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13. ABSTRACT (Maximum 200 Words) Antiestrogens and selective estrogen receptor modulators (SERMs) such as tamoxifen are effective in controlling the progression of estrogen receptor (ER)- positive breast tumors to more invasive phenotypes. However, over time, many patients acquire resistance to tamoxifen. The mechanisms underlying tamoxifen resistance remain elusive. Proline-, glutamic acid-, and leucine-rich protein-1 (PELP1) is a novel ER coactivator that plays a role in both genomic and nongenomic actions of ER. PELP1 is predominantly localized in the nucleus in hormonally responsive tissues, but recent studies suggest that PELP1 may be exclusively localized in the cytoplasm in cancer cells. We found that MCF-7 cells engineered to specifically express PELP1 in the cytoplasm (PELP1-cyto), exhibited hypersensitivity to estrogen but resistance to tamoxifen. In addition, PELP1-cyto cells exhibited increased association of PELP1 with Src, enhanced MAPK activation, and constitutive stimulation of AKT. Altered localization of PELP1 was sufficient to trigger PELP1's interaction with the p85 subunit of PI3 kinase and PI3K activation. Consistent with these findings, human tumors expressing PELP1 in the cytoplasm showed increased AKT activation. Our results suggest that altered localization of coactivators such as PELP1 promotes resistance to hormonal therapy, presumably via stimulation of nongenomic estrogen responses such as activation of MAPK and AKT.				
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Title: Significance of Localization of Coactivators in Tamoxifen Resistance

PI: Ratna K Vadlamudi

INTRODUCTION

The estrogen receptor (ER) has been implicated in the progression of breast cancer. About 60-70% of breast tumors are ER positive at presentation. Antiestrogens and selective estrogen receptor modulators (SERMs) are effective in controlling the progression of ER-positive breast tumors to more invasive phenotypes. However, many patients with metastatic breast tumors eventually develop resistance to antiestrogens and SERMs. Several mechanisms have been proposed for resistance to hormonal therapy, including expression of variant or mutant ER, ligand-independent activation of ER, adaptation of tumors to lower concentrations of estrogen, and pharmacological alterations. Emerging data also suggest that ER coregulators might play a role in hormonal responsiveness and tumor progression. ER, a ligand-dependent transcription factor, modulates transcription of a number of genes and contributes to genomic responses. Recent studies suggest that in addition to its well-studied nuclear functions, ER participates in cytoplasmic and membrane-mediated signaling events (nongenomic signaling). Nongenomic signaling has been linked to the generation of rapid responses to estrogen and generally involves stimulation of the Src, MAPK, PI3K and PKC pathways in the cytosol. Proline-, glutamic acid-, and leucine-rich protein-1 (PELP1),¹ also known as modulator of nongenomic activity of estrogen receptor (MNAR)^{2,3} is a novel ER coregulator that has been implicated in both genomic activity of ER via histone interactions⁴ and nongenomic functions of ER via activation of Src/MAPK pathways. Although PELP1 is predominantly localized in the nucleus in hormonally responsive tissues⁴, recent studies suggest that under certain conditions, PELP1 could be exclusively localized in the cytoplasm of a subset of cancer cells⁵. In this study we examined the **hypothesis that altered localization of PELP1 may contribute to the hormonal independence and tamoxifen resistance via excessive activation of nongenomic signaling pathways.**

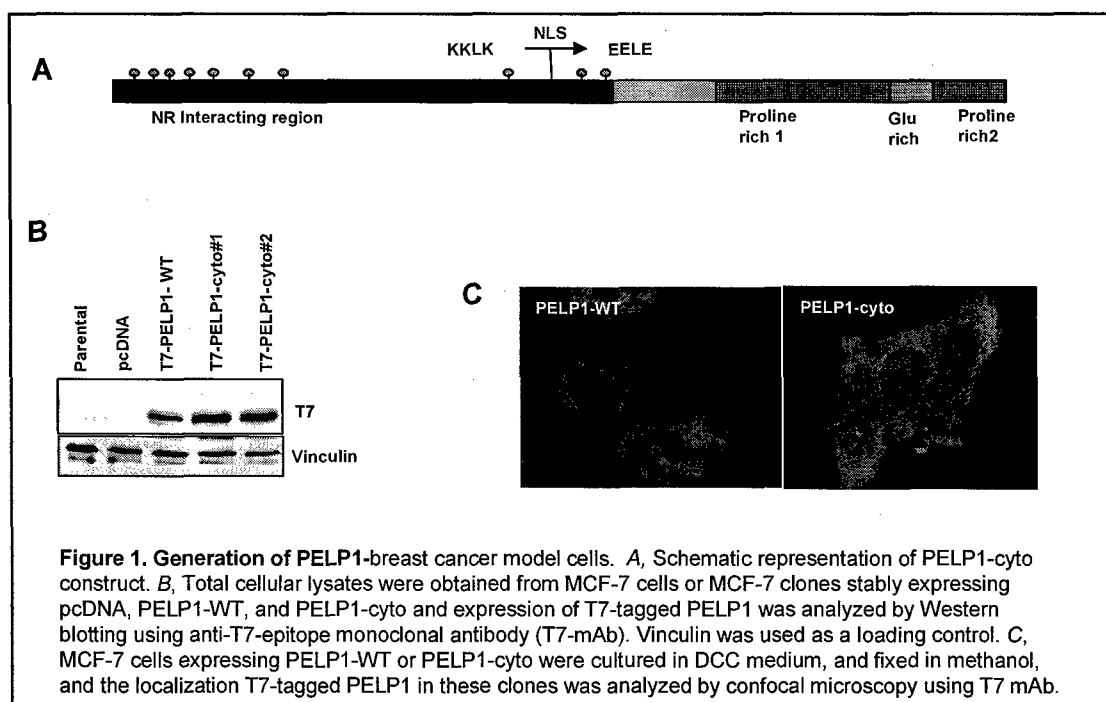
BODY

The scope of this proposal is to undertake the following two tasks outlined in the approved statement of work

Task1. Generation of MCF-7 model cells

Task2. Testing and characterization of model cells for hormonal responsiveness

Task 1. Generation of MCF-7 model cells expressing PELP1-cyto mutant: To examine the functional consequences of cytoplasmic localization of ER coactivator PELP1, we generated a mutant of PELP1 (PELP1-cyto) in which the nuclear localization signal was mutated from KKLK to EELE. Using this construct, we generated MCF-7 cells stably expressing PELP1-cyto mutant. To avoid potential clonal variations, we developed two pooled clones. Western blotting showed expression of T7-tagged-PELP1-cyto mutant in these clones. As a control, we have used an earlier characterized wild type PELP1 (PELP1-WT) pooled clone. Confocal scanning analysis showed that PELP1-cyto mutant was localized to the cytoplasm while PELP1-WT was localized to the nucleus.



Task 2. Testing and characterization of PELP1-cyto model cells for hormonal

responsiveness : Since earlier studies have shown that PELP1 interacts with Src kinase, we examined whether altered localization of PELP1 to the cytoplasmic compartment promotes PELP1 interaction with Src. Results from confocal analysis suggested that a substantial amount of PELP1 was localized in the nuclear compartment in PELP1-WT-expressing cells and that E2 treatment stimulated colocalization of PELP1 with Src at the membrane in some cells (Fig. 2A,

top panels). In contrast, in PELP1-cyto-expressing cells, E2 treatment promoted substantial colocalization of PELP1-cyto with Src both at the membrane and in the cytoplasm (Fig. 2A, bottom panels). Since PELP1 interaction with Src has been shown to promote Src activation, these findings raised the possibility that cytoplasmic localization of PELP1 might provide an opportunity for PELP1 to interact with Src, leading to enhanced Src kinase activation. To confirm that PELP1-cyto mutant indeed interact with Src kinase, immunoprecipitation was performed using total cellular lysates from pcDNA and PELP1-cyto mutant expressing cells. Results showed that PELP1-cyto interacts with Src kinase (Fig. 2B). To confirm that overexpression of PELP1-cyto activates Src kinase, we have used western blotting with phospho specific antibody (phospho-Src Tyr 416) which recognizes activated Src kinase. PELP1-cyto expressing cells showed increased activation of Src kinase compared to pcDNA expressing clones (Fig. 2C, *right panel*). These results suggest that PELP1 cytolocalization promotes its interactions with Src kinase.

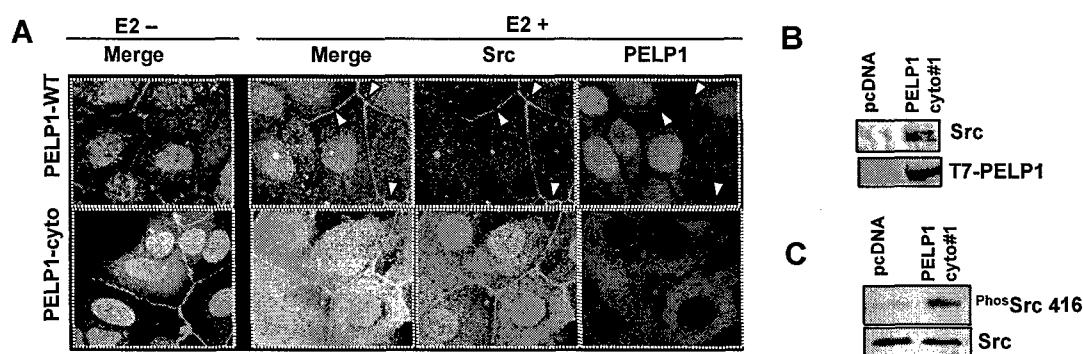


Fig. 2. PELP1 Cytoplasmic Retention Promotes PELP1-Src kinase Interaction and MAPK Activation. A, PELP1-WT and PELP1-cyto cells were treated with or without E2 (10^{-9} M) for 10 min and the colocalization of PELP1 (red) and Src (green) was analyzed by confocal microscopy. B, PELP1-cyto interacts and activates Src kinase.

Cytoplasmic Retention of PELP1 Promotes Activation of MAPK Pathway. We next

examined whether the PELP1 localization with Src observed in the PELP1-cyto clones led to functional activation of MAPK, a well-established downstream endpoint of ER nongenomic signalin. When these clones were cultured in DCC serum, PELP1-cyto-expressing clones exhibited increased basal MAPK activity compared to pcDNA-vector-overexpressing clones, and E2 treatment substantially enhanced MAPK activation in PELP1-cyto clones (Fig. 3A). PELP1-cyto expressing cells also showed increased MAPK activity upon tamoxifen treatment compared to pcDNA-expressing clones (Fig. 3B). To confirm that the increased MAPK activation seen in PELP1-cyto clones is due to PELP1-cyto interactions with Src kinase, we have used a specific Src kinase inhibitor PP2. Pretreatment of cells with Src kinase inhibitor PP2 reduced the basal as

well as E2 induced MAPK activation seen in the PELP1-cyto clones (Fig. 3C). These results suggest that PELP1 cytoplasmic localization enhances MAPK activation via its interactions with Src kinase. Since MAPK activation has been shown to phosphorylate ER on Ser-118, we next examined whether ER was phosphorylated on ER-Ser 118 in the PELP1-cyto clones. Western analysis of cells grown in 5% serum, revealed that PELP1 cyto expressing clones have increased ER-Ser118 phosphorylation compared to pcDNA expressing clones (Fig. 3D).

