Nonproteasomal Targets of the Proteasome Inhibitors Bortezomib and Carfilzomib: a Link to Clinical Adverse Events

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Abstract

**Purpose:** Bortezomib (Velcade), a dipeptide boronate 20S proteasome inhibitor and an approved treatment option for multiple myeloma, is associated with a treatment-emergent, painful peripheral neuropathy (PN) in more than 30% of patients. Carfilzomib, a tetrapeptide epoxyketone proteasome inhibitor, currently in clinical investigation in myeloma, is associated with low rates of PN. We sought to determine whether PN represents a target-mediated adverse drug reaction (ADR).

**Experimental Design:** Neurodegenerative effects of proteasome inhibitors were assessed in an in vitro model utilizing a differentiated neuronal cell line. Secondary targets of both inhibitors were identified by a multifaceted approach involving candidate screening, profiling with an activity-based probe, and database mining. Secondary target activity was measured in rats and patients receiving both inhibitors.

**Results:** Despite equivalent levels of proteasome inhibition, only bortezomib reduced neurite length, suggesting a nonproteasomal mechanism. In cell lysates, bortezomib, but not carfilzomib, significantly inhibited the serine proteases cathepsin G (CatG), cathepsin A, chymase, dipeptidyl peptidase II, and HtrA2/Omi at potencies near or equivalent to that for the proteasome. Inhibition of CatG was detected in splenocytes of rats receiving bortezomib and in peripheral blood mononuclear cells derived from bortezomib-treated patients. Levels of HtrA2/Omi, which is known to be involved in neuronal survival, were upregulated in neuronal cells exposed to both proteasome inhibitors but was inhibited only by bortezomib exposure.

**Conclusion:** These data show that bortezomib-induced neurodegeneration in vitro occurs via a proteasome-independent mechanism and that bortezomib inhibits several nonproteasomal targets in vitro and in vivo, which may play a role in its clinical ADR profile.

Introduction

Polypharmacology is the study of identifying secondary targets for an active pharmaceutical agent (drug promiscuity). In some disease etiologies, targeting multiple proteins may improve the therapeutic efficacy, such as for antipsychotic drugs or kinase inhibitors with anticancer activity (1). However, off-target activity can lead to adverse drug reactions (ADR) such as the case with fenfluramine and torcetrapib, 2 agents halted in development because of fatal hypertension that was the consequence of secondary target inhibition (1–3). Highlighting the promiscuity of small molecule inhibitors, a recent in silico screening of 213 Roche compounds revealed an average of 6.3 target proteins per compound (1). The standard approach for detecting off-target activity consists of screening compounds against panels of targets, often involving recombinant proteins in cell-free systems, and in silico prediction tools (1, 4). More recently, whole genome- and proteome-based analyses have been utilized to infer potential off-target activity, but these methods identify affected pathways rather than direct protein targets for a drug (3).

Bortezomib (Velcade) is a dipeptide boronate proteasome inhibitor that has been approved for the treatment of multiple myeloma (MM) and mantle cell lymphoma (5). The proteasome is a multicatalytic protease whose active sites are unique N-terminal threonine proteases (6, 7). Proteasome inhibition results in the modulation of many cellular activities, including alterations in signal transduction pathways, modifications in antigen presentation, and induction of an apoptotic response (5). Despite robust
Bortezomib has validated the proteasome as a target for the treatment of myeloma but is hindered by a painful peripheral neuropathy (PN) that may occur because of proteasome inhibition. Carfilzomib, a second in class proteasome inhibitor, is in late-stage clinical trials and displays minimal signs of PN. These observations suggest a proteasome-independent mechanism (off-target activity) underlying PN.

In this study, we utilized an in vitro model of neurodegeneration in which, despite equivalent proteasome inhibition, only bortezomib induced neurite degeneration. Utilizing a multifaceted approach to identify off-targets that included activity-based probe (ABP) profiling in cells and in silico database mining, we found that bortezomib targets several serine hydrolases. We report the first demonstration of inhibition of a nonproteasomal target (cathepsin G) in bortezomib-treated patients using ABPs. Also, bortezomib induces a prosurvival protease, HtrA2/Omi, in neuronal cells, supporting the hypothesis that bortezomib-induced PN results from off-target activity.

