Cell Cycle, Cell Death, and Senescence

Bortezomib Sensitizes Human Renal Cell Carcinomas to TRAIL Apoptosis through Increased Activation of Caspase-8 in the Death-Inducing Signaling Complex

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Abstract

Bortezomib (VELCADE) could sensitize certain human renal cell carcinoma (RCC) lines to the apoptotic effects of tumor necrosis factor–related apoptosis-inducing ligand (TRAIL). Analysis of seven human RCC showed a clear increase in the sensitivity of four of the RCC to TRAIL cytotoxicity following bortezomib (5-20 nmol/L) treatment, whereas the remaining three remained resistant. Tumor cell death following sensitization had all the features of apoptosis. The enhanced antitumor activity of the bortezomib and TRAIL combination was confirmed in long-term (6 days) cancer cell outgrowth assays. The extent of proteasome inhibition by bortezomib in the various RCC was equivalent. Following bortezomib treatment, neither changes in the intracellular protein levels of various Bcl-2 and IAP family members, nor minor changes in expression of TRAIL receptors (DR4, DR5), correlated well with the sensitization or resistance of RCC to TRAIL-mediated apoptosis. However, enhanced procaspase-8 activation following bortezomib pretreatment and subsequent TRAIL exposure was only observed in the sensitized RCC in both cell extracts and death-inducing signaling complex immunoprecipitates. These data suggest that the molecular basis for bortezomib sensitization of RCC to TRAIL primarily involves early amplification of caspase-8 activity. In the absence of this increased caspase-8 activation, other bortezomib-induced changes are not sufficient to sensitize RCC to TRAIL-mediated apoptosis. Mol Cancer Res; 8(5); OF1–10. ©2010 AACR.

Introduction

Tumor necrosis factor–related apoptosis-inducing ligand (TRAIL or Apo2L) preferentially causes apoptosis in cancer cells rather than normal cells. Therefore, there has been much interest in the use of TRAIL, or other TRAIL receptor agonists such as antibodies, as a novel approach to cancer therapy. Currently, humanized agonist antibodies to TRAIL R1 (DR4), TRAIL R2 (DR5), and TRAIL itself are in clinical trials in a wide variety of cancer patients (1). Encouragingly, these agents can be administered to cancer patients in the absence of major toxicities. Nonetheless, in vitro studies indicate that only a minority of cancer cell lines undergo apoptosis in response to TRAIL. This has prompted intense interest in the identification of compounds that can sensitize cancer cells to TRAIL.

Bortezomib, a proteasome inhibitor that has been approved for the therapy of multiple myeloma, has been investigated extensively for its therapeutic effects in many types of cancer (2). Earlier studies also suggested that bortezomib could dramatically sensitize not only multiple myeloma cells (3), but also a variety of human and mouse solid tumor cells to the apoptotic effects of TRAIL in vitro (4, 5). Subsequent studies confirmed that a combination of bortezomib with TRAIL could overcome the resistance to TRAIL cytotoxicity in vitro in a wide variety of cancer cells including leukemias and lymphomas (6, 7), prostate (8, 9), colon (5), bladder (8), thyroid (10), ovarian (11), lung (12), sarcoma (13), hepatoma (14, 15), and glioma (16). Encouragingly, normal cells still seem to remain relatively resistant to TRAIL, even in combination with bortezomib (14, 15). However, the molecular events responsible for the sensitizing effect of bortezomib on TRAIL-induced apoptosis in cancer cells remain controversial and unclear.

