all immune cell types examined were present in osteosarcoma samples, only infiltration by dendritic cells (28.3% vs. 83.9%, $p = 0.001$) and macrophages (45.5% vs. 84.4%, $p = 0.031$) were associated with worse five-year-EFS. PD-L1 expression was significantly associated with poorer five-year-EFS (25.0% vs. 69.4%, $p = 0.014$). Further studies in osteosarcoma are needed to determine if targeting the PD-L1:PD-1 axis improves survival.

Osteosarcoma is the most common primary malignant bone tumor in children and young adults^{1–3}. Although there have been great advances in the treatment of osteosarcoma, survival rates have remained stagnant over the past three decades. With the introduction of multimodal therapy, including surgery and chemotherapy, 5-year event-free survival (EFS) has remained less than 70%, which has remained unchanged over the past few decades^{4,5}. Patients with metastatic or relapsed disease have particularly dismal outcomes, with a less than 30% 5-year EFS^{6–10}. There is an urgent need for new therapeutic approaches to improve outcomes for these patients.

In recent years, numerous studies have sought to define the interplay between the immune system and tumor, with the potential to target this interaction for therapeutic intervention. In many malignancies, both innate and adaptive immune cells play a role in the tumor microenvironment, with communication between natural killer (NK) cells, antigen presenting cells (APCs) such as macrophages and dendritic cells (DC), and lymphocytes allowing for effective tumor control¹¹. Aberrations in the function of immune cells can lead to ineffective tumor surveillance and tumor elimination. Abnormal immune function can also contribute to enhanced tumor growth by increasing local inflammation, creating an environment conducive to tumor growth^{12–14}.

Immune cells are initially attracted to tumor cells by the presence of tumor specific antigens. Cancers with higher mutational loads present greater numbers of tumor specific neoantigens and are frequently associated with

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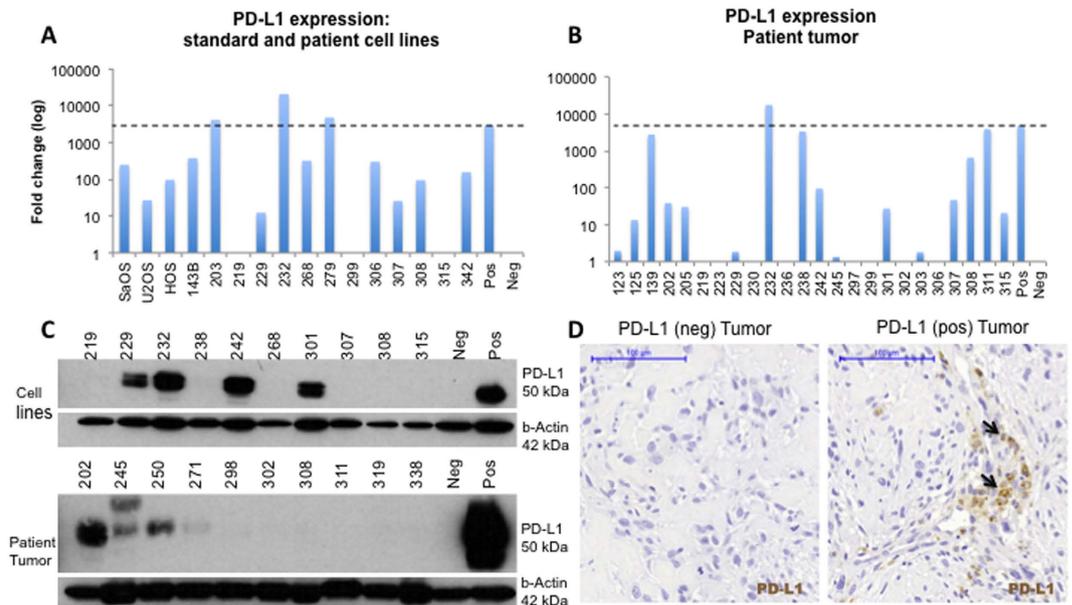


