Antitumor Effects of Proteasome Inhibition in Anaplastic Thyroid Carcinoma

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The ubiquitin-proteasome pathway has been identified as a potential molecular target for cancer therapy. In this study, we investigated the effect of the proteasome inhibitor bortezomib on anaplastic thyroid carcinoma (ATC) characterized by complete refractoriness to multimodal therapeutic approaches. Methods: The ATC cell lines C643 and SW1736 were treated with bortezomib (1 nM to 1 µM) for 12–72 h. Thereafter, growth inhibition was analyzed by thymidine uptake experiments and determination of the viable cell number. Apoptosis was measured and a cell cycle analysis was done. Using gene chip analysis and the real-time quantitative PCR system, we measured transcriptional changes. The activity of the nuclear factor (NF)-κB and p53 signal transduction pathways was monitored using the reporter constructs pNF-kB-TA-Luc and pp53-TA-Luc in the luciferase activity assay. Uptake measurements using ³H-FDG, ¹⁴C-aminoisobutyric acid, and Na¹²⁵iodide were performed to investigate metabolic changes and iodide symporter activity in vitro. Moreover, the ¹⁸F-FDG uptake was evaluated in ATC tumor-bearing nude mice 1 or 2 d after treatment with bortezomib. Results: Bortezomib induced growth inhibition, apoptosis, and G2-M cell cycle arrest associated with upregulation of p21^{CIP1/WAF1} expression in SW1736 and C643 cells. Moreover, the glucose metabolism and aminoisobutyric acid uptake significantly decreased in vitro in both of the ATC cell lines in vivo only in SW1736 tumors at 2 d after the bortezomib treatment. The transcriptional profile in bortezomib-treated SW1736 and C643 cells revealed increased expression of genes involved in stress response, apoptosis, regulation of the cell cycle, and differentiation. Using real-time quantitative PCR for the quantification of gene expression, we additionally noticed upregulation of the tumor necrosis factor-related apoptosis-inducing ligand and the thyroid-specific transcription factors Pax8 and TTF-1, leading to expression of the thyroid-specific target genes thyroglobulin, sodium iodide symporter, thyroperoxidase, and thyroid-stimulating hormone receptor and to a moderate accumulation of iodide in ATC cells. Conclusion: On the basis of our data, bortezomib represents a promising antineoplastic agent

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for the treatment of ATC. To improve the clinical outcome, further investigation into the potential of bortezomib therapy of thyroid cancer is clearly warranted.

Key Words: anaplastic thyroid carcinoma; proteasome inhibition; transcriptional profile; redifferentiation; iodide transport

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A hyroid carcinomas mostly appear as well-differentiated papillary and follicular tumors that are successfully treated by thyroidectomy, radioiodide therapy, and thyroid-stimulating hormone (TSH)–suppressive L-thyroxine medication. Occasionally, however, some patients develop more aggressive disease with local recurrence and distant metastasis due to dedifferentiation of the tumor, which is associated with a decrease or even loss of thyroid-specific gene expression and the inability to take up radioiodide (*I*). The prognosis is even worse for anaplastic thyroid carcinoma (ATC), which completely lacks thyroid-specific gene expression (*2*) and is not accessible to any therapeutic regimes (*3*). Because the overall median survival of patients with ATC is limited, there is a need for new therapeutic strategies for the treatment of these tumors.

As a major component for the degradation of proteins involved in cell survival, proliferation, and regulation of apoptosis and transcription, the ubiquitin–proteasome pathway has been identified as a potential molecular target for cancer therapy (4). Proteasome inhibitors have been shown to interfere with key steps of tumor cell regulation, leading to the inhibition of proliferation and death of neoplastic cells both in vitro and in vivo (5). Accordingly, the proteasome inhibitor bortezomib (PS-341, Velcade; Janssen-Cilag) provides strong antiproliferative effects in a variety of hematologic and solid cancer cells (6,7) and has marked clinical activity even in the setting of relapsed refractory multiple myeloma (8).