Bortezomib has been reported to affect levels of TRAIL receptors (12, 17), c-FLIP (4), NF-κB (18), p21 and p27 (8), and pAkt (19), and these changes have been implicated in promoting tumor cell apoptosis on subsequent exposure to TRAIL. In addition, other studies suggested that the effects of bortezomib on members of the Bcl-2 family (20), particularly Bik and Bim (21), could also be crucial in promoting TRAIL apoptosis. Nonetheless, proteasome inhibition results in changes in the levels of many cellular
proteins. The extent to which some of the observed changes are directly involved in amplifying TRAIL-mediated apoptosis has not been stringently addressed. In this study, to investigate the molecular mechanisms whereby bortezomib sensitizes human renal cell carcinoma (RCC) to TRAIL, we have done a thorough molecular analysis of bortezomib sensitization using a panel of human RCC. We have used concentrations of bortezomib that could be realistically achieved in vivo. All these human RCC were quite resistant to TRAIL alone. Interestingly, some human RCC were dramatically sensitized to TRAIL-mediated apoptosis by bortezomib, whereas others remained resistant. This allowed us to systematically test which molecular changes induced by bortezomib were crucial for TRAIL sensitization and attempt to distinguish these from changes that were merely a consequence of proteasome inhibition. Results show that a crucial mechanism of bortezomib sensitization of human RCC to TRAIL-mediated apoptosis involves increased activation of caspase-8 in the death-inducing signaling complex (DISC).

Materials and Methods

Reagents and cell lines. Antibodies were purchased from the following companies: anti-FLIP Dave-2 (Kamiya Biomedical); anti-FLIP NF6 (Axxora); anti-clAP-1 AF818 and anti-Survivin AF886 (R&D Systems); anti-β-actin AC74 and anti-DR5 D3938 (Sigma); anti-DR4 32A242 (Stratagene); anti-FADD 1F7 (Millipore); anti-bax sc-493, anti-bik sc1710, anti-ubiquitin sc8017, and anti-Bcl-xS/L sc-1041 (Santa Cruz); anti–bcl-2 PC68 (EMD Chemicals, Inc.) and anti-Noxa 114C307 (EMD Chemicals, Inc.); anti-AKT, anti–phoso-AKT (Ser473; D9E), anti-bid (2002), anti-Bim C34C5, anti-Puma (4976), anti–cleaved caspase-8 18C8, and anti–cleaved caspase-9 (Asp315; Cell Signaling); anti–Mcl-2 599027, anti-XIAP 610716, p21 SX118, and anti–bak G317-2 (BD Biosciences), horseradish peroxidase (HRP)–conjugated anti-goat (Santa Cruz); HRP-anti-rabbit and HRP-anti-mouse (Thermo Fisher Scientific, Inc.); HRP-anti-rat (Cell Signaling); and rhTRAIL (Peprotech). Cell lines were obtained for nontreated cells.

Flow cytometric analysis. TRAIL receptor 1 to 4 Flow Cytometry Set (Axxora) was biotinylated using the PlatinumLink Antibody Labeling kit (Kreatech Diagnostics). For immunoprecipitation and immunoblotting, rhTRAIL (Peprotech) was biotinylated using the PlatinumLink Antibody Labeling kit (Kreatech Diagnostics). For immunoprecipitation, cells were treated overnight with either 20 nmol/L bortezomib or media treated with or without 500 ng/mL biotinylated TRAIL for 60 minutes. Cells were washed in cold PBS and lysed in immunoprecipitation (IP) DISC lysis buffer [30 mmol/L Tris-HCl (pH 7.5), 150 mmol/L NaCl, 10% glycerol, 1% Triton X-100] supplemented with the Halt Protease Inhibitor Cocktail (Thermo Fisher Scientific Inc.) and 20 μmol/L zVAD-FMK (Biomol). Samples were equalized for protein concentration and incubated with MagnaBind Streptavidin resin (Thermo Fisher Scientific, Inc.) overnight at 4°C with end over end rotation, washed thrice in IP DISC lysis buffer, and eluted in 75 μL 2x loading buffer (NuPAGE LDS Sample Buffer and NuPAGE Reducing Agent, Invitrogen). Samples for immunoblot only were lysed in lysis buffer [50 mmol/L Tris-Cl (pH 8.0), 0.5% Triton X-100, 300 mmol/L NaCl, and 5 mmol/L EDTA] supplemented with Halt Protease Inhibitor Cocktail and 20 μmol/L zVAD-FMK. Western blotting was done as previously described (22).