Figure 1. PD-L1 is expressed in some primary osteosarcoma tumors. (A) PD-L1 mRNA expression in cell lines derived from primary osteosarcoma tumors. The PD-L1 expressing breast cancer cell line MDA-MB-231 was used as the positive control. (B) PD-L1 mRNA expression in osteosarcoma tumor specimens. Dashed lines represent the level of PD-L1 expression in the positive control. (C) Western blot validation confirms that PD-L1 is expressed in primary osteosarcoma tumors (30%) (D) IHC on selected osteosarcoma tumor specimens show areas of heterogeneous PD-L1 expression. Scale bar represents 100 μ m. Neg = negative, pos = positive.

robust immune infiltration^{11,15}. Osteosarcoma demonstrates significant genetic complexity, with the majority of tumors displaying loss of both p53 and RB^{16–18}. In addition, 33% of primary osteosarcoma shows evidence of chromothripsis, or chromosome shattering^{19,20}, and over 50% exhibit kataegis, or localized areas of hypermutation^{18,21}. The high mutational load in osteosarcoma, along with the regular interaction between immune cells and bone cells in normal tissue, suggests that osteosarcoma may be an immunogenic tumor and evasion of the immune response may be an important component of its pathogenesis. Although previous studies have suggested a potential role for immune cells in osteosarcoma, to date no study has provided a comprehensive picture of the immune microenvironment^{22,23}.

Tumors escape immunosurveillance by expressing immune checkpoints, such as programmed death ligand-1 (PD-L1)^{24,25}. The interaction between PD-L1, expressed on tumor cells and its receptor, programmed death-1 (PD-1), expressed on immune cells, leads to immune cell apoptosis, anergy, and tolerance. Recent studies have demonstrated that cancers like melanoma, despite being highly immunogenic tumors, are not effectively cleared, partially due to their ability to express PD-L1²⁶. A number of studies have already demonstrated that blockade of the PD-L1:PD-1 interaction in patients with Hodgkin lymphoma, non-small-cell lung cancer, and melanoma leads to improved outcomes^{27,28}, and anti-PD-1 antibodies have been FDA approved for the treatment of melanoma^{29–31}. In addition, studies have reported that tumor response to PD-L1 or PD-1 inhibition is directly related to the level of PD-L1 expression and lymphocytic infiltration of the tumor^{32–35}.

Previous studies have begun to examine the role of PD-L1 in osteosarcoma. Utilizing quantitative real-time PCR (qRT-PCR), PD-L1 mRNA expression was associated with lymphocyte infiltration²³. Subsequent studies, utilizing immunofluorescence (IF), reported PD-L1 was not expressed in primary specimens and was only expressed in metastatic tissue²². Further studies examining PD-L1 blockade in a mouse model of osteosarcoma showed initial regression of the tumor followed by growth of PD-L1 antibody resistant clones. In response to treatment with anti-PD-L1 antibody, tumor cells downregulated PD-L1 and upregulated CD80 and CD86. In addition, neighboring T-cells exhibited decreased expression of PD-1 and increased expression of CTLA4, suggesting that alternate immune checkpoints may play a role in osteosarcoma resistance to PD-L1 blockade³⁶. Combination therapy targeting PD-L1 and CTLA-4 showed control of osteosarcoma growth in the majority of tumors in a mouse model^{22,36}. In this study, we determined the potential prognostic and therapeutic utility of PD-L1 by examining its expression and characterizing the immune microenvironment of human osteosarcoma.

Results

PD-L1 is expressed in primary osteosarcoma tissue. PD-L1 expression was quantified in osteosarcoma patient and standard cell lines by qRT-PCR. Three quarters of patient derived cell lines, all the osteosarcoma standard cell lines (HOS, HOS-143b, SaOS, and U2OS), and two thirds of tumor specimens demonstrated expression of PD-L1 mRNA greater than the negative MCF7 control cell line; however, only a small subset expressed PD-L1 at the same level as the positive control, breast cancer cell line MDA-MB-231. Specifically, three of the sixteen cell lines (203, 232, and 279), 19% (Fig. 1A), and four of the 24 tumors (311, 139, 232, 238), 17%, were highly positive for PD-L1 mRNA, with expression levels near or exceeding the positive control (Fig. 1B).

