Cytoplasmic PML promotes TGF-β-associated epithelial–mesenchymal transition and invasion in prostate cancer

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Epithelial–mesenchymal transition (EMT) is a key event that is involved in the invasion and dissemination of cancer cells. Although typically considered as having tumour-suppressive properties, transforming growth factor (TGF)-β signalling is altered during cancer and has been associated with the invasion of cancer cells and metastasis. In this study, we report a previously unknown role for the cytoplasmic promyelocytic leukaemia (cPML) tumour suppressor in TGF-β signalling-induced regulation of prostate cancer-associated EMT and invasion. We demonstrate that cPML promotes a mesenchymal phenotype and increases the invasiveness of prostate cancer cells. This event is associated with activation of TGF-β canonical signalling pathway through the induction of Smad proteins and Mad related family 2 and 3 (SMAD2 and SMAD3) phosphorylation. Furthermore, the cytoplasmic localization of promyelocytic leukaemia (PML) is mediated by its nuclear export in a chromosomal maintenance 1 (CRM1)-dependent manner. This was clinically tested in prostate cancer tissue and shown that cytoplasmic PML and CRM1 co-expression correlates with reduced disease-specific survival. In summary, we provide evidence of dysfunctional TGF-β signalling occurring at an early stage in prostate cancer. We show that this disease pathway is mediated by cPML and CRM1 and results in a more aggressive cancer cell phenotype. We propose that the targeting of this pathway could be therapeutically exploited for clinical benefit.

INTRODUCTION

The gene encoding the promyelocytic leukaemia (PML) tumour suppressor is a member of the Ring-B box-Coiled Coil family, originally identified at the breakpoint of the t(15;17) translocation found in acute promyelocytic leukaemia (APL).1–3 PML is involved in the regulation of many cellular pathways such as apoptosis, senescence, response to DNA damage and resistance to viral infection.4–7 Although the function of PML has been extensively investigated because of its essential role in the formation of PML-nuclear bodies (PML-NBs), the subcellular localization of PML is not restricted to the PML-NBs and is observed in other cellular compartments including the nucleoplasm, the nucleolus, the nuclear envelope and the cytoplasm.8,9 The roles of PML in these cellular compartments remain unclear, and may involve mechanisms that are independent of PML-NBs. There are seven main isoforms of PML (I–VIIb), which appear to have cellular functions that are related to their expression pattern in physiological and pathological cellular contexts. Most of the isoforms contain the nuclear localization signal (NLS), which enables them to localize in the nucleus (PML I to PML VI), whereas PML VIIb lacks the NLS and is found in the cytoplasm.9 The PMLI isoform also possesses a nuclear export signal (NES), which allows its nuclear export in a CRM1-dependent manner.10

Recent studies suggest the involvement of cytoplasmic PML in tumorigenesis, glycolysis, anti-viral responses, laminopathies and cell cycle regulation.8 In a physiological context, cytoplasmic PML has been shown to activate transforming growth factor (TGF)-β signalling, a cellular pathway involved in tumour suppression, apoptosis and senescence.11–13 Furthermore, defects in TGF-β signalling in Pml−/− mouse embryonic fibroblasts have also been reported.14 TGF-β binding to its receptors results in the phosphorylation of the SMAD proteins, a cellular event that is facilitated by the TGF-β-receptor adapter SARA (SMAD Anchor for Receptor Activation). This process results in the translocation of SMADs to the nucleus and the activation of target genes.15 In this context, cytoplasmic PML appears to favour the interaction of SARA and the SMAD protein members, SMAD2 and SMAD3.14 Although TGF-β signalling is a potent inhibitor of cell proliferation, this function seems to be altered in tumour cells, in that it acts as an oncogene.16–21 In particular, defects in normal TGF-β signalling have been shown to trigger tumorigenic events including cancer-associated epithelial–mesenchymal transition (EMT), an important process that is involved in the progression of advanced tumours.22,23 Based on this background, we hypothesized that cytoplasmic PML promotes cancer-associated EMT and local tumour invasion via the induction of TGF-β signalling. Furthermore, we reveal a new role for cytoplasmic PML in the progression of prostate cancer. To better understand the role of cytoplasmic PML in tumour progression, we investigated its influence on EMT and cell invasion using prostate cancer cell lines and mutant constructs. Furthermore, we showed the translational importance of this proposal in a clinical setting by showing that cPML expression correlates with the expression of the nuclear export factor CRM1 (Exportin-1), and is associated with a worse prognosis.
These studies demonstrate that cPML induces cell invasion by promoting the transdifferentiation of epithelial cells into mesenchymal cells involving TGF-β signalling, via SMAD2 and SMAD3 phosphorylation. Collectively, these results support the concept that cPML promotes TGFβ-induced EMT promoting a more aggressive cancer cell phenotype resulting in increased cell invasion and reduced survival for patients with prostate cancer.

