Therapeutic Effects of XPO1 Inhibition in Thymic Epithelial Tumors

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Abstract

Exportin 1 (XPO1) mediates nuclear export of many cellular factors known to play critical roles in malignant processes, and selinexor (KPT-330) is the first XPO1-selective inhibitor of nuclear export compound in advanced clinical development phase for cancer treatment. We demonstrated here that inhibition of XPO1 drives nuclear accumulation of important cargo tumor suppressor proteins, including transcription factor FOXO3a and p53 in thymic epithelial tumor (TET) cells, and induces p53-dependent and -independent antitumor activity in vitro. Selinexor suppressed the growth of TET xenograft tumors in athymic nude mice via inhibition of cell proliferation and induction of apoptosis. Loss of p53 activity or amplification of XPO1 may contribute to resistance to XPO1 inhibitor in TET. Using mass spectrometry–based proteomics analysis, we identified a number of proteins whose abundances in the nucleus and cytoplasm shifted significantly following selinexor treatment in the TET cells. Furthermore, we found that XPO1 was highly expressed in aggressive histotypes and advanced stages of human TET, and high XPO1 expression was associated with poorer patient survival. These results underscore an important role of XPO1 in the pathogenesis of TET and support clinical development of the XPO1 inhibitor for the treatment of patients with this type of tumors. Cancer Res; 77(20); 5614–27.

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Introduction

Almost all targeted therapies developed in the modern era work through inhibition of particular oncogenic pathways that confer cancer cells the ability to proliferate and survive independent of exogenous growth signals (1). However, to undergo dysregulated proliferation and survival, cancer cells also need to circumvent internal negative regulatory programs often governed by tumor suppressor proteins (TSP; ref. 2). Although inactivation of TSPs is paramount in tumorigenesis, no cancer therapy has yet been developed aiming to restore their activity.

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TSPs can be inactivated by several mechanisms, including mutation, deletion, and epigenetic silencing (2). However, many TSPs such as p53 and p27 execute their tumor suppressor functions mainly in the nucleus and are in part regulated by nuclear–cytoplasmic shuttling for a rapid on/off switch. Dysregulation of nuclear–cytoplasmic shuttling affects nuclear activity of various TSPs and can contribute to abnormal cell survival, tumor progression, and drug resistance (3, 4). Although many molecules are involved in the shuttling process, alteration of exportin 1 (XPO1) plays a prominent role in tumor pathogenesis. XPO1, also known as chromosome region maintenance 1 (CRM1), mediates nuclear export of ~200 leucine-rich-nuclear export signal (LR-NES)-containing proteins (4). Importantly, XPO1 is the sole nuclear export receptor for a large number of TSPs involved in apoptotic signaling and cell-cycle regulation. Overexpression of XPO1 is reported in both solid tumors and leukemias and correlates with poor prognosis of several tumor types (4).

Consequently, XPO1 inhibition has emerged as a cancer therapeutic strategy (4). The first XPO1 inhibitor discovered was Leptomycin-B (LMB), a natural compound with antitumor activity (5), but severe toxicity profile prevented further clinical development (6). Latterly developed small-molecule selective inhibitors of nuclear export (SINE) represent a different class of XPO1 inhibitors with better specificity and efficacy. Selinexor (KPT-330) is one of such SINE compounds that is currently in clinical development (4, 7, 8). Unlike LMB that forms an irreversible covalent bond with cysteine-528 residue located in the vicinity of LR-NES-binding domain in XPO1, selinexor binds XPO1 through the same residue in a covalent but slowly reversible manner (4).

In this study, we investigated the role of XPO1 in thymic epithelial tumors (TET) and used selinexor to mechanistically

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explore the feasibility of XPO1-targeted therapy for TET treatment. We demonstrated that XPO1 is active in TET cells, and selinexor induced p53-dependent and independent antitumor activity and suppressed TET xenograft tumor growth. We also demonstrated that p53 loss and XPO1 amplification are potential mechanisms of resistance to selinexor. Furthermore, we globally profiled the factors and pathways/networks that are affected by selinexor using a proteomics approach and identified several important changes including those on the transcriptional programs. Finally, we demonstrated that targeting XPO1 in TETs is relevant by providing evidence of XPO1 overexpression in this tumor type and its correlation with the disease outcome.

