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Immunosensitization of Tumor Cells to Dendritic Cell-Activated Immune Responses with the Proteasome Inhibitor Bortezomib (PS-341, Velcade)\textsuperscript{1}

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Proteasome inhibition results in proapoptotic changes in cancer cells, which may make them more sensitive to immune effector cells. We established a murine model to test whether the proteasome inhibitor bortezomib could sensitize established B16 melanoma tumors to dendritic cell (DC)-activated immune effector cells. Day 3-established s.c. B16 tumors had significantly decreased tumor outgrowth when treated with a combination of bortezomib and DC, regardless of whether the DC were loaded or not with a tumor Ag. In vivo Ab-depletion studies demonstrated that the effector cells were NK and CD8\textsuperscript{+} cells, but not CD4\textsuperscript{+} cells. NF-κB nuclear transcription factor assay and gene-expression profiling of B16 treated with bortezomib was consistent with inhibition of NF-κB target genes leading to a proapoptotic phenotype. In vitro lytic assays demonstrated that TNF-α, but not perforin, Fas-ligand, or TRAIL, was responsible for bortezomib-sensitized B16 cytotoxicity. In conclusion, the proteasome inhibitor bortezomib can pharmacologically sensitize tumor cells to the lytic effects of DC-activated immune effector cells. \textit{The Journal of Immunology}, 2006, 176: 4757–4765.

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We hypothesized that the pharmacological inhibition of tumor proteasome enzymatic activity would sensitize tumor cells to immune effectors. To test this hypothesis, we developed an in vivo murine model treating established B16 melanoma with a combined therapy of dendritic cell (DC)-based immunization/activation and the proteasome inhibitor bortezomib. Tumor growth was significantly delayed in mice treated with the combined therapy as opposed to either therapy alone. Each group received either AdVMART1/DC or untransduced DC, in both cases in conjunction with bortezomib following the same schedule as in A, C. Lack of adverse effects of bortezomib on MART-1-specific T cells. Splenocytes from mice treated with or without AdVMART1/DC combined with bortezomib, as indicated, were harvested on day 14 after study initiation. RBC-depleted splenocytes were restimulated in vitro with EL4 cells stably transfected with MART-1 (EL4-MART1) or parental cells not expressing MART-1 (EL4) for 48 h, after which cells were added to an IFN-γ ELISPOT plate. D. Treatment of established murine tumors with combined DC and bortezomib. Kaplan Meier plot of percent tumor-free mice over time combining data from eight independent replicate studies (p < 0.0001).

Materials and Methods

**Mice**

C57BL/6, and CD8 α-chain knockout mice (CD8KO) in a C57BL/6 back- 
ground (B6.129S2-Cd8α KO B6.129S2-Cd8α KO backcrossed >28 generations), were purchased from The Jackson Laboratory and were bred and kept under defined-flora pathogen-free conditions at the Association for Assessment of Laboratory Animal Care-approved Animal Facility of the Division of Experimental Radiation Oncology (University of California, Los Angeles, CA; UCLA). Mice were handled in accordance with the UCLA animal care policy.

**Cell lines and MHC surface staining**

B16-F0 (B16), a murine melanoma, EL4, a murine lymphoma, and Yac-1, a murine lymphoma, were obtained from the American Type Culture Collection (ATCC). B16 and EL4 were maintained in vitro in DMEM (Invitrogen Life Technologies), and Yac-1 in RPMI 1640 (Invitrogen Life Technologies), in both cases supplemented with 10% FCS (Gemini) and 1% (v/v) penicillin, streptomycin, and amphotericin (Gemini). Generation of EL4 (MART1) stable transfectants has been previously described (13). Stably transfected cells were maintained in vitro under constant G418 se- 
lection (0.5 mg/ml; Invitrogen Life Technologies). Determination of sur- 
face expression of MHC class I molecules was performed by cell surface staining with a PE-conjugated anti-H2-Kd Ab (BD Pharmingen). Staining was analyzed by flow cytometry.