Methods

Compounds

The biotinylated general serine hydrolase fluorophosphonate-biotin (FP-biotin) probe was synthesized as previously described (20). Bortezomib (Fig. 1A) was purchased from a pharmacy. Cbz-Leu-Leu-Leu-boronic acid was purchased from AG Scientific, Inc. MG-132 (Cbz-Leu-Leu-Leu-aldehyde) was purchased from Boston Biochem.

Carfilzomib (Fig. 1A) was synthesized as previously described (US 07232818). Cbz-Leu-Phe-ketoepoxide and Cbz-Leu-Leu-Leu-ketoepoxide were synthesized as previously described (WO 2005111008) with more than 95% purity by high-performance liquid chromatography (HPLC) and nuclear magnetic resonance (NMR) methods. Cbz-Leu-Phe-aldehyde was synthesized as previously described (21), and the aldehyde version of carfilzomib was synthesized by the reduction of the Weinreb amide of carfilzomib to an aldehyde with lithium aluminum hydride in tetrahydrofuran with more than 75% purity.
Cells

PBMCS from normal healthy volunteers were obtained from AllCells. HepG2 (hepatocellular carcinoma), THP-1 (acute monocytic leukemia), 786-O (renal cell adenocarcinoma), and SH-SY5Y (neuroblastoma) cells were obtained from the American Type Culture Collection and were cultured in media recommended by the supplier at 37°C in 5% CO2.

Animals

Male Sprague-Dawley rats (275–300 g) were purchased from Charles River Laboratories and housed for 1 week before experimentation. For all experiments, animals had access to food and water ad libitum. All experiments were done under protocols approved by an institutional animal care and use committee.

SH-SY5Y neurodegeneration assay

This assay was conducted as previously described (22, 23). Briefly, SH-SY5Y cells were differentiated in multi-well plates by culturing in the presence of retinoic acid for 5 days, followed by exposure to brain-derived neurotrophic factor for 3 days. The cells were then treated with compound for 24, 48, and/or 72 hours, fixed, and stained with Hoechst (blue) for identification of cell nuclei staining to identify live cells and neuronal-specific βIII-tubulin (green) for the cytoskeleton to visualize cell bodies and neurites. Endpoint analysis was done by high-content imaging and quantitative image analysis using GraphPad Prism determined on the basis of the reaction velocity measured between 60 and 75 minutes by using GraphPad Prism 4.01. Antibodies used for Western blotting were mouse monoclonal anti-human CatA and anti-human chymase obtained from R&D Systems, rabbit polyclonal anti-human CatG and anti-human DPPII purchased from Abcam, and horseradish peroxidase–conjugated (HRP) secondary antibodies obtained from Jackson ImmunoResearch Laboratories, Inc. ImmunoPure Streptavidin resin and HRP–streptavidin overlay were purchased from Thermo Scientific.

Purified enzyme assays

Substrate-based assays were initiated by enzyme addition and monitored for aminocetyl 7-amino-4-methylcoumarin amide (MCA) and 7-amino-4-methylcoumarin (AMC) product formation with a plate-based spectrofluorometer (Tecan). Chymase (EMD Chemicals) activity was measured using Suc-Ala-Ala-Pro-Phe-AMC (MP Biomedicals) in assay buffer consisting of 200 mmol/L HEPES, pH 8.0, 2 mol/L NaCl, and 0.01% Triton X-100 at 27°C. CatG (EMD) activity was measured using Suc-Ala-Ala-Pro-Phe-AMC (MP Biomedicals) in assay buffer consisting of 50 mmol/L NaOAc, pH 5.5, 2 mmol/L EDTA, and 1 mmol/L dithiothreitol (DTT) at 30°C. CatA (R&D Systems) activity was measured using MCA-Arg-Pro-Pro-Gly-Phe-Ser-Ala-Phe-Lys(Dnp) in assay buffer consisting of 50 mmol/L MES, pH 5.5, and 5 mmol/L DTT at 27°C. DPPII (R&D Systems, Inc.) activity was measured using Suc-Lys-Pro-AMC (MP Biomedicals) in assay buffer consisting of 25 mmol/L MES, pH 6.0, at 27°C. IC50 (50% inhibitory concentration) values were determined by GraphPad Prism 4.01. Antibodies used for Western blotting were mouse monoclonal anti-human CatA and anti-human chymase obtained from R&D Systems, rabbit polyclonal anti-human CatG and anti-human DPPII purchased from Abcam, and horseradish peroxidase–conjugated (HRP) secondary antibodies obtained from Jackson ImmunoResearch Laboratories, Inc. ImmunoPure Streptavidin resin and HRP–streptavidin overlay were purchased from Thermo Scientific.