**Materials and Methods**

**Cell lines, drug, and antibodies**

T1889 and T1682 cell lines were kindly provided by Marco Breining (Heidelberg University Hospital, Heidelberg, Germany; ref. 9). Ty82 was purchased from Japan Health Science Foundation (Tokyo, Japan). IU-TAB1 was kindly provided by George Sledge Jr (Indiana University School of Medicine, Indianapolis, IN; ref. 10). TEC84 was kindly provided by Phong Le (Loyola University, Chicago, IL; ref. 11), and TEC41.2 was kindly provided by Julian Sage and Brian Condie (University of Georgia, Athens, GA; ref. 12). MP57 was established in our lab as recently described (13). Except TEC84, which was cultured in TE media as previously described (11), all other cell lines were cultured in RPMI media containing 1% penicillin/streptomycin and 10% heat-inactivated bovine serum (Invitrogen). All cell lines were passaged less than 6 months between thawing and use in the described experiments.

**Generation of selinexor-resistant cell line**

IU-TAB1 cells (1 × 10^6) were seeded in T75 flasks in growth medium supplemented with selinexor (starting from 20 nmol/L). Medium was replaced every 2 to 3 days, and surviving cells were allowed to grow to 70% confluence before trypsination and reseded in medium containing (Pierce) according to the manufacturer's protocol (100 μL of CER1 and 5.5 μL of the CER2 reagents for the cytoplasmic extraction and 50 μL of the NER reagent for nuclear extractions).

**Immunofluorescence microscopy**

Cells were fixed with 4% paraformaldehyde and permeabilized with 0.1% Triton X-100 for 10 minutes. After blocking in 3% bovine serum albumin for 1 hour, cells were incubated with primary antibody for 1 hour, followed by 1 hour incubation with Alexa Fluor 488 dye (Thermo Fisher Scientific) and DAPI at room temperature. The staining was visualized under an immunofluorescence microscope.

**Cell viability assay**

Cells were seeded in 96-well plates (2.5 × 10^3 cells/well), allowed to adhere overnight, and exposed to DMSO or selinexor for 72 hours. Cell viability was evaluated using the CellTiter-Glo viability assay (Promega). Each assay was performed in triplicate. Data were plotted and analyzed using GraphPad Prism and represent mean ± SD. IC_{50} values were obtained by curve fitting to the Hill equation.

**Cell-cycle analysis**

Cells were exposed to DMSO or selinexor for 24 hours and harvested. After propidium iodide (PI) staining of the harvested cells, DNA content was measured by FACStar Plus Dual Laser System and FACSort System (Becton Dickinson). The percentage of cells in G1, S, and G2–M phases was analyzed by the ModFit LT program (Verify Software Home).

**Apoptosis assay**

Cells were exposed to DMSO or selinexor for 72 hours. Apoptotic cells were detected by Annexin V/PI staining assay (Life Technologies) and measured by flow cytometry. Results represent mean ± SD of triplicates.

**Caspase-3/7 assay**

Caspase activity was detected using the Caspase-Glo 3/7 assay kit (Promega). Cells (2.5 × 10^4 cells/well) were seeded in a luminometer white 96-well plate (Thermo Scientific) and treated with DMSO or selinexor for 48 hours. Caspase-3/7 reagents were then added to each well and incubated for 1 hour at room temperature. Luminescence was recorded using the GLOMAX+ instrument (Promega).

**Transfection of siRNAs and plasmids**

Cells were transfected with ON-TARGET plus siRNAs targeting XPO1, p53, and FOXO3a (Dharmacon) using Lipofectamine RNAiMAX transfection reagent (Invitrogen). AllStars Negative Control siRNA (Qiagen) was used as negative control. Transfection of H1682 cells with pIRE2-EGFP-p53WT or pIRE2-EGFP plasmids (Addgene) was carried out using Lipofectamine 2000 (Invitrogen).