In the attempt to use proteasome inhibition for the treatment of undifferentiated thyroid tumors, we investi-

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gated the cellular response of the ATC cell lines SW1736 and C643 after exposure to bortezomib and applied combined gene expression profiling and real-time quantitative PCR (qPCR) to characterize the corresponding molecular pathways. Here, we show that in these cells bortezomib induced apoptosis, cell cycle arrest, changes in metabolism, and upregulation of genes involved in thyroid-specific functional activity.

MATERIALS AND METHODS

Cell Cultures and Treatment

The ATC cell lines SW1736 and C643 were cultured in RPMI 1640 medium (Gibco) supplemented with 10% fetal calf serum at 37°C in an atmosphere of 95% air and 5% CO₂. For the experiments, SW1736 and C643 cells were seeded in 6-well plates (cell cycle analysis, cell uptake measurements), in 12-well plates (apoptosis measurement), or in tissue culture flasks (RNA extraction) and cultured until 70% confluence was reached. Thereafter, the cells were treated with 10-fold dilutions (10 nM, 100 nM, and 1 μ M) of bortezomib for 12, 24, 48, and 72 h, respectively, before experimental procedures were performed. The viable cell number was determined in a Vi-CELL XR Cell Viability Analyzer (Beckmann Coulter).

Total RNA Extraction

The cells were lysed in Trizol (Invitrogen), and the RNA was purified by chloroform extraction and centrifugation, followed by isopropanol precipitation for 10 min at room temperature. The quality and the concentration of the RNA dissolved in diethylpyrocarbonate– H_2O was photometrically determined by the 280-to-260 ratio in the Nano Drop ND 100 (Peqlab).

Real-Time qPCR

After digestion of the RNA with RQ1 RNase-Free DNase I (Promega), reverse transcription of 2 µg of RNA was performed using the high-capacity RNA-to-complementary DNA kit (Life Technologies) as described by the manufacturer. The real-time qPCR of samples (n = 2) was performed on the Step One Plus qPCR system (Life Technologies) using the TaqMan Fast Univ PCR Master Mix and the TaqMan gene expression assays (Life Technologies) Hs01121172_m1 (p21^{CIP/WAF1}), Hs00153349_m1 (p53), Hs00247591_m1 (Pax8), Hs00201121_m1 (TTF-1), Hs00794359 m1 (thyroglobulin), Hs01053846 m1 (thyroidstimulating hormone receptor [TSHr]), Hs00892519_m1 (thyroperoxidase), Hs00166567_m1 (iodide symporter [NIS]), Hs00234355_m1 (tumor necrosis factor[TNF]-related apoptosis-inducing ligand [TRAIL]), and Hs01043166_m1 (TRAIL receptor DR5/TRAIL-R2) as specific probes and glyceraldehyde 3-phosphate dehydrogenase (GAPDH) (Hs99999905_m1) as an endogenous control. The relative quantification of transcription was determined using the formula $2^{-\Delta\Delta Ct}$ (9).

Measurement of Apoptosis

Apoptosis was measured by the Caspase-Glo 3/7 assay (Promega) as recommended by the manufacturer. For modulation, the cells were lysed with Luciferase Cell Culture Lysis Reagent (200 μ L/well; Promega). Thereafter, the lysate was centrifuged at 12,000g for 15 s, and 50 μ L of supernatant were incubated with 50 μ L of Caspase-Glo Reagent (Promega) for 1 h at room temperature before the caspase activity was quantified in the luminometer.

Cell Cycle Analysis

SW1736 and C643 cells were washed twice with phosphatebuffered saline (PBS), trypsinized, and centrifuged in culture medium. The cell pellet was resuspended in PBS and diluted 10-fold in 70% ethanol overnight. After being washed and centrifuged in PBS, the pellet was resuspended in 0.5 mL of 2.1% citric acid/0.5% polysorbate 20, and 3 mL of 7.1% Na₂HPO₄/0.5% 4',6-diamidino-2phenylindole solution (10 mg of 4',6-diamidino-2-phenylindole in 28.57 mL of H₂O) was added for fluorescence-activated cell sorting analysis.