**Reagents**

Bortezomib (Velcade), a dipeptide boronic acid proteasome inhibitor, was purchased or was a gift (in some experiments) from Millennium Pharmacuticals. Bortezomib was prepared as a stock solution in 0.9% saline. For
administration to animals, stock solutions were thawed and further diluted in 0.9% saline to a dilution to yield 0.02 mg/0.1 ml (1 mg/kg/mouse) for i.p. injection. Purified hamster anti-mouse Fas mAb (Jo2) and recombinant mouse TNF-α were purchased from BD Pharmingen. Stock solutions of reagents were routinely prepared in PBS or medium as appropriate.

Proteasome function assay

Proteasome function was measured as described previously (14) with some minor modifications. Briefly, B16 melanoma cells were treated with or without 150 nM bortezomib for 3 h. The cells were then washed with 1× PBS (pH 7.2), then with buffer I (50 mM Tris (pH 7.4), 2 mM EDTA, 5 mM MgCl₂, 2 mM ATP), and pelleted by centrifugation. Homogenization buffer (50 mM Tris (pH 7.4), 1 mM DT, 5 mM MgCl₂, 2 mM ATP, 250 mM sucrose) was then added and cells underwent three freeze-thaw vortex cycles with liquid nitrogen and a 37°C water bath. Cell debris was next removed by centrifugation at 4°C, 1,000 g for 15 min, and supernatant was transferred to new Eppendorf tubes. Twenty micrograms of protein of each sample was diluted with buffer I to a final volume of 200 μl. The fluorogenic proteasome substrates SucLLVY-MCA (chymotrypsin-like; Sigma-Aldrich), Z-Leu-Leu-Leu-AMC (Calbiochem), and Boc-Val-Leu-Lys-AMC (trypsin-like; Sigma-Aldrich) were dissolved in DMSO and added in a final concentration of 80 μM (0.8% DMSO). Proteolytic activity was monitored continuously by the release of the fluorescent group 7-amido-4-methylcoumarin (AMC) measured in a fluorescence plate reader (Spectrafluor (Tecan) and Spectra Max Gemini XS (Molecular Devices), 37°C) at 380/460 nm. In control experiments using MG-132, the chymotryptic activity of the lysates was always inhibited by >90%, indicating that the observed cleavage activity was mainly based on proteasome function.

Cell viability assay

Cell viability upon treatment with bortezomib was determined by the MTT assay, as described (15). A total of 25 × 10⁴ B16 cells/well was incubated in 96-well plates in the presence or in the absence of bortezomib at the specified concentrations. The blue MTT formazan precipitate was dissolved in 100 μl of MeSO₄, and the absorbance values at 565 nm were determined on a multiwell plate reader. Murine melanoma B16 cell cultures used in this study were free of mycoplasma, tested using a Mycoplasma PCR ELISA kit (Roche Diagnostic).

Recombinant adenoviruses

The replication-deficient adenoviral vector AdVMART1 is a previously described E1-deleted vector based on human type 5 adenovirus (16). It contains the 400-bp human MART-1 cDNA driven by the CMV early enhancer/promoter.

Preparation of DC and adenoviral transduction

DC were differentiated from bone marrow progenitor cells derived from C57BL/6 mice by in vitro culture in murine GM-CSF (100 ng/ml; Amgen) and murine IL-4 (500 Units/ml; R&D Systems) as described by Inaba et al. (17) with minor modifications (16). DC were harvested as loosely adherent cells, washed twice in PBS (Mediatec), and immediately prepared for injection in 0.2 ml of PBS/mouse. When indicated, DC were transduced in RPMI 1640 with 2% FCS transduction media at a multiplicity of infection of 100–200 viral PFU/each DC. Transduction was conducted for 2 h at 37°C, washed twice in PBS and resuspended in 0.2 ml of PBS/animal for injection (16, 18).}