**Generation of selinexor-resistant cell line**

IU-TAB1 cells (1 × 10^6 cells) were seeded in growth medium supplemented with selinexor (starting from 20 nmol/L). Medium was replaced every 2 to 3 days, and surviving cells were allowed to grow to 70% confluence before trypsination and reseded in medium containing...
gradually increased selinexor concentrations. This process was repeated for 2 months until selinexor reached 500 nmol/L. The resistant cells were maintained in growth medium containing 500 nmol/L of selinexor.

Gene copy-number variation analysis

Genomic DNA was extracted using the DNA/RNA mini kit (Qiagen) and subjected to a real-time PCR-based XPO1 gene copy-number analysis using TaqMan copy number assay on the ABI 7900HT system (Applied Biosystems). Primers for the XPO1 gene (XPO1: Hs03075013_c1) were purchased from ThermoFisher Scientific, and the RPPH1 gene was used as an (endogenous reference control) was purchased from ThermoFisher Scientific. Gene copy number was analyzed by CopyCaller software v1.0 (Applied Biosystems), and the copy number of the XPO1 gene was calibrated to that of the RPPH1 reference gene.

Xenograft study

Cells (1.5 × 10⁶/mouse in 0.1 mL of 50% Matrigel) were subcutaneously injected into the flanks of 6- to 8-week-old athymic nude mice (Charles River Laboratories). Tumor volume was measured once a week: Volume = 1/2 (Length × Width²). When average tumor volumes reached 100 mm³, mice were randomized into three groups (8 mice/group) and treated with DMSO or selinexor (0.5 or 1 mol/L) for 48 hours. Subcellular fractions were obtained using NE-PER Nuclear and Cytoplasmic Extraction Reagents (Thermo Scientific) and subjected to a real-time PCR-based Gene copy-number variation analysis (95% CI) for each cell line. NR

Statistical analysis

To determine the relevance of XPO1 inhibition in TETs, we first examined XPO1 expression in two thymoma cell lines (IU-TAB1 and T1682) and three thymic carcinoma cell lines (Ty82, MP57, and T1889). We observed moderate to high levels of XPO1 in TET cell lines, treated with 1 mol/L of selinexor resulted in a dose-dependent decrease or complete loss of XPO1 protein, localized mainly in the nucleus in these cells (Fig. 1A and B; Supplementary Fig. S1A). Treatment of TET cells with selinexor resulted in a significant shift in the balance of FOXO3a, p53, and p27 proteins in the nucleus and cytoplasm, leading to nuclear accumulation of these TSPs (Fig. 1E; Supplementary Fig. S1B). The concomitant robust increase of p27 cytoplasmic levels, observed after 24 hours of

Mass spectrometry–based proteomics

IU-TAB1 cells were cultured for at least five passages in SILAC media containing L-arginine and L-lysine (light), or L-¹³C₆-arginine and L-¹³C₆-lysine (heavy; Cambridge Isotope Laboratories). Cells were then treated with DMSO (“light” labeled) or selinexor (400 nmol/L; “heavy” labeled) for 6 hours. Subcellular fractions were obtained using NE-PER Nuclear and Cytoplasmic Extraction Reagents (Thermo Scientific) and validated by Western blot using anti-tubulin (cytoplasmic marker) and anti-histone H3 (nuclear marker). Equal amounts of protein lysates were mixed together for either nuclear or cytoplasmic fraction. The combined proteins were subjected to trypsin digestion, followed by purification of digested peptides, basic RPLC fractionation, LC-MS/MS, and data analyses as described elsewhere (15).

Results

XPO1 is expressed in TET cell lines and its activity can be inhibited by selinexor

To determine the relevance of XPO1 inhibition in TETs, we examined XPO1 expression in two thymoma cell lines (IU-TAB1 and T1682) and three thymic carcinoma cell lines (Ty82, MP57, and T1889). We observed moderate to high levels of XPO1 protein, localized mainly in the nucleus in these cells (Fig. 1A and B; Supplementary Fig. S1A). Treatment of TET cells with selinexor resulted in a significant shift in the balance of FOXO3a, p53, and p27 proteins in the nucleus and cytoplasm, leading to nuclear accumulation of these TSPs (Fig. 1E; Supplementary Fig. S1B). The concomitant robust increase of p27 cytoplasmic levels, observed after 24 hours of
Inhibition of XPO1 impairs proliferation and survival of TET cells

To determine the biological effect of XPO1 inhibition, we performed cell viability assay to test the sensitivity of TET cells to selinexor. Selinexor induced a dose-dependent inhibition of cell viability assay to test the sensitivity of TET cells. Inhibition of XPO1 impairs proliferation and survival (Fig. 1E; Supplementary Fig. S1C and S2A).