	Osteosarcoma patients (n = 33)
Median age (SD)	16 (5)
Huvos Response	
I/II standard	22
III/IV poor	11
Metastasis	
Present	4
Absent	29

Table 1. Patient demographic information.

PD-L1 protein expression was validated with western blotting: 3 of 10 patient samples (30.0%) and 4 of 10 osteosarcoma patient derived cell lines (40%) demonstrated detectable levels of PD-L1 protein (Fig. 1C). mRNA and protein expression did not correlate as some tumors that had mRNA expression as detected by qPCR but no detectable protein expression, specifically cell lines 307 and 308 and tumors 308 and 311.

PD-L1 protein expression was subsequently examined using IHC (Fig. 1D) on an osteosarcoma TMA containing 54 patient samples (Table S1). PD-L1 expression was quantified as the percentage of positive tumor cells on the slide compared to the tumor volume. Seven percent of all specimens on the TMA stained positive for PD-L1, with the majority of positive tumors demonstrating less than 25% staining. No significant differences in PD-L1 expression were seen between primary specimens and metastatic specimens (6.8% vs 15.4%, $p = 0.24$). Due to the low percentage of PD-L1 positive primary tumors detected by IHC compared to qPCR and western blotting, PD-L1 staining was performed on a second cohort of samples using whole slides, which included increased area of osteosarcoma tissue per sample. In the whole slide cohort, 12 patients (25.0%) had positive PD-L1 expression, of which eight had associated clinical data (Table 1). All of the positive samples demonstrated <25% PD-L1 staining.

PD-L1 positive tumors are associated with the presence of tumor infiltrating immune cells in the tumor microenvironment.

The presence of six groups of tumor infiltrating immune cells was examined by IHC in whole slides (Figures S1–S6). Presence of the immune cell was classified as a tumor that had greater than 1% of the tumor volume consisting of positively stained cells. Tumor infiltrating lymphocytes (TILs) examined included multiple T-lymphocyte subsets as well as B-lymphocytes. Additionally, slides were stained for the presence of macrophages, DCs, and PD-1 positive cells. Ninety eight percent of all osteosarcoma tumors examined demonstrated infiltration by at least one immune cell type, including 18.8% that demonstrated infiltration by all immune cell types assessed (Fig. 2).

PD-L1 positive tumors were associated with infiltration by multiple immune cell types as compared to PD-L1 negative tumors, which demonstrated infiltration by fewer immune cell types. Tumors expressing PD-L1 were significantly more likely to demonstrate infiltration by all immune cell types compared with PD-L1 negative tumors (50.0% vs 5.6%, $p = 0.002$). Tumors expressing PD-L1 compared to those that were PD-L1 negative were more likely to demonstrate infiltration by CD3+ T-cells (91.7% vs 55.6%, $p = 0.002$), CD56+ NK cells (100% vs 66.7%, $p = 0.02$), CD68+ macrophages (91.7% vs 50%, $p = 0.02$), and CD1a+ DCs (83.3% vs 30.6%, $p = 0.002$). In addition, immune cells were significantly more likely to express PD-1 in PD-L1 positive tumors compared with PD-L1 negative tumors (91.7% vs 47.2%, $p = 0.002$) (Table S2).

Expression of PD-L1 is consistent within the tumor mass. IHC indicated that PD-L1 expression and immune cell infiltration is heterogeneous within the whole slide samples examined. Two osteosarcoma tumor maps, containing slides from multiple areas of tumor, were examined in order to assess the consistency of PD-L1 expression and tumor infiltrating immune cells across whole tumors. Four slides were selected from the map for patient A, containing tissue from four locations across the tumor mass (Fig. 3A). Six slides were selected from the map for patient B, containing tissue from three locations across the tumor mass and three locations containing the resection margin (Fig. 3B).

Patient A's tumor was PD-L1 negative across all sections examined (Fig. 3C) and patient B's tumor was PD-L1 positive in five out of six sections, with only one region located outside of the tumor mass demonstrating negative PD-L expression (Fig. 3D). In both tumors, presence of specific tumor infiltrating cells varied only slightly across multiple sections of the tumor, with the exception of CD56+ NK cells and CD8+ T-regulatory cells, which were universally present. Overall, the PD-L1 positive tumor contained more tumor infiltrating immune cell types than did the PD-L1 negative tumor.