RESULTS

PML cytoplasmic localization is mediated by CRM1-dependent nuclear export

PML expression is altered in cancers of various histological origins.24 To investigate the expression status of PML, we performed immunostaining on prostate adenocarcinoma tissues using an antibody against PML. In normal prostate tissue, PML was expressed in the nucleus of cells within the prostate gland. The expression was typical of that of PML-NBs (Figure 1a). In contrast, PML expression was decreased (Figure 1b, upper panel) or exhibited nucleo-cytoplasmic and/or cytoplasmic expression patterns (Figure 1b, lower panel) in prostate adenocarcinoma cells. Similar results were obtained by immuno-fluorescence (IF), in that PML is expressed in the nucleus of the normal prostate cell line PNT2, and in the nucleo-cytoplasmic and/or cytoplasm of the prostate cancer cell lines DU145, PC3 and LNCaP (Figures 1c and d). Although the decrease of PML expression in cancer tissues has been previously reported, PML localization in the cytoplasm and its role in that cellular compartment are not understood. To investigate this, we performed immunoprecipitation (IP) experiments using a PML antibody and whole-cell extracts from DU145 human prostate cancer cells. The efficiency of the IP was determined by investigating the presence of PML in PML-immunoprecipitates and using immunoblotting (IB) with PML antibody. PML immunoprecipitates were also used to identify PML-interacting partners using MALDI-TOF mass spectrometry. PML and several molecules were identified as PML-interacting partners (Figure 1e). Interestingly, one of the interacting proteins identified was CRM1 (Exportin-1), a molecule which is involved in nuclear export. Therefore, the cytoplasmic localization of PML within prostate cancer cells might be a consequence of its nuclear export in a CRM1-dependent manner. To confirm the interaction of PML with CRM1, IP of DU145 extracts with a PML antibody was performed (Figure 1f). In addition to PML, CRM1 was also pulled down as demonstrated by IB of the immunoprecipitates with PML and CRM1 antibodies. Finally, the nuclear export of endogenous PML in DU145 and PC3 cells was markedly decreased after treatment with Leptomycin B also affected cell migration (Supplementary Figure 2c). Furthermore, knockdown of endogenous PML in DU145 and PC3 resulted in a decreased cell migration. Treatment of these cells with Leptomycin B also affected cell migration (Supplementary Figure 2c). Finally, CRM1 knockdown in DU145 expressing PML ΔNL5 did not significantly affect cell migration (Supplementary Figure 3d). The increased cell migration and invasion observed in the presence of PML ΔNL5 was not caused by increased proliferation (Supplementary Figure 3e). These results suggest that the cytoplasmic fraction of PML promotes the migration and invasive capacities of prostate cancer cells.