Selinexor induces p53-dependent and -independent cytotoxic effects in TET cells

Selinexor treatment resulted in nuclear accumulation of p53 and FOXO3a (Fig. 1), two important TSPs that control the expression of many crucial genes and regulate various key cellular processes (4, 7). To definitively demonstrate the contribution of these two TSPs to the antitumor activity of selinexor, we assessed the viability of TET cells treated with selinexor after siRNA knockdown of p53 and/or FOXO3a. Knockdown of p53 significantly reduced selinexor cytotoxicity in T1889 and IU-TAB1 cells (IC50 from 150 nmol/L to 600 nmol/L in T1889, P < 0.001 and from less than 150 nmol/L to more than 300 nmol/L in IU-TAB1, P < 0.05; Fig. 3A and B); the effect was also reflected by increase of the area under the dose–response curve (AUC; from 30 to 52 in T1889, P = 0.002, and from 29 to 47 in IU-TAB1, P = 0.006; Fig. 3C). However, knockdown of FOXO3a in these two cell lines had almost no effect on selinexor cytotoxicity (Fig. 3A–C). Interestingly, FOXO3a knockdown resulted in a significantly reduced selinexor cytotoxicity in MP57 cells (IC50 from <150 nmol/L to ~600 nmol/L, P < 0.001, and AUC from 24 to 42, P = 0.02; Fig. 3A–C), a p53-deficient thymic carcinoma cell line (13). Nonetheless, no effect was observed in the Ty82 cells with the knockdown of p53 and/or FOXO3a (Fig. 3A and B), indicating the involvement of other factors. Together, these data suggest that selinexor depends upon different XPO1 cargo proteins to exert its antitumor activities under different cellular contexts (i.e. p53-dependent and p53-independent fashions).

Loss of p53 activity in T1682 cells contributes to selinexor resistance

T1682 cells are primarily resistant to selinexor (Figs. 1 and 2), even though XPO1 is functional and its inhibition by selinexor led to a nuclear accumulation of FOXO3a in this TET line (Fig. 3D). Given that no p53 was detected in T1682 cells (Fig. 3D; refs. 21–23), we asked whether p53 loss contributes to the resistance. We ectopically expressed p53 in T1682 cells and reassessed their sensitivity to selinexor. Reinstallation of p53 significantly increased the sensitivity of T1682 cells to selinexor (at least 2.5-fold decrease of IC50, compared to that of Mock control; AUC: 73 versus 23 in T1682 cells treated with 0.5 μmol/L). These results clearly indicate a significant contribution of p53 loss to the survival of T1682 cells.
resistance of T1682 cells to selinexor. Nevertheless, we cannot completely rule out the potential role of other XPO1 cargoes to this resistance.

**XPO1 gene amplification confers acquired resistance to selinexor**

To further explore potential mechanisms of resistance to the XPO1 inhibitor, we analyzed experimentally established selinexor-resistant IU-TAB1 cells (referred to as IU-TAB1-R; Fig. 4A). The resistant cells displayed a 5-fold increase of selinexor IC₅₀ (400 nmol/L for IU-TAB1-R versus 80 nmol/L for the parental cells; Fig. 4B) and significantly less selinexor-induced apoptosis than the IU-TAB1 cells (Fig. 4C). While no acquired mutation was found in the XPO1 coding region in the IU-TAB1-R cells (data not shown), XPO1 protein levels were significantly elevated and higher concentrations of selinexor were required to inhibit XPO1 and to deregulate p53 and p27 in this resistant line (Fig. 4D). Gene copy-number analysis revealed at least 3.5-fold more copies of the XPO1 gene in the resistant cells than in the parental cells (Fig. 4E). A targeted exome sequencing of more than 200 cancer genes (13) confirmed XPO1 gene amplification but identified no additional alteration in the IU-TAB1-R cells (data not shown). Importantly, partial knockdown of XPO1 by siRNA (to a level comparable to that in the parental cells) was able to restore the sensitivity of IU-TAB1-R cells to selinexor (Fig. 4F and G). These data indicate that amplification of the XPO1 gene is one of the potential mechanisms of resistance to the XPO1 inhibitor.