PD-L1 expression or APC infiltration is associated with event free survival. In order to assess the prognostic relevance of PD-L1 and tumor infiltrating immune cells in osteosarcoma, Kaplan Meier curves were generated. EFS, with an event being defined as relapse or death, was calculated based on presence or absence of PD-L1 protein and presence of specific immune cells as determined by IHC on both TMA and whole slide samples. PD-L1 expression was associated with significantly poorer five-year EFS in the whole slide samples (25% vs. 69.4%, $p = 0.01$). In contrast, although the TMA cohort showed a trend for poorer survival with PD-L1 expression (33.3% vs. 71.4%, $p = 0.18$), this result did not achieve statistical significance (Fig. 4A,B).

In the whole slide specimens, tumor infiltration by APCs was associated with poorer 5 yr-EFS, including CD1a+ DCs (28.3% vs. 83.9%, $p = 0.001$) and CD68+ macrophages (45.5% vs. 84.4%, $p = 0.032$) (Fig. 5). Infiltration by the other immune cells assessed was not associated with differences in the 5-year EFS. (Figures S1–S6).

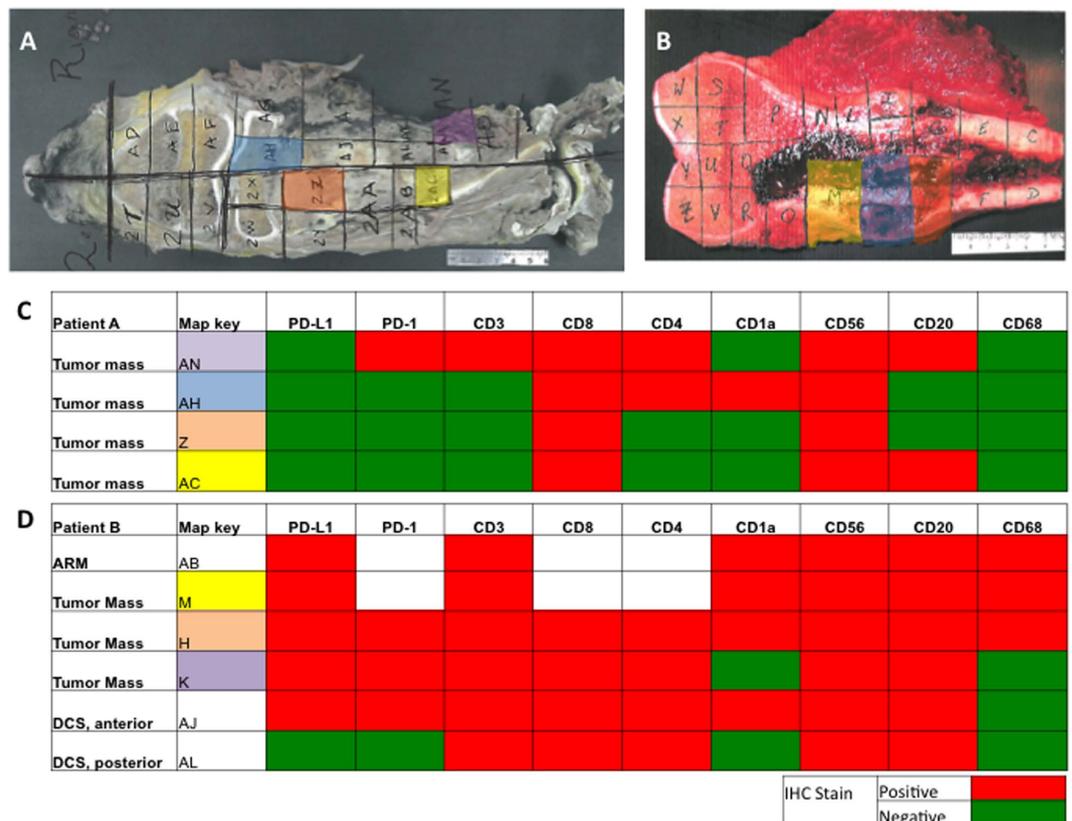