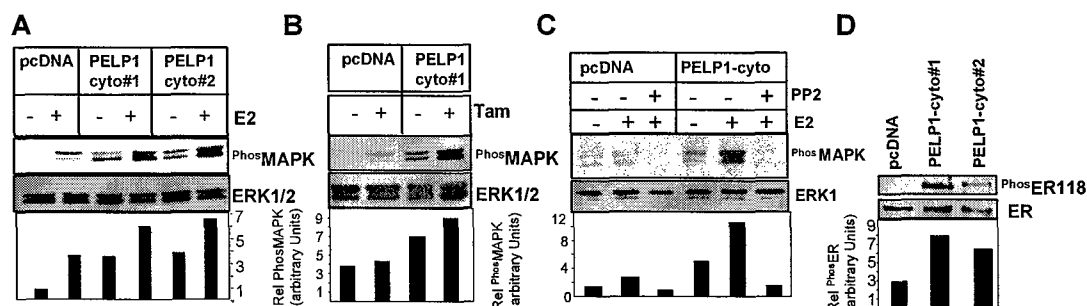


Fig 3. PELP1 Cytoplasmic Retention Promotes MAPK activation. A,B, pcDNA or PELP1-cyto were treated with E2 (1-9 M) or tamoxifen (10-8 M), and total lysates were subjected to Western blotting by using a phospho-MAPK antibody. C, pcDNA and PELP1-cyto clones were pretreated with Src kinase inhibitor PP2 for 30 min before the addition of E2 (10-9 M). Activation status of MAPK pathway was analyzed by Western blotting. D, MCF-7 clones expressing pcDNA and PELP1-cyto were cultured in 5% serum, and total lysates were subjected to Western blotting using phospho-ER Ser118.

Cytoplasmic Retention of PELP1 Promotes Constitutive Activation of AKT Pathway. Since activation of PI3K is another recognized nongenomic signaling of ER, we next examined the functionality of PI3K signaling in PELP1-cyto clones using phospho-AKT as a downstream marker of this pathway. Interestingly, PELP1-cyto clones showed excessive constitutive activation of the AKT pathway, which could not be further induced by E2 treatment (Fig. 4A). Because the AKT pathway has also been shown to promote tamoxifen resistance, we next examined the status of the AKT pathways in PELP1-cyto clones after treating them with tamoxifen. PELP1-cyto expressing cells showed increased AKT activity compared to pcDNA-expressing clones (Fig. 4B) and the status of already elevated levels of AKT in PELP1-cyto clones was not affected by tamoxifen treatment. Our finding that PELP1-cyto clones exhibit significant constitutive stimulation of the AKT pathway even in the absence of E2 treatment raised the possibility that PELP1 activation of the AKT pathway may be independent of PELP1's ability to interact with ER. To examine this possibility, we treated pcDNA and PELP1-cyto clones with ICI 182,780 for three days to downregulate ER expression and activity. The ICI 182,780 treatment did not affect the levels of constitutive AKT stimulation observed in the PELP1-cyto clones (Fig. 4C) suggesting the increased activation seen in the PELP-cyto clones is independent of ER. We then examined the possibility that increased AKT was due to activation of PI3K, an upstream activator of AKT. Treatment of PELP1-cyto clones with LY294002, a

widely used inhibitor of PI3K, blocked the constitutive activation of AKT (Fig. 4D). These findings suggested that cytoplasmic localization of PELP1 might contribute towards the observed enhanced AKT signaling by activating PI3K. We next examined whether PELP1 directly interacts with the p85 subunit of PI3K and whether the observed altered PELP1 localization facilitates constitutive interaction of PI3K. Immunoprecipitation of pcDNA, and PELP1-cyto lysates with p85 and T7-antibody showed that PELP1-cyto interacted with the p85 subunit of PI3K (Fig. 4E). Immunoprecipitation and *in vitro* PI-3 kinase assay also confirmed the noted constitutive activation of the PI3K pathway in the PELP1-cyto clones (Fig. 4F). Western analysis of PELP1WT clones also revealed mild activation of AKT however is significantly lower than the magnitude of AKT activation seen in PELP1-cyto clones (Fig. 4G). Since AKT activation has been shown to phosphorylate ER on Ser-167, we next examined whether ER was phosphorylated in the PELP1-cyto clones. Western blotting using phosphospecific antibody recognizing ER Ser-167 confirmed that, ER was constitutively phosphorylated on Ser-167 in PELP1-cyto clones (Fig. 4H). Collectively, these findings suggested that cytoplasmic localization of PELP1 promotes constitutive activation of the PI3K-AKT pathway which can phosphorylate ER.

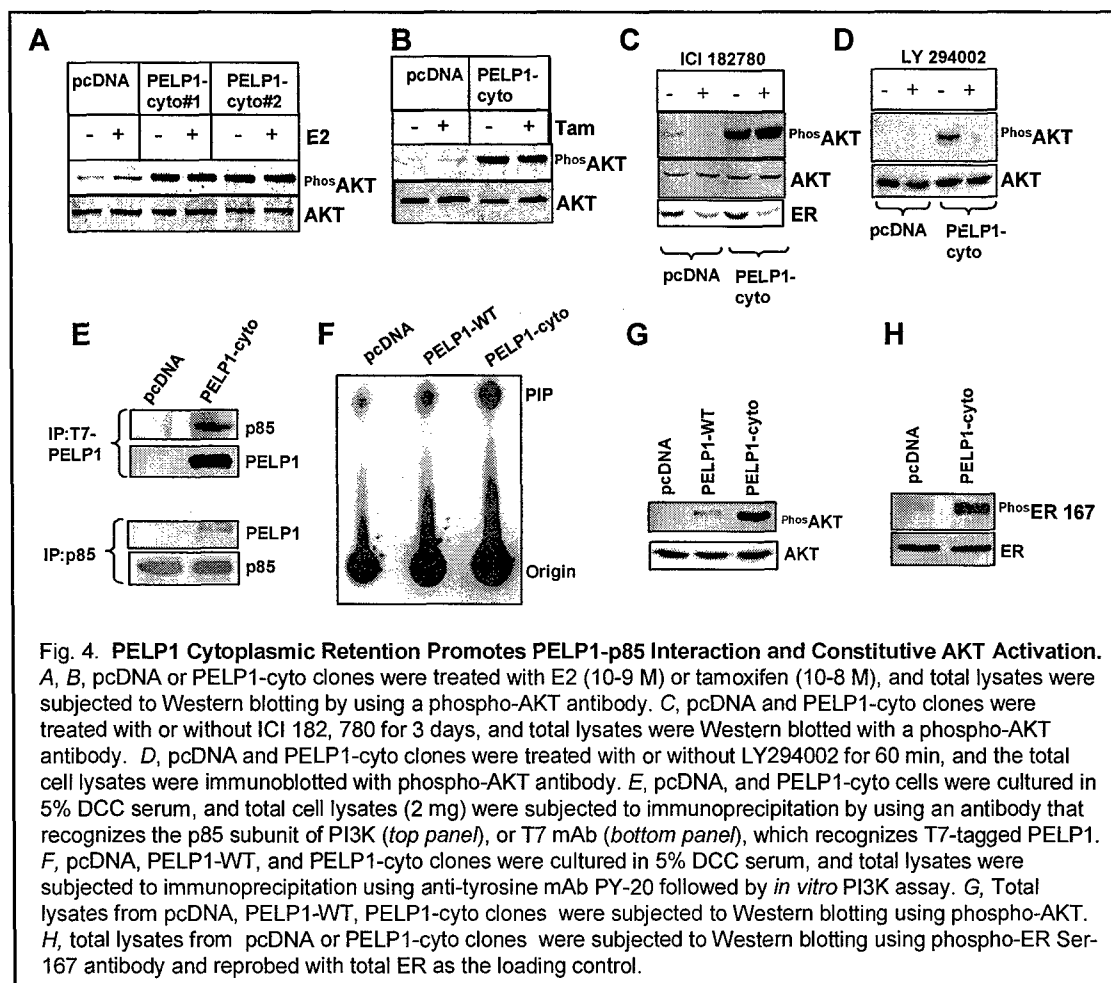
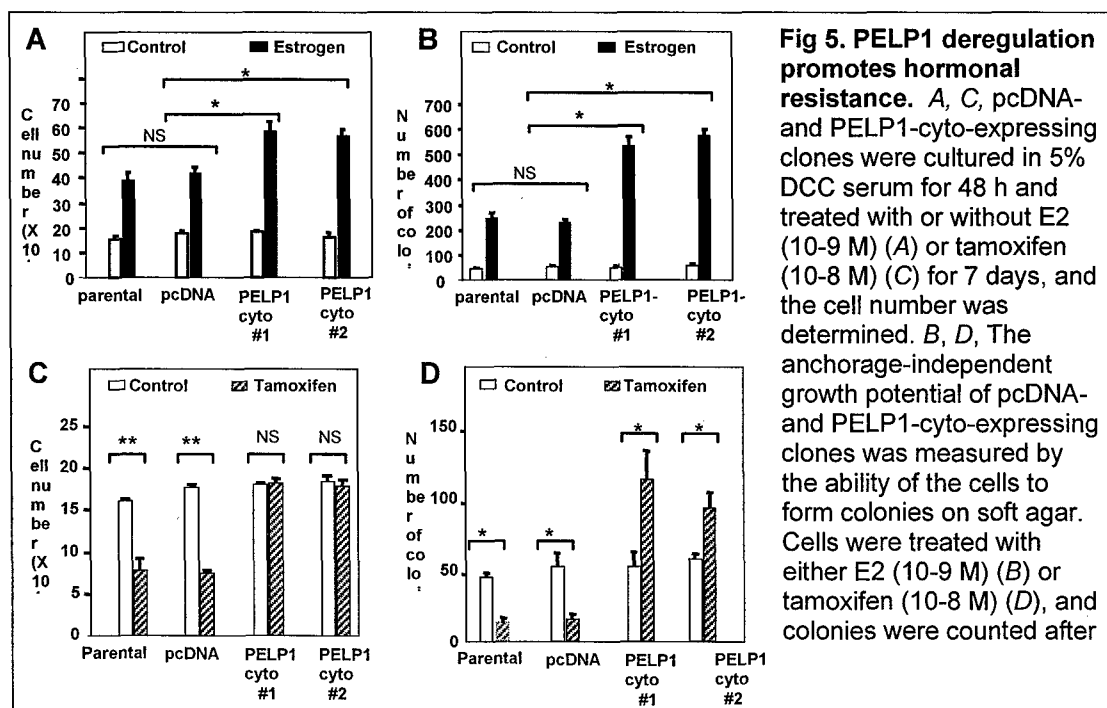


Fig. 4. PELP1 Cytoplasmic Retention Promotes PELP1-p85 Interaction and Constitutive AKT Activation. A, B, pcDNA or PELP1-cyto clones were treated with E2 (10-9 M) or tamoxifen (10-8 M), and total lysates were subjected to Western blotting by using a phospho-AKT antibody. C, pcDNA and PELP1-cyto clones were treated with or without ICI 182, 780 for 3 days, and total lysates were Western blotted with a phospho-AKT antibody. D, pcDNA and PELP1-cyto clones were treated with or without LY294002 for 60 min, and the total cell lysates were immunoblotted with phospho-AKT antibody. E, pcDNA, and PELP1-cyto cells were cultured in 5% DCC serum, and total cell lysates (2 mg) were subjected to immunoprecipitation by using an antibody that recognizes the p85 subunit of PI3K (top panel), or T7 mAb (bottom panel), which recognizes T7-tagged PELP1. F, pcDNA, PELP1-WT, and PELP1-cyto clones were cultured in 5% DCC serum, and total lysates were subjected to immunoprecipitation using anti-tyrosine mAb PY-20 followed by *in vitro* PI3K assay. G, Total lysates from pcDNA, PELP1-WT, PELP1-cyto clones were subjected to Western blotting using phospho-AKT. H, total lysates from pcDNA or PELP1-cyto clones were subjected to Western blotting using phospho-ER Ser-167 antibody and reprobated with total ER as the loading control.