Animal studies

B16 cells used for tumor challenge were obtained from single-cell suspensions of progressively growing tumors in syngeneic mice to avoid the confounding effects of the presentation of medium- and serum-deprived epitopes. To generate single-cell suspensions, tumors were surgically removed, decapsulated, and minced. Minced tumors underwent enzymatic digestion (2 h with DNase 0.1 mg/ml; Sigma-Aldrich) and collagenase D (1 mg/ml; Roche Applied Sciences) in 50 ml of AIM-V medium (In-vitrogen Life Technologies). Viable cells were washed three times in PBS and resuspended in 0.1 ml of PBS/animal (16, 18). Tumors were injected s.c. into the left flank (1 × 10⁶ cells/injection) on day 0. Injected cells were >70% viable, as determined by trypan blue exclusion. On days 3 and 10, mice received untransduced DC or DC transduced with AdVMART1 (0.5–1.0 × 10⁶ cells/injection) s.c. in the right flank. Bortezomib was administered at 1 mg/kg/mouse i.p. on days 7, 10, and 14. Each treatment group typically contained five mice.

FIGURE 2. CD8 and NK cells mediate the antitumor effects of DC plus bortezomib. Groups of five mice were depleted of CD4 cells (A), CD8 cells (B), or NK cells (C) using specific Abs as described in Materials and Methods. B16 tumors were then implanted s.c. in a flank and mice received combined DC plus bortezomib therapy. In all studies, tumor outgrowth after administration of DC or bortezomib alone was not different from control mice at the p = 0.05 significance level, while DC plus bortezomib was significantly different in all studies. Depletion of CD4 cells did not abrogate the antitumor effects of the combination, while depletion of CD8 and NK cells did abrogate the antitumor activity.

Immune cell subset depletion in vivo

In vivo Ab ablation of CD8⁺ (clone 2.3), CD4⁺ (clone GK1.5) T cell subsets, and NK1.1⁺ (clone PK136) cells was performed by i.p. injection of 100–200 μg of purified endotoxin-free Ab/mouse/injection (BioExpress) on days −5, −3, and −1 before tumor inoculation, and every 6 days thereafter. Monitoring of successful depletion was performed by flow cytometry on splenocytes harvested on the day of tumor inoculation for CD4⁺ and CD8-depleting Abs. NK depletion was confirmed by NK cell activity microcytotoxicity assays using the NK-sensitive Yac-1 cell line as target cells and IL-2-activated lymphocyte-activated killer (LAK) cells as effectors (Yac-1 assay) (19).

DC phenotyping

Bone marrow-derived DC were evaluated after an 8-day culture in GM-CSF and IL-4 and were treated for 18 h with a dose titration of bortezomib at 200, 100, 10, and 1 nM. DC were then stained with H2-Kb PE anti-mouse (MHC class I molecule), CD86 PE anti-mouse, CD80 FITC anti-mouse, CD80 PE anti-mouse, CD80 FITC
anti-mouse, CD11c PE anti-mouse (all from BD Pharmingen), and I-A<sup>b</sup> FITC anti-mouse (MHC class II molecule; Caltag Laboratories). Stained cells were analyzed by flow cytometry. DC were also stained with annexin V (BD Pharmingen) according to the manufacturer’s instructions.

**Cytokine profile by ELISPOT**

For the detection of MART-1 Ag-specific responses, splenocytes depleted of RBC by hypotonic lysis, cultured in vitro with irradiated EL4 or EL4 (MART-1) at a 25:1 ratio for 48 h in the presence of 10 U/ml IL-2, were added in duplicate 3-fold dilutions to 96-well mixed cellulose plates (MultiScreen filtration system; Millipore) precoated with anti-IFN-γ Ab (BD Pharmingen). For Ag-nonspecific responses, RBC-depleted splenocytes were added directly to IFN-γ-coated plates with or without the addition of PMA (Sigma-Aldrich). Plates were incubated for 24 h at 37°C. Spots were developed using a secondary biotin-labeled Ab and counted under a dissecting microscope as previously described (20).