Growth inhibition assay. RCC (5 × 10³) were plated, incubated overnight, treated with 20 nmol/L bortezomib for 2 hours, then treated with rhTRAIL (100 ng/mL) and incubated overnight. Growth inhibition was measured using the CellTiter 96 AQueous Non-Radioactive Cell Proliferation (MTS) Assay (Promega). In some experiments, the caspase inhibitors zVAD-FMK (Axxora) or zIETD-FMK (BD Biosciences) were added 2 hours before the addition of bortezomib. For TRAIL wash kill experiments, TRAIL was added to cells for 90 minutes; all wells were washed thrice with 200 μL media to remove unbound TRAIL. Wells were refilled with fresh media and bortezomib was added for an overnight treatment.

Long-term cell survival assays. RCC (5 × 10⁴) were plated in 24-well plates, incubated overnight, treated with bortezomib for 2 hours, then with rhTRAIL (1,000 ng/mL), and incubated overnight. Treatments and dead cells were washed off and replaced with fresh media. Surviving cells were cultured for an additional 6 days, then fixed with methanol and stained with a 0.1% crystal violet solution.

CaspaSE-8 activity assays. Cells were plated in white 96-well plates at 1 × 10⁴ cells per well, incubated 24 hours, and then treated overnight with 20 nmol/L bortezomib or media. RhTRAIL (500 ng/mL) was added for 60 minutes before the addition of Caspase-8 glo reagent (Promega) containing MG-132 and readings were taken 1 hour after the addition of substrate. Duplicate samples were run in standard 96-well plates to measure growth inhibition. Luminescence values were then adjusted based on cell number and 1 luminescence unit was defined as the value obtained for nontreated cells.

Proteasome inhibition assays. Cells were plated in white 96-well plates at 1 × 10⁵ cells per well, incubated 24 hours, and then treated overnight with 1, 10, or 20 nmol/L bortezomib or media. Cells were assayed using the Proteasome-Glo Chymotrypsin-Like Cell-Based Assay (Promega) according to the manufacturer’s protocol. Duplicate samples were run in standard 96-well plates to measure growth inhibition.

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Results

Bortezomib sensitization of RCC lines to TRAIL. The sensitivity of a panel of human RCC to TRAIL in the presence or absence of bortezomib was tested after overnight incubation. Bortezomib alone resulted in a 20% to 45% reduction in the number of viable cells for these RCC. TRAIL treatment alone had little effect on viable cell number, whereas the combination of bortezomib and TRAIL treatment for A498, ACHN, TK-10, and UO-31 resulted in a 60% to 95% reduction in the number of viable cells (Fig. 1A). Therefore, for these RCC, bortezomib treatment dramatically sensitized cells to the effects of TRAIL. By contrast, CAKI-1, SN12C, and 786-0 were not sensitized to TRAIL by bortezomib and the combination resulted in a reduction in viable cell number no different to that induced by the single agents alone. Thus, we consider these lines to be resistant to bortezomib sensitization for TRAIL cytotoxicity. Interestingly bortezomib sensitized these RCC to FasL-mediated apoptosis in a similar manner (Supplementary Fig. S1). Furthermore, for the resistant RCC, bortezomib failed to sensitize over a range of TRAIL concentrations (Supplementary Fig. S2). The pan-caspase inhibitors VAD-FMK and the more caspase-8–specific zIETD-FMK both abolished the reduction of cell number that occurred in response to bortezomib plus TRAIL.
(Fig. 1B). This suggested that the combination enhanced cell death by apoptosis. In addition, because zVAD-FMK had no effect on the reduction of cell number observed in response to bortezomib alone, this suggested that the effects of bortezomib on RCC were predominantly cytostatic rather than cytotoxic. This was confirmed by Annexin V and propidium iodide staining of cells (Fig. 1C). However, the combination of bortezomib and TRAIL did not further increase the number of apoptotic cells in the resistant CAKI-1 cells.