Thymidine Incorporation into DNA

The cells were pulsed for 2 h with 74 kBq of methyl-³Hthymidine (American Radiolabeled Chemicals; specific activity, 370 GBq [10 Ci]/mmol) per milliliter of medium and 0.05 mM cold thymidine. After the cells had been washed 3 times with ice-cold PBS, lysis was initially performed with 0.5 mL of 0.5 M perchloric acid using a cell scraper. After 30 min on ice, the lysate was stirred in a vortex mixer and centrifuged at 1,500g for 5 min at 0°C. The supernatant (acid-soluble fraction) was removed, and the pellet (acid-insoluble fraction) was lysed in 0.5 μ L of 0.3N NaOH. The radioactivity in 300- μ L aliquots of the acid-insoluble fraction was quantified by scintillation counting.

³H-FDG Uptake

The uptake experiments were performed in glucose-free RPMI 1640 medium. After preincubation for 30 min, 37 kBq of 2-fluoro-2-deoxy-D-glucose, [5,6-³H] (ART 0105 [37 MBq (1 mCi)/mL]; specific radioactivity, 740 GBq [20 Ci/mmol]; radioactive concentration, 36,963 kBq [999 μ Ci]/vial), per milliliter of medium and cold FDG (final concentration, 0.1 mM) were added. After incubation for 10 min, the cells were washed twice with ice-cold PBS. The lysis was done in 0.5 μ L 0.3N NaOH. The radioactivity in 300- μ L aliquots was quantified by scintillation counting.

Iodide Uptake and Efflux

The iodide uptake was determined as described previously (10). Briefly, ATC cells were incubated with 74 kBq of Na¹²⁵I (Amersham Buchler; specific activity, 625.3 MBq/µg; radioactive concentration, 3.7 GBq/mL; radiochemical purity, 99.3%) per well for 1 h. The cells were washed twice with ice-cold PBS and lysed with 0.3N NaOH on ice. Thereafter, the radioactivity in the lysates was measured using a Cobra II γ -Counter (Packard). For modulation, the cells were incubated with or without TSH (50 µU/mL) for 4 h before Na¹²⁵I was added. To determine the iodide efflux, the Na¹²⁵I-containing medium was exchanged after 1 h by nonradioactive medium, and the cells were incubated again for 20 min and prepared for the measurement of radioactivity as described.

Animal Studies

Female BALB/c (*nu/nu*) mice (age, 6 wk) were supplied by Charles River Laboratories. All animal studies were performed in compliance with the national laws relating to the conduct of animal experimentation. After inoculation of cell suspensions (4×10^6 cells per animal) into the thigh of the mice, tumor growth was measured using calipers. ¹⁸F-FDG PET was performed when the tumors reached diameters of 7–13 mm (8–10 wk after transplantation). The animals (weight, 18–23 g) were studied before and at 2 d after treatment with bortezomib (2.25 mg/m²), with no statistically significant difference in size of tumors before and after treatment. Before PET, the animals were kept fasting for 4 h. The animals were kept in an inhalation narcosis with sevorane



FIGURE 1. Proliferation of SW1736 (A) and C643 (B) cells was evaluated by ³H-thymidine incorporation in presence of 0.05 mM cold thymidine into acid-insoluble cell fraction after exposure to bortezomib (10 nM, 100 nM, and 1 μ M) graded for 12, 24, and 48 h. Specific activity (pmol) of each sample was calculated and normalized to 10⁵ cells. Data represent mean values and SD of triplicate samples.

(0.5 volume-%) and oxygen (flow, 500 mL/min) during the PET examination. The plasma glucose level of the animals was determined using a blood glucose sensor electrode (MediSense), revealing values between 78 and 145 mg/dL. A transmission scan was obtained for 10 min before tracer administration with 2 rotating germanium pin sources to obtain cross sections for attenuation correction. After the acquisition of transmission data for attenuation correction, 6 MBq of ¹⁸F-FDG in 0.3 mL were administered into the tail vein, and PET data were acquired for 1 h in list mode on a Siemens Inveon scanner using a matrix of 256×256 (pixel size, $0.3882 \times 0.3882 \times 0.796$ mm). Thereafter, images were reconstructed at definite times after tracer administration $(2 \times 15, 8 \times 10^{-5})$ 30, 5×60 , 4×120 , 2×210 , and 7×300 s). The images were reconstructed iteratively using the space-alternating generalized expectation maximization method (16 subsets, 4 iterations), applying a median root before correction, and were converted to standardized uptake value (SUV) images on the basis of the formula SUV = tissue concentration (Bq/g)/(injected dose[Bq]/body weight [g]). The individual tumor SUV was calculated for each study as the mean of the last 2 time frames from 50 to 60 min after injection.