Detection of protease activity in lysates

PMBC lysozyme IC50 determinations. A total of 2 × 10⁶ to 3 × 10⁶ cells were lysed per condition in 50 to 100 μL of lysis buffer [PBS with 1% Nonidet P40 (NP40)] for 1 hour on ice. Lysates were cleared by centrifugation and treated with inhibitor at various concentrations for 30 minutes, followed by 1-hour incubation with 10 μmol/L FP-biotin at room temperature (RT). SDS was added to a final concentration of 1% (5 μL in a 50 μL reaction volume) and the samples were boiled for 5 minutes. The SDS was diluted to 0.1% with PBS (Vf = 500 μL) and incubated with rotating overnight with 20 μL of prewashed Pierce Streptavidin UltraLink Resin (Thermo Scientific) at RT. FP-biotin–bound proteome resin was washed 3 times in lysis buffer, and after the final wash, any remaining liquid was extracted with a 30-gauge needle. The resin was resuspended in 10 μL PBS, 20 μL 4 × SDS-PAGE loading dye (Bio-Rad), and 3 μL of 10× reducing agent (Bio-Rad). The FP-biotin–bound proteome was eluted by 3 cycles of 5 minutes boiling and vortexing. The eluate (10 μL) was loaded onto each lane for Western blot analysis. The signal was quantitated by densitometry using ImageJ software (http://rsbweb.nih.gov/ij/) and IC50 values were determined by GraphPad Prism 4.01. Antibodies used for Western blotting were mouse monoclonal anti-human CatA and anti-human chymase obtained from R&D Systems, rabbit polyclonal anti-human CatG and anti-human DPPII purchased from Abcam, and horseradish peroxidase–conjugated (HRP) secondary antibodies obtained from Jackson ImmunoResearch Laboratories, Inc. ImmunoPure Streptavidin resin and HRP–streptavidin overlay were purchased from Thermo Scientific.

Profiling tumor cell lines

Tumor cells (1.5 × 10⁶–3 × 10⁶ cells per condition) were lysed in lysis buffer for 30 minutes on ice and cleared by centrifugation. The lysates were then incubated with 10 μmol/L bortezomib or carfilzomib for 30 minutes at RT and then probed with 5 μmol/L of FP-biotin at RT for 1 hour. Proteins were separated by SDS-PAGE chromatography and FP-bound targets detected by Western blotting with HRP-conjugated streptavidin.

Ex vivo treatment of whole blood

Whole blood (10 mL) was treated with 100 mmol/L of bortezomib or carfilzomib for 40 minutes and transferred to BD Vacutainer cell preparation tubes (CPT; BD Biosciences) to isolate PBMCs. After centrifugation, PBMCs were collected, erythrocytes were removed by hypotonic lysis (BD PharmLyse; BD Biosciences), and the cells were lysed and cleared as described above. Cleared lysate was probed with 5 μmol/L of FP-biotin at RT for 1 hour. CatG was detected as described above.
Pharmacodynamics *in vivo*

Rats received intravenous bolus administration of bortezomib at 0.3 mg/kg and carfilzomib at 7 mg/kg. Fifteen minutes after dosing, animals were sacrificed by CO₂ narcosis, and blood and spleens were collected and processed as described previously (24). Proteasome CT-L activity was measured as described previously (24), and CatG activity in lysates from erythrocyte-depleted splenocytes was measured by FP-biotin probe binding as described above by using an anti-rodent CatG antibody (Santa Cruz Biotechnologies).

Whole blood (10 mL) was collected from consenting patients receiving 1.0 (n = 1), 1.3 (n = 2), or 1.6 mg/m² (n = 1) bortezomib into CPT to isolate PBMCs as described above. Proteasome and CatG activity was measured as described above. The study was approved by the independent cantonal Research Ethics Review Board and carried out in accordance with good clinical practice guidelines, the Declaration of Helsinki, and the applicable national law.