**Selinexor inhibits TET xenograft tumor growth**

To evaluate antitumor efficacy of selinexor, two xenograft tumor models (MP57 and T1889) were treated with two different doses of selinexor (10 and 15 mg/kg, 3 times weekly) or vehicle for 4 weeks. Both dosages were effective and led to 47% and 43% growth inhibition of MP57 tumors, and 42% and 53% growth inhibition of T1889 tumors, respectively (Fig. 5A and B). Note-significantly higher concentrations of selinexor was required to inhibit MP57 tumors, and 42% and 53% growth inhibition of MP57 tumors, and 42% and 53% growth inhibition of T1889 tumors, respectively (Fig. 5A and B). Note-significantly less selinexor-induced apoptosis than the IU-TAB1 cells (Fig. 4C). While no acquired mutation was found in the XPO1 coding region in the IU-TAB1-R cells (data not shown), XPO1 protein levels were significantly elevated and higher concentrations of selinexor were required to inhibit XPO1 and to deregulate p53 and p27 in this resistant line (Fig. 4D). Gene copy-number analysis revealed at least 3.5-fold more copies of the XPO1 gene in the resistant cells than in the parental cells (Fig. 4E). A targeted exome sequencing of more than 200 cancer genes (13) confirmed XPO1 gene amplification but identified no additional alteration in the IU-TAB1-R cells (data not shown). Importantly, partial knockdown of XPO1 by siRNA (to a level comparable to that in the parental cells) was able to restore the sensitivity of IU-TAB1-R cells to selinexor (Fig. 4F and G). These data indicate that amplification of the XPO1 gene is one of the potential mechanisms of resistance to the XPO1 inhibitor.

**Proteomic profiling of factors and pathways affected by selinexor**

To extend the understanding of selinexor antitumor activity, we performed proteomic analyses of the cytoplasmic and nuclear fractions of IU-TAB1 cells treated with DMSO (light state) and selinexor (heavy state) by SILAC approach. Subcellular fractions of nucleus and cytoplasm from each state were subjected to proteomics by digestion with trypsin/LystC, basic RPLC fractionation, and LC-MS/MS analysis. In total, 72 LC-MS/MS runs from three biological replicates resulted in the identification of 3,246 and 1,617 proteins in the nucleus and cytoplasm, respectively (Fig. 6A; Supplementary Tables S1 and S2). With 1.5 fold-change cutoff, abundance of 303 proteins was decreased and of 63 proteins increased in the nucleus, whereas abundance of 185 proteins decreased and of 19 proteins increased in the cytoplasm upon selinexor treatment (Fig. 6A). The volcano plots indicate the affected proteins in the nucleus and cytoplasm with a minimum of 1.5-fold change combined with a P < 0.05 (Fig. 6B).

Interestingly, hierarchical clustering of 78 quantified transcription factors based on the SILAC ratio of selinexor over DMSO treatment (H/L) showed specific clusters of transcription factors whose abundance significantly changed in the opposite direction between the nucleus and cytoplasm (Fig. 6C). In particular, NPM1, CCAR1, GTF2I, and MYBBP1A decreased in cytoplasm and increased in the nuclear fraction (Fig. 6C), suggesting potential enhancement of transcription of genes regulated by these transcription factors upon XPO1 inhibition. Gene set enrichment analysis using a published transcription factor protein–protein interaction library (24) identified several significantly enriched transcription factor networks, including the NF-kB (Fig. 6D). In addition, a list of p53 network proteins was also significantly altered by selinexor (Supplementary Tables S3 and S4). Furthermore, Ingenuity Pathway Analysis (IPA) identified various enriched networks and canonical pathways affected by selinexor (Supplementary Tables S3 and S5). The effect of XPO1 inhibition on these transcriptional programs and functional networks/pathways likely contributes to the antitumor activity of selinexor cumulatively.

