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BMI1 silencing enhances docetaxel activity and impairs antioxidant response in prostate cancer

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The BMI1 oncogene promotes prostate cancer (PC) progression. High B-cell-specific Moloney murine leukemia virus integration site 1 (BMI1) expression predicts poor prognosis in PC patients. Recent evidence suggests that BMI1 may also play a role in docetaxel chemoresistance. However, mechanisms and clinical significance of BMI1-related chemoresistance have not been investigated. For this purpose, BMI1 was silenced in 2 PC cell lines (LNCaP and DU 145). Cell proliferation and apoptosis after docetaxel treatment were measured. Guanine oxidation was assessed by in-cell western. Global gene expression analysis was performed on BMI1 silenced cells. Oncomine database was used to compare in vitro data with gene expression in PC samples. BMI1 silencing had no effect on cell proliferation but significantly enhanced docetaxel-induced antitumor activity. Gene expression analysis demonstrated that BMI1 silencing downregulates a set of antioxidant genes. Docetaxel treatment increased guanine oxidation, whereas the antioxidant N-acetyl cysteine rescued docetaxel-induced cell death. Examination of clinical datasets revealed a positive correlation of BMI1 and antioxidant gene expression. BMI1-controlled antioxidant genes were predictive of poor prognosis in PC patients. In conclusion, BMI1 enhances antioxidant response, thereby allowing PC survival after docetaxel-based chemotherapy. BMI1-controlled antioxidant genes are overexpressed in aggressive PC and should be tested as predictors of chemotherapy failure.

Prostate cancer (PC) is not a single disease, but an umbrella under which a plethora of heterogeneous diseases is hidden. These range from indolent localized tumors, to aggressive metastatic diseases. Metastatic hormone-refractory prostate cancer (MHRPC) is the most aggressive form and is generally associated with very poor prognosis. Docetaxel is currently the only effective drug for MHRPC. Docetaxel is a microtubule-targeting drug that suppresses spindle microtubule dynamics and, thus, blocks mitosis and induces apoptosis. It has been shown that docetaxel disrupts centrosome organization in the late S, G2 and M phases of the cell cycle. This results in incomplete mitosis, accumulation of cells in G2/M phase and cell death. Two large Phase III trials showed that docetaxel treatment significantly improves survival and quality of life in MHRPC patients. Unfortunately, the survival advantage conferred by docetaxel is limited to 2–3 months, and life expectancy for MHRPC is only 18–19 months. Indeed, after an initial response to docetaxel, 80% of patients experience rapid disease progression. For this reason, insights into the mechanisms of docetaxel chemoresistance in PC cells are warranted.

Mechanisms of resistance to docetaxel are still not clear, although several hypotheses have been proposed. Docetaxel-induced microtubule damage inactivates the antiapoptotic protein BCL2, with subsequent triggering of the apoptotic machinery. For this reason, BCL2 overexpression in PC has been linked to chemoresistance. Keeping with this observation, activation of the antiapoptotic Akt pathway leads to docetaxel resistance. An additional mechanism of docetaxel resistance could be due to overexpression of efflux drug transporters. Some members of the ATP-binding cassette (ABC) transporter family are expressed by PC cells. ABC protein activity is associated with resistance to several
chemotherapeutic drugs, including docetaxel. Recent evidence shows that docetaxel antitumor activity is not limited to its antimitotic effects. Docetaxel is able to trigger reactive oxygen species (ROS) formation in cancer cells, thus inducing deoxyribonucleic acid (DNA), protein and cell membrane damage. ROS seem to play a major role in docetaxel antitumor activity. Indeed, head and neck carcinoma cells resistant to docetaxel displayed an abnormal mitochondrial DNA content and an enhanced antioxidant response. Interestingly, blocking the antioxidant response was able to restore docetaxel sensitivity.