Statistical Analysis

Data were analyzed using the paired 2-tailed Student t test, and significance was assumed at a P value of less than 0.05.

RESULTS

Effect of Bortezomib on Proliferation and Cell Cycle

After incubation with increasing concentrations of bortezomib for 12, 24, and 48 h, a time- and dose-dependent growth inhibition of SW1736 and C643 cells was measured by [Fig. 1] ³H-thymidine uptake (Fig.1) and determination of the cell number (Supplemental Fig. 1; supplemental materials are available online only at http://jnm.snmjournals.org). In SW1736 cells (Fig. 1A) treated with 1 µM bortezomib, a decrease of the ³H-thymidine uptake between 81% and 96% was measured. The same concentration reduced the proliferation of C643 cells between 76% and 95% (Fig. 1B). In addition, the fraction of viable cells was reduced to 25% and 6.5% for SW1736 (Supplemental Fig. 1A) and C643 (Supplemental Fig. 1B), respectively. The growth inhibitory effect of bortezomib was associated with cell cycle [Table 1] arrest (Table 1) and upregulation of the cyclin-dependent [Fig. 2] kinase inhibitor p21^{CIP1/WAF1} (Fig. 2). In cells treated

(Fig. 2) kinase inhibitor p21cm (Fig. 2). In cells treated with 100 nM bortezomib, an up to 2-fold accumulation

in the G_2 -M phase and a decrease of both the G_1 phase cell fraction by 21% (SW1736) and 45% (C643) and the S phase cell fraction by 25% (SW1736) and 17% (C643) was observed (Table 1). Simultaneously, a more than 11fold and 5.5-fold increased p21^{CIP1/WAF1} transcription was measured in SW1736 and C643 cells (Fig. 2), respectively. Previous studies have shown that in thyroid carcinoma cells, bortezomib inhibits the activity of nuclear factor (NF)- κ B (11), which has been implicated in tumor development (12). Instead, we measured an up to 9.6-fold $(P \le 0.05)$ and up to 3.7-fold $(P \le 0.05)$ increase of NF-KB activity in SW1736 (Supplemental Fig. 2A) and C643 cells (Supplemental Fig. 2B), respectively, after the treatment with 100 nM bortezomib. In contrast, neither p53 transcription (Fig. 2) nor p53 activity (Supplemental Fig. 2) was measured using qPCR and the reporter gene assay.

Induction of Apoptosis

Although induction of p21CIP1/WAF1 expression and G2-M phase arrest are implicated in resistance to apoptosis (13), we detected a concentration-dependent increase of the apoptotic cell fraction in both cell lines using the annexin V/ propidium iodide assay (Supplemental Fig. 3) and the measurement of the caspase-3 and caspase-7 activity (Fig. 3). In [Fig. 3] SW1736 cells, an up to 27-fold increase of caspase-3 and caspase-7 activity was measured 24 h after the treatment with bortezomib (Fig. 3A) ($P \le 0.01$). In C643 cells, an up to 15-fold increase of the caspase-3 and caspase-7 activity was observed at 48 h after exposure to the proteasome inhibitor (Fig. 3B) ($P \le 0.01$). We also found a significant increase of annexin V-labeled SW1736 cells at 24 and 48 h, whereas annexin V-labeled C643 cells were detected only at 48 h after exposure to bortezomib (Supplemental Fig. 3). Simultaneously, a 19- and 10-fold increased expression of TRAIL and a 2.1- and 2.6-fold increased expression of the TRAIL receptor DR5/TRAIL R-2 were measured by qPCR in SW1736 and in C643 cells (Fig. 3C) exposed to 100 nM bortezomib for 24 h. The expression pattern obtained by the gene chip analysis, however, differed between the ATC cell lines. C643 cells provided increased expression of the growth-arrest and DNA damage-inducible α-transcripts and the growth-arrest and DNA damage-inducible (GADD)34 (Supplemental Table 1), which is induced in certain cell lines and correlates with p53-independent apoptosis after