**High XPO1 expression is associated with aggressive histotypes and advanced stages of TETs and poor patient survival**

To evaluate whether targeting XPO1 is clinically relevant, we assessed XPO1 expression by IHC in all 6 different histological subtypes of TETs using a tissue microarray consisting of 132 TETs and 16 histologically normal thymic tissues (Fig. 7; Supplementary Fig. S3A–S3C; ref. 14). TET samples were mainly collected from primary surgery, and survival was assessable in all patients with a median follow-up of 82 months (95% CI, 68–94 months). Fourteen samples were excluded due to poor staining quality, and a total 118 TETs were evaluated for association study (Fig. 7A). Analysis of XPO1 staining took into consideration both the percentage of positive cells and intensity of the staining, as previously described (25). We defined samples with final IHC scores of 0 to 6 as XPO1-low and 7 to 12 as XPO1-high. Based on this cutoff, we found a significant association between high XPO1 and more aggressive histological subtypes of TETs (χ² test, P = 0.003), as higher percentages of thymomas type B2, B3, and thymic carcinoma (TC) expressed high XPO1 (Fig. 7A–C). High XPO1 were also significantly correlated with advanced stages of disease (10%–15% cases in stages I–II versus 50% cases in stages III–IV; P = 0.0006; Fig. 7A and D). Furthermore, a significant association between high XPO1 and poorer overall survival (OS) was observed (Wilcoxon P = 0.04; HR 2.025; 95% CI, 0.9188–4.463; Fig. 7E). Interestingly, higher XPO1 expression was a significant predictor of worse survival for stage I–II patients (median OS not reached in XPO1-low vs. 58 months in XPO1-high; P < 0.005; Fig. 7F), but not for patients at stage III–IV (Supplementary Fig. S3C). Finally, XPO1 expression retained its prognostic value in a Cox multivariate analysis of main known prognostic factors in TETs, including histologic grade, stage
Figure 3.
Selinexor confers p53-dependent and -independent antitumor activity in TET cells. A, Western blot detection of p53 and FOXO3a in the indicated TET cells after siRNA knockdown. B, Viability of T1889, IU-TAB1, MP57, and Ty82 cells treated with increasing concentrations of selinexor with/without control siRNA, p53 siRNA, and/or FOXO3a siRNA for 72 hours. C, Comparison of the AUC between cells treated with selinexor with and without p53 and/or FOXO3a siRNAs. Data represent mean values ± SD of triplicates. **, P < 0.01. D, Western blot detection of XPO1, FOXO3a, and p53 in the cytoplasmic (CE) and nuclear extraction (NE) of T1682 cells treated with DMSO or selinexor for the indicated times. E, Western blot confirmation of ectopic p53 expression in T1682 cells transfected with pIRES2-EGFP-p53 vector. Cells transfected with pIRES2-EGFP empty vector (Mock) were used as the control. F, Cell viability of Mock- or p53-transfected T1682 cells treated with increasing concentrations of selinexor for 72 h. IC_{50} concentrations were determined using GraphPad Prism program. G, Comparison of AUCs between p53- and Mock-transfected T1682 cells treated with selinexor. Data represent mean values ± SD of triplicates. **, P < 0.01.
of disease, and completeness of surgery (HR 3.6; 95% CI, 1.3–10.2; \( P = 0.01 \)).

**Discussion**

TETs are rare tumors originated from thymic epithelia, and therapeutic options are limited for advanced-stage diseases (26–28). Development of targeted therapy for TETs has been hampered by insufficient knowledge of their tumor biology and scarcity of targetable oncogenic driver abnormalities (28). In this study, we took an unconventional targeted approach to explore a potential therapeutic strategy for TETs by targeting XPO1. The concept of this strategy is to boost tumor suppressor activity by inhibiting XPO1-mediated nuclear export of key TSPs, thereby eliciting antitumor activity. XPO1 is highly expressed in TETs, and its inhibitor selinexor significantly increased nuclear p53, p27, and FOXO3a and induced antitumor activities at clinically achievable concentrations (29). Importantly, selinexor displayed less cytotoxicity in normal thymic epithelial cells, indicating its specificity against tumor cells (7).