Nonetheless, it remained important to determine whether bortezomib sensitized some RCC to TRAIL-mediated apoptosis independent of any proapoptotic effects of bortezomib alone. Indeed, we have observed that the cytotoxic effects of bortezomib as a single agent may not become apparent until 48 to 72 hours after exposure to the drug. However, using longer term cell survival assays following removal of the drugs after overnight exposure (Fig. 2), we observed that the combination of bortezomib and TRAIL resulted in an ∼100% reduction of cell number 6 days later, for at least one concentration of bortezomib (between 5 and 20 nmol/L) for A498, ACHN, TK-10, and UO-31. The transient exposure of the cells to bortezomib in this assay did initially reduce cell number due to the cytostatic effects of the drug, but most of the RCC recovered and then started proliferating. As seen in the shorter term assays, 786-0 and CAKI-1 were highly resistant to the combination treatment; so in this case, the combination of bortezomib and TRAIL did not result in increased loss of cells due to apoptosis. SN12C was highly sensitive to bortezomib alone even at quite low doses of the drug. This only became apparent in the longer term cell survival assay. This makes it difficult to determine whether there is a combined effect of bortezomib and TRAIL in reducing cell survival for SN12C.

Proteasome inhibition in various human RCC. One possible explanation for the differences in the effects of bortezomib on TRAIL sensitization for different RCC could be the differential effects of this inhibitor on proteasome enzyme activity. However, assays on proteasome activity using either a luminescence-based assay for proteasome enzyme activity (Fig. 3A), or increases in ubiquitinated proteins as assessed by Western blotting (Fig. 3B), suggested that the proteasome activity was inhibited to a similar degree in all RCC, irrespective of whether they were sensitized or remained resistant to TRAIL apoptosis. Because the proteasome inhibition appeared constant, it seemed that specific changes that occurred downstream of this inhibition were most likely to account for sensitization or resistance to TRAIL-mediated apoptosis.

Effect of bortezomib on TRAIL DISC components. One possible effect of bortezomib, which might sensitize cells to TRAIL apoptosis, could be on altering the level of cell surface expression of TRAIL receptors, or components of the TRAIL DISC. We therefore assessed the effects of bortezomib on the cell surface expression of various TRAIL receptors on some of our panel of human RCC (Fig. 4A). Interestingly, slight increases in DR4 expression were observed only on sensitized cells (ACHN and A498). However, the resistant CAKI-1 cells had high endogenous cell surface levels of DR4 expression. The expression of DR5 was increased not only in sensitized A498 and ACHN but also in the resistant CAKI-1. Therefore, there was no clear correlation between the cell surface expression of DR4 and DR5 and bortezomib sensitization of RCC to TRAIL. Cell surface levels of the decoy receptors DcR1 and DcR2 did not change in any of the RCC following bortezomib treatment (data not shown).