**Lactate dehydrogenase (LDH) in vitro cytotoxicity assay**

The LDH-based CyTox 96 Assay (Promega) was used to determine in vitro cytotoxicity as described previously. Briefly, 5 × 10<sup>3</sup> to 7 × 10<sup>3</sup> cells/sample, in quadruplicate, were cultured into a 96-well flat-bottom microtiter plate (Costar) and cultured at low serum concentration (0.1% FBS) 18 h before each treatment. The medium was then changed to complete DMEM with 5% heat-inactivated FBS. B16 or EL4 cells were then cultured in the presence of bortezomib at the specified concentrations. After incubation for each different experimental condition, released LDH into the culture supernatants was measured with a 30-min coupled enzymatic assay, which results in the conversion of a tetrazolium salt (INT) into a red formazan product that is read at 490 nm in an automated plate reader (Axxora). The assay plates were incubated for 24 h at 37°C in 5% CO<sub>2</sub>. After incubation for each different experimental condition, released LDH into the culture supernatants was measured with a 30-min coupled enzymatic assay, which results in the conversion of a tetrazolium salt (INT) into a red formazan product that is read at 490 nm in an automated plate reader (Emax; Molecular Devices). Percentage cytotoxicity was calculated using the spontaneous release-corrected OD as follows: percent cytotoxicity = (OD of experimental well/OD of maximum release control well) × 100.

**NF-κB-related subunits DNA-binding assay**

The nuclear DNA-binding activity of the NF-κB-related subunits p50, p52, p65 (RelA), c-Rel, and RelB was assessed using a nonisotopic quantitative DNA-binding assay (BD TransFactor NF-κB family kit; BD Clontech). A total of 20 μg of nuclear extract (prepared as we have reported previously. Ref. 22) from B16 mouse melanoma cells, treated in the absence or presence of the proteasome inhibitor bortezomib at 10 nM, were bound to microwells coated with NF-κB consensus oligonucleotide. Wells were incubated with specific Abs against p50, p52, p65 (RelA), c-Rel, or RelB, followed by the secondary HRP-conjugated and horseradish peroxidase-conjugated streptavidin, as per manufacturer’s recommendations. Chemiluminescence signals were visualized by autoradiogram and their relative intensity was assessed by densitometric analysis of the digitized image (ScanAlalyze Microarray Image Analysis, [http://rana.lbl.gov/Eisen-Software.htm](http://rana.lbl.gov/Eisen-Software.htm)) after housekeeping gene-based normalization, background subtraction, and statistical significances determination (GEArray software; SuperArray Bioscience).

**Results**

**Bortezomib treatment permits tumor suppression of established B16 melanoma by DC-based immunotherapy**

C57BL/6 mice immunized with DC engineered to express MART-1 melanoma Ag are partially protected from a challenge of B16 melanoma, which express MART-1. This model has been extensively studied by our group (13, 19–21, 23, 24). MART-1-engineered DC can partially or completely protect mice from a subsequent B16 challenge, but we have never been successful in having any impact on even microscopically established tumors (13).

We set up a treatment model to determine whether systemic bortezomib treatment of microscopically established B16 would render them more sensitive to DC-based immunotherapy. Mice received 10<sup>6</sup> B16 implanted s.c. in one flank. On days 3 and 10, they received 10<sup>6</sup> immature DC (7-day cultures of GM-CSF/IFN-γ) transduced with an adenoviral vector expressing MART-1 (AdVMART1, multiplicity of infection 1:100) administered s.c. in the contralateral flank. Mice also received bortezomib at 1 mg/kg i.p. on days 3, 7, 10, and 14. Neither AdVMART1/DC nor bortezomib alone had any impact on B16 tumor development or growth kinetics (Fig. 1A). However,
their combination significantly delayed tumor appearance and retarded tumor growth kinetics. Three replicate studies were performed with similar results.