XPO1 regulates nuclear–cytoplasmic shuttling of key cell-cycle regulators with different roles in G1–S and G2–M transition (7). Indeed, XPO1 inhibition induced cell-cycle arrest, albeit with varies of G1 and/or G2 arrest in different lines, likely due to different status and/or regulation of these cell-cycle regulators under different cellular contexts. The ability of selinexor to block cell cycle is intriguing in light of the encouraging antitumor activity of the CDK-inhibitor milciclib in TETs (30).

Along with the cell-cycle effect, selinexor induced apoptosis of TET cells, a crucial aspect of the antitumor activity resulted from XPO1 inhibition. Selinexor-induced apoptosis could be explained in part by the significant induction of the proapoptotic proteins BIM and BAX, which are not directly regulated by XPO1 but transcriptionally controlled by XPO1 cargoes such as p53 and/or FOXO3a proteins (4, 7, 19, 20). Interestingly, p53 and FOXO3a appeared to be differently needed for selinexor cytotoxicity in different cell lines, a likely reflection of different statuses of p53/FOXO3a and/or of their networking molecules in the cells. Importantly in the selinexor-resistant T1682 cell line, p53 deficiency is the primary cause of resistance, and p53 reinstallation...
Figure 5.
Selinexor exhibits in vivo antitumor activity against TET xenograft tumors. A and B, Growth curves of MP57 and T1889 xenograft tumors in athymic nude mice treated with vehicle or selinexor (10 mg/kg or 15 mg/kg) three times a week for 2 weeks. Arrows, when treatment started. Data represent the mean ± SD of tumor volumes (n = 8). *, P < 0.05; **, P < 0.01. C–E, IHC staining of Ki67 (C), cleaved caspase-3 (D), and p27 in MP57 and T1889 tumors (E) after treatment with/without selinexor for 2 weeks. Representative IHC images are shown in the left panels, and quantification of IHC staining results is shown on the right. Histograms represent mean ± SD of percentage of IHC positive cells in three tumors from each group. *, P < 0.05; **, P < 0.01; ***, P < 0.001.
leads to selinexor resensitization. Notably, about 20% human TETs harbor p53 inactivating mutations, indicating the importance of p53 loss in TET pathogenesis (21–23). Unlike p53, the tumor suppressor activity of FOXO3a could be diminished by other mechanisms, for instance, overactivation of the PI3K–AKT pathway, which causes XPO1-dependent cytoplasmic entrapment of FOXO3a (4, 7). It is clear, however, that the antitumor activity of selinexor in TETs may depend on p53, FOXO3a, and/or other XPO1 cargo proteins, even though p53 seems to be particularly important in most cases. While loss of p53 activity can cause resistance to the XPO1 inhibitor under certain circumstances, other alterations such as XPO1 amplification may also contribute...
Figure 7.
High XPO1 expression is associated with aggressive histotypes and advanced stages of human TETs as well as poorer patient survival. A, Characteristic of patient population and XPO1 expression statuses. B, Representative images of XPO1 IHC staining in different histotypes of thymomas and thymic carcinomas. C, Percentage of XPO1-high cases relative to the cases of each histological subtype. For comparison, normal thymic tissues were also examined. D, Percentage of XPO1-high cases relative to the analyzed cases of TETs in each corresponding stage. E and F, Kaplan–Meier patient survival curves plotted according to the XPO1 expression levels (XPO1-high versus XPO1-low) in the TETs at all stages (E) or at early stages (I–II; F).
to the resistance. Our conclusions are also supported by other studies that indicated that in certain tumors, expression of certain TSPs is needed for selinexor sensitivity, whereas in other tumors, inactivation of these TSPs does not affect drug sensitivity (18, 31–34).