Figure 3. PD-L1 expression is consistent throughout the tumor mass. (A,C) All four slides from the tumor mass of patient A did not demonstrate PD-L1 staining. Few immune cells are present in this tumor map. (B,D) All of the slides from the tumor mass of patient B stained PD-L1 positive with the exception of one section outside the tumor mass (not shown). Numerous immune cells are present throughout all sections of this tumor map. ARM = anterior resection margin, DCS = distal cross section.

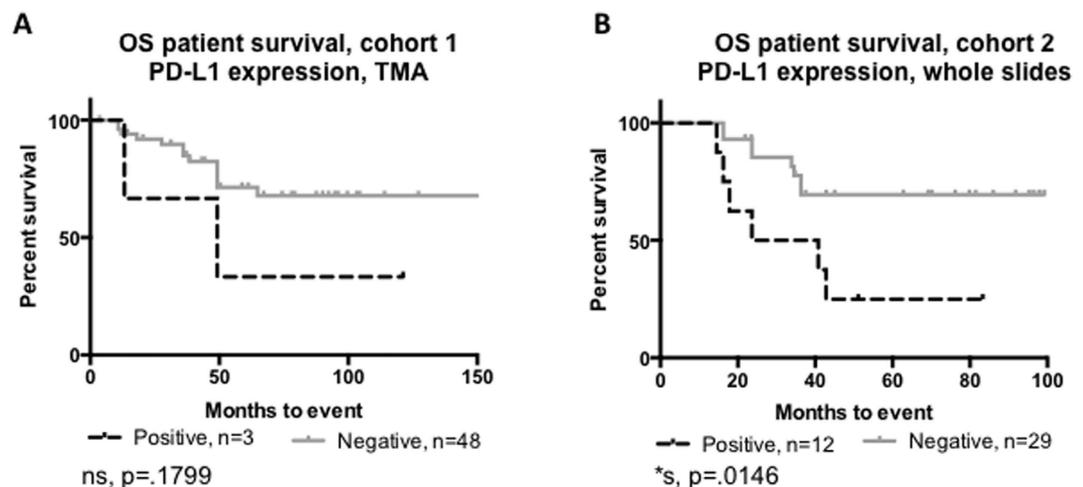


Figure 4. PD-L1 expression is prognostic in osteosarcoma. (A) PD-L1 expression in the TMA trends with poorer EFS. (B) PD-L1 expression in the whole slide specimens is significantly associated with poorer EFS.

Tumor associated macrophages (TAMs) are present in multiple solid tumors, including breast, gastric, and ovarian cancers³⁸. Multiple studies have shown that TAMs play an important role in breast cancer pathogenesis by promoting angiogenesis, tumor cell invasion, and migration^{39–42}. Unlike the presence of TAMs in other solid tumors, the presence of CD14 positive immune cells, which include macrophages, monocytes, neutrophils, and dendritic cells⁴³, was previously found to be associated with decreased metastasis but increased angiogenesis in osteosarcoma⁴⁴. The presence of M1 macrophages, which are pro-inflammatory but not pro-tumor, has been