 TABLE 1

 Bortezomib Induces G2-M Phase Arrest in ATC Cell Lines

		Phase		
Cell line	Concentration (nM)	G ₁	S	G ₂ –M
SW1736				
12 h	0	47.97 ± 0.78	36.77 ± 1.07	15.20 ± 0.29
	100	38.28 ± 1.43	32.90 ± 1.45	28.51 ± 0.42
24 h	0	42.83 ± 1.5	34.77 ± 0.35	22.31 ± 1.02
	100	34.94 ± 2.02	38.68 ± 3.35	26.38 ± 1.33
48 h	0	50.16 ± 1.42	31.44 ± 1.4	18.40 ± 0.02
	100	48.25 ± 0.78	23.42 ± 0.75	28.33 ± 0.03
C643				
12 h	0	49.57 ± 0.03	31.25 ± 0.21	19.17 ± 0.18
	100	27.36 ± 0.13	33.70 ± 1.29	38.94 ± 1.42
24 h	0	35.90 ± 1.06	40.71 ± 0.59	23.39 ± 1.66
	100	22.14 ± 0.03	33.97 ± 0.34	43.89 ± 0.37
48 h	0	58.27 ± 1.94	29.32 ± 0.39	12.41 ± 1.55
	100	46.09 ± 0.72	27.50 ± 1.33	26.41 ± 0.35

Cell cycle analysis of SW1736 and C643 cells incubated without (0 nM) or with bortezomib (100 nM) for indicated times was performed using fluorescence-activated cell sorting analysis. Data represent percentage of cells in G_1 , S, and G_2 –M phases, indicated as mean values of triplicate samples and SD.

ionizing radiation (14). In contrast, in SW1736 cells, the transcripts of the death-associated protein kinase3, the growth arrest and DNA damage–inducible β -protein, the TRAIL receptor DR4/TRAIL-R1 and the Fas-associated factor (FAF)1 were upregulated (Supplemental Table 2). When exposed to 10 nM bortezomib, SW1736 cells predominantly provided increased expression of members of the TNF superfamily and the TNF receptor member 6 (Supplemental Table 3).



FIGURE 2. Transcription of p21^{CIP/WAF1} and p53 in SW1736 and C643 cells exposed to graded bortezomib concentrations for 24 h was determined by qPCR using p21^{CIP/WAF1-} or p53-specific probe and GAPDH-specific probe as endogenous control. Values represent relative quantification of bortezomib-treated vs. untreated cells.

Determination of Metabolic Changes

Uptake studies performed to evaluate the effect of bortezomib on the metabolism of the ATC cells revealed a dose- and time-dependent decrease in tracer uptake (Fig. 4). [Fig. 4] The ³H-FDG accumulation declined to 1.5% in SW1736 (Fig. 4A) and to 3.4% in C643 (Fig. 4B) cells. Similarly, the ³H-aminoisobutyric acid uptake decreased to 1.7% in SW1736 (Supplemental Fig. 4A) and 2.2% in C463 (Supplemental Fig. 4B) cells.

After transplantation of SW1736 and C643 tumor cells in nude mice in vivo, studies of ¹⁸F-FDG uptake were done using PET before and at 1 and 2 d after bortezomib therapy (Fig. 5; Supplemental Fig. 5). No significant changes were [**Fig. 5**] seen on day 1 for either tumor. However, a significant decrease (45%) of tracer uptake (P < 0.05) was observed for SW1736 tumors on day 2 after therapy.