A putative mediator of PC chemoresistance is the B-cell-specific Moloney murine leukemia virus integration site 1 (BMI1) oncogene. BMI1 belongs to the family of Polycomb group (PcG) genes. PcG proteins are organized in multimeric complexes, which mediate histone modifications of specific set of genes during cell development. BMI1 is a part of the Polycomb repressive complex 1, which triggers histone H2A ubiquitylation and gene silencing. BMI1 is responsible for locus silencing, thus contributing to prostate carcinogenesis. Although a mechanistic link has not been established, BMI1 is thought to silence many other oncosuppressors, particularly in PC cells. For example, BMI1 is essential for anchorage-independent growth and metastatic spreading of PC cells. This effect is likely mediated by silencing of several cell adhesion genes. In PC samples, BMI1 overexpression is associated with high Gleason score and increased risk of recurrence after prostatectomy. In addition, BMI1 is overexpressed in a subpopulation of PC cells with tumor-initiating capabilities. Microarray data analysis by Glinsky et al. identified a BMI-1-pathway signature with concordant profiles in normal stem cells and PC metastasis. In the same study, expression of the BMI1 signature was strongly associated with poor survival and therapy failure in five different types of epithelial neoplasms, including PC.

Recent studies showed that BMI1 silencing enhanced 5-fluorouracil antitumor activity in nasopharyngeal carcinoma. This effect seems to be dependent on the inactivation of antiapoptotic mechanisms, namely a reduced Akt phosphorylation. In addition, Hedgehog (HH) signaling activation enhanced ABC transporter expression and docetaxel resistance in PC cells. BMI1 is a well known downstream effector of HH signaling. Finally, BMI1 silencing strongly impairs antioxidant defense in different cell types. Given its prominent role in PC carcinogenesis, progression and prognosis, we sought to investigate the role of BMI1 in PC response to docetaxel.

Thus, we hypothesized that BMI1 silencing in PC cell could enhance docetaxel antitumor activity by at least one of three mechanisms: (i) inactivating antiapoptotic pathways (Akt phosphorylation); (ii) downregulating ABC transporter expression; (iii) impairing antioxidant defenses. For this purpose, we silenced BMI1 in 2 MHRPC cell lines: LNCaP (derived form androgen receptor-positive tumor) and DU 145 (derived from androgen receptor-negative tumor). We investigated putative mechanisms of BMI1-dependent chemoresistance, and we queried Oncomine database to test the clinical relevance our in vitro findings.

Our results show that BMI1 silencing impairs antioxidant defense and sensitizes PC cells to docetaxel. Examination of clinical datasets confirmed the relationship between BMI1 expression, antioxidant response and PC aggressiveness.

**Material and Methods**

**Cell culture**

The MHRPC cell lines LNCaP and DU 145 were obtained from American Type Culture Collection (Manassas, VA). According to ATCC, LNCaP cells are derived from a lymph node metastasis and DU 145 cells from a brain metastasis. Both cell lines are derived from androgen-independent PCs, although LNCaP still expresses the androgen receptor. Cells were maintained in RPMI-1640 medium with 10% fetal bovine serum, glutamine (1%), and penicillin-streptomycin (1%). Docetaxel (Sigma-Aldrich, St. Louis, MO) was dissolved in dimethyl sulfoxide (DMSO) and diluted in culture medium immediately before use. Final DMSO concentration never exceeded 0.1%. N-Acetyl cysteine (NAC; Sigma) was dissolved in sterile water and α-tocopherol (Sigma) was dissolved in ethanol and diluted in culture medium immediately before use. Final concentration for both NAC and α-tocopherol were 20 mM.

**Generation of ShBMI1 LNCaP and DU 145 cells**

BMI1-silenced cells were generated using the TRIPZ lentiviral doxycycline inducible Tet-On shRNA system (Open Biosystems, Huntsville, AL), after the protocols provided by the company. They are referred as DU145ShBMI1 and LNCaPShBMI1 from therein. Nonsilencing-TRIPZ lentiviral inducible ShRNAmir expressing cell lines (DU145NS and LNCaPNS) were generated and used as controls in all the experiments. Experiments were performed after at least 3 days of doxycycline (1 μg/ml) induction.