Western blotting for HtrA2 substrates

SH-SY5Y cells were seeded at 3 × 10⁵ cells/mL and cultured for 2 days. The cells were then treated for 12 hours with carfilzomib or bortezomib (both at 100 nmol/L) and harvested for Western blotting analysis. Cell pellets were lysed in PBS + 1% NP40 + protease inhibitor cocktail (Roche Diagnostics) for 1 hour on ice and the lysates were cleared by centrifugation. Proteins were resolved by SDS-PAGE, transferred to nitrocellulose, and probed with antibodies for HtrA2, eIF4G1, and pyruvate dehydrogenase. All the antibodies were purchased from Abcam.

Statistical analysis

For comparisons of treatment groups, a 1-way ANOVA followed by Bonferroni post hoc analysis using GraphPad Prism (version 4.01) was done. Statistical significance was achieved when P < 0.05.

Results

**Bortezomib induces neurite degeneration *in vitro by a proteasome-independent mechanism***

To investigate the role of proteasome inhibition in neurite degeneration, an *in vitro* assay was employed. This assay involved the differentiation of the human neuroblastoma cell line SH-SY5Y in the presence of retinoic acid and brain-derived neurotrophic factor, followed by exposure to proteasome inhibitors (22). Following treatment, cells were stained with Hoechst dye (blue) to detect nuclei and fluorescein isothiocyanate (FITC) anti-βIII-tubulin (green) to detect neurite outgrowth and analyzed using a high-content image analysis.
Bortezomib inhibits multiple serine proteases in vitro

Bortezomib has been shown to inhibit the serine proteases chymotrypsin, CatG, and chymase but not proteases with a preference for basic residues at P1, such as thrombin (19). We used a candidate approach to compare the activity of bortezomib and carfilzomib against a panel of 21 serine, cysteine, aspartyl, and metalloproteases (Supplementary Fig. S1). At 10 μmol/L, bortezomib inhibited serine proteases with a P1 selectivity of large hydrophobic or aromatic amino acids (chymotrypsin, CatG, and chymase) by 95% and elastase, rennin, and angiotensin-converting enzyme by 30% to 40%. On the other hand, carfilzomib displayed modest inhibition of chymotrypsin (~40%) but had minimal to no activity against the other 20 proteases. These data show that bortezomib has a broader off-target activity profile against nonproteasomal proteases than was previously considered.

To broadly profile the activity of bortezomib as an inhibitor of serine hydrolases, we utilized an ABP approach using a general serine hydrolase probe FP-biotin (refs. 19, 20; Supplementary Fig. S2). Lysates from 3 PBMC samples, an easily accessible primary human tissue that is a rich source of proteases (Supplementary Table S1), were exposed to both proteasome inhibitors prior to the addition of the FP-biotin probe. Isolated streptavidin-bound proteins were identified by liquid chromatography and tandem mass spectrometry (LC/MS-MS). As a validation of the method, we found that the broad-based protease inhibitors difluorophosphonate and diisocoumarin resulted in the inhibition of several serine hydrolases (Supplementary Table S1). Further validating this approach, CatG inhibition by bortezomib was detected. By the ABP approach, we also detected inhibition of CatA and DPPII by bortezomib, 2 proteases that were not evaluated in the candidate screen (Supplementary Table S1). Carfilzomib did not result in significant inhibition of serine proteases in PBMC lysates as measured by ABP profiling.

We extended these findings by determining the relative potency of bortezomib by using fluorescent substrate assays on purified CatG, CatA, chymase, and DPPII. Bortezomib had varying potencies against these purified enzymes, with IC50 values ranging from 0.3 to 9 μmol/L (Table 1). To investigate whether the activity of the enzymes might differ in a complex cellular milieu, we determined the IC50 values by using the FP-biotin probe and a gel-based assay (Table 1 and Fig. 2A). In each case, bortezomib was more potent against serine proteases in cell lysates versus purified enzymes. For CatA and chymase, IC50 values were equivalent to those of the proteasome. Carfilzomib did not result in significant inhibition of these serine proteases in either assay system. These data show potent inhibition of several serine protease targets of bortezomib in physiologically relevant conditions.