**FIGURE 2.** Long-term cell survival assays also show bortezomib sensitization of RCC to TRAIL apoptosis. RCC were treated overnight with bortezomib and rhTRAIL alone or in combination. The following morning drugs were washed out and replaced with fresh media. After 6 d, cells were fixed with methanol and stained with crystal violet.
The DISC is important for proximal signaling events in TRAIL-mediated apoptosis. We therefore analyzed changes in cellular levels of DISC components in sensitized and resistant RCC following bortezomib treatment (Fig. 4B). No major changes in the cellular levels of DR4, DR5, FADD, cFLIPs, or caspase-8 were observed following Western blotting in either sensitized or resistant RCC following bortezomib treatment, whereas increases in cFLIPS were observed in both sensitized and resistant. Bik increases following bortezomib treatment were only detected in two of the RCC, one sensitized and one resistant. Overall, there was no direct correlation between any changes in levels of Bcl-2 family members and sensitization to TRAIL apoptosis. Concerning the IAP family, no changes in IAP-1 or XIAP levels occurred following bortezomib treatment. Survivin decreased in both the sensitized ACHN and A498 and the resistant CAKI-1, making it unlikely that decreases in survivin alone were crucial for TRAIL sensitization. Levels of p21 increased in all RCC tested. It has also been proposed that bortezomib effects on Akt phosphorylation or caspase activation could help promote TRAIL-mediated apoptosis. However, we did not observe any effect of sensitizing doses of bortezomib on Akt phosphorylation in either sensitized or resistant cells. Furthermore, at these concentrations of bortezomib, we did not see activation of caspase-9 in most of the RCC tested. This would be consistent with the fact that the tested concentrations of bortezomib were not directly triggering any apoptosis in these cell lines. Interestingly, the SN12C line was the exception, in which bortezomib alone caused caspase-9 activation consistent with the cytotoxic effects of bortezomib observed with this cell line in long-term growth assays (Fig. 2). Because we observed no obvious changes in downstream apoptotic signaling that correlated with sensitization to TRAIL apoptosis, we then focused on determining if bortezomib induced any changes in the activation of procaspase-8.

**Bortezomib effects on procaspase-8 activation.** Measuring caspase-8 activity using a synthetic substrate assay (Fig. 6A), the combination of bortezomib and TRAIL resulted in increased caspase-8 activation in all the sensitized cells (ACHN, A498, UO-31, and TK-10) over and above that of either bortezomib or TRAIL alone. However, for the resistant CAKI-1 and 786-0, no such enhancement of caspase-8 activation was observed. In addition, Western blotting using an antibody that only recognized the cleaved forms of procaspase-8 showed increased enzyme processing...
following TRAIL exposure in response to bortezomib in the sensitized ACHN, A498, UO-31, and TK-10, but not in the resistant CAKI-1 or 786-0 (Fig. 6C). Interestingly, no processing of procaspase-8 in response to bortezomib alone was observed in any of the RCC with the exception of SN12C. This illustrates that for most RCC, the sensitizing effect of bortezomib to TRAIL can occur in the absence of any direct apoptotic effects produced by the bortezomib alone. Because caspases can be activated during the process of apoptosis, it was necessary to clarify whether observed caspase-8 activation following bortezomib treatment was due to enhanced proximal processing of procaspase-8, or merely a consequence of ongoing apoptosis. Using the MCF-7 breast cancer cell line that is resistant to apoptosis due to a lack of caspase-3, we observed that bortezomib treatment still dramatically enhanced caspase-8 activation. A slight increase in the activation of caspase-9 was also observed (Fig. 6D). No apoptosis was observed in MCF-7 cells in response to bortezomib and TRAIL treatment (data not shown). Thus, bortezomib treatment still increased caspase-8 activation even in the absence of subsequent apoptosis.

Since all of the molecular parameters that we had assessed only increased activation of caspase-8 activity correlated