PELP1 Cytoplasm Retention Promotes Hormonal Independence. To examine the effect of cytoplasmic localization of PELP1 on biological functions, we performed cell proliferation and anchorage independence assays. PELP1-cyto-expressing clones were hypersensitive to E2 treatment compared to parental MCF-7 or pcDNA-vector-expressing clones (Fig. 5A). Further, PELP1-cyto clones showed an increased ability to form colonies in an anchorage-independent manner compared to pcDNA-expressing cells (Fig. 5B). Since AKT activation has been shown to promote tamoxifen partial agonist action in breast cancer cells, we repeated these biological assays in the presence or absence of tamoxifen for 7 days. MCF-7 parental and pcDNA-expressing clones showed a 50% reduction in cell number. Interestingly, PELP1-cyto-expressing clones showed resistance to tamoxifen treatment, and the cell number was not altered (Fig. 5C). In anchorage independence assays, PELP1-cyto-expressing cells exhibited an increased ability to form colonies in the presence of tamoxifen compared to parental or pcDNA clones (Fig. 5D). These findings suggest a close relationship between PELP1 cytoplasmic localization promotes hormonal resistance possibly by excessively activating MAPK and AKT signaling pathways.



LIST OF PERSONEL

Neeta Sharma, SS# 093 72 8379, 100% effort

KEY RESEARCH ACCOMPLISHMENTS

- Altered localization of PELP1 promotes resistance to hormonal therapy
- Altered localization of PELP1 promotes excessive activation of MAPK and AKT
- PELP1 can directly interact and activate PI3K-AKT pathway

REPORTABLE OUTCOMES

This study has resulted in the following publications

Balasenthil S, Broaddus, R, Kumar R, Gustafsson, J-A, and Vadlamudi R,K. Potential involvement of PELP1/MNAR in tamoxifen resistance. **Breast Cancer Research and Treatment**. 82, supplement 1, abstract 27, 2003.

Vadlamudi, R.K., Manavathi, R., Balasenthil, S., Yang, Z., Sahin, A.¶, and Kumar, R. Altered Localization of PELP1 Promotes Estrogen Hypersensitivity, and Tamoxifen resistance. In review **Cancer Research**, 2005 (revised and resubmitted on 05/15/05).

CONCLUSIONS

Our findings suggest that cytoplasmic localization of Er coactivator PELP1 may play a role in the tamoxifen resistance. The ability of PELP1 to interact with and modulate the PI3K-AKT pathways may represent one potential mechanism by which PELP1-cyto cells might have developed resistance to hormonal treatment. Altered localization of the ER coactivator PELP1, which has the potential to stimulate nongenomic ER functions, may alter the ratio of genomic and nongenomic signaling in breast cancer cell and thus might promote hormonal independence by modulating ER-transactivating functions and SERM actions. Since PELP1 expression is deregulated in breast cancer, alteration in the localization of PELP1 is expected to contribute towards the excessive activation of the MAPK and PI3K-AKT pathways, leading to follow-up modifications of ER and other coregulators phosphorylation. Collectively, the findings from this study suggested that cytoplasmic localization of PELP1 promotes ligand-mediated enhanced MAPK activation and that constitutive activation of the PI3K-AKT pathway can phosphorylate ER. The ability of PELP1 to activate the MAPK and AKT signaling pathways, its potential to enhance tamoxifen resistance, and its distinct localization in a subset of breast tumors suggest that PELP1 plays a role in the biology of a subset of breast cancers and that its deregulation might contribute to hormonal independence.

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from the start of TAM treatment, was significantly prolonged in the trastuzumab-treated tumors ($M=146$ days) compared to gefitinib-treated tumors ($M=98$ days). Combining gefitinib and trastuzumab did not further increase tumor regression, but may further delay the onset of resistance, since median time to resistance has not yet been reached in this group ($M>153$ days). Interestingly, immunohistochemical analysis of phosphorylated (active) ERK1/2 showed that the high levels induced by E2 and maintained by TAM are markedly decreased by gefitinib but not by trastuzumab, even in regressing tumors in the TAM group. This observation suggests that trastuzumab's effect on tamoxifen sensitivity in this xenograft model does not occur via the ERK1/2 pathway. Together, our results suggest that gefitinib and trastuzumab modulate the growth of HER2-overexpressing breast tumors through distinct mechanisms and that combining such agents, in combination with ER-targeted therapy is a valid strategy to test in the clinic. ['Iressa' is a trademark of the AstraZeneca group of companies.]

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Molecular classification of tamoxifen-responsive and -resistant breast carcinomas by gene expression profiling.

Jansen M, Foekens J, Van Staveren I, Dirkwager-Kiel M, Look M, Meijer-Van Gelder M, Dorssers L, Klijn J, Berns E, JN1, Rotterdam, ZH, Netherlands

Background: Anti-estrogen resistance is one of the major challenges in the treatment of breast cancer. Approximately half of the patients with advanced disease does not respond to tamoxifen, whereas in those who respond the tumor ultimately becomes resistant. We aim to assess whether high-throughput gene expression profiling can lead to the discovery of a new set of markers for therapy resistance that may be predictive for the type of response to anti-estrogen treatment.

Material and Methods: The study was performed on a series of 70 ER-positive primary breast cancer samples from patients with advanced disease that showed the extreme types of response (objective response, $n=30$, versus progressive disease, $n=40$, from start of treatment). None of the patients received adjuvant systemic tamoxifen therapy. The main clinical endpoint of the study is the measurable effect of this endocrine therapy on tumor size and on time until tumor progression. RNA from these breast tumor samples and pool of control cell-lines was isolated and amplified and co-hybridized to cDNA micro-arrays, containing 18,500 transcripts in the human genome, produced by the NKL/AvL Micro-Array Facility (<http://microarrays.aki.nl>). After normalization, per sub grid, and a spot quality check, 8600 potentially relevant genes were selected.

Results: Using a training set of 50 tumors and two distinct bioinformatics tools (Spotfire and BRB, <http://linus.nci.nih.gov/BRB-ArrayTools.html>) and a leave-one out approach, a set of 143 overlapping genes was found to be differentially expressed ($P<0.01$) between tamoxifen-responsive and -resistant breast tumor samples. Using this gene-set, the cases clustered into two major cluster branches. This 143-gene-set predicted correctly the outcome of response to tamoxifen treatment in 82% of the cases (with a specificity of 84% and a sensitivity of 80%). This set includes gene associated with cell proliferation (e.g. CFP1), signal transduction (e.g. PTHR1), RNA and/or DNA binding activity (e.g. NSAP1 and TCFL1), extracellular matrix (e.g. Osteonectin) and protein serine-threonine kinases (e.g. PCK1). Using an add 4 more gene analysis, the classifier set could be optimized and was reduced to 40 genes which correctly predicted the outcome for 96% of the training-cases. This 40-gene predictive set was then used in an independent test set of 15 tamoxifen resistant tumors, and correctly predicted the outcome of treatment resistance in 80% of the cases validated. This predictive gene set is currently validated in a larger set of breast tumors.

Discussion: Our data suggest that gene expression profiling can be used to discriminate between breast cancer patients with an objective response to tamoxifen or patients with progressive disease. Further validation studies will prove whether this gene-set allows for the identification of the individual patient who could benefit or fail from (adjuvant) tamoxifen therapy.

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Potential involvement of PELP1/MNAR in tamoxifen resistance.

Balasenthil S, Broaddus R, Kumar R, Gustafsson J-A, Vadlamudi RK, UT MD Anderson Cancer Center, Houston, TX; Karolinska Institute, Huddinge, Sweden

Background: Selective estrogen receptor modulators (SERMs) including Tamoxifen are effectively used to retard the progression of hormone-dependent breast tumors to more invasive phenotypes. However such therapies have been closely linked with an acquired resistance in breast tumors and an increased incidence of endometrial cancer. The molecular basis of tamoxifen-resistance of breast cancer and -associated endometrial cancer remains elusive. In the present study, we tested the hypothesis that the recently discovered novel ER-regulatory protein proline glutamic acid leucine rich protein (PELP1, also referred to as MNAR, modulator of nongenomic activity of estrogen receptor), with functions in both nuclear and cytoplasmic compartments, plays any role in the action of tamoxifen.

Methods: To test the above hypothesis, we used ER-positive breast cancer cells with or without PELP1 deregulation, or sensitive or resistant to tamoxifen, and endometrial cancer cells. Sensitivity to estrogen and tamoxifen was determined by using reporter gene assays and protein, confocal and siRNA based biochemical and biological methods. The significance of these findings was established by PELP1 analysis in normal and cancerous endometrium tissues.

Results: Tamoxifen-sensitive and -resistant cells expressed similar levels of PELP1 but exhibited distinct subcellular localization in tamoxifen resistant cells. PELP1 deregulation led to a potentiation of agonist activity of tamoxifen in endometrial cancer cells. Expression of PELP1 mutant deficient in activation domain or PELP1 siRNA effectively blocked tamoxifen mediated agonist responses. Deregulation of PELP1 expression was observed in both breast and endometrial tumors. Immunohistological examination of PELP1 in tumors revealed its predominant localization in the cytoplasm as opposed to nuclear localization in the normal tissues.

Discussion: Our results suggest that altered localization of PELP1/MNAR, which could also activate nongenomic signaling, may be associated with tamoxifen resistance. Since PELP1 expression is deregulated in hormonal tumors and because PELP1 modulates SERM action, we will discuss results implying a role of PELP1/MNAR in hormone resistance. (This work was supported by Department of Army Grant BC023845)

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Drug metabolizing enzyme polymorphisms and outcome in node-positive breast cancer patients: differential impact of glutathione-S-transferase subtypes.

DeMichele A, Aplenc R, Botbyl J, Gor P, Foulkes A, Stadtmayer E, Rebbeck T, University of Pennsylvania, Philadelphia, PA

Background: The efficacy of adjuvant chemotherapy for breast cancer may depend, in part, on individual differences in drug metabolizing enzymes (DMEs), reflecting the balance of phase I (activating) DMEs and phase II (detoxifying) DMEs. Polymorphisms in genes encoding cytochrome p450 (CYP) enzymes may increase activation of some chemotherapeutics (CYP 3A4_*1A/*1A or CYP3A5_*1/*1; "favorable" phase I DME genotypes) while others may impair activation (CYP3A4_*1B or CYP 3A5_*3; "unfavorable" phase I DME genotypes). Meanwhile, improved outcome may be seen with homozygous null deletions of the glutathione-S-transferase (GST) genes μ and θ (GSTM1 and GSTT1) or point mutations in GST-p (GSTP1; "favorable" phase II DME genotypes) due to their inhibition of drug metabolism leading to a prolonged half-life of reactive oxygen species. We hypothesized that combinations of "unfavorable" phase I and phase II DME genotypes would impact recurrence of breast cancer among patients receiving adjuvant chemotherapy. **Material and Methods:** We tested this hypothesis by examining cyp3A4/3A5 and GSTM1, GSTT1 and GSTP1 polymorphisms in a cohort of 80 women with lymph-node positive breast cancer who received high-dose cyclophosphamide (1500mg/m²) and thiotepa (200mg/m²) in addition to standard anthracycline-based adjuvant chemotherapy. Genotypes were obtained from subject stem cells using PCR-based methods. We evaluated disease-free survival (DFS) by genotype (individually and in combination) using Kaplan-Meier and Cox proportional hazards models.