**Assay of cell viability and caspase activity**

Number of viable cells and caspase activity were measured though CellTiter-Glo- and CaspaseGlo luminescent assay (Promega, Madison, WL) and caspase. Both assays were previously described. For cell viability, three kinds of experiments were performed:

a. To assess cell proliferation after BMI1 silencing, LNCaP and DU 145 cells (NS and ShBMI1) were plated in triplicate in 96-well plates (1,000 cells/well). After 1, 3, 5 and 7 days, cell numbers were measured.

b. To assess cell viability after docetaxel treatment, LNCaP and DU 145 cells (NS and ShBMI1) were plated in triplicate in 96-well plates (5,000 cells/well). The following day, cells were exposed to different concentrations of docetaxel (1, 10, 100 and 1,000 nM) for 1 hr. After treatment, cells were allowed to grow in drug-free medium for 7 days. Cell viability was assessed using the CellTiter-Glo method.
medium for an additional 3 days, as previously described.\textsuperscript{25} Cell viability was measured at the end of the 3 days. Docetaxel concentrations used in our study are clinically achievable.\textsuperscript{26} The 1-hr treatment has been chosen in analogy with the 1-hr clinical infusion of docetaxel.\textsuperscript{26}

c. To assess the effect of ROS on docetaxel antitumor activity, cells were pretreated with NAC (20 mM, as described\textsuperscript{26}), then exposed to the same docetaxel concentration for 1 hr and grown in NAC-containing medium for 3 days.

For (b) and (c), the fraction of proliferating cells and IC\textsubscript{50} values relative to untreated cultures were calculated by non-linear leastsquares curve-fitting.

**Western blot**
Total protein was isolated from LNCaP and DU 145 (NS and ShBMI1) cells using RIPA lysis buffer (Thermo Scientific, Waltham, MA) and quantified using the BCA protein assay kit (Pierce) kit. A total of 30 µg of protein extract was loaded per lane into a 4 to 20% Tris-glycine gel (Invitrogen, Carlsbad, CA). For the experiment testing the effects of docetaxel and antioxidants on caspase and PARP cleavage, the experiment was set up identical to the experiment testing cell viability. Proteins were transferred to a polyvinylidene fluoride membrane, blocked in 10% nonfat dry milk, 0.1% Tween-20 PBS, incubated with primary (anti-BMI1; Millipore, 1/1,000 dilution; anti actin, Abcam 1/5,000 dilution; anticleaved Caspase 3 (1:1,000), anticleaved Caspase 8 (1:1,000), anticleaved Caspase 9 (1:1,000), anti-PARP (1:1,000), Akt (1:2,000), and phospho-Akt (1:1,000) Cell Signaling Technology) and secondary (LI-COR Biosciences, Lincoln, NE) antibodies and scanned by the LI-COR Odyssey IR Imaging System.

**Gene expression analysis**
Total RNA was extracted from NS and ShBMI1 (LNCaP and DU 145) cells using RNeasy kit. Total RNA from LNCaP and DU 145 (NS and ShBMI1) cells was labeled by reverse transcription with Superscript II (Invitrogen) and oligo-dT in the presence of Cy3-dUTP for Universal RNA reference control (Universal Human Reference RNA Stratagene p/n 740,000) or Cy5-dUTP for the samples. Whole Human Genome Oligo Microarray 4 × 44K format gene expression arrays (Agilent, p/n G4112F) were used for gene expression studies. The hybridization and washes were performed using Agilent reagents and following related protocols, and the slides were scanned on a GenePix 4000B scanner (Molecular Devices, Sunnyvale, CA). Differentially expressed genes were identified by ANOVA analysis of the replicates using Partek Genomic Suite Package (Partek). Data were further analyzed through the use of Ingenuity Pathways Analysis, with statistical methods and threshold described by the manufacturer (Ingenuity\textsuperscript{8} Systems, www.ingenuity.com). Unless otherwise indicated, genes that are up- or downregulated more than twofold and genes modulated with a \( p < 0.01 \) are referred as “significantly modulated genes.”