Transcriptional Profile of Bortezomib-Treated ATC Cells

A summary of bortezomib-induced transcriptional changes in SW1736 and C643 cells is presented in Supplemental Tables 1, 2, and 3. Besides the induction of both proapoptotic and antiapoptotic protein transcripts of components of the ubiquitin–proteasome pathway—most notably the ubiquitinconjugating enzyme E2H and ubiquitin-specific proteases transcripts implicated in growth signaling and survival cascades were upregulated. These include the insulin-like growth factor 1 and 2, serine–threonine protein kinase, serine–threonine kinase 17b, tyrosine kinase transcripts, protein kinase and proteinase inhibitors in C643 cells (Supplemental Table 1), and the insulin-like growth factor receptors 2 and 1 in SW1736 cells (Supplemental Table 2). In addition, in both cell lines bortezomib induced a significant increase of heat



FIGURE 3. Determination of apoptosis in SW1736 (A) and C643 (B) cells treated with increasing bortezomib concentrations for 12, 24, and 48 h. Caspase-3 and caspase-7 activity was measured by Caspase-Glo 3/7 assay (Promega). Data evaluated by luminescence (RLU) and normalization to 10⁵ cells represent mean values and SD of triplicate samples. Expression of TRAIL and DR5/TRAIL-R2 in SW1736 and C643 cells (C) exposed to bortezomib for 24 h was quantified by qPCR using TRAIL- and DR5/TRAIL-R2 probe and GAPDH probe as endogenous control. Values represent relative quantification of bortezomib-treated vs. untreated cells. ** $P \le 0.01$.

shock protein (hsp) transcripts, including the hsp-70, hsp-90, hsp-40, and hsp-27 families. Interestingly, the transcriptional profile of the ATC cells differed with respect to the expression of thyroid-specific transcription factors because in SW1736 cells treated with 10 nM bortezomib, but not in C643 cells, TTF-2 expression was initiated and transcription of Pax8 and Pax8 isoforms was increased (Supplemental Table 3).

Induction of Thyroid-Specific Gene Expression

In ATC expression of thyroid-specific genes and of the transcription factor TTF-1 is completely absent (15). The fact that increased Pax8 transcription was detected by the gene chip analysis, at least in bortezomib-treated SW1736 cells (Supplemental Table 3), prompted us to use qPCR for the investigation of thyroid-specific gene expression in the ATC cell lines. In fact, after incubation with 100 nM bortezomib for 24 h, a more than 4,500- and 711-fold increase of Pax8 transcription and a 5- and a 3.4-fold induction of TTF-1



thyroglobulin and thyroperoxidase (16, 17), these data provide evidence for their functional activity. The transcription of NIS was most affected by proteasome inhibition, ranging from 23- to 184-fold in SW1736 and 63- to 826-fold in C643 cells; the induction of NIS expression was achieved by the incubation with 100 nM and 1 µM bortezomib. Iodide uptake experiments performed to investigate functional activity of NIS revealed a 4.5-fold increase of iodide uptake ($P \leq$ 0.01) in C643 cells exposed to 100 nM bortezomib for 72 h; this increase was followed by an efflux of more than 90% radioactivity within 20 min (Fig. 7B, $P \le 0.05$). In SW1736 [Fig. 7] cells (Fig. 7A), a 1.6-fold increase of iodide accumulation was achieved ($P \le 0.01$), followed by an efflux of less than 43% of the intracellular radioactivity ($P \le 0.01$), pointing to functional activity of the thyroperoxidase and presence of thyroglobulin. Because NIS activity is affected by TSH (18), and TSHr expression was initiated on bortezomib treatment (Fig. 6), we performed iodide uptake experiments in the presence of TSH. However, in none of the ATC cell lines was an augmentation of iodide accumulation achieved by the addition of TSH (data not shown).

DISCUSSION

This study demonstrated an antiproliferative effect of bortezomib on the ATC cell lines SW1736 and C643 involving the induction of cell cycle arrest and apoptosis and changes in metabolism and gene expression. Bortezomib induced a dose- and time-dependent growth inhibition, which was



FIGURE 4. Uptake of ³H-FDG in SW1736 (A) and C643 (B) cells after exposure to different doses of bortezomib for 12-48 h. ATC cell lines were incubated with 37 kBq of ³H-FDG and 0.1 mM cold FDG for 10 min, and radioactivity in lysates was quantified by scintillation counting. Specific activity (pmol) of each sample was calculated and normalized to 10⁵ cells. Data represent mean values and SD of triplicate samples.