Functional Implications of Altered Subcellular Localization of PELP1 in Breast Cancer Cells*

Ratna K. Vadlamudi,^{1,3} Bramanandam Manavathi,¹ Seetharaman Balasenthil,¹ Sujit S. Nair,³ Zhibo Yang,¹ Aysegul A. Sahin,² and Rakesh Kumar¹

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Running Title: PELP1 in Cytoplasmic Signaling

Key Words: Nuclear Receptors, Coregulator, Cytoplasmic Signaling, Cancer

ABSTRACT

It is increasingly accepted that steroidal receptor coregulators may also function in the cytoplasmic compartment. PELP1 (Proline-, glutamic acid-, and leucine-rich protein-1) is a novel coregulator, which plays a role in both the genomic and extranuclear actions of estrogen receptors (ER) in hormonally responsive tissues. In this study using breast tumor arrays, we found PELP1 was localized only in the cytoplasm in 58% of the PELP1-positive breast tumors. To help explain the significance of the cytoplasmic localization of PELP1 in human breast tumors, we created a mutant protein that was expressed only in the cytoplasm (PELP1-cyto) and then generated a model system wherein MCF-7 breast cancer cells were engineered to specifically express this mutant. We found that PELP1-cyto cells were hypersensitive to estrogen but resistant to tamoxifen. PELP1-cyto cells, but not parental MCF-7 cells, formed xenograft tumors in nude mice. In addition, PELP1-cyto cells exhibited increased association of PELP1 with Src, enhanced mitogen-activated protein kinase (MAPK) activation, and constitutive activation of AKT. The altered localization of PELP1 was sufficient to trigger PELP1's interaction with the p85 subunit of phosphatidylinositol-3-kinase (PI3K), leading to PI3K activation. In addition, PELP1 interacted with epidermal growth factor receptors and participated in growth factor-mediated ER transactivation functions. Our results suggest that the altered localization of PELP1 modulates sensitivity to antiestrogens, potentiates tumorigenicity, presumably via the stimulation of extranuclear estrogen responses such as the activation of MAPK and AKT, and also enhance cross-regulation of ER transactivation activity by growth factors.

Introduction

The estrogen receptor (ER), a ligand-dependent transcription factor that modulates the transcription of a number of genes and contributes to genomic responses, has been implicated in the progression of breast cancer, and this is borne out by the finding that about 60-70% of breast tumors are ER-positive at presentation (1, 2). Although antiestrogens and selective estrogen receptor modulators (SERMs) are effective in curbing the progression of ER-positive breast tumors to more invasive phenotypes (3), many patients with metastatic breast tumors eventually become resistant to this treatment (4). Several mechanisms have been proposed to explain this resistance to hormonal therapy, including the expression of variant or mutant ER, the ligand-independent activation of ER, the adaptation of tumors to lower concentrations of estrogen, and pharmacological alterations (4, 5). Emerging data also suggest that ER coregulators play a role in hormonal responsiveness and tumor progression (6, 7). However, the causes of ER coregulator-linked resistance to hormonal therapy and ways to interfere with this phenomenon remain elusive.

The answer may be in the fact that, in addition to its well-studied nuclear functions, ER also participates in cytoplasmic and membrane-mediated signaling events (nongenomic signaling; 8, 9). Such nongenomic signaling has been linked to rapid responses to estrogen and generally involves the stimulation of the Src kinase, mitogen-activated protein kinase (MAPK), phosphatidylinositol-3-kinase (PI3K), and protein kinase C pathways in the cytosol (10, 11). Further, ER-activated nongenomic pathways have been shown to modify ER or its coactivators by phosphorylation, resulting in the altered topology of ER and its coregulator proteins and eventually leading to ligand-independent activation or differential responses to SERMs (9, 12).

Other factors implicated in the development and progression of breast cancer are, deregulated epidermal growth factor receptor (EGFR) signaling, and the constitutive activation of cytosolic pathways (MAPK; PI3K; AKT) (13). Now there is also emerging evidence that resistance to endocrine therapies may stem from complex interactions between ER and EGFR signaling components (14). The phosphorylation of ER and its associated coregulatory proteins has been suggested as one such mechanism by which growth factor signaling contributes to hormonal resistance (15). However, very little is known about the molecular mechanisms that lead to EGFR-ER crosstalk and the molecular adaptors that facilitate EGFR-ER signaling crosstalk.

PELP1 (proline-, glutamic acid-, and leucine-rich protein-1) / modulator of nongenomic activity of estrogen receptor (MNAR) is a novel ER coactivator (16) plays an important role in the genomic (17), nongenomic actions of ER (18) and promotes cell proliferation by sensitizing cells to G1>S progression (19). PELP1 expression is up regulated by estradiol (E2)-ER signaling and differentially modulated by SERMs (20). PELP1 is widely deregulated in hormone-responsive cancers including breast and endometrial cancers (16, 21). Although PELP1 is predominantly localized in the nucleus in hormonally responsive tissues (16, 17), recent studies have suggested that under certain conditions, PELP1/MNAR could be exclusively localized in the cytoplasm of cancer cells (21). However, the functional implications of the localization of PELP1 are unknown. Therefore in the present study, we investigated the localization and functional consequences of PELP1 in human breast tumor specimens. We found that, PELP1 was localized in the cytoplasm only in 58% of PELP1-positive tumors. To mimic this situation in order to gain further insight into the functional consequences of this event, we generated a novel MCF-7 model cell line to specifically express PELP1 in the cytoplasm. We found that the cytoplasmic localization of PELP1 made these cells, hypersensitive to estrogen but resistant to

SERMs and promoted tumorigenesis in nude mice. Our findings suggest that the altered localization of the ER coactivator PELP1 might lead to the excessive stimulation of cytosolic signaling pathways, thereby causing hormonal independence and increasing tumorigenic potential of breast cancer cells.

Materials and Methods

Cell Cultures and Reagents. MCF-7, HeLa and COS-1 cells were maintained in Dulbecco's modified Eagle's medium (DMEM)-F12 (1:1) supplemented with 10% fetal calf serum.

Antibodies against vinculin, actin and the steroid hormone 17 β -estradiol were purchased from Sigma Chemical Co. (St. Louis, MO). Antibodies against ER α and the phospho-tyrosine (pY20) were purchased from Lab Vision (Fremont, CA). Anti-T7-epitope antibody was purchased from Novagen (Milwaukee, WI). Antibodies against AKT, phospho-AKT, MAPK, phospho-MAPK, Phospho-Src Tyr 416 and phospho-ER-Ser 167, ER-Ser 118 were purchased from Cell Signaling (Beverly, MA). Antibody for Src and specific Src kinase inhibitor PP2 was purchased from Calbiochem (La Jolla, CA). Antibodies for EGFR were purchased from SantaCruz Biotechnology (SantaCruz, CA). Charcoal-stripped serum (DCC serum), ICI 182,780, and LY294002 were purchased from Sigma Chemical Co.

Human Samples and Tumor Array Staining. Tissue sample from patients who had undergone routine surgery for breast cancer were snap frozen in liquid nitrogen and stored at -80°C in the M.D. Anderson Breast Core Pathology Laboratory. Paraffin-embedded archival breast tissue specimen array was obtained from the pathology core at The University of Texas M. D. Anderson Cancer Center. The utilization of human tissue samples and array used in this study was approved by M.D. Anderson Cancer Center institutional human research committee. All

tissue classifications were verified by light microscopic examination of H&E stained slides by a breast pathologist (AAS). For immunohistochemical detection of PELP1 sections were deparaffinized with xylene and rehydrated using graded ethanol. The tumor array section was boiled for 10 minutes in 0.01 M citrate buffer and cooled for 30 minutes at room temperature to expose antigenic epitopes. The tumor array was blocked with 2% normal goat serum in 1% bovine serum albumin, PBS for 30 minutes. The tumor array section was incubated with polyclonal rabbit antihuman-PELP1 antiserum at a dilution of 1:500 and incubated overnight at room temperature. PELP1 antibody was developed in our laboratory, well characterized, and used in immunohistochemistry in earlier published studies (16, 21). The sections were washed three times with 0.05% Tween in PBS for 10 minutes, incubated with secondary antibody for 1 hour, and washed three times with 0.05% Tween in PBS for 10 minutes, developed with DAB-H₂O₂, and then counter-stained with Mayer's hematoxylin. Negative controls were performed by replacing the primary antibody with control rabbit IgG or peptide absorbed PELP1 antibody. The finding that no cells or less than 10% of cells immunoreactive was considered as a negative and the finding that more than 10% of the cells were immunoreactive was considered positive.

Generation of PELP1-cyto Cells. PELP1 cytoplasmic mutant (PELP1-cyto) lacking nuclear localization signal was generated using a Quick Change kit (Stratagene, La Jolla, CA) by mutating PELP1 aa 495KKLK498 to EELE using T7-tagged wild type PELP1 as a template. The sequence of the primer used for mutagenesis was 5'-cctagcgccccgaggagctagagctggatgtg-3'. MCF-7 cells stably expressing PELP1-cyto were generated by transfecting PELP1-cyto using FuGENE-6 transfection reagent (Roche Molecular Biochemicals, Indianapolis, IN). Stable clones were selected using G418 selection (1 mg/ml).

Tissue and Cell Extracts, Immunoblotting, Immunoprecipitation, Kinase Assays. Tissue and cell lysates were prepared using Triton X100 buffer (50 mM Tris-HCl, pH 7.5, 100 mM NaCl, 0.5% Triton X100, 1X protease inhibitor mixture, 1 mM sodium vanadate) for 15 minutes on ice. The lysates were centrifuged in an Eppendorf centrifuge at 4° C for 15 minutes. Cell lysates containing equal amounts of protein (~200 µg) were resolved on SDS-polyacrylamide gels (8% acrylamide), transferred to nitrocellulose membranes, probed with the appropriate antibodies, and developed using either the enhanced chemiluminescence method (ECL) or the alkaline phosphatase-based color reaction method. Immunoprecipitation was performed for 2 hours at 4°C using 1 µg of antibody per milligram of protein. For PI3K assay, cell lysates (1 mg) were immunoprecipitated with anti-phosphotyrosine mAb PY20 (Labvision) and subjected to *in vitro* kinase reaction in 50 µl of kinase buffer containing 0.2 mg/ml phosphatidylinositol (Sigma) and 20 µCi of [γ -³²P] ATP and 20 mM MgCl₂. The reaction products were separated on thin layer chromatography plates using chloroform: methanol: ammonium hydroxide: water (87:76:10:14) buffer and visualized by autoradiography.