**Quantitative immunofluorescence**
To measure ROS effects after docetaxel treatment, LNCaP and DU 145 (NS and ShBMI1) cells were plated on 96-well plates. For each cell line, cells were treated with docetaxel at IC\textsubscript{50} concentration of NS cells, as described in “Assay of Cell Proliferation.” After treatment, cells were fixed in methanol for 15 min, washed in a solution of PBS and 1% fish oil gelatin (PBSG) and incubated in the same solution for 60 min to block. After blocking, cells were incubated overnight with primary antibodies (mouse antioxygenase, Millipore, 1/200 dilution; rabbit antiactin, Abcam, 1/500 dilution) in PBSG. After washing with PBSG + 0.1% Tween 20, cells were incubated with IRDye 800CW labeled donkey antimouse and IRDye 680CW labeled donkey antirabbit antibody (LI-COR Biotechnology, 1/200 dilution) for 2 h. After washing with PBSG + 0.1% Tween 20, cells were scanned with the LI-COR Odyssey IR Imaging System.

**Cell cycle analysis**
Cells were seeded at 50% confluence to ensure logarithmic growth and treated with docetaxel and antioxidants in the same manner as described for cell viability assays. Following treatment, one million cells were fixed in ice cold 70% ethanol overnight. Following fixation, cells were centrifuged and resuspended in PBS containing 40 µg/mL propidium iodide and 100 µg/ml RNAs A and incubated at 37°C for 1 hr.

**Meta-analysis for correlation of in vitro data with clinical data**
*In vitro* data on BMI1 silencing were compared to PC expression profiles derived from Oncomine 4.2 database analysis tool (http://www.oncomine.org). Oncomine database was also used to investigate the correlation between antioxidant genes and PC prognosis. For this purpose, we identified 22 PC studies on Oncomine database. We selected genes positively
correlated to BMI1 based on a correlation coefficient higher than 0.5. To avoid a duplicated analysis on same samples, we identified studies with the same first author and analyzed the one with the highest number of BMI1-correlated genes (based on correlation coefficient). All statistical values relative to this meta-analysis were calculated as described by Chinnaian and coworkers.27

Statistical analysis
Unless otherwise specified, all experiments were done in triplicate and were repeated at least twice. Data were expressed as mean values SE or SD and were analyzed by Student’s t-test, ANOVA followed by Bonferroni’s multiple comparison though GraphPad Prism software. The level of significance was set at $p < 0.05$.

Results
BMI1 silencing in PC cells
To test the effectiveness of our silencing system, we measured BMI1 mRNA and protein expression in NS and ShBMI1 cells. Both LNCaP and DU 145 ShBMI1 cells exhibited a more than threefold reduction in BMI1 mRNA levels (Fig. 1a) and a nearly complete BMI1 protein silencing (Fig. 1b).

Cell viability and caspase activity after BMI1 silencing
We first tested the hypothesis that BMI1 silencing affects PC cell proliferation. Cell growth was not significantly affected in BMI1-silenced cells (data not shown). Given these results, we investigated the effect of BMI1 silencing on docetaxel antitumor activity. As expected, treatment with docetaxel induced a dose-dependent growth inhibition, with IC$_{50}$ concentrations of 6.53 ± 1.59 and 22.60 ± 2.64 nM (LNCaP NS and DU 145 NS, respectively). BMI1 silencing caused a significant IC$_{50}$ reduction in both cell lines (2.01 ± 0.76 and 9.96 ± 1.02 nM, in LNCaP and DU 145, respectively; Fig. 2a). These results were consistent with cell cycle and apoptosis measurements. Docetaxel-treated NS and ShBMI1 LNCaP cells exhibited a similar cell cycle distribution (Fig. 2b), but BMI1 silencing increased both the number of apoptotic cells, as measured by flow cytometry (from 41 to 51%) and caspase 3/7 activity (Fig. 2d). On the contrary, BMI1 silencing in DU145 cells did not enhance docetaxel-induced apoptosis (data not shown) but increased the percentage of cells arrested in G2/M phase (from 12 to 23%; Fig. 2c).