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FIGURE 5. (A) Changes (in percentage of pretreatment value in same animal) in ¹⁸F-FDG uptake (maximum SUV) before and 1 and 2 d after therapy with bortezomib in animals bearing SW1736 and C643 tumors in right thigh (mean and SEM, n = 4). (B) Coronal and transaxial slices of the SW1736 tumor-bearing animal before therapy (left) and 2 d after therapy (right). * $P \le 0.05$.

associated with an increased G₂–M phase cell fraction. Given the essential role of p21^{CIP1/WAF1} in the regulation and maintenance of G₂–M phase arrest in response to anticancer drugs (*19,20*), it is not surprising that we measured a significant upregulation of p21^{CIP1/WAF1} expression in both cell lines.

Although the induction of p21^{CIP1/WAF1} expression and G2-M phase arrest has been demonstrated to impede caspase-3-mediated apoptosis in MCF7 cells (13), we measured an increase of the caspase-3 and caspase-7 activity in the ATC cells. In addition, the transcriptional profile revealed an increased expression of the TNF superfamily members including TNF superfamily 10 (TRAIL) and of the TRAIL receptor DR4/TRAIL-R1 in SW1736 cells. Using qPCR for investigation of gene expression, we detected the upregulation of TRAIL expression and, although to a lesser extent, of DR5/TRAIL-R2 expression in both cell lines. Although a caspase-3-independent apoptotic pathway cannot definitively be excluded, we suggest that in ATC cells activation of the caspase-3 pathway is initiated through the interaction of TRAIL with the specific receptors DR5/TRAIL-R2 or DR4/TRAIL-R1 as already observed for follicular thyroid (21) and renal carcinoma (22).

The determination of metabolic changes after therapeutic intervention has been proven to be valuable for the assessment of early therapy effects (23). Using ³H-aminoisobutyric acid and ³H-FDG as tracer, we observed a significantly decreased glucose metabolism and amino acid uptake in both bortezomib-treated ATC cell lines. However, in vivo only SW1736 tumors responded with a decrease of the ¹⁸F-FDG uptake on day 2 after

treatment. Given that we did not observe any differences in growth and size of the ATC tumors or in the plasma glucose levels of the mice, we assume that the tumor perfusion and, therefore, the drug availability—differed in SW1736 and C643 tumors. Besides differences in perfusion, additional factors might have contributed to the different effect of bortezomib on the ¹⁸F-FDG uptake in SW1736 and C643 tumors. Generally, we observed a more pronounced response of SW1736 cells to proteasome inhibition in vitro with respect to gene expression (including that of p21^{CIP/WAF1}), differentiation, and induction of apoptosis. Moreover, differences in the tumor environment recruitment of benign cells such as fibroblasts, immune cells, and endothelial cells and the secretion of protecting factors may additionally affect tumor response to therapeutic agents.

NF-κB has been shown to play an important role in the anticancer activity of proteasome inhibitors. On the basis of the stabilization of IκB, the inhibitor of the NF-κB, proteasome inhibition prevents nuclear translocation and transcriptional activity of NF-κB in different tumor cells including medullary thyroid carcinoma cells (*5*,*11*). Our results instead demonstrated an increased NF-κB activity in SW1736 and C643 cells. This discrepancy might be explained by different experimental approaches used for the investigation of NF-κB activity. Furthermore, tumor cells often differ in sensitivity and response to therapeutic agents even when they originate from the same tumor entity (*24*,*25*). Thus, although bortezomib mediates similar effects on apoptosis and cell cycle in carcinoma cell lines, the NF-κB signaling pathway might be differently affected in ATC cells. More importantly, we

FIGURE 6. Quantification of thyroid-specific gene expression in SW1736 (A) and C643 (B) cells after bortezomib treatment for 24 h. Values represent relative quantification of bortezomib-treated vs. untreated cells evaluated by qPCR using Pax8-, TTF-1-, NIS-, thyroglobulin-, thyroperoxidase-, and TSHr-specific probes. GAPDH was used as endogenous control.