The expression of PML constructs also influenced the cellular morphology of DU145 and PC3 prostate cancer cells. Notably, cells expressing the PML ΔNL5-HA construct exhibited a mesenchymal phenotype, which was characterized by the presence of individual elongated cells. In contrast, DU145 and PC3 cells expressing PML ΔNES-HA exhibited a more epithelial phenotype. The cells expressing the wild-type construct presented an intermediate phenotype, which was probably a result of its dual nuclear and cytoplasmic expression (Figures 4a and b). To confirm these observations, we performed IF on these cells using antibodies that recognize key markers of epithelial cells (E-Cadherin) and mesenchymal cells (Vimentin and N-Cadherin). Lower E-cadherin expression and increased expression of Vimentin and N-Cadherin were observed in cells expressing PML ANL5. On the contrary, cells expressing PML ΔNES-HA, expressed higher levels of E-Cadherin, but a lower expression of Vimentin and N-cadherin. Some of the cells that expressed PML WT-HA exhibited a higher expression of E-Cadherin, whereas others expressed higher levels of Vimentin and N-Cadherin. To confirm these observations, we performed IB of lysates from cells expressing the empty vector, PML WT-HA, PML ΔNL5-HA and PML ΔNES using expression of PML, nor the expression of cytoplasmic CRM1 protein alone was found to be predictive of reduced disease-specific survival or metastasis (Supplementary Table 2). To test the hypothesis that PML is transported by CRM1, the capacity of PML/CRM1 expression to predict clinical outcome was assessed. Eighty percent (40/50) of patients exhibited either complete loss of PML and CRM1 in tumours or simultaneous expression of cytoplasmic PML and CRM1. This patient group showed reduced cancer-specific survival (χ² = 6.47, P = 0.011) over a 10-year term (Figure 2e), but not significantly increased metastasis over the same term (χ² = 3.108, P = 0.078). A Cox proportional hazards model demonstrated that simultaneous expression of cytoplasmic PML/CRM1 predicted reduced disease-specific survival that was independent of prostate-specific antigen levels (>4 ng/ml) and the Gleason score (>7; hazard ratio = 0.264, P = 0.009, 95% confidence interval = 0.097–0.720).
Figure 1. PML expression in normal and cancer prostate cells. (a and b) Immunohistochemistry/fluorescence micrographs showing PML expression (green) in normal prostate tissue and prostate adenocarcinoma. The nucleus is stained in blue (4,6-diamidino-2-phenylindole (DAPI)). Scale bar = 100 μm. (c and d) Immunofluorescence micrograph showing PML expression (red) in normal prostate cell line (PNT2) and in prostate cancer cell lines (DU145, PC3 and LNCaP). Scale bar = 20 μm. (e) Schematic representation of the methodology used to identify PML-interacting partners from DU145 cell extracts using immunoprecipitation (IP) with PML antibody and MALDI-TOF mass spectrometry. (f) Immunoprecipitation of DU145 cell extract with PML antibody and immunoblotting (IB) with PML and CRM1 antibodies. (g) Immunofluorescence micrograph showing PML expression in DU145 and PC3 treated with or without Leptomycin B (15 ng/ml). Scale bar = 20 μm.
antibodies recognizing key markers of epithelial and mesenchymal phenotypes (Figures 4c and d). PMLΔNLS cells expressed lower levels of epithelial markers and higher levels of mesenchymal markers (Figures 4c and d). PMLΔNES expressed higher levels of epithelial markers and lower levels of mesenchymal markers. These results demonstrate that cytoplasmic PML promotes EMT, whereas nuclear PML promotes an epithelial phenotype.

Cytoplasmic PML-induced EMT and cell invasion are mediated by TGF-β signalling

To investigate the potential involvement of TGF-β signalling pathway in PML-mediated EMT and cell migration and invasion, we performed IB of lysates from DU145 and PC3 cells expressing the empty vector, PMLWT-HA, PMLΔNLS-HA and PMLΔNES using an antibody against the phosphorylated forms of SMAD2 and 3. Interestingly, higher levels of phospho-SMAD2/3 were observed in...
PML ΔNLS-expressing cells, when compared with cells expressing the other constructs (Figure 5a). To confirm this result, we performed IB using the phospho-SMAD2/3 antibody on cell lysates from DU145 and PC3 cells expressing PML ΔNLS that had been treated with a specific TGFβRI inhibitor (untreated cells as controls). Treatment with the inhibitor significantly reduced the...
level of phospho-SMAD2/3 expression, but did not affect cPML expression (Figures 5b and c). E-Cadherin expression levels were increased following treatment with the inhibitor (Figures 5b and c), whereas the expression of N-Cadherin was decreased (Figure 5c). As the inhibition of TGF-β signalling in this experimental setting may also affect cell migration, we tested the influence of the TGFβRI inhibitor on DU145 and PC3 cells expressing PML mutant constructs using antibodies against molecules expressed by epithelial cells (E-Cadherin, β-Catenin, ZO-1 and Claudin1) and mesenchymal cells (Vimentin, N-Cadherin, SNAIL, SLUG and ZEB1). β-Actin represents the loading control.