Gene expression analysis
Because one possible explanation for BMI1-dependent chemoresistance may be drug transporter modulation, we investigated if BMI1 silencing downregulated three ABC transporters involved in docetaxel chemoresistance28 and expressed by PC cells.29 As shown in Supporting Information Figure 1 A, LNCaP cells expressed much lower ABC transporter mRNA, compared to DU 145 cells. Interestingly, LNCaP are more sensitive to docetaxel and express lower BMI1 than DU 145 cells (Figs. 1 and 2). However, BMI1 silencing did not significantly modulate ABC transporter expression. Because docetaxel resistance has also been linked to induction the Akt pathway, we also looked at Akt phosphorylation following BMI1 silencing. Similarly, we found that BMI1 silencing did not significantly reduce Akt phosphorylation in both cell lines (Supporting Information Fig. 1B).

To dissect the molecular pathways involved in BMI1-dependent chemosensitivity, we compared gene expression profiles of ShBMI1 and NS cells. BMI1 silencing produced significant modulation of 564 (LNCaP) and 880 (DU 145) genes (Supporting Information Tables 1 and 2). Our array data show that BMI1 silencing upregulated 280/564 (49.6%) and 492/879 (55.9%) genes in the two cell lines. To gain functional insights into the consequences of BMI1 silencing, we analyzed gene categories modulated by BMI1 (Supporting Information Table 3), using Ingenuity Pathway Analysis (IPA) software. Interestingly, we found that most of the genes modulated in both cell lines were significantly correlated to cancer ($9.3e^{-15} < p < 3.9e^{-2}$). In particular, 150 and 141 genes in LNCaP and DU 145 cells respectively were correlated to tumorigenesis. We also found that “cell death” and “cell growth and proliferation” were significantly affected (Supporting Information Table 3). As docetaxel sensitivity...
may be mediated by modulation of pro- and antiapoptotic pathways, we further assessed function and classification of these genes. In both LNCaP and DU 145, we were unable to identify a significant modulation of pro- or anti apoptotic pathways after BMI1 silencing. These data are consistent with the similar Akt phosphorylation level in NS and Sh BMI1 cells. Because analysis of classical death pathways was not able to explain our data on docetaxel activity, we investigated TOX pathways modulated after BMI1 silencing. This IPA analysis determines genes expression changes in response to several kinds of cell stress, including DNA damage, hypoxia and xenobiotics. Interestingly, BMI1 silencing was able to modulate ROS-related genes in both cell lines. In particular, “Oxidative Stress Response” genes ranked first among significantly affected TOX categories (Supporting Information Table 4). Keeping with our cell cycle analysis, G2/M transition genes were also significantly modulated. More interestingly, we found that BMI1 silencing was able to downregulate an antioxidant gene set in each of the cell lines (Fig. 3a). Among these, we found classical antioxidant genes (gamma glutamyl transferase), different heat shock protein (HSP) 40 isoforms and ferritin light chain (FLT). Notably, BMI1 silencing never led to an upregulation of antioxidant genes. In DU 145 cells, this effect was coupled to upregulation of genes involved in ROS production (MAOA)30 and ROS-dependent apoptosis (caspase 8)31. Interestingly, BMI1 silencing also upregulated KEAP1, which is known to inhibit antioxidant response. In addition, we confirmed a significant downregulation of FTL and HSP40 genes by qRT-PCR.