and others (8,26) suggest that NF- κ B inhibition may not be the sole mediator of the anticancer activity of bortezomib. Our gene chip analysis demonstrated a specific coordinated pattern of transcriptional changes in bortezomib-treated ATC cells similar to that already observed for multiple myeloma cells (8). Besides the upregulation of genes involved in apoptosis, cell cycle, and growth signaling, we noticed a significant increase of hsp transcripts in both ATC cell lines. Especially, inductions of the hsp-70, hsp-90, hsp-40, and hsp-27 families were the most pronounced responses to bortezomib, likely reflecting a stress response to the therapeutic agent, and are consistent with the welldocumented role of these molecular chaperones in conferring protection against therapeutic agents (27). Furthermore, bortezomib triggered transcription of components of the proteasome-ubiquitin pathway, including the proteasome 26s subunits in C643 cells and enzymes involved in ubiquitination in both ATC cells. Obviously, tumor cells attempt to compensate for the loss of proteasome activity by synthesizing new proteasomes. Although these cytoprotective responses are insufficient to rescue ATC cells from apoptosis, they indicate that proteasome activity is required for cell survival. In SW1736 cells, bortezomib also induced the upregulation of FAF1, which is involved in protein degradation (28), negative regulation of NF- κ B activity (29), chaperone activity of hsps (30), and induction of apoptosis (31). Therefore, FAF1 might be considered an important factor in therapeutic response to the treatment with bortezomib.

Finally, we demonstrated that bortezomib upregulates transcription of Pax8, TTF-1, and thyroid-specific target genes in the ATC cells. In addition, we measured an increased uptake of radioiodide in both cell lines, which, however, washed out of C643 cells within 20 min to more than 90% but less than 43% of SW1736 cells. These data implicate that bortezomib initiates a redifferentiation process in both cell lines, restoring to a low to moderate extent the functional activity of NIS. The intracellular accumulation of iodide additionally depends on the coupling to tyrosine residues of thyroglobulin by thyroperoxidase. The fact that SW1736 cells failed to do so implies that the ATC cells differed with respect to expression and functional activity of thyroperoxidase and thyroglobulin.

FIGURE 7. Uptake of ¹²⁵iodide within 1 h in SW1736 (A) and C643 (B) cells untreated or treated with indicated bortezomib concentrations for 72 h (uptake experiment). In efflux experiment, intracellular radioactivity of tumor cells was evaluated at 20 min after Na¹²⁵I-containing medium had been replaced by nonradioactive medium. Counts (cpm) for each sample were normalized to 10⁵ cells. Data represent mean values and SD of triplicate samples from representative experiment. * $P \leq 0.05$. ** $P \leq 0.01$

CONCLUSION

The present study showed that bortezomib provides strong antitumor effects, including cycle arrest, apoptosis, and changes in metabolism in ATC. In patients, the maximal concentration for bortezomib is around 112 ng/ mL, with a half-life of 9-15 h. In our cell experiments, we used concentrations between 4 and 400 ng/mL. Therefore, our experimental procedures underlying the in vitro and in vivo data correspond to the clinical situation. We also observed that the upregulation of thyroid-specific gene expression restores to some extent the capacity of iodide accumulation in ATC cells. However, the effectiveness of radioiodide therapy critically depends on the local concentration and the biologic half-life of the isotope in the tumor. To achieve therapeutically useful absorbed doses in vivo, a longer exposure with low doses of bortezomib may result in a more pronounced redifferentiation and higher iodide uptake as has been observed in patients with non-iodideconcentrating differentiated thyroid carcinoma after longterm treatment with retinoic acid (32). Thus, combining bortezomib with agents that additionally affect metabolism, sensitivity, and redifferentiation of ATC cells could improve the clinical outcome. For instance, in thyroid carcinoma cells, the histone deacetylase inhibitor MS-275 was shown to enhance the functional activity of NIS (25), and the combination of histone acetylase inhibitors and proteasome inhibitors synergistically enhanced TRAIL-mediated apoptosis (33). Thus, bortezomib may be considered a promising therapeutic agent for the treatment of anaplastic thyroid carcinoma in the setting of a combination therapy, but further investigation is clearly warranted to improve clinical outcome.

DISCLOSURE STATEMENT

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