**DISCUSSION**

EMT is a key cellular event that is involved in embryonic development, wound healing and cancer progression. The baso-apical polarity of epithelial cells is maintained through lateral cell–cell junctions that include desmosomes, adherens junctions, tight junctions and gap junctions. During EMT, epithelial cells, which begin their transdifferentiation via the loss of lateral junctions, start to express molecules associated with the mesenchymal phenotype and this results in the generation of mesenchymal cells. These later have an essential role in the spread of disease to adjacent tissues and other organs via a process called metastasis. Upon reaching the target organ, these cells begin a reverse cellular process (mesenchymal–epithelial transition), which results in the generation of cancer epithelial cells, which integrate into the structure of target tissue and therefore ‘ignite’ its oncogenic transformation. EMT can be naturally induced by several cellular pathways such as growth factors, TGF-β, WNT, HH and Notch signalling. In cancer, these pathways can be altered and result in cancer-associated EMT. Notably, TGF-β signalling is a well-known tumour-suppressive pathway, and cancerous cells circumvent this function by inactivating the anti-proliferative arm of the pathway or by altering the function of downstream components of the pathway.27

EMT is induced as a consequence of TGF-β signalling dysfunction, to result in an increase in the invasion and dissemination of cancer cells. In this regard, cytoplasmic PML appears to activate the phosphorylation of SMAD2 and SMAD3, two main components of the canonical SMAD pathway in...
Figure 5. Cytoplasmic PML induces SMAD2/3 phosphorylation, EMT and cell invasion in vitro. (a) Immunoblotting of whole-cell extracts from DU145 and PC3 cells expressing PML mutant constructs using antibodies against phosphorylated forms of SMAD2/3 and β-actin (loading control). (b) Immunoblotting analysis of whole-cell extracts from DU145 and PC3 cells expressing the PML ΔNLS construct, treated with DMSO (control) or TGFβ-RI inhibitor (3 μM for 48 h) using antibodies against HA, phosphorylated forms of SMAD2/3, E-Cadherin and β-actin (loading control). (c) Immunoblotting of whole-cell extracts from DU145 and PC3 cells expressing PML mutant constructs using antibodies against total SMAD2, SMAD3 and β-actin (loading control). (d) Immunofluorescence analysis of DU145 and PC3 cells expressing the PML ΔNLS construct, treated with DMSO (control) or TGFβ-RI inhibitor (3 μM for 48 h) using antibodies against phosphorylated forms of SMAD2/3, E-Cadherin and N-Cadherin. Scale bar = 20 μm. (e, f) Wound healing assay (Scratch assay) using DU145 and PC3 cells expressing the PML ΔNLS construct, treated with DMSO (control) or TGFβ-RI inhibitor (3 μM for 48h) and the data relating to the percentages of gap closure at time points 0 and 24 h following scratching. Two-tailed paired test: *P = 0.0451 and **P = 0.0088.
response to TGF-β. In this study, we observe a similar mechanism, in that cPML induces TGF-β signalling through cPML induction of SMAD2 and SMAD3 phosphorylation. However, and in comparison with Lin et al. study which used a physiological model (mouse embryonic fibroblasts), our work investigated the role of cPML in the regulation TGF-β signalling within a pathological setting, namely prostate cancer. In this context, we showed that cPML promotes TGF-β-induced prostate cancer-associated EMT and invasion, a mechanism that was previously unknown (Figure 7). This difference is not surprising given that the consequences of TGF-β signalling are dependent on whether the cellular context is physiological or pathological. Moreover, cPML is mainly found in the cytoplasm of tumour tissue for patients with prostate cancer that exhibited a poor clinical prognosis. In addition, several reports have reported on the oncogenic potential of cPML mutants. Two different PML mutations (1272delAG and IVS3–1G-A) have been identified in aggressive APL patients. The mutations generated a stop codon before the NLS domain leading to the generation of cPML mutants. Other studies have shown that cPML is upregulated in hepatocellular carcinoma. In the present study, we showed increased cPML and cytoplasmic expression of Exportin-1 (CRM1) in tumour tissue of patients with prostate cancer correlates with a poor clinical prognosis, suggesting a functional link between the two molecules. Indeed, the PMLI isoform is the only known isoform that contains a NES, and thereby could be exported to the cytoplasm.
cytoplasm. This possibility has been demonstrated, in that we have shown that the mutant PMLI which lacked the NES promoted a more epithelial phenotype, which is associated with an increased expression of epithelial markers. This suggests that nuclear PML may prevent cancer-associated EMT and may have a role in promoting mesenchymal–epithelial transition. Taken together, these observations highlight the involvement of cPML in prostate cancer progression and provide further insights on the role of deregulated TGF-β signaling in pathogenesis of cancer invasion.