**FIGURE 4.** Bortezomib causes an increase in the surface expression of some death receptors and cFLIPs in some RCC lines. A, death receptor expression on tumor cell lines was analyzed after overnight treatment with 20 nmol/L bortezomib or media alone by flow cytometry using anti-TRAIL Receptor 1 to 4 Flow Cytometry Set (Axxora). B, DISC protein expression was measured by Western blotting of cell lysates. C, MTS assay; cells were treated with TRAIL for 90 min and then washed thrice with media to remove unbound TRAIL. Media were replaced in wells and bortezomib was added for an overnight treatment. For the control plate, cells were washed with media then TRAIL and/or bortezomib were added for the overnight incubation.
with the sensitizing effects of bortezomib, we examined the DISC components following ligand-induced immunoprecipitation from the bortezomib-sensitized ACHN and resistant CAKI-1 (Fig. 6B). Higher levels of processed caspase-8 were associated with the ACHN DISC following pretreatment of the cells with bortezomib, whereas for the CAKI-1 DISC, no effect of bortezomib was observed. In the DISC of both cell lines, there seemed to be someewhat higher levels of DR5 following bortezomib in both sensitized and resistant RCC. We failed to detect DR4 in our DISC immunoprecipitates from either ACHN or CAKI-1 cells. A more pronounced increase in FADD at the DISC following bortezomib treatment of ACHN, as opposed to CAKI-1 cells, was observed. The levels of cFLIP in the immunoprecipitates were somewhat difficult to determine, due to what seemed to be nonspecific antibody binding in the region where full-length cFLIP should be detected. However, levels of the cleaved fragment of cFLIP (p43) were increased in ACHN following bortezomib treatment, whereas no change was seen with CAKI-1. This would also be consistent with an increased caspase-8 activity at the ACHN DISC. Increased cFLIP, was associated with DISC immunoprecipitates particularly with ACHN cells; however, this also could have been due to nonspecific binding events rather than TRAIL-mediated recruitment because it was also present in DISC immunoprecipitates of ACHN treated with bortezomib alone. Overall, the data suggest that bortezomib treatment in sensitized RCC promotes a more efficient DISC formation that results in enhanced activation of caspase-8. This increase in proximal apoptotic signaling then results in apoptosis of these cells. By contrast, in the resistant RCC lines, no enhanced caspase-8 activation was observed following bortezomib treatment.

Discussion

Bortezomib can sensitize certain human RCC to TRAIL-mediated apoptosis, whereas others remain resistant. We have previously shown that all these RCC can be sensitized to TRAIL-mediated apoptosis by cycloheximide (22); thus, the TRAIL apoptotic signaling pathway is intact. Consequently, the ability of bortezomib to sensitize RCC cells to TRAIL at the concentrations we used is selective. Previous studies have suggested that increases in the surface expression of the TRAIL death receptors (particularly DR5 or TRAIL-R2) could play an important role in sensitization of cancer cells to TRAIL-apoptosis following proteasome inhibition (12, 17). From our data, we conclude that increases in cell surface expression of TRAIL death receptors in response to bortezomib are not absolutely required for sensitizing RCC to TRAIL apoptosis. Several other studies have also concluded that changes in cell surface levels of DR5 do not play an essential role in the effects of bortezomib in sensitizing cancer cells to TRAIL (15, 23).

Concerning human RCC, we could find no direct evidence for an absolute requirement for the intrinsic apoptotic signaling pathway in bortezomib sensitization to TRAIL apoptosis. No obvious changes in cellular levels of most Bcl-2 family members were observed. Overall, it was noteworthy that all the bortezomib-induced changes to Bcl-2 family members (Bik and Mcl-1) as well as changes in survivin and p21 that occurred in the sensitized ACHN cells were paralleled in Caki-1 cells that remained TRAIL resistant. Furthermore, neither the overexpression of Bcl-2 nor Bcl-xL had any influence on bortezomib sensitization of ACHN to TRAIL (data not shown). In addition, Mcl-1 increased in all RCC tested. Mcl-1 is an antiapoptotic molecule that efficiently blocks the intrinsic or mitochondrial apoptosis pathway during TRAIL-mediated apoptosis (24), yet its increase in response to bortezomib in RCC does not block TRAIL apoptosis. Indeed others have reported that Mcl-1 increases following bortezomib treatment can limit the TRAIL sensitizing activity of bortezomib (25). Therefore, it is very unlikely that effects of bortezomib on
the intrinsic signaling pathway are required to promote TRAIL-mediated apoptosis in human RCC.