**Role of ROS in docetaxel activity**

Because our array data strongly suggested that BMI1 silencing disrupts antioxidant response in PC cells, we analyzed ROS production after docetaxel treatment. For this purpose, we measured oxoguanine levels in NS and ShBMI1 cells. Oxoguanine is the main product of DNA oxidative damage and is widely used to measure oxidative stress.32 As expected, docetaxel treatment induced a 1.5- to 2-fold increase in oxoguanine levels (Fig. 3b). More importantly, BMI1 silenced cells exhibited a significantly higher oxoguanine concentration in both cell lines (Fig. 3b). Finally, to show that ROS play a crucial role in docetaxel antitumor activity, we treated BMI1-silenced cells with the antioxidant NAC. As shown in Figures 3c and 3d, NAC treatment greatly reduced docetaxel-dependent cell death. These results show that BMI1 silencing impairs PC antioxidant response, thus enhancing docetaxel antitumor activity.

To further corroborate our data, we measured PARP and caspase cleavage in docetaxel treated and untreated cells. In addition, we evaluated the effect of two antioxidant molecules (NAC and α-tocopherol) on docetaxel treated cells. Consistent with our previous results, LNCaP cells exhibited greater PARP and caspases 9 and 7 cleavage after BMI1 silencing (Fig. 4). In DU 145 cells, we did not detect caspase 9 cleavage but instead detected caspase 8 cleavage. However, there was no change in either caspase or PARP cleavage in the BMI1-silenced cells as compared to the NS DU 145 cells. Importantly, in both LNCaP and DU 145 cells both antioxidants were able to rescue docetaxel-treated cells form apoptosis.
Gene expression analysis in clinical samples

Finally, we investigated the possible clinical significance of our in vitro findings. First, we tested the hypothesis that BMI1 expression is associated with ROS-related genes in PC samples. We used Oncomine database to investigate global gene expression data from 22 independent studies involving 20 to 596 PC samples of different stage and grade. We identified 1207 genes positively correlated to BMI1. We then analyzed, through IPA, the TOX pathways associated to BMI1 in clinical samples. As shown in Table 1, "Oxidative Stress Response" genes were significantly correlated with BMI1 expression. Among these genes, we found two isoforms of HSP40, EPHX1 and GST. When we compared oxidative stress related genes modulated by BMI1 in PC cell lines (Supporting Information Table 5) and in clinical samples (Supporting Information Table 6) we found a very significant overlap (odds ratio: 488.2, \( p < 0.05 \), t-test (NS vs. ShBMI1)). In addition, BMI1 expression was significantly correlated with "Hypoxia-Inducible Factor-Signalling," which is also involved in ROS metabolism.

Both in vitro and clinical data indicated that BMI1 controls a set of antioxidant genes in PC cells. Our IPA analysis generated a BMI1-dependent antioxidant signature (BAS) composed of 19 genes (Supporting Information Table 5). Given the prominent role of BMI1 in PC pathogenesis and progression, we investigated if this antioxidant signature had a prognostic significance. Using the Oncomine database, we found 3 studies in which the BAS was significantly correlated with PC clinical features (Table 2). In these studies, the signature was highly predictive of metastasis, shorter recurrence after prostatectomy and shorter survival. Interestingly, Oncomine data showed that high BMI1 expression is predictive of shorter relapse-free survival in PC (Supporting Information Fig. 2). These results show that BMI1 controls a set of antioxidant genes in PC and that these genes are activated in poor prognosis PC.

Discussion

In this article, we showed that BMI1 silencing enhances docetaxel antitumor activity in PC and that this effect is mediated by ROS production. As a part of Polycomb repressive complex 1, BMI1 can silence several genes in different cellular contexts. In PC, BMI1 is crucial for tumor progression and metastatic spreading.\(^{11,12}\) In addition, BMI1 expression is higher in a small fraction of PC cells with tumor initiating abilities, which may be the seeds of chemoresistance, metastasis and recurrence.\(^{15}\) Clinical data strongly suggest that BMI1 expression is a poor prognostic indicator in PC patients.\(^{14}\) Despite all these indications, very little is known about the pathways controlled by BMI1 in PC. In addition, it has been shown that BMI1 enhances chemo- and radio-resistance in nasopharyngeal carcinoma cells.\(^{17,33}\) For this reason, we
investigated the relationship between BMI1 expression and PC sensitivity to docetaxel. We also compared our in vitro findings to clinical data, in an attempt to evaluate its relevance for the development of new strategies to detect and overcome PC chemoresistance.