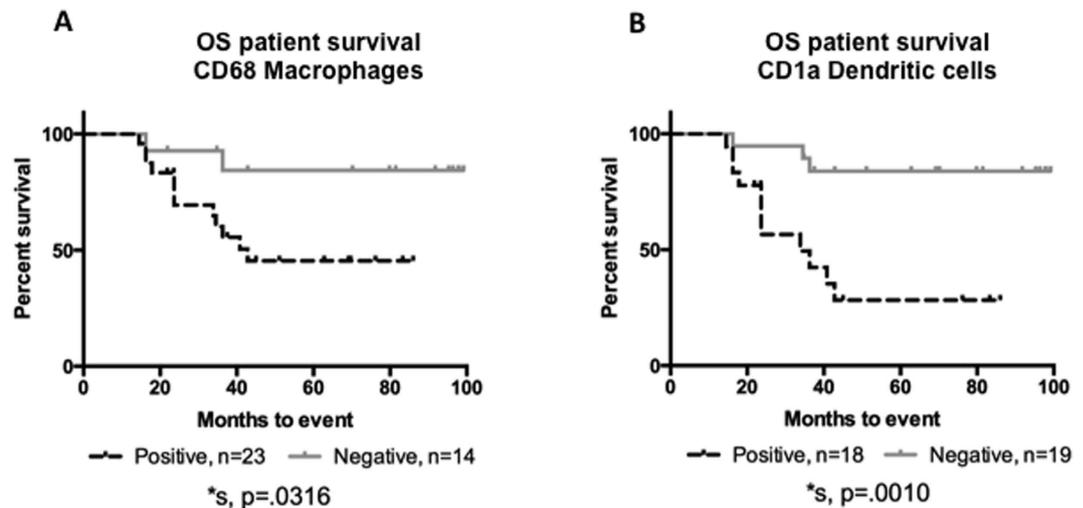


Figure 5. APC infiltration is prognostic in osteosarcoma. Presence of APCs, specifically (A) CD68 positive macrophages or (B) CD1a positive dendritic cells are significantly associated with poorer EFS in osteosarcoma. Over 50% of osteosarcoma samples have APCs present.

associated with better outcomes in osteosarcoma. Treatment of osteosarcoma with muramyl-tripeptide, which appears to activate monocytes to become M1 macrophages, causing a cytokine cascade⁴⁵, leads to osteosarcoma cell death *in vitro*⁴⁶.

Although previous studies may suggest that the opposite may be the case, we found that the presence of CD68+ TAMs in primary tissue may be significantly associated with worse survival in osteosarcoma. In previous studies, by capturing both monocytes and mature (M1 and M2) macrophages with CD14 staining, the role of mature macrophages may have been confounded. In our current study CD68 staining captures both pro-inflammatory M1 and pro-tumor M2 macrophages. In order to clarify the role of macrophages in osteosarcoma, future studies should correlate the presence of CD68+/CD80+ M1 and CD68+/CD163+ M2 macrophages to osteosarcoma survival.

It is also possible that osteoclasts, a subset of CD68+ bone specific macrophages, may impact osteosarcoma patient survival. Presence of osteoclasts in osteosarcomas have been linked to conflicting outcomes in past studies⁴⁷. Although a recent study shows that pulmonary osteosarcoma metastases have fewer osteoclasts⁴⁸, earlier work has shown that patients with aggressive osteosarcomas often have increased osteoclast activity⁴⁹ and that inhibition of osteoclast function via a RANK-ligand antagonist leads to favorable outcomes in mice⁵⁰.

Presence of a second APC, CD1a+ DCs, is also associated with poor survival in osteosarcoma. With the recent advent of DC vaccines as a cancer treatment, multiple groups have shown that DC vaccines have varying levels of effectiveness in animal models of osteosarcoma⁵¹⁻⁵⁴. It has been previously shown that human osteosarcoma cells in culture inhibit the ability of DC to present antigens⁵⁵. Furthermore, human bone does not contain lymphatic vessels, implying that DCs cannot function in bone as they do in other tissues⁵⁶. However, osteosarcomas frequently have a soft tissue component which may contain lymphatic vessels. The relevance of lymphatic vessels in osteosarcoma is not clear, however, the primary mode of metastasis in osteosarcoma is via hematogenous spread, with fewer than 4% of patients displaying lymph node metastasis^{57,58}. It seems that despite showing the potential of vaccines in animal models, DCs may be unable to function properly in human osteosarcoma.

A major limitation of this study is the small number of samples available in this single institution analysis of the tumor immune microenvironment in osteosarcoma. Our sample size was further limited by the lack of associated clinical data for a subset of tumors. Although the non-significant correlative outcome results in the TMA may be explained by the limited sample assessed in each core, and the limited sample number, these represent two different patient cohorts and we cannot rule out the possibility that the prognostic correlation is purely a random occurrence in a small patient population. Further studies are necessary to validate that these results can be recapitulated in a larger multi-institutional cohort.