Cell Proliferation, Soft-Agar, and Tumorigenesis Assays. For the cell proliferation assays, cells were grown in phenol red-free medium supplemented with 5% DCC serum for 48 hours, and then estrogen or tamoxifen was added. The proliferation rate of the cells was measured by counting them in a Beckman Coulter Counter as previously described (19). Soft-agar colony-growth assays were performed as previously described (22). Briefly, 1 ml of 0.6% DIFCO agar in DMEM supplemented with 5% DCC serum and insulin was layered onto tissue culture plates. Test cells (1×10^4) mixed with 1 ml of 0.36% bactoagar solution in DMEM were layered on top of the 0.6% bactoagar layer. The plates were incubated at 37°C in 5% CO₂ for 21 days. For tumorigenesis studies, 5×10^7 cells were implanted subcutaneously into the mammary fat pads of

8 nude mice as previously described (23) and allowed to grow for 8 weeks. Tumor size was then measured.

Reporter Gene Assay. For reporter gene transient transfections, COS-1 cells were cultured for 48 hours in minimal essential medium without phenol red-containing 5% DCC serum. ERE-luciferase reporter constructs were cotransfected with ER along with or without pcDNA vector, PELP1-WT or PELP1-cyto expression plasmids using FuGENE-6 according to the manufacturer's instructions. Twenty-four hours later, cells were treated with EGF for 12 h. Cells were then lysed with passive lysis buffer, and the luciferase assay was performed using a luciferase reporter assay kit (Promega). The total amount of DNA used in the transfections was kept constant by adding a parental vector. Each transfection was carried out in six-well plates in triplicate wells.

Immunofluorescence and Confocal Microscopy. The cellular localization of PELP1 was determined using indirect immunofluorescence as described previously (17).

Statistical Analysis. Statistical analysis was done using Student's *t* test, and values with $p < 0.05$ were considered statistically significant.

Results

PELP1 Localization is altered in a Subset of Breast Tumors. To examine the localization of PELP1 in human breast tumors, we stained 62 breast tumor specimens using a well-characterized PELP1 antibody (16, 21; Fig. 1A). The staining intensity was scored in both nuclear and cytoplasmic compartments. Normal breast epithelium ($n=8$) was used as a control. All normal breast epithelium stained positively; PELP1 staining was strong in the nucleus and modest in the cytoplasm. In tumors, however, PELP1 staining intensity and localization was

variable. PELP1 staining was observed in 38 (61%) of the 62 analyzed tumors. In 22 (58%) of the 38 PELP1-positive tumors, PELP1 was localized in the cytoplasm only; in one tumor (3%), intense nuclear staining was observed, and in 15 tumors (39%), both nuclear and cytoplasmic staining were observed (Table 1). We then correlated the expression and localization of PELP1 with the clinical ER status of these tumors. ER staining was present in 19 (50%) of 38 PELP1-positive tumors. ER expression was observed in the one tumor that showed nuclear PELP1 expression, in 8 of 22 (36%) cytoplasmic PELP1-expressing tumors, and in 10 of 15 (66.6%) nuclear and cytoplasmic PELP1-expressing tumors. These results suggest that in a subset of breast tumors, PELP1 is localized in the cytoplasm only and that a substantial number of breast tumors coexpress of PELP1 and ER.

Generation of PELP1-cyto Model Cells. To examine the functional consequences of the cytoplasmic localization of PELP1, we next generated a mutant of PELP1 (PELP1-cyto) in which the nuclear localization signal was mutated from KKLK to EELE (Fig. 1B). Preliminary analysis using confocal microscopy showed that PELP1-cyto was predominantly localized in the cytosol. Using this construct, we generated MCF-7 cells that stably expressed PELP1-cyto. To avoid potential clonal variations, we developed two pooled clones. Western blotting showed expression of T7-tagged PELP1-cyto in these clones (Fig. 1C). We used a previously characterized wild-type PELP1 (PELP1-WT) pooled clone as the control (19). Confocal scanning analysis showed that PELP1-cyto was localized to the cytoplasm while PELP1-WT was localized to the nucleus, as expected (Fig. 1D). To examine whether the cytoplasmic PELP1 observed in the tumors corresponded to the full-length PELP1, we performed Western blot analysis of the lysates of the tumors that expressed PELP1 in the cytoplasm only. Lysates from MCF-7 cells expressing both PELP1-WT and PELP1-cyto were used as controls. Results showed

that PELP1 isolated from lysates of the tumors that exclusively expressed PELP1 in the cytoplasm migrated in a similar fashion as PELP1-WT and PELP1-cyto that were expressed in MCF-7 cells (Fig. 1E). These findings suggest that PELP1 expressed in the cytoplasm of tumors represents the full-length PELP1.

Cytoplasmic Retention Enhances PELP1 Interactions with Src Kinase. Recent studies have shown that in addition to its suggested nuclear functions (17), PELP1 participates in nongenomic signaling activities, such as activation of Src (18, 24). Therefore, we examined whether the localization of PELP1 to the cytoplasmic compartment, a physiologic situation existing in a subset of breast tumors, promotes PELP1 interaction with Src. Results from confocal microscopy showed that a substantial amount of PELP1 was localized in the nuclear compartment of PELP1-WT-expressing cells and that E2 treatment stimulated colocalization of PELP1 with Src at the membrane in some cells (Fig. 2A, *top panels*). In contrast, in PELP1-cyto-expressing cells, E2 treatment promoted substantial colocalization of PELP1-cyto with Src both at the membrane and in the cytoplasm (Fig. 2A, *bottom panels*). Another study found that PELP1 interaction with Src promotes Src activation (24); this raised the possibility that the cytoplasmic localization of PELP1 might provide an opportunity for PELP1 to interact with Src, thereby leading to enhanced Src kinase activation. To test whether that PELP1-cyto interacts with Src kinase, immunoprecipitation was performed by using total cellular lysates from pcDNA- and PELP1-cyto-expressing cells. Results showed that PELP1-cyto interacts with Src kinase (Fig. 2B, *left panel*). To confirm that overexpression of PELP1-cyto activates Src kinase, we performed Western blotting with a phospho-specific antibody (phospho-Src Tyr 416) that recognizes activated Src kinase. This showed that PELP1-cyto-expressing cells increased the activation of Src kinase compared with pcDNA-expressing clones (Fig. 2B, *right panel*). These

findings suggest that PELP1 cytolocalization promotes PELP1 interaction with Src kinase. Since PELP1-WT functions as a scaffolding protein for the coupling of ER with Src kinase, we then investigated whether PELP1-cyto also interacts with ER. The immunoprecipitation of PELP1-cyto showed that PELP1-cyto could interact with ER (Fig. 2C).

Cytoplasmic Retention of PELP1 Promotes Activation of the MAPK Pathway. Next we examined whether the interaction we observed between PELP1 and Src in the PELP1-cyto expressing clones leads to the functional activation of MAPK, a well-established downstream endpoint of ER nongenomic signaling (10, 11). Western blot analysis of cells grown in 5% serum revealed that PELP1-WT and PELP1-cyto-expressing cells showed increased activation of the MAPK pathway compared with pcDNA-expressing clones. However, the PELP1-cyto-expressing clones exhibited significantly more MAPK activation than did the PELP1-WT clones (Fig. 2D). When these clones were cultured in DCC serum, the PELP1-cyto-expressing clones exhibited increased basal MAPK activity compared with pcDNA-vector-overexpressing clones, and E2 treatment substantially enhanced MAPK activation in PELP1-cyto clones (Fig. 2E). PELP1-cyto-expressing cells also showed greater MAPK activity in response to tamoxifen treatment than did pcDNA-expressing clones (Fig. 2F). To confirm that the increased MAPK activation seen in PELP1-cyto clones was due to PELP1-cyto interactions with Src kinase, we pretreated cells with PP2, a specific Src kinase inhibitor. PP2 reduced the basal as well as E2-induced MAPK activation seen in the PELP1-cyto clones (Fig. 2G). These results suggests that PELP1 cytoplasmic localization enhances MAPK activation via its interactions with Src kinase. Since MAPK activation has been shown to phosphorylate ER on Ser-118, we then examined whether ER was phosphorylated on ER-Ser 118 in the PELP1-cyto clones. Western blot analysis

of cells grown in 5% serum revealed that PELP1-cyto-expressing clones showed increased ER-Ser118 phosphorylation compared with pcDNA-expressing clones (Fig. 2H)

Cytoplasmic Retention of PELP1 Promotes Constitutive Activation of the AKT

Pathway. Since the activation of PI3K is another recognized nongenomic signaling activator of ER (13), we examined the functionality of PI3K signaling in PELP1-cyto clones by using phospho-AKT as a downstream marker of this pathway. Interestingly, PELP1-cyto clones showed excessive constitutive activation of the AKT pathway that could not be further induced by E2 treatment (Fig. 3A). Because the AKT pathway has also been shown to promote tamoxifen resistance (25), we examined the status of the AKT pathway in PELP1-cyto clones after treating the clones with tamoxifen. PELP1-cyto-expressing cells showed increased AKT activity compared with pcDNA-expressing clones (Fig. 3B), and the status of already elevated levels of AKT in PELP1-cyto clones was not affected by tamoxifen. Our finding that PELP1-cyto clones exhibit significant constitutive stimulation of the AKT pathway even in the absence of E2 treatment raised the possibility that PELP1 activation of the AKT pathway might be independent of PELP1's ability to interact with ER. To examine this possibility, we treated pcDNA and PELP1-cyto clones with the antiestrogen ICI 182, 780 for 3 days to down regulate ER expression and activity. The ICI 182, 780 did not affect the levels of constitutive AKT stimulation observed in the PELP1-cyto clones (Fig. 3C), suggesting that the increased activation seen in the PELP1-cyto clones is independent of ER. We then examined the possibility that increased AKT was due to the activation of PI3K, an upstream activator of AKT. Treatment of PELP1-cyto clones with LY294002, a widely used inhibitor of PI3K, blocked the constitutive activation of AKT (Fig. 3D). These findings suggested that the cytoplasmic localization of PELP1 might activate PI3K, thereby contributing to enhanced AKT signaling.

PELP1 Interacts with the P85 Subunit of PI3K. Analysis of the motif scan

(<http://scansite.mit.edu>) of the PELP1 amino acid sequence revealed that PELP1 contains two potential p85-SH3 model 1 sites (HPPNRSAPHLPLGLMC; PSPFRAPPFHPPGPM) and one potential p85-SH3 model 2 site (QTGKPSAPKKLKLDV). Therefore, we examined whether PELP1 directly interacts with the p85 subunit of PI3K and whether the observed altered PELP1 localization facilitates constitutive interaction of PI3K. Immunoprecipitation of pcDNA and PELP1-cyto lysates with antibodies p85 and T7 showed that PELP1-cyto interacts with the p85 subunit of PI3K (Fig. 3E). Immunoprecipitation and *in vitro* PI3K assays also confirmed the constitutive activation of the PI3K pathway in the PELP1-cyto clones (Fig. 3F). Western blot analysis of PELP1-WT clones also revealed mild activation of AKT; however the activation was significantly lower than the magnitude of AKT activation seen in PELP1-cyto clones (Fig. 3G). Since AKT activation has been shown to phosphorylate ER on Ser-167, we examined whether ER was phosphorylated in the PELP1-cyto clones. Western blotting using a phospho-specific antibody that recognizes ER Ser-167 confirmed that ER was constitutively phosphorylated on Ser-167 in PELP1-cyto clones (Fig. 3H). Collectively, these findings suggested that cytoplasmic localization of PELP1 promotes constitutive activation of the PI3K-AKT pathway, which can phosphorylate ER.