The SILAC-based LC-MS/MS proteomic analysis has enabled us to profile the global impact of XPO1 inhibition on cellular factors/pathways/networks in the TET cells. In line with the changes of hundreds of XPO1 cargo proteins (4, 7), we found significant enrichment of several important pathways/networks that are functionally relevant to XPO1 inhibition. Among them include networks of "DNA replication, recombination, and repair," "cellular growth and proliferation," and "cell death and survival" (Supplementary Table S3), and the sum of these changes likely reflects the overall antitumor effect of selinexor. Nonetheless, the most interesting observation from the proteomics analysis is the deep impact of XPO1 inhibition on the transcriptional programs. Selinexor treatment causes a prominent shift in the abundance of nearly 80 different transcription factors in the nucleus and cytoplasm, with NPM1, CCAR1, GTF2I, and MYBBP1A being the most affected ones.

Both MYBBP1A and NPM1 are known XPO1 cargoes. MYBBP1A can bind to p53 and enhance the transcription of p53 target genes (35, 36), and its inhibition significantly enhanced tumorigenesis (35). MYBBP1A can also interact with NF-kB and inhibit NF-kB–dependent transcription (36). On the other hand, NPM1 is involved in many cellular functions, which require continuous nuclear–cytoplasmic shuttling of NPM1 (37). Notably, NPM1 has been characterized as the most frequently mutated gene in acute myeloid leukemia and the most frequent frameshift mutation at its exon 12 results in an additional nuclear export signal recognized by XPO1, leading to aberrant nuclear export and stable cytosolic localization of NPM1 (37). Evidence suggests that when mislocalized in the cytoplasm NPM1 acquires oncogenic properties and alters cell cycle and apoptotic regulation (37).

Unlike NPM1 and MABB1A, however, GTF2I and CCAR1 are not known cargoes of XPO1. The identification of GTF2I as one of the top transcription factors in TET cells affected by XPO1 inhibition is of particular interest. In our recent study, we reported the identification of a single highly frequent mutation in the GTF2I gene in a large cohort of human TETs (22). GTF2I is a general transcription factor that regulates many target genes expression, and its two major splicing isoforms have been shown to exert opposite transcriptional activities due to differences in their subcellular localizations (38). The shift in the subcellular location of GTF2I from the cytoplasm to the nucleus after XPO1 inhibition reported in this study as well as recently documented (39) suggests that the nuclear–cytoplasmic shuttling of GTF2I is somehow regulated by XPO1. Further study will be required to determine whether nuclear entrapment of these transcription factors contributes to the antitumor activity of selinexor.

Even though the impact of selinexor on the p53 and NF-kB proteins was evident in Western blot and immunofluorescence analyses, both proteins were not identified by the LC-MS/MS approach. This is not a surprise as protein identification with mass spectrometry is a stochastic process and depends on multiple variables. Nevertheless, the proteomics analysis revealed a significant enrichment of a numbers of proteins in the networks of p53 or NF-kB. NF-kB is an important transcription factor that regulates many cellular processes including cell proliferation and survival. Although not functionally examined in this study, inhibition of NF-kB by selinexor likely plays a role in the selinexor-induced cytotoxicity in TETs as well. Recently, it was shown that a synthetic–lethal interaction of XPO1 inhibition with KRAS mutation in NSCLC cells requires its inhibition of NF-kB transcription factor activity (34).

Most importantly, targeting XPO1 in TETs is not only biologically achievable and effective but also clinically relevant. A recent phase I study indicates clinical activity of selinexor in patients with TETs. Four patients with TETs were enrolled in this trial, of which 1 patient obtained partial response, 2 patients had stable disease lasting more than 4 months, and 1 patient had stable disease lasting less than 4 months (8). Here, we demonstrated that XPO1 is highly expressed in a subset of human TETs, and overexpression of XPO1 is significantly associated with aggressive histological subtypes, advanced stage of disease, and poor patient survival. These data together with the antitumor activity of selinexor in the preclinical TET models support clinical development of XPO1 targeted therapy for the treatment of TETs. In fact, we are on the process of launching a phase II trial of selinexor in TET patients.

Disclosure of Potential Conflicts of Interest
No potential conflicts of interest were disclosed.

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