In conclusion, we have shown that tumor infiltration by either CD68+ TAMs or CD1a+ DCs may be associated with poorer survival in osteosarcoma. Furthermore, PD-L1 is expressed in a subset of primary osteosarcoma samples at the time of biopsy, and it may be prognostic for increased mortality. PD-L1 expression is associated with TILs and APCs, including PD-1 positive immune cells. Due to its expression in a significant number of osteosarcoma tumors and an abundant presence of immune cells, the PD-L1:PD-1 axis may be a promising target for immunotherapy in osteosarcoma, particularly given osteosarcoma known genetic complexity. Further evaluation of PD-L1 protein expression is warranted in a larger cohort of patients with osteosarcoma. Currently a phase II study of pembrolizumab (anti-PD-1 antibody) in patients with advanced sarcomas, including osteosarcomas, is ongoing, the results of which may be informative.

Materials and Methods

Cell Culture and Human Materials. Both osteosarcoma standard cells lines (HOS, HOS-143b, SaOS, and U2OS) and isolated cells were cultured in Eagle minimum essential medium supplemented with 20% fetal bovine

serum (FBS) and antibiotics. PD-L1 negative MCF7 breast cancer cells, PD-L1 positive MDA-MB-231 breast cancer cells, and PD-L1 negative 3T3 cells were cultured in Dulbecco's modified Eagle's medium supplemented with 10% FBS. Primary osteosarcoma cultures were generated using standard collagenase disaggregation of surgical specimens from seven osteosarcoma patients, as has been described previously⁵⁹. All cell lines were grown in 5% CO₂ humidified atmosphere at 37 °C.

Tumors collected from two cohorts of osteosarcoma patients were used to construct a TMA or whole slides, as previously described^{60,61}. Tissue was decalcified overnight, formalin-fixed, and paraffin embedded. Four to five-micron thick sections were cut from representative areas of the tumor and used for immunohistochemistry (IHC). In total, the TMA contained cores from 62 samples representing 51 unique patients, of which 54 cores were usable (Table S1). The cores that detached from the slide during processing were deemed unusable. Slides with an area of 50–135 cm² were constructed using tumors collected from a second cohort of patients (Table 1), and were used to validate the TMA results. This second cohort included 48 total patients, of which 37 patients had associated clinical outcomes and 29 had clinical outcomes and additional demographic information.

The Ethics Committees and Institutional Review Boards at Montefiore Hospital, Memorial Sloan Kettering Cancer Center, and the Center for Cancer Research approved the study. Written informed consent was obtained from the patients and their parents/guardians prior to tissue collection. All procedures were conducted in accordance with guidelines provided by the Ethics Committees and Institutional Review Boards.

Protein extraction and Western Blotting. Protein was extracted from 15 frozen patient tumors using RIPA buffer (ThermoScientific). The tumors were collected in accordance with an IRB approved research protocol with patient/guardian informed consent as described previously⁶². Proteins were denatured in a reducing sample buffer and run on a 12% tris gel using tris-glycine buffer. Proteins were transferred to a nitrocellulose membrane, incubated with 5% milk block, probed with antibody for PD-L1 and subsequently with either a goat anti-rabbit HRP conjugated secondary antibody (Cell Signaling) diluted in 5% bovine serum albumin. Horse radish peroxidase conjugated secondary antibodies were detected using autoradiography using chemoluminescence (Amersham Enhanced Chemiluminescence Prime Western Blotting Detection Reagent). 3T3 cells and MDA-MB-231 cells were used as a negative and positive control for PD-L1 expression, respectively.

RNA extraction and quantitative PCR (qPCR). RNA was extracted from flash frozen osteosarcoma tumor samples from 15 patients using Trizol reagent (Invitrogen) and converted to cDNA using Superscript II transcriptase (Life Technologies). qPCR was performed using commercially available TaqMan primers for PD-L1 (Life Technologies, Hs01125301_m1). Relative gene expression was normalized to an internal control, GAPDH, and calculated using the $2^{-\Delta\Delta CT}$ method. MCF7 and MDA-MB-231 cell lines were used as a negative and positive control for PD-L1 mRNA expression, respectively. Due to lack of detectable amplification of PD-L1 in the negative control, the CT value for MCF7 was set to 40 for calculations.