BMI1 silencing significantly increased docetaxel-dependent growth arrest (Fig. 2a). In the p53 positive LNCaP cells, this effect was due to an increased apoptotic activity. In the p53-mutated DU145 cells, BMI1 silencing enhanced G2/M phase arrest induced by docetaxel. Interestingly, in both cell lines, antioxidant drugs like NAC or tocopherol counteracted the effects of BMI1 silencing (Figs. 3c and 3d).

The chemosensitivity acquired by both cell lines after BMI1 silencing was not mediated by direct apoptotic pathway modulation, as shown for nasopharyngeal carcinoma cells, or by ABC transporter downregulation (Supporting Information Fig. 1). Our results suggest that BMI1 antioxidant role is crucial in PC cells. Previous studies have shown that BMI1 plays a key role in maintaining the oxidative stasis of the cell. In these models, BMI1 function has been linked to p16 and p53 suppression, with no association to antioxidant genes. In this article, we showed that BMI1 silencing significantly increased docetaxel-dependent oxidative stress (Fig. 3b) and that the antioxidants NAC and α-tocopherol can completely abrogate this effect (Figs. 3c and 3d). Gene expression analysis showed BMI1 silencing downregulated antioxidant genes and upregulated pro-oxidant genes (Fig. 3a). These results were supported by clinical samples data, showing a significant association between genes classified by IPA as playing a role in “Oxidative Stress Response” and BMI1. This set comprises genes that protect cells from oxidative stress, including epoxide hydrolase, superoxide dismutase and gamma glutamyl transferase. Interestingly, terminal caspase activity was increased after BMI1 silencing.

Although there is a diversity of oxidative stress-related genes regulated in these samples, HSP 40 genes are common to all datasets. HSP 40 is a family of co-chaperones comprising 41 members. Recent evidence suggests that some HSP 40 isoforms are involved in various aspects of cancer biology and that their expression is deregulated in many neoplasms. In addition, HSP 40 members contribute to inhibition of apoptosis, protection from oxidative stress, and maintenance of the mitochondrial membrane potential. In view of our results, it would be interesting to further explore the relationship between HSP 40, BMI1 and PC progression.

The putative relationship between BMI1 antioxidant genes and PC aggressiveness is further supported by clinical data, showing a correlation between BMI1-dependent antioxidant signature (BAS) and PC aggressiveness. In particular, three independent studies show that BAS is predictive of shorter disease-free survival after prostatectomy. This relationship is confirmed by Oncomine data (Supporting Information Fig. 2). A recent meta-analysis on PC patients found that a shorter time of PC progression is predictive of poor response to docetaxel. Our findings suggest that a subset of
PC is more resistant to docetaxel because of higher BMI1 expression and more effective antioxidant response. These tumors are also characterized by faster progression. Our in vitro results call for clinical studies on BMI1 expression in docetaxel-treated PC patients. According to our data, BMI1 should be overexpressed in chemoresistant tumors. Unfortunately, docetaxel is currently used to treat metastatic disease, which is always unresectable. Thus, clinical specimens from docetaxel-treated PC patients are hard to get.

These results could pave the way to new approaches to treat and detect chemoresistant PC. In particular, BMI1 expression, as well as the expression of BAS could prove to be valid predictive indicators of chemosensitivity. These genes could be used to personalize docetaxel regimens in MHRPC patients. Recently, some drug targeting histone modifications showed an interesting anticancer effect, and several new compounds are being tested. In the future, it is conceivable that specific inhibitors of Polycomb repressive complex 1 will be developed. Our results raise the possibility that docetaxel and Polycomb-targeting drugs could be synergistic in PC cells and that a combined regimen could improve docetaxel activity and prevent tumor recurrence.

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