Bortezomib-facilitated DC tumor suppression is Ag nonspecific

We and others have shown that DC-based immunotherapy can induce both adaptive and innate antitumor responses (19, 24–29). Administration of unmodified GM-CSF/IL-4 DC to B16-bearing, bortezomib-treated mice provided a comparable degree of tumor suppression as was observed with MART-1-transduced DC (Fig. 1B). We confirmed by IFN-γ ELISPOT analysis that the administration of AdVMART1/DC expanded MART-1-specific T cells in these mice (Fig. 1C), but these tumor Ag-specific effectors were not required for the antitumor effect of DC-bortezomib combined therapy. Pooled data from eight replicate studies using a total of 157 mice demonstrated that tumor appearance was consistently delayed by \(~10\) days, but essentially all mice developed tumors (Fig. 1D).

NK and CD8+ T cells, but not CD4+ T cells, are required

CD8, CD4, or NK cells were depleted by Ab before tumor implantation and DC-bortezomib treatment. As shown in Fig. 2, CD4 depletion had no impact on tumor protection, while depletion of CD8 or NK1.1-positive cells completely abrogated tumor suppression. In summary, bortezomib treatment of B16-bearing mice facilitates tumor suppression by CD8 and NK cells driven by immature DC.

B16 melanoma treated with bortezomib acquires a proapoptotic phenotype and is sensitized to TNF-α but not perforin or FasL killing

Our hypothesis is that bortezomib sensitizes B16 to immune effectors. We sought to confirm the work of others that tumor cells acquired a proapoptotic phenotype with proteasome inhibition. In preliminary studies, we determined that the IC_s of B16 treated for 24 h with bortezomib was 20 nM in an in vitro MTT assay (Fig. 3A). Also, a range of concentrations of bortezomib (1, 10, 50, 100, 200 nM) was tested to define its ability to induce apoptotic death in B16 cells in vitro. There was a concentration- and time-dependent increase in annexin V and propidium iodine single- and double-positive cells, demonstrating apoptotic death of B16 with increasing bortezomib exposure (Fig. 3B).

Using a sublethal concentration of 10 nM of bortezomib, we sought to determine the effects of proteasome inhibition on the differential gene expression pattern from bortezomib-untreated (Fig. 4A) and -treated (Fig. 4B) mouse melanoma B16 cells, using a mouse pathway-focused gene expression microarray corresponding to 96 key apoptosis-related genes. The following transcripts were up-regulated with bortezomib (Fig. 4B, solid boxes): Bcl2/adenovirus E1B 19kDa-interacting protein 1, NIP3 (Bnip3), a proapoptotic member of the Bcl-2 family (30, 31); transformed mouse 3T3 cell double minute 2 (Mdm2), a p53-related apoptosis regulator with both pro- and antiapoptotic activities (32); and nucleolar protein 3 (Nol3), an apoptosis repressor with a caspase recruitment domain (33). Four genes were found to be significantly down-regulated by bortezomib (Fig. 4B, dotted boxes): baculoviral inhibitor of apoptosis protein (IAP) repeat-containing 5 (Birc5, Survivin), a critical antiapoptosis regulator (34); replication protein A3 (Rpa3), a DNA stability/cell cycle checkpoint regulator (35); TNF-receptor-associated factor 4 (Traf4), a NF-κB-associated antiapoptosis gene (36); and transformation-related protein 53 (Trp53), a cell cycle-related anti-apoptosis gene (37).