Next, we utilized the FP-biotin probe to detect protease inhibition in 3 tumor cell lines. Lysates of HepG2

Table 1. IC50 values of serine proteases in purified enzymes and PBMC lysates

<table>
<thead>
<tr>
<th>Target</th>
<th>P1 selectivity</th>
<th>Purified enzyme</th>
<th>PBMC lysatea</th>
<th>Purified enzyme</th>
<th>PBMC lysatea</th>
</tr>
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<tbody>
<tr>
<td>CatA</td>
<td>F, Y, V, E</td>
<td>&gt;10</td>
<td>2.5 (1.6–4.1)</td>
<td>0.0047 (0.0022–0.0008)</td>
<td></td>
</tr>
<tr>
<td>CatG</td>
<td>L, F, H, Y, N, I, V</td>
<td>&gt;10</td>
<td>0.95 (0.77–1.2)</td>
<td>0.3 (0.12–0.76)</td>
<td></td>
</tr>
<tr>
<td>Chymase</td>
<td>F, Y</td>
<td>&gt;10</td>
<td>0.28 (0.26–0.31)</td>
<td>0.0016 (0.00071–0.0036)</td>
<td></td>
</tr>
<tr>
<td>DPPII</td>
<td>P, A, R</td>
<td>&gt;10</td>
<td>8.7 (5.6–14)</td>
<td>0.26 (0.17–0.4)</td>
<td></td>
</tr>
<tr>
<td>HtrA2/Omi</td>
<td>L, V</td>
<td>&gt;10</td>
<td>0.003 (0.001–0.009)</td>
<td>N/A</td>
<td></td>
</tr>
<tr>
<td>Proteasome (LMP7)</td>
<td>L, F</td>
<td>0.025 (0.015–0.041)</td>
<td>0.022 (0.016–0.030)</td>
<td>0.031 (0.021–0.047)</td>
<td>0.00099 (0.00063–0.0016)</td>
</tr>
</tbody>
</table>

NOTE: 95% CIs are shown within parenthesis.
Abbreviation: N/A, not available.

aLysate IC50 values were generated by using the ABP method except LMP7 activity which was measured by using a fluorogenic substrate assay.
(liver adenocarcinoma), THP-1 (monocytic leukemia), and 786-O (renal adenocarcinoma) were exposed to bortezomib or carfilzomib. In all samples, we detected one or more FP-biotin–reactive protein species whose binding to FP-biotin could be blocked by prior exposure to 10 mM bortezomib but was not affected by carfilzomib (Fig. 2B). We identified CatA as a bortezomib target in HepG2 lysates and CatG in THP-1 cells lysates (data not shown). Taken together, these data show a broad and variable profile of nonproteasomal targets for bortezomib.

Peptide boronic acids are potent serine protease inhibitors

Two possible explanations for the striking difference in serine protease inhibition between bortezomib and carfilzomib are the peptide length (dipeptide vs. tetrapeptide) and the reactive pharmacophore (boronate vs. epoxyketone). Indeed, peptide boronates have been described as potent serine protease inhibitors (25, 26). To address the roles of peptide length and pharmacophore on off-target activity, we generated a matrix of 9 compounds related to the peptide backbones of carfilzomib (hPhe-Leu-Phe-Leu), MG-132 (Leu-Leu-Leu), and bortezomib (Phe-Leu), in which each of the peptide backbones were synthesized with each of the 3 pharmacophores, epoxyketone, aldehyde, and boronate (Fig. 2C). PBMC lysates were exposed to each compound and probed with FP-biotin and Western blotting for 4 target proteases (CatG, CatA, DPPII, and chymase). Boronate-containing compounds, regardless of the peptide backbone, were potent inhibitors of serine proteases, whereas none of the epoxyketone containing inhibitors resulted in detectable inhibition (Fig. 2D). Aldehyde-based compounds were also unable to inhibit FP probe binding, further suggesting that serine protease inhibition is driven by the boronate pharmacophore. These data were further validated by fluorogenic substrate assays with CatG and the 3 warhead versions of Leu-Leu-Leu (compounds 4–6) in which only the boronic acid analogue (compound 6) was capable of inhibiting enzymatic activity (IC$_{50}$ = 78 nmol/L).