**MATERIALS AND METHODS**

**Antibodies**

For this study, we used antibodies to PML (1:250 for IB, 1:100 for IP, sc-966, Santa Cruz Biotechnology, Delaware Ave., Santa Cruz, CA, USA), PML (1:500 for immunohistochemistry (IHC), sc5621, Santa Cruz Biotechnology, Delaware Ave., HA (1:1000 for IB, 2:200 for IF, A190-108 A, Bethyl Laboratories Inc., Montgomery, TX, USA), β-actin (1:5000 for IB, A5441, Sigma-Aldrich, St Louis, MO, USA), Exportin-1 (CRM1; 1:5000 for IB, 1:200 for IHC, A300-469A, Bethyl Laboratories Inc.), E-Cadherin (1:500 for IF, 610181, BD Transduction Laboratories, San Jose, CA, USA), EMT antibody Sampler Kit (9782, Cell Signaling Technology, Danvers, MA, USA), which contains rabbit antibodies each used at 1:300 (IB), N-Cadherin and Vimentin antibodies were used at 1:500 for IF, Phospho-Smad2/Smad3 (1:300 for IB, 8828, Cell Signaling Technology), Phospho-Smad2/3 (1:100 for IHC, sc-11769, Santa Cruz Biotechnology), Anti-rabbit IgG, HRP-linked Antibody (1:1000 for IB, 7074, Cell Signaling Technology), Anti-TGFβ Receptor I (1:1000 for IB, 3712, Cell Signaling Technology), Anti-Smad2 (1:1000 for IB, 3122, Cell Signaling Technology), Anti-Smad3 (1:1000 for IB, 9513, Cell Signaling Technology), Anti-phospho-Smad2/3 (1:500, SAB4504207, Sigma-Aldrich), Anti-phospho-Smad3 (1:500, SAB4300253, Sigma-Aldrich), Anti-mouse IgG, HRP-linked antibody (1:1000 IB, 7076, Cell Signaling Technology).

**Cell lines, growth conditions, cell fractionation and cell proliferation assay**

DU145 and PC3 prostate cancer cell lines were purchased from American Type Culture Collection (Manassas, VA, USA) and cultured in Eagle’s Minimum Essential Medium (BE12-6621, Lonza, Basel, Switzerland) and F-12k Nut Mix (1x; 21127-022, Gibco Life Technologies, Waltham, MA, USA) respectively. The media were supplemented with 10% (v/v) fetal calf serum and 1% (w/v) L-glutamine (Lonza). Cells were incubated at 37 °C in 5% (v/v) CO2 and 100% (v/v) humidity. For the relevant experiments, cells were treated for 48 h with 3 μM of TGF-βR1 inhibitor SB431542 (1614, Tocris Bioscience, Moorend Farm Avenue, Bristol, UK) and for 20 h with 15 ng/ml of Leptomycin B. TGF-β1 was purchased from PeproTech (London, UK) (100-22) and used at 10 ng/ml. Cell proliferation was quantified using CyQUANT NF Cell Proliferation Assay Kit (C35006, Invitrogen, Waltham, MA, USA).