Tumor microarray (TMA) construction. Tumors, collected from osteosarcoma patients were used to construct a TMA, as has been described previously^{60,61}. The TMA contains cores from osteosarcoma tumors collected at the time of biopsy and definitive surgery, as well as from metastatic tissues. Tumor specimens obtained at the time of biopsy or definitive surgery were all obtained from the patients' primary site of disease. A pathologist reviewed the tumors in order to confirm the diagnosis of osteosarcoma. The TMA was constructed using 1-millimeter (mm) cores that were acquired from the formalin-fixed paraffin embedded tissue block from each individual tumor. Five-micron thick sections were cut and used for subsequent analysis⁶³.

Tumor Maps. Multiple slides were selected from tumor maps that were previously generated for clinical purposes using osteosarcoma specimens obtained with parental consent from two patients enrolled in an IRB-approved bio-banking protocol at Montefiore Medical Center. Samples were analyzed from multiple locations within the tumor mass and surrounding tissue at the time of definitive surgery following neoadjuvant chemotherapy. A set of slides was generated for each patient and this was related to the specimen map, which provided each slide's location within the tumor.

Immunohistochemistry. Immune staining was optimized for proper antigen retrieval and antibody concentrations using immune tissue for positive controls and brain tissue for negative controls. Osteosarcoma tumor slides were baked at 60 °C for one hour and antigen retrieval was performed using either tri-citrate buffer, pH 6.0 or tris-EDTA buffer pH 8.0 at 100 °C for 20 minutes. Slides were de-paraffinized using xylene, rehydrated along an ethanol gradient, and blocked using dual endogenous enzyme-blocking reagent (Dako) and a subsequent normal goat serum block (Santa Cruz). Slides were incubated overnight at 4 °C in primary antibody followed by a one hour room-temperature incubation with secondary antibody. Antibody concentrations and information can be found in Supplementary Table 1. Antigen expression was detected by addition of VECTASTAIN Elite ABC Kit (Vector Laboratories) and DAB (3,3-diaminobenzidine) HRP substrate (Vector Laboratories). Slides were counterstained with hematoxylin (Harris). Human placenta was used as positive control of PD-L1 expression. A full list of antibodies and their positive controls for all antibodies can be found in Supplementary Table 1.

Quantification of IHC. Slides were scanned and graded by three independent reviewers without previous knowledge of patient outcomes or demographics. Slides were graded with a quartile score based on the percentage of positive staining cells present in the core/slide (0 = 0% staining, 1 = <25%, 2 = 25–50%, 3 = 50–75%, and 4 = >75%). If scores between the three graders were discordant the average, rounded to the nearest whole integer, was used. A tumor with a score of 1 or higher, representing $\geq 1\%$ staining, was considered positive for PD-L1 expression or immune cell infiltration. Images of IHC staining were obtained at 20 \times magnification using the Panoramic Scanner and captured using Panoramic Viewer (3DHISTECH Ltd).

Statistical analysis. Patient demographic data from each cohort was reported as frequencies for categorical variables and means with standard deviations for continuous variables. The association between the expression of PD-L1 and the presence of immune cells was assessed using Fisher's exact test or chi square analysis. Kaplan Meier curves were generated assessing the association between EFS and PD-L1 expression or immune cell infiltration, and the Log-rank (Mantel-Cox) test was utilized (GraphPad Prism software). Survival curves were generated using expression thresholds of negative, defined as no staining (grade 0) and positive defined as at least 1% staining (grade 1, 2, 3, or 4).

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Author Contributions

P.K. participated in study design, performed the study, and prepared the manuscript. M.E.R. and J.G. participated in study design, data analysis, and manuscript editing. J.M.C., S.P., A.P. and M.A.F. helped with data analysis. D.S.G. and B.H.H. participated in study design and manuscript editing. R.G. and X.Z. conceived of study and participated in its design and coordination, and manuscript editing. All the authors read and approved this manuscript.

Additional Information

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