Bortezomib inhibits serine proteases in intact cells and \textit{in vitro}

Following exposure of intact PBMCs to bortezomib, we could detect CatG inhibition via ABP profiling, suggesting
that this approach could be applied to in vivo settings (data not shown). Subsequently, we incubated whole blood for 1 hour with 100 nmol/L of bortezomib, a concentration/exposure period maintained in patients receiving a dose of 1.3 mg/m² (19, 27, 28), and observed potent inhibition of CatG by ABP detection in isolated PBMCs. Carfilzomib, on the other hand, did not affect CatG activity despite equivalent levels of proteasome inhibition to bortezomib (Fig. 3A).

To determine whether bortezomib results in serine protease inhibition in vivo, bortezomib or carfilzomib were administered intravenously to rats and splenocytes were isolated 15 minutes after dosing and analyzed for inhibition of CatG and the proteasome. Bortezomib inhibited splenocyte CatG activity by 50% and proteasome activity by 70% (Fig. 3B). On the other hand, carfilzomib resulted in approximately 85% inhibition of the proteasome but no inhibition of CatG activity. Interestingly, when splenocytes were isolated 1 hour after administration of bortezomib, CatG activity had recovered to basal levels whereas proteasome inhibition remained at approximately 70% (data not shown).

Next, we investigated the inhibition of nonproteasomal proteases in PBMCs derived from myeloma patients treated with either agent. We found a time- and dose-dependent inhibition of CatG in patients receiving 1.0, 1.3, and 1.6 mg/m² bortezomib (Fig. 3C). One hour after administration of 1.6 mg/m², both CatG and proteasome CT-L and LMP2 activities were inhibited by 60% to 70% in PBMC samples (Fig. 3C and Supplementary Fig. S4). In the PBMC samples from patients treated with 20 mg/m² carfilzomib, no inhibition of CatG was detected in PBMCs of 2 solid tumor patients treated with 20 mg/m² bolus administration of carfilzomib despite proteasome inhibition of more than 75% (Supplementary Fig. S3). These findings show that nonproteasomal proteases are inhibited in vivo following administration of bortezomib.

By Western blot analysis, we could not detect expression of the bortezomib-targeted proteases identified in PBMCs (CatG, CatA, chymase, and DPPII) in SH-SY5Y cells.
(data not shown). Furthermore, probing lysates from these cells with FP-biotin did not yield discernible proteins on streptavidin blots, indicating that target identification by LC/MS-MS would not be fruitful (data not shown). Therefore, we utilized a database mining approach to identify potential bortezomib targets that could be involved in the neurite degeneration in differentiated SH-SY5Y cells described previously (Fig. 1). MEROPS (http://merops.sanger.ac.uk/) is a comprehensive database of all known proteases and gene sequences predicted to be proteases. Of the 200 serine proteases in the database, 100 are annotated for a P1 preference. Of these, 14 proteases met our search criteria of having a P1 preference of Leu, Phe, or Tyr (Supplementary Table S2), with 5 of them already described above (CatG, CatA, chymase, and chymotrypsin B and C). We conducted literature searches on the remaining 9 candidates to determine whether a role in neurodegeneration or neuronal cell survival had been reported. One of these proteases, HtrA2/Omi, a stress-inducible mitochondrial protease involved in neuronal cell survival (29), was identified as a potential target. We utilized a gel-based assay to measure processing of the substrate β-casein by purified enzyme and found that bortezomib was an extremely potent inhibitor of HtrA2/Omi (IC50 by purified enzyme and found that bortezomib was an extremely potent inhibitor of HtrA2/Omi (IC50 by purified enzyme and found that bortezomib was an extremely potent inhibitor of HtrA2/Omi (IC50 by purified enzyme). We also utilized a candidate screen approach to determine specific protease families and target profiles for bortezomib. From this, we utilized ABP profiling with a general serine hydrolase probe to survey primary cells for other nonproteasomal targets of bortezomib. When converted to a quantitative assay, we were able to show that bortezomib results in inhibition of multiple serine proteases with relative levels similar to the CT-L and LMP2 subunits of the proteasome. Importantly, we found that bortezomib administration to myeloma patients resulted in a potent and dose-dependent inhibition of one of these proteases (CatG). Future studies will explore the onset and recovery of CatG and other serine proteases in patient tissue samples following bortezomib exposure.