**Retroviral expression of PML constructs**

The PMLI gene fragment was obtained by PCR amplification from pcDNA3.1-PMLI (a kind gift from Dr Mounira Chelbi-Alix, University PARIS V) using forward primer SpeI/PML F: 5’-CATCTAACTAGTATGGAGAGCCCTGCG-3’. The resulting PCR product was cloned into the SpeI site of pCMV-SPORT6 (Invitrogen, Waltham, MA, USA) and sequenced for correct orientation. The clones were linearized by NotI and transfected into the packaging cell line 293 T. The supernatants were collected 48 h post transfection and used to transduce the prostate cancer cell lines. The transduced cells were selected with puromycin (2 μg/ml) and stable clones were isolated by limiting dilution. The stability of the PML constructs was verified using Western blotting with antibodies specific to PML (1:250 for IB, sc-966, Santa Cruz Biotechnology).
ACCCGCC-3′ and reverse primer PML R′/BanHII 5′-CATCTAGATCATCCTACGC
TCTGCTCGAGGACC-3′ (Eurofins MWG Operon). The fragment was obtained when clonned into the plKOpuro vector using the Spel and BanHII restriction sites (SHC001, Sigma, St Louis, MO, USA). An HA-tag was introduced in the C-terminal end of PML by PCR using QuikChange Site-Directed Mutagenesis Kit (200518, Stratagene, Cedar Creek, TX, USA) and the following primers: HA-tag-sense 5′-CCCCCAAGCAGACCTACCTATGATGCAGATT
ACGCTTGAGATCACCAGG-3′ and HA-tag-antisense 5′-CCGGTGGATCCTCA
AGCGTAATCTGGAACATCGTATGGGTAGCTCTGCTGGGAGG-3′. The deletions of PML NLS and PML NES were also performed by PCR using QuikChange Site-Directed Mutagenesis Kit using ∆NLS-sense 5′-GCCCAAGGAGTTGGCG
GAAGAAGGACAG-3′; ∆NLS-antisense 5′-CTTGCTCTCTGCCAGGCTTCTGCG
GCC-3′; ∆NES-sense 5′-ACATTTAAACAGCTGGAAGGAAATGCGCCGGG-3′; ∆NES-antisense 5′-GCCGGGCGACCTTTCCAGCTTAATG-3′. The lenti-viral particles used to infect DU145 and PC3 with plKO-PML constructs were produced according to the manufacturer’s recommendations. The lentiviral packaging mix was also purchased from Sigma (SHP001).

Stable and transient transfection of cell lines
The transient knockdown of CRM1 and TGFβRII was carried out using an
CRM1- or TGFβRII-specific small interfering RNA molecule (CRIS1 small
interfering RNA (h), sc-35116, Santa Cruz Biotechnology, Inc; TGFβRII small
interfering RNA (h), sc-40222, Santa Cruz Biotechnology, Inc) and Interferon
transfection reagent (Polyplus, Woodley, Reading, UK) following the
manufacturer’s recommendations. PML lentiviral plasmids short hairpin
RNA control and PML short hairpin RNA 1 and 2 were purchased from Sigma (TRCN0000008866, TRCN000000869 and SHC001).