Immune effector cells, including T, NK, and LAK cells, kill targets either through the release of perforin/granzyme B resulting in target cell death in short-term (4 h) in vitro assays, or the engagement of the TNFR family of death receptors that require longer assays to detect cytotoxicity. Both pathways activate the extrinsic and intrinsic apoptotic machinery, eventually resulting in the activation of effector caspases and target cell apoptotic death. We tested whether bortezomib would sensitize target cells to perforin-mediated killing in 4-h chromium release assays. We tested the lytic activity of DC and IL-2-stimulated lytic cells against B16, with or without bortezomib treatment, compared with the lytic activity against Yac-1 as a positive control. Splenocytes from DC-administered mice, regardless of exposure to high concentrations of IL-2 in vitro to generate LAK cells with NK lytic activity, had low lytic activity to B16 in a short-term microcytotoxicity assay. Only splenocytes cultured in IL-2 ex vivo had lytic activity to the
NK-preferred target Yac-1, while splenocytes from DC-administered mice did not demonstrate NK lytic activity under these assay conditions (Fig. 5A). To further confirm the lack of sensitization to perforin-mediated killing, we turned to a CD8null environment with enhanced NK responses to DC administration. We have previously reported that administration of DC to CD8KO mice results in the activation of much higher levels of NK effector cell lytic activity than when administered to wild-type mice. These NK effectors lyse B16 through the perforin pathway (19). Therefore, we administered DC to CD8KO mice and tested their ability to lyse chromated B16 targets with or without prior bortezomib treatment. Again, killing of B16 treated with bortezomib, if anything, was decreased compared with untreated B16 targets and Yac-1 (data not shown). In conclusion, no target cell sensitization by bortezomib to NK lytic activity could be detected in short-term microcytotoxicity assays, which predominantly detect perforin-mediated target cell killing.

We then tested the in vitro sensitivity to the three death receptor ligands in long-term LDH release microcytotoxicity assays. The death receptor pathway requires longer exposure than when assessing perforin-mediated killing in standard 4-h chromium release assays. We used EL4 lymphoma as a positive control for Fas, TRAIL, and TNF-α-mediated killing in this assay. Sublytic doses of FasR-activating Ab or TRAIL ligand, with or without concurrent bortezomib at 10 nM, a concentration below the IC50 for B16, did not result in significant killing of B16 (Fig. 5, C and D). On the contrary, the combination of TNF-α and bortezomib resulted in increased killing of B16 (Fig. 5B), a cell line that has been previously shown to be resistant to TNF-α-mediated killing (38). In conclusion, these functional assays suggest that bortezomib sensitizes B16 cells to killing through TNF-α receptor engagement. These findings confirm those of Murphy and colleagues (3, 4) and collectively provide circumstantial support for our immunosensitization hypothesis.

**Bortezomib has no significant impact on several immune effector functions**

An alternative hypothesis is that bortezomib enhances immune effector cell function to produce this antitumor response. We could find no evidence to support this. As already shown in Fig. 1C, the expansion and activation of Ag-specific T cell responses are unaffected by the dose and treatment schedule of the proteasome inhibitor we used. In vitro treatment of DC with bortezomib did decrease MHC class I expression, less markedly for MHC class II, and had no effect on expression of costimulatory molecules CD80 and CD86 or the adhesion molecule CD11c (Fig. 6A). There was evidence of early apoptosis in DC starting at 10 nM concentrations of bortezomib using the marker annexin V (Fig. 6B). Finally, spleen cells from bortezomib-treated mice were not enhanced nor impaired in their ability to be stimulated by PMA to produce...
IFN-γ or to generate IL-2-activated lytic NK cells (Fig. 6, C and D). In summary, there appeared to be a modest impact of the proteasome inhibitor on cells from the innate and adaptive responses in the models we studied.