In contrast to our findings with RCC, several publications using other human cancer cell lines have shown that bortezomib sensitization of cells to TRAIL apoptosis involves effects on Bcl-2 family members (21, 26, 27). However, caution should be used when invoking a contribution of signaling through the intrinsic apoptotic pathway for bortezomib sensitization to TRAIL. In some of these studies, quite high concentrations of bortezomib (50-1,000 nmol/L) were used to show the sensitizing effect. In our experience, bortezomib cytotoxicity may only become apparent for most cancer cells 48 to 72 hours after exposure to the drug. Therefore, we feel it is crucial that survival of the cancer cells should be assessed 5 to 6 days after the removal of bortezomib and TRAIL. Otherwise, the increased cytotoxic effects of the bortezomib and TRAIL combination observed at early time points might just reflect an acceleration of apoptosis in cells already destined to die.

Bortezomib has also been proposed to overcome TRAIL resistance in hepatoma cells through inhibition of the phosphoinositide 3-kinase/Akt pathway (28). For human RCC, we did not observe any effect of bortezomib on levels of pAkt at the concentrations we used. In addition, inhibitors of NF-κB that did not block proteasome activity were unable to sensitize RCC to TRAIL (data not shown). This would be consistent with other studies suggesting that inhibition of NF-κB activation is not required for bortezomib sensitization of RCC to TRAIL apoptosis (29). The presence or absence of mutations in p53 in the RCC did not correlate with the effects of bortezomib on TRAIL
sensitization (data not shown), suggesting a lack of p53 involvement in the apoptotic signaling in this instance.

For RCC, the amplification of the caspase-8 signal in response to TRAIL following bortezomib treatment is the one molecular change that always correlated with enhanced apoptosis. This does not rule out the possibility that other changes in apoptosis proteins induced by bortezomib may further amplify apoptotic signaling. Indeed, the relative importance of such additional changes may vary in a cell type–specific manner. However, for RCC, the increased caspase-8 activation seems to be the crucial event. In contrast to the studies with hepatoma and glioma cells (15, 16), and previous findings from our laboratory and others (4, 30), we did not consistently observe a major reduction in cellular levels of cFLIPL following bortezomib treatment of RCC. However, the TRAIL DISC contained a significantly increased level of caspase-8 and FADD in bortezomib-sensitized RCC. In resistant RCC, there were no obvious effects of bortezomib on the composition of the DISC, following exposure to TRAIL. Our data in agreement with others (14–16) suggest that a crucial component of bortezomib sensitization of cancer cells to TRAIL apoptosis involves increased generation of caspase-8 at the DISC. The molecular basis for this increased activity of caspase-8 in RCC previously exposed to bortezomib is not yet known. Recent findings suggest that the glycosylation state of the TRAIL receptors (31) and ubiquitination of caspase-8 (32) can influence the strength of the apoptotic signal. Therefore, the effects of bortezomib on these processes are worthy of further investigation.

Although numerous studies have shown that bortezomib can sensitize tumor cells to TRAIL apoptosis in vitro, few studies have attempted to extend this to the in vivo setting. We have shown improved therapeutic efficacy in the absence of toxicity in a mouse model of renal cancer when bortezomib is combined with an agonist antibody to mouse DR5 (33). Furthermore, combinations of bortezomib together with natural killer cells have also shown some therapeutic benefits in vivo in the same mouse renal cancer model (34). More recently, therapeutic benefit of the bortezomib and TRAIL combination was observed in a xenograft model of human prostate cancer (35). These studies need to be extended. Because bortezomib has many side effects, and may be somewhat immunosuppressive in vivo (36), identification of the molecular mechanism(s) whereby it sensitizes tumor cells to TRAIL could be very beneficial. This may allow for the development of tumor biomarkers for prior identification of specific tumors that are most likely to respond to TRAIL-based therapies. Furthermore, this could also allow for the identification of components of the apoptotic pathway that could be targeted using agents more specific, and maybe more potent and less toxic than proteasome inhibitors.

Disclosure of Potential Conflicts of Interest

None of the authors have any competing financial interests in relation to the work described.

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