Immunoblotting and immunoprecipitation
The experimental procedures were performed as described previously.30
For IB, the cells lysed in 1× solution containing 50 mM Tris-HCl (pH 6.8),
100 mM dithiothreitol, 2% (w/v) SDS, 0.1% (w/v) bromophenol blue and 10% (v/v) glycerol, and loaded onto Tris/glycine SDS–polyacrylamide gels
for electrophoresis. The proteins were transferred onto Amersham
Hybond-P PVDF membranes (GE Healthcare, Chalfont St Giles, Buck-
inghamshire, UK). Membranes were blocked with 10% (w/v) Marvel milk
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buffer and suspended in lysis buffer, after which samples were analysed by
overnight at 4 °C followed by washing and incubation with secondary
washes in tris-buffered saline solution with 0.1% (v/v) Tween-20,
inghamshire, UK). Membranes were blocked with 10% (w/v) Marvel milk

step at 4 °C followed by washing and incubation with secondary
antibodies for 1 h at room temperature before visualization using Rapid
Step ECL reagent (Calbiochem, San Diego, CA, USA) and a CCD camera
overnight at 4 °C followed by washing and incubation with secondary
washes in tris-buffered saline solution with 0.1% (v/v) Tween-20,
inghamshire, UK). Membranes were blocked with 10% (w/v) Marvel milk

stable extracts for overnight IP. The beads were washed four times in IP
buffer and suspended in lysis buffer, after which samples were analysed by

using the software programme X-tile to determine the optimal cutoff.32
The cutoff values used were PML: nuclear (230), cytoplasm (80); CRM1: cytoplasm (110). Patients were classified according to PML and
CRM1 expression patterns (Supplementary Table 2) and a Kaplan–Meier
survival plot performed to compare the different classes according to
disease-specific survival. Subsequently, a group of patients exhibiting a poor prognosis was defined on the basis of an absence of
PML and CRM1 expression, or simultaneous cytoplasmic expression of
PML and CRM1.

REMARK guidelines for reporting on prognostic biomarkers were
followed and a flow diagram showing the final number of cases that
were analysed as a proportion of the whole cohort is shown in
Supplementary Figure 1.31 Following immunostaining, the number of
samples available for analysis was reduced for several reasons: the amount
of available malignant tissue was considered too small to be representative
of the tumour, tissue cores were detached during processing, cases had
data for only one biomarker/case or clinical data (survival or metastasis)
were not available for follow-up.

Liquid chromatography-matrix-assisted laser desorption/ ionization time-of-flight mass spectrometry identification of
PML-interacting partners
Following IP, identification of the potential interacting partner proteins
was carried out by LC-MALDI mass spectrometry. Samples were tryptically
digested and fractionated by nano-LC (C18 PepMap100, 150 mm × 75 μm;
5 μm particle size) before spotting on 10 s fractions over a 384-spot target
plate. Analysis was carried out on a Bruker UltraflaXtreme MALDI-TOF/TOF
mass spectrometer (Bruker Daltonics, Banner La, Coventry, UK) in
reflectron-positive mode over a m/z range of 650–3500. Tandem mass
spectrometry fragmentation of the top 10 precursors per spot was carried out and results exported using Biotools 3.1 software (Bruker Daltonics)
for searching via Mascot 2.1 against the human Swissprot database.

Cell migration and invasion assays
For wound healing assay (Scratch assay), DU145 and PC3 cells expressing the
empty vector, PMLI WT-HA, PMLI ∆NLS-HA and PMLI ∆NES-HA were
cultured to 80% confluence and serum starved for 24 h, after which a
scratch was made in the middle of each well using a 10-μl pipette tip. Images
taken of replicate experiments at 0 and 24 h were taken and the distances
between the edges of the scratch were measured at three
different points using Carl Zeiss Axiowision software (Carl Zeiss,
Oberkochen, Germany). The measurements were expressed as percentages
of gap closure. For the well cell migration and invasion assay, we used
Cultrex 96-Well Cell Invasion Assay (3465-96-K, Cultrex) and Cultrex
96-Well BME Cell Invasion Assay following the manufacturer’s recom-
endations (Trevigen, Gaithersburg, MD, USA). The cells in the bottom
chamber were labelled by Calcein AM (Trevigen) and the quantification
of the number of cells was performed using the plate reader Infinite
M200 Pro TECAN (Tecan Group Ltd, Männedorf, Switzerland) at 585 nm
excitation and 520 nm emission.

CONFLICT OF INTEREST
The authors declare no conflict of interest.

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