Discussion

Tumor cells acquire a variety of mechanisms for resistance to apoptosis induced by immune effector cells (39, 40). The improved understanding of these pathways, together with the development of specific inhibitors for critical molecules responsible for tumor resistance, may facilitate the reversal of tumor escape from the immune system (40, 41). The ubiquitin-proteasome pathway plays an important role in regulating the degradation of proteins involved in the cell cycle, cell survival, transcriptional activation, and tumor growth. Many proteasome substrates are known mediators of pathways that are dysregulated in neoplasia. Proteasome substrates include signaling molecules, tumor suppressors, cell-cycle regulators, transcription factors, inhibitory molecules (degrading these molecules in turn activate other molecules), and antiapoptotic proteins (42). Altering the function of the proteasome and disrupting the degradation of these proteins has profound effects on cell growth and survival, particularly in rapidly dividing cancer cells, which require increased availability of growth-promoting proteins (43). Inhibiting the proteasome modulates the function of critical cellular molecules, such as the tumor suppressor p53 and the cyclin-dependent kinase inhibitors p21 and p27 (2). Its best-characterized effect is in the modulation of the transcriptional factor NF-κB. The activity of NF-κB is regulated by proteasome-mediated degradation of the inhibitor protein IκB. In response to stress such as neoplasia, IκB becomes phosphorylated and deactivated by the proteasome thus promoting NF-κB-induced prosurvival pathways. Therefore, when the proteasome is inhibited, IκB in turn inhibits the activation of NF-κB thus affecting the role of NF-κB in antiapoptosis, tumor metastasis, and angiogenesis (44). The inhibition of NF-κB via proteasome inhibition has also been shown to sensitize melanoma cells to undergo apoptosis when treated with TRAIL and other TNF family members (3, 45). Therefore, bortezomib has favorable characteristics as a potential immune sensitization drug.

Bortezomib is licensed for the treatment of multiple myeloma and is currently in trials for melanoma, ovarian, glioma, lung, lymphomas, renal cell, breast, sarcoma, colorectal, and neuroendocrine cancers (2). Preliminary data from a multicenter trial in patients with melanoma suggests that bortezomib lacks single agent activity for this malignancy (46). Melanoma is well recognized to be inherently resistant to multiple proapoptotic agents such as chemotherapy and irradiation. However, a small subset of patients with metastatic melanoma respond to a variety of immunotherapies, including high dose IL-2, whole tumor cell-based vaccines, DC vaccines, and adoptive transfer of Ag-restricted T cells (47). Lack of response to immunotherapy in the majority of patients...
with melanoma may be due to the use of inefficient immunotherapy strategies, a lack of adequate recognition of tumor cells by activated immune effector cells, or a tumor cell insensitivity to the cellular cytotoxic apoptotic signals (40).

Several laboratories have demonstrated that DC can induce antitumor Ag-nonspecific cell-mediated cytotoxicity responses (19, 25–29). In the current study, we confirm that administration of Ag-unloaded DC to mice results in tumor responses, in our case, when target cells have been cosensitized using systemic treatment with bortezomib. It is not surprising that this is an Ag-independent phenomenon in this model, because bortezomib would be expected to inhibit endogenous peptide processing by the proteasome, and thereby result in decreased surface expression of tumor Ag peptides in MHC molecules. Strategies that could enhance the participation of innate effectors in antitumor responses have the obvious advantage of marshaling a large pool of rapidly inducible, Ag-nonspecific cells. Our in vitro correlative studies suggest an increased sensitivity of bortezomib-treated B16 cells to the lytic activity of TNF-α receptor ligation, which may be the molecular basis for the immunosensitization phenomenon. However, the role of TNF-α was not tested in vivo using specific knockout mice or soluble blocking reagents, and therefore may not have a role in the combined ability of DC vaccines and bortezomib to treat established B16 tumors.

In conclusion, our studies highlight that strategies aimed at increasing the activation state of immune effectors, together with pharmacological target modulation by intracellular alteration of sensitivity to apoptotic signals, may result in increased antitumor therapeutic effects. The approach used in the current studies was aimed at testing this mechanism by activating innate effectors with DC and modulating targets with the proteasome inhibitor bortezomib. Our data demonstrate that the antitumor activity of bortezomib can be enhanced by innate immune cell activation induced by DC administration, by virtue of this small molecule targeted inhibitor to sensitize cancer cells to innate effector cell-mediated lysis.

**Disclosures**

The authors have no financial conflict of interest.

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