The use of the ABP approach to target profiling reveals several new and clinically relevant findings. By measuring off-target activity in physiologically relevant conditions (i.e., in cells and cell lysates), we found that bortezomib potently inhibited several serine proteases at clinically relevant concentrations. This suggests that the utilization of purified proteins may underestimate the potency of a drug against secondary targets. Furthermore, by using the ABP approach, we obtained an unbiased and broad survey for target profiling. The identification of DPPII, which has a strong preference for proline-containing dipeptides, as a target shows that bortezomib is a potent and promiscuous inhibitor of multiple classes of proteases. Finally, we show that secondary target inhibition can be monitored in vitro by the ABP approach. Inhibition of CatG in the spleen of bortezomib-treated rats shows tissue inhibition of nonproteasomal targets. Potent inhibition of CatG in PBMCs of patients receiving bortezomib is the first demonstration of the use of ABP profiling to monitor off-target activity in patients with an approved drug.

We also applied the ABP approach to show that serine protease inhibition is a generalized property of peptide boronates. When boronates are added to the tri- and tetrapeptide backbone of MG-132 and carfilzomib, respectively, the resulting compounds are potent serine protease inhibitors. Peptide epoxyketones, even the epoxyketone version of bortezomib, had no activity against serine proteases. Inhibition of DPPII by the tri- and tetrapeptide boronates suggests that the promiscuity of boronic acid–containing compounds may be broader than originally anticipated. However, we cannot rule out a role for the peptide backbone, as MG-132 could mediate potent inhibition of chymase as determined by the use of a fluorogenic substrate (data not shown). Although the ABP approach enables the detection of multiple targets in a physiologically relevant condition, such as primary cells, it is limited by the relative abundance of the probe to individual targets. This may explain...
why chymase was not found in PBMCs via LC/MS-MS target identification but was detectable in Western blot analysis of PBMC lysates. Another limitation to the assay is the cellular source of targets. Indeed, we found distinct patterns of nonproteasomal target inhibition in individual tumor cell lines exposed to bortezomib. Further exploration with the ABP approach will involve different cell types and new active site probes. However, by utilizing the P1 preference for bortezomib against serine proteases we determined by using purified enzymes and ABP profiling, we were able to carry out database mining as a screen for other proteases that may be potential targets of bortezomib in cells but either lack affinity for the probe or are underexpressed in PBMCs. From this approach, we found that HtrA2/Omi, a serine protease with weak affinity for the FP-biotin probe (data not shown) and whose expression is induced following cellular stress, is potently inhibited by bortezomib.

HtrA2/Omi is expressed as mitochondrial and endoplasmic reticulum forms, each with distinct sets of substrates (35–37). Mitochondrial HtrA2/Omi has been shown to play a protective role in both in vitro and in vivo models of neurodegeneration and lymphocyte survival (29, 38–40). Following exposure to either bortezomib or carfilzomib, HtrA2/Omi expression increased in SH-SY5Y cells to levels similar to that reported for MG-132 (35), which may be the result of an unfolded protein response leading to oxidative stress. In these cells, processing of 2 substrates of HtrA2/Omi, eIF4G1, a translation factor, and pyruvate dehydrogenase, a Kreb cycle regulatory enzyme involved in generating oxidative species, are inhibited by bortezomib but not carfilzomib. Because both of these substrates have been implicated in neurodegenerative diseases (35, 41), inhibition of their processing may represent downstream mechanisms by which bortezomib alters neuronal cell function. Our data are consistent with a model in which bortezomib reduces neurite length by dual inhibition of the proteasome (resulting in oxidative and proteotoxic stress) and the neuronal prosurvival protease HtrA2/Omi.

In conclusion, we have utilized a multifaceted approach, including the first use of an ABP technique in clinical samples, to show that 2 structurally and mechanistically distinct proteasome inhibitors, bortezomib and carfilzomib, have dramatically different profiles for nonproteasomal targets in vitro and in vivo. Together with data from our in vitro model of neurite length inhibition, we propose that a specific ADR of bortezomib, PN, may be caused by distinct, off-target activities. Further exploration of the target profile for both agents by the approaches such as those described here is warranted.

Disclosure of Potential Conflicts of Interest

S. Arastu-Kapur, S.J. Lee, T. Muchamuel, and C.J. Kirk have ownership interest (including patents) in Onyx Pharmaceuticals, Inc.

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