PELP1 associates with Epidermal Growth Factor Receptor. Previous studies have suggested bidirectional signaling between ER and EGFR pathways and growth factors promotes ER phosphorylation through the PI3K-AKT pathway (13). Since we observed that PELP1 interacts with PI3K, whose activity is modulated by growth factor signaling, we examined whether PELP1 interacts with growth factor receptors as well. To determine whether there is physiological evidence of endogenous PELP1-EGFR interactions, we initially examined this in

HeLa cells, which abundantly express EGFR and PELP1. HeLa cells were treated with EGF, and the colocalization of PELP1 and EGFR was analyzed by confocal microscopy. Our results showed that EGF stimulation caused a small portion of PELP1 to associate with membranous regions, and in some areas, the colocalization of PELP1 with EGFR was observed (Fig. 4A, left panel). We next examined whether PELP1-cyto also interacts with EGFR. Confocal microscopy examination of EGF treated PELP1-cyto clones revealed colocalization of PELP1-cyto with EGFR (Fig. 4A, right panel). To confirm these results, we performed immunoprecipitation experiments in COS-1 cells, which were treated with or without EGF. Immunoprecipitation of EGFR followed by Western blot analysis with PELP1 confirmed that PELP1 indeed associates with EGFR upon EGF stimulation (Fig. 4B).

Since EGF has been shown to promote ligand-independent activation of ER transactivation functions (26, 27), we examined whether PELP1 promotes EGF-mediated ER transactivation functions by performing estrogen response element (ERE) reporter gene assays using HeLa cells cotransfected with ER along with or without pcDNA vector, PELP1-WT or PELP1-cyto constructs. Results showed that pcDNA transfected clones exhibited 2.3 fold activation of ERE reporter upon addition of EGF. PELP1 WT transfection modestly enhanced the basal activity of the ERE reporter, however EGF treatment induced a 3.2 fold induction of ERE reporter gene activity over untreated control. Interestingly, PELP1-cyto transfection substantially increased the ERE reporter gene activity in the absence of ligand to a level similar to EGF treated, PELP1-WT transfected cells and EGF treatment further increased ERE reporter activity in PELP1-cyto transfected cells (Fig. 4C). Increased basal activity of the ERE reporter in PELP1-cyto transfected cells could be due to the constitutive activation of AKT pathway seen in the PELP1-cyto cells. These results suggest that PELP1-WT has a potential to enhance EGF mediated ER

transactivation functions and mislocalization of PELP1 in the cytoplasm can potentially activate ER basal activation by excessively coupling EGFR-PI3K signaling pathways.

Growth factor signaling has been shown to modulate coactivator functions by post-translation modifications (14). Since PELP1 interacted with EGFR and promoted EGF-mediated ERE transactivation functions, we next examined whether EGF stimulation promotes the phosphorylation of PELP1. COS-1 cells were transfected with T7-tagged PELP1, and after 48 hours, cells were treated with or without EGF. Total lysates were immunoprecipitated with T7-epitope antibody (Fig. 4D, *left panel*), and immunoprecipitates were blotted with T7 and phosphotyrosine antibodies. Results showed that EGF promotes tyrosine phosphorylation of PELP1. Similarly, immunoprecipitation of PELP1-cyto from PELP1 stable clones also showed that EGF promotes tyrosine phosphorylation of PELP1-cyto (Fig. 4D, *right panel*).

Since PELP1 interacts with ER and promotes E2-mediated nongenomic signaling pathways, we examined whether E2 can promote the tyrosine phosphorylation of PELP1. MCF-7 cells stably expressing PELP1-WT were treated with or without E2. Immunoprecipitation experiments showed that E2 also promoted the tyrosine phosphorylation of PELP1 (Fig. 4E). We confirmed these results with confocal microscopic analysis. As shown by phosphotyrosine antibody staining, E2 promoted the membrane localization of PELP1 and its colocalization with phosphotyrosine (Fig. 4F). These results suggest that both EGF and E2 signaling modulate PELP1 functions and that the cytoplasmic retention of PELP1 excessively promotes the interactions of PELP1 with components of the growth factor signaling axis.

PELP1 Cytoplasm Retention Promotes Hormonal Independence. To examine the effect of the cytoplasmic localization of PELP1 on biological functions, we performed cell

proliferation and anchorage independence assays. Results showed that PELP1-cyto-expressing clones were hypersensitive to E2 compared with parental MCF-7- or pcDNA-vector-expressing clones (Fig. 5A). Further, PELP1-cyto clones showed an increased ability to form colonies in an anchorage-independent manner compared with pcDNA-expressing cells (Fig. 5B). Since AKT activation has been shown to promote tamoxifen partial agonist action in breast cancer cells, we repeated these biological assays in the presence or absence of tamoxifen for 7 days. MCF-7 parental and pcDNA-expressing clones showed a 50% reduction in cell number. Interestingly, PELP1-cyto-expressing clones showed resistance to tamoxifen, in that the cell number was not altered (Fig. 5C). In anchorage-independence assays, PELP1-cyto-expressing cells exhibited an increased ability to form colonies in the presence of tamoxifen compared with parental or pcDNA clones (Fig. 5D). Consistent with these results, PELP1-cyto clones exhibited tumorigenic potential in nude mice, and these tumors also exhibited constitutive activation of the AKT pathway (Fig. 5E). These findings suggest that a close relationship exists between PELP1 cytoplasmic localization, AKT activation, and tumorigenesis.

Discussion

In this study, we observed that the localization of the ER coactivator PELP1 is altered in a subset of breast tumors. Specifically, model cells that mimic PELP1 localization in tumors, we observed that the cytoplasmic localization of PELP1 promotes its interactions with EGFR, Src, and PI3K, thereby leading to the activation of the MAPK and AKT pathways. In addition, the altered localization of PELP1 promoted hypersensitivity to estrogen and anchorage independence. Our results thus indicate that the ER coactivator PELP1 may contribute to

tumorigenesis in breast tumors by serving as a molecular adaptor between EGFR-ER pathways and by promoting excessive crosstalk between these pathways.

The altered localization of the ER coactivator PELP1, which has the potential to stimulate nongenomic ER functions (18, 24), may alter the ratio of genomic and nongenomic signaling in breast cancer cells and thus might promote hormonal independence by modulating ER-transactivating functions and SERM actions. As our study and other studies indicate, PELP1 expression is deregulated in breast and endometrial cancers (16, 21); therefore, the altered localization of PELP1 is expected to contribute toward the excessive activation of the MAPK and PI3K-AKT pathways, leading to follow-up modifications of ER. Such modifications of the ER pathway may lead to the activation of ER target genes in a ligand-independent manner. Alternatively, the modified ER may allow the recruitment of a different set of coregulator proteins, and thus may exhibit resistance to hormonal therapy. In our study, the cytoplasmic localization of PELP1 promoted increased activation of MAPK, leading to enhanced ER-Ser118 phosphorylation. Similarly, these clones also exhibited constitutive activation of the AKT pathway, which was not further induced by E2. Therefore, the enhanced E2-mediated growth and anchorage independence seen in these clones is likely to be caused by excessive activation of the MAPK pathway, while tamoxifen-mediated resistance could be due to the cumulative effect of the activation of the MAPK and AKT pathways, which leads to the modification of ER and its associated proteins as well as the activation of downstream pathways by activated MAPK and AKT.

ER coregulatory proteins have been suggested to play a role in the generally observed tissue-specific effects of tamoxifen (29, 30). However, these ER coregulators are targeted by excessive ER-HER2 crosstalk leading to hormonal resistance in a subset of breast tumors (15).

High levels of the ER coactivator AIB1 and HER2 in breast cancer contribute to tamoxifen resistance (31). Our findings that PELP1 interacts with EGFR, and that PELP1 has the ability to enhance EGF-mediated ER transactivation functions suggest that PELP1 potentiates growth factor receptor-mediated hormonal resistance in a subset of tumors that have deregulated growth factor signaling.

Growth factor signaling promotes ER phosphorylation (both Ser and tyrosine; 14, 32). Recent evidence suggests that ER coactivators; in addition to ER, are targets of growth factor signaling (33). Growth factor-mediated activation of nongenomic pathways and phosphorylation of ER and ER-coregulatory proteins have been shown to have a role in tamoxifen resistance (28, 34). Our findings that PELP1 enhances tamoxifen resistance and that EGF promotes the phosphorylation of PELP1 suggest that possibility that growth factor-mediated posttranslational modification of PELP1 may play an important role in PELP1-mediated hormonal resistance functions.

Activation of the PI3K-AKT pathway has been shown to be an essential step in the estrogenic action of growth factors (35). Previous studies also demonstrated that forced expression of constitutively active AKT in MCF-7 cells promotes E2-independent growth as well as tamoxifen response (25). A recent study found that overexpression of the ER coactivator AIB1 promoted high tumor incidence, which is associated with the activation of the PI3K-AKT pathway (36). Our findings suggest that cytoplasmic localization of PELP1 plays a role in tamoxifen resistance and that the ability of PELP1 to modulate the PI3K-AKT pathways may represent one mechanism by which PELP1-cyto cells develop resistance to hormonal treatment. Further, our analysis of human breast tissues provided the proof-of-principle that PELP1 localization is altered in a subset of human breast tumors.

Our results suggest that the cytoplasmic localization of PELP1 may play a role in the constitutive activation of AKT in tumor cells. Our findings also suggest the existence of a close relationship between cytoplasmic PELP1 localization and increased nongenomic signaling. Additionally, our findings suggest that the cytoplasmic localization of PELP1 might be sufficient to promote tumorigenic phenotypes and hormonal independence. Further, the ability of PELP1-cyto cells to form tumors in nude mice and the presence of increased AKT signaling in these tumors also supports the potential role of PELP1 localization in tumorigenesis. Since these tumors were formed in the absence of estrogen pellet implantation, PELP1-mediated constitutive activation of the AKT pathway rather than the MAPK pathway might play a role in PELP1-cyto-mediated *in vivo* tumorigenic functions. It will be interesting to discover the mechanisms by which PELP1 localization is altered in pathologic conditions. Since cytoplasmic PELP1 observed in the tumors migrated in a similar fashion as PELP-WT, it is likely that it represents a full-length protein and that its deregulation in localization could be because of posttranslational modifications or mutations that affect its localization to the nuclear compartment. Our future studies will be directed at identifying the mechanisms of PELP1 deregulation, including possible posttranslational modifications / mutations in tumors.

In summary, our results suggest that the localization of ER coactivators such as PELP1 could activate nongenomic signals and may play an important role in the hormonal responses of ER-positive cancer cells (Fig. 5F). Furthermore, the ability of PELP1 to activate MAPK and AKT signaling pathways, its potential to enhance tamoxifen resistance, and its distinct localization in a subset of breast tumors suggest that PELP1 plays a role in the biology of a subset of breast cancers and that its deregulation might contribute to hormonal independence.

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FIGURE LEGENDS

Figure 1. Localization of PELP1 in human breast tissues and breast cancer model cells. *A*, A tissue array containing samples of 62 breast tumors was stained with a PELP1 antibody. Three representative PELP1 localizations (nuclear, cytoplasmic, and nuclear and cytoplasmic) observed in breast tumors are shown. Magnification, X200. Normal breast tissue was used as a control. *B*, Schematic representation of PELP1-cyto construct. *C*, Total cellular lysates were obtained from MCF-7 cells or MCF-7 clones stably expressing pcDNA, PELP1-WT, and PELP1-cyto. Expression of T7-tagged PELP1-cyto was analyzed by Western blotting using anti-T7-epitope monoclonal antibody (T7-mAb). Vinculin was used as a loading control. *D*, MCF-7 cells expressing PELP1-WT or PELP1-cyto were cultured in DCC medium and fixed in methanol. The localization of T7-tagged PELP1 in these clones was analyzed by confocal microscopy using T7 mAb. *E*, Total lysates obtained from breast tumors or MCF-7 clones that stably express PELP1-WT and PELP1-cyto were analyzed by Western blotting using PELP1 antibody (*top*) and subsequently reprobed with an actin antibody, which served as a loading control (*bottom*).

Figure 2. PELP1 cytoplasmic retention promotes PELP1-Src kinase interaction and MAPK activation. *A*, MCF-7 cells stably expressing T7-tagged PELP1-WT and PELP1-cyto were treated with or without E2 (10^{-9} M) for 10 min. Cells were fixed, and the colocalization of PELP1 (red) and Src (green) was analyzed by confocal microscopy. Colocalization of PELP1 and Src is indicated by yellow. *B*, pcDNA- and PELP1-cyto clones were cultured in 5% DCC serum. Total cellular lysates (2 mg) were subjected to immunoprecipitation using T7 antibody followed by Western blot analysis with Src antibody (*left panel*). Total lysate (100 μ g) was analyzed by Western blotting using phospho-Src Tyr 416 antibody (*right panel*). *C*, Total

cellular lysate (2 mg) from pcDNA or PELP1-cyto clones was subjected to immunoprecipitation by using T7 antibody followed by Western blot analysis with ER α antibody. *D*, MCF-7 clones expressing pcDNA, PELP1-WT, PELP1-cyto were cultured in 5% serum, and total lysates were subjected to Western blotting by using phospho-MAPK antibody. Blots were stripped and reprobed with total MAPK as the loading control. *E*, *F*, MCF-7 clones expressing pcDNA or PELP1-cyto were treated with E2 (10^{-9} M) or tamoxifen (10^{-8} M), and total lysates were subjected to Western blotting by using a phospho-MAPK antibody. Blots were stripped and reprobed with total MAPK as the loading control. *G*, pcDNA and PELP1-cyto clones were pretreated with Src kinase inhibitor PP2 30 minutes before the addition of E2 (10^{-9} M). Activation status of MAPK pathway was analyzed by Western blotting. *H*, MCF-7 clones expressing pcDNA and PELP1-cyto were cultured in 5% serum, and total lysates were subjected to Western blotting using phospho-ER Ser118 and subsequently reprobed with total ER.

Figure 3. PELP1 cytoplasmic retention promotes PELP1-p85 interaction and constitutive AKT activation. *A*, *B*, MCF-7 clones expressing pcDNA or PELP1-cyto were treated with E2 (10^{-9} M) or tamoxifen (10^{-8} M), and total lysates were subjected to Western blotting by using a phospho-AKT antibody. Blots were stripped and reprobed with total AKT as a loading control. *C*, pcDNA and PELP1-cyto clones were treated with or without ICI 182, 780 for 3 days, and total lysates were Western blotted with a phospho-AKT antibody. Blot was reprobed with total AKT and ER antibody. *D*, pcDNA and PELP1-cyto clones were treated with or without LY294002 for 60 min, and the total cell lysates were immunoblotted with phospho-AKT antibody. *E*, pcDNA, and PELP1-cyto cells were cultured in 5% DCC serum, and total cell lysates (2 mg) were subjected to immunoprecipitation by using an antibody that recognizes the p85 subunit of PI3K

(*top panel*), or T7 mAb (*bottom panel*), which recognizes T7-tagged PELP1. *F*, pcDNA, PELP1-WT, and PELP1-cyto clones were cultured in 5% DCC serum, and total lysates were subjected to immunoprecipitation by using anti-tyrosine mAb PY-20 followed by *in vitro* PI3K assay. *G*, MCF-7 clones expressing pcDNA, PELP1-WT, PELP1-cyto were cultured in 5% DCC serum, and total lysates were subjected to Western blotting using phospho-AKT followed by total AKT as the loading control. *H*, pcDNA or PELP1-cyto clones were cultured in 5% DCC serum and total lysates were subjected to Western blotting using phospho-ER Ser-167 antibody and reprobed with total ER as the loading control.

Figure 4. PELP1 interacts with EGFR and promotes EGF-mediated ligand-independent ER transactivation functions. *A*, HeLa cells were treated with EGF (50 ng/ml). The cells were then fixed and costained with antibodies against PELP1 (green) and EGFR (red). The images were analyzed by confocal microscopy. Arrows point to the area of colocalization (yellow) of PELP1 and EGFR (left panel). PELP1-cyto cells were treated with EGF (50 ng/ml). Cells were stained with antibodies against T7-tag (to detect T7-tagged PELP1-cyto, green color) and EGFR (red). (right panel). A small portion of the image showing the colocalization (yellow) was enlarged and shown in the side panels *B*, COS-1 cells were serum starved for 24 h, and treated with or without EGF. EGFR was immunoprecipitated by using an EGFR antibody; and the presence of PELP1 in the immunoprecipitates was analyzed by Western blotting by using an anti-PELP1 antibody. *C*, HeLa cells were cotransfected with ER and ERE-luc reporter gene along with pcDNA or PELP1-WT or PELP1-cyto. After 48 h, cells were stimulated with EGF for 12 h and ERE-luc reporter activity was measured. *D*, COS-1 cells were transfected with T7-tagged PELP1-WT and after 72 h, treated with or without EGF. Total lysates were immunoprecipitated with T7-epitope specific

antibody and blotted with phosphotyrosine and T7 antibodies (*left panel*). MCF-7 cells that stably express T7-tagged PELP1-cyto were treated with or without EGF, and T7-PELP1-cyto was immunoprecipitated, followed by Western blotting with the phosphotyrosine antibody (*right panel*). *E*, MCF-7 cells that stably express T7-tagged PELP1-WT were treated with or without E2 for indicated periods of time and T7-PELP1 was immunoprecipitated; this was followed by Western blotting with the phosphotyrosine antibody. *F*, MCF-7 cells that stably express T7-tagged PELP1-WT were treated with or without E2, and colocalization of PELP1 (green) and phosphotyrosine (red) was analyzed by confocal microscopy. Arrow point to the areas of colocalization of PELP1 and phosphotyrosine at the membrane.

Figure 5. PELP1 deregulation promotes hormonal resistance and tumorigenesis. *A, C*, pcDNA- and PELP1-cyto-expressing clones were cultured in 5% DCC serum for 48 hours and treated with or without (*A*) E2 (10^{-9} M) or (*C*) tamoxifen (10^{-8} M) for 7 days, and the cell number was determined. *B, D*, The anchorage-independent growth potential of pcDNA- and PELP1-cyto-expressing clones was measured by the ability of the cells to form colonies on soft agar. Cells were treated with either (*B*) E2 (10^{-9} M) or (*D*) tamoxifen (10^{-8} M), and colonies were counted after 21 days. Similar results were obtained in three independent experiments. Error bars represent standard error of mean. **, $p < 0.01$; ***, $p < 0.001$; NS. Nonsignificant (Student's *t* test). *E*, Nude mice were injected in the mammary fat pad with 5×10^7 MCF-7 cells that stably express either pcDNA or PELP1-cyto, and tumor growth was measured after 8 weeks. Morphology of tumor was evaluated by hematoxylin-eosin staining, and activation of AKT pathway in the tumors was confirmed by immunohistochemistry with a phospho-AKT antibody. *F*, Model for PELP1 regulation of ER functions. Cytoplasmic localization of PELP1 promotes its

interactions with EGFR, Src kinase and PI3K, thus leading to the activation of nongenomic pathways including MAPK and AKT. Such excessive activation of nongenomic pathways promotes hypersensitivity and tamoxifen resistance, which could in part be due to the modification of ER phosphorylation. PELP1 serves as an adaptor protein because it interacts with growth factor receptors, signaling kinases, and nuclear receptors, its deregulation may perturb ER signaling network toward tumorigenesis.

Table 1. Summary of the immunoreactive staining of PELP1 in breast tumors

No staining	Positive staining	Nuclear staining	Cytoplasmic staining	Nuclear and cytoplasmic staining
38.7% (24/62)	61.3% (38/62)	2.63% (1/38)	57.9% (22/38)	39.5% (15/38)

Fig 1

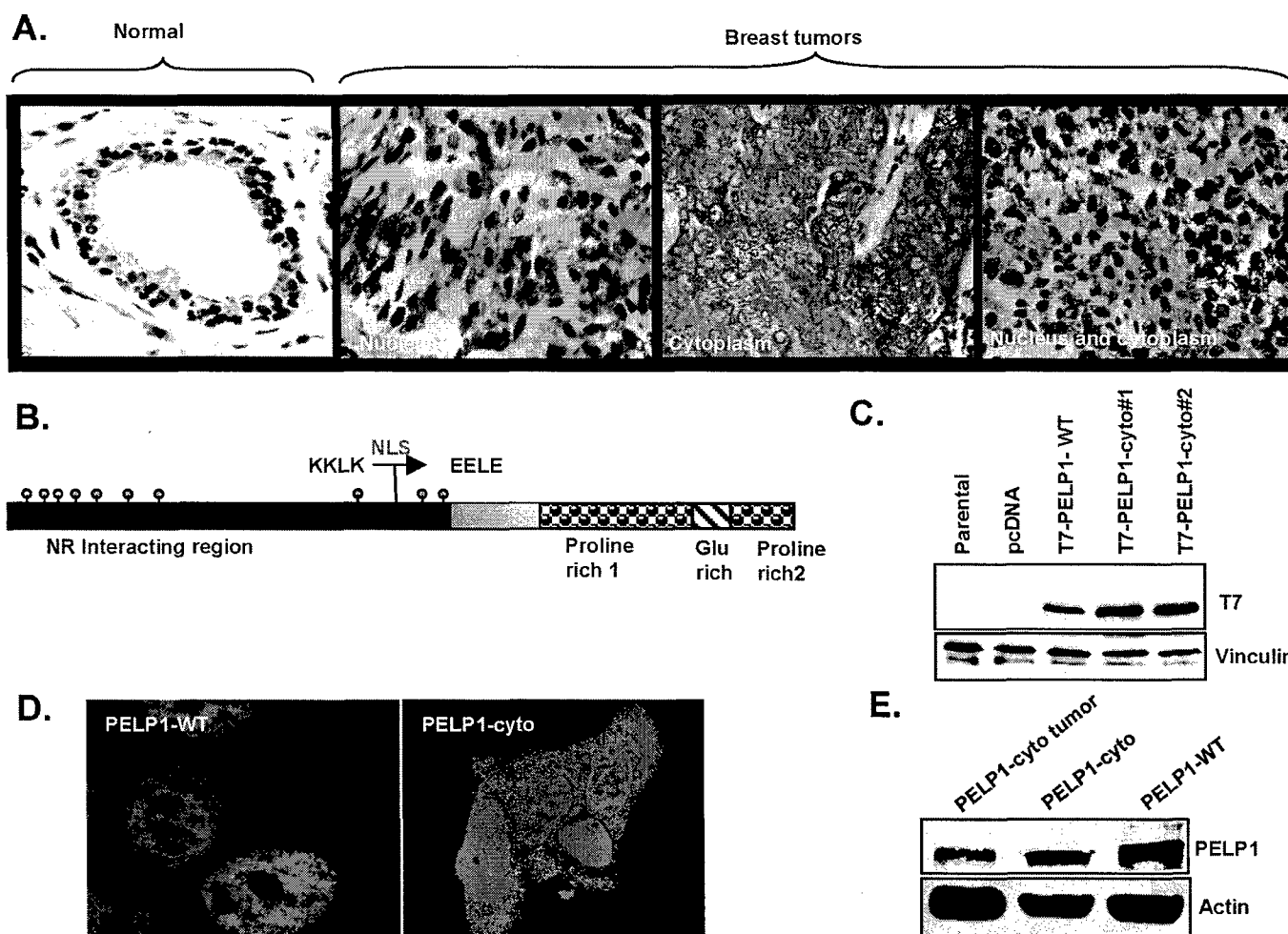


Fig 2

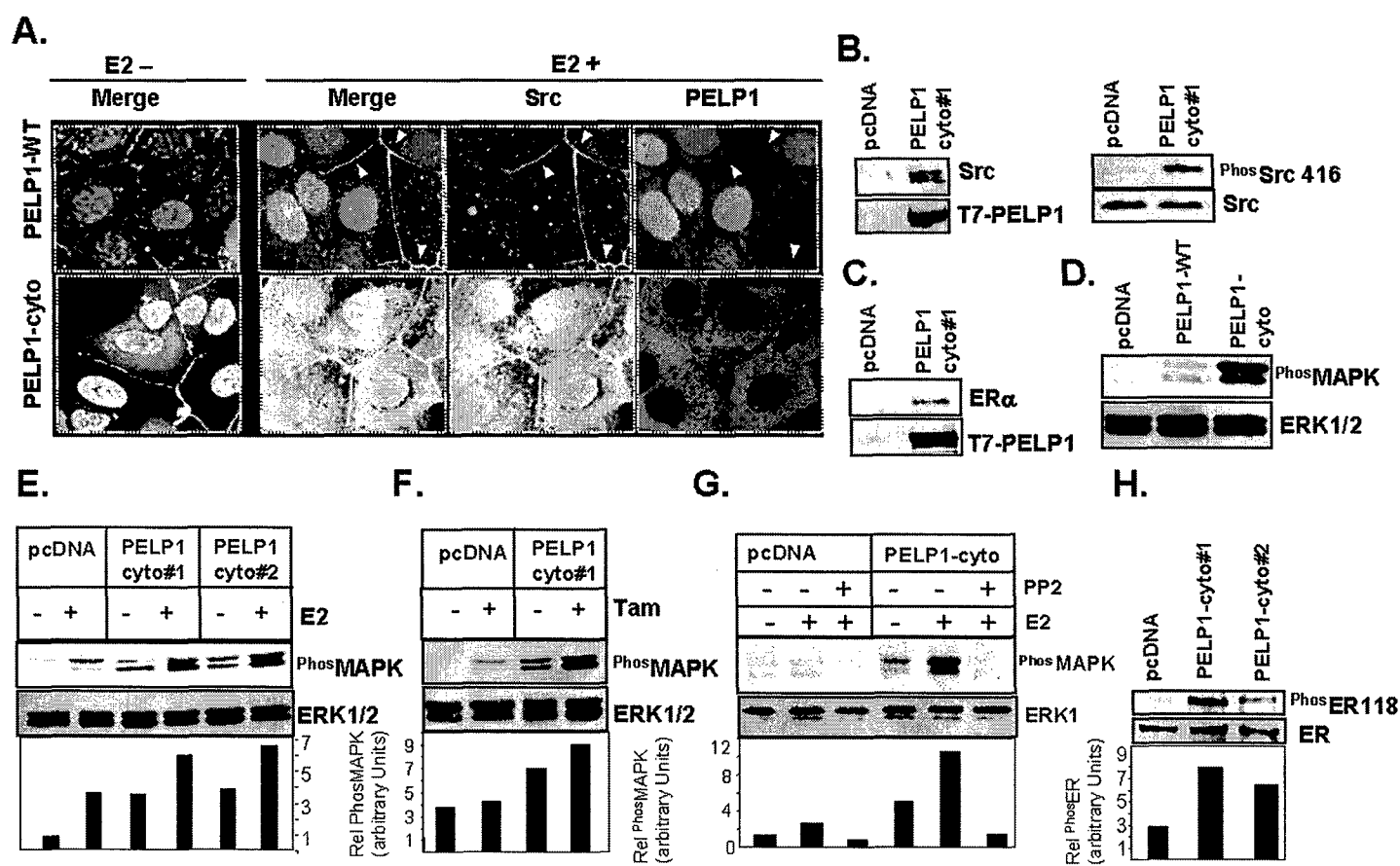


Fig 3

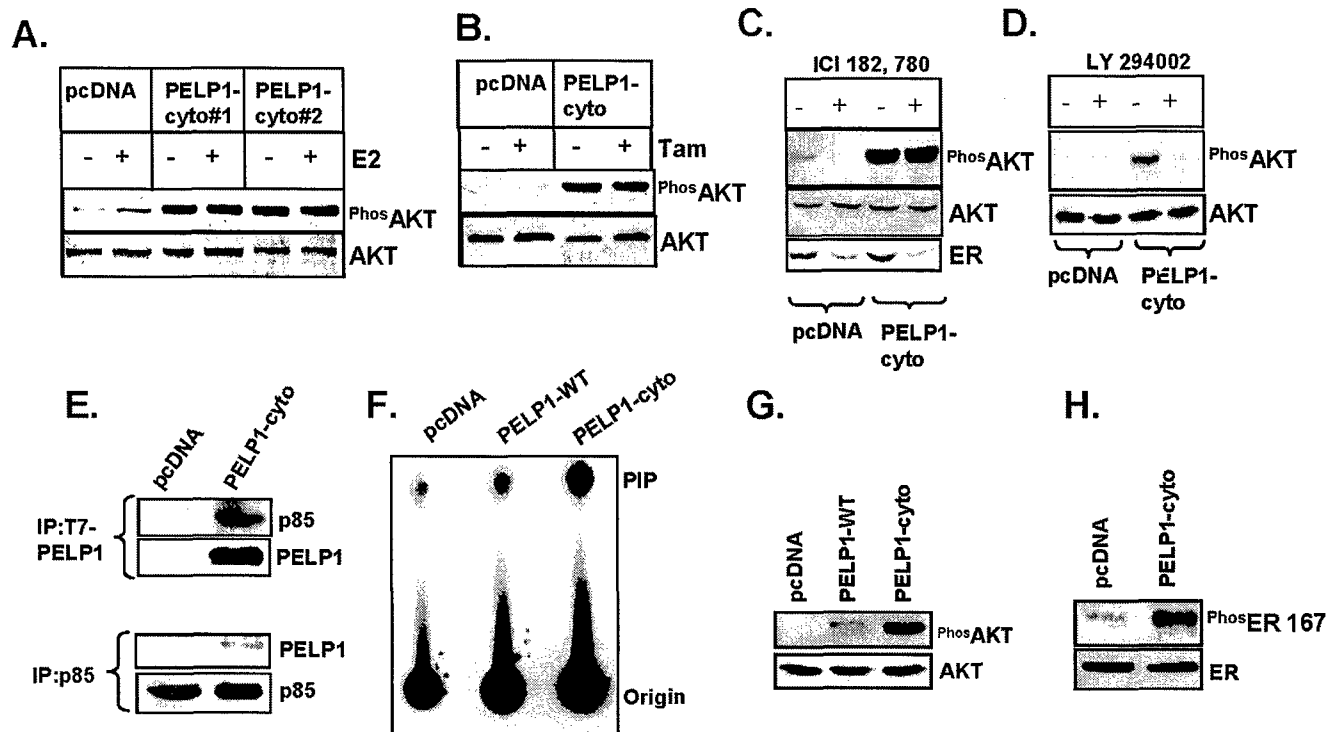


Fig 4

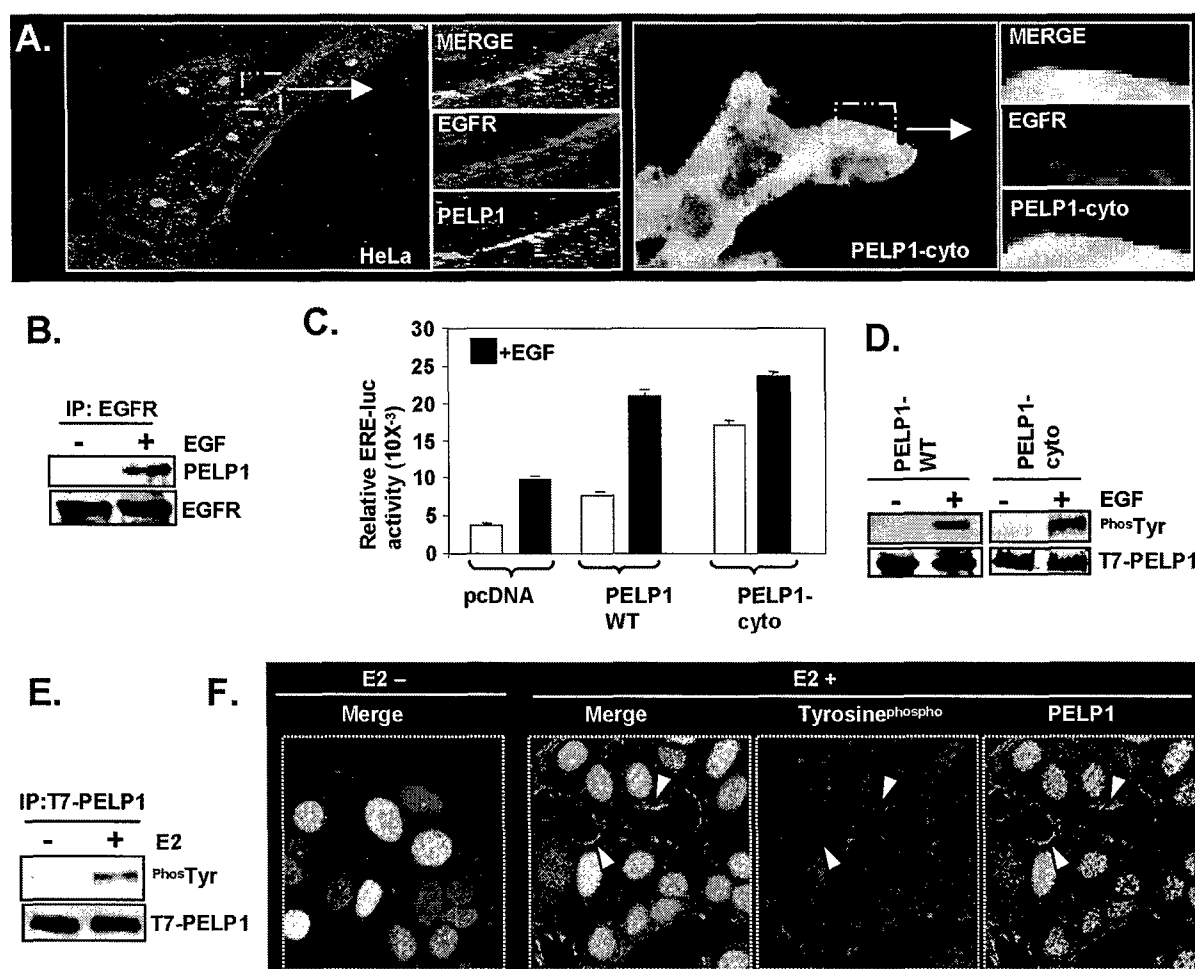


Fig 5

