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Epidermal Growth Factor Promotes Protein Degradation of Epithelial Protein Lost in Neoplasm (EPLIN), a Putative Metastasis Suppressor, during Epithelial-mesenchymal Transition*

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Background: The mechanism of EGF signaling in the regulation of prostate cancer (PCa) metastasis remains unclear.

Results: EGF promotes epithelial-mesenchymal transition (EMT) and induces degradation of epithelial protein lost in neoplasm (EPLIN), a putative suppressor of PCa metastasis.

Conclusion: EGF activates ERK1/2-dependent phosphorylation, ubiquitination, and protein turnover of EPLIN.

Significance: This study suggested that blockade of EGF signaling could retard EMT and inhibit invasiveness of PCa cells.

Aberrant expression of EGF receptors has been associated with hormone-refractory and metastatic prostate cancer (PCa). However, the molecular mechanism for EGF signaling in promoting PCa metastasis remains elusive. Using experimental models of PCa metastasis, we demonstrated that EGF could induce robust epithelial-mesenchymal transition (EMT) and increase invasiveness. Interestingly, EGF was found to be capable of promoting protein turnover of epithelial protein lost in neoplasm (EPLIN), a putative suppressor of EMT and tumor metastasis. Mechanistic study revealed that EGF could activate the phosphorylation, ubiquitination, and degradation of EPLIN through an extracellular signal-regulated kinase 1/2 (ERK1/2)-dependent signaling cascade. Pharmacological inhibition of the ERK1/2 pathway effectively antagonized EGF-induced EPLIN degradation. Two serine residues, i.e., serine 362 and serine 604, were identified as putative ERK1/2 phosphorylation sites in human EPLIN, whose point mutation rendered resistance to EGF-induced protein turnover. This study elucidated a novel molecular mechanism for EGF regulation of EMT and invasiveness in PCa cells, indicating that blockade of EGF signaling could be beneficial in preventing and retarding PCa metastasis at early stages.

Aberrant overexpression of EGF receptors (EGFRs) has been associated with hormone-refractory and metastatic prostate cancer (PCa)2. Activation of EGFR signaling cascade could promote the proliferation, survival, and invasion of PCa cells. As the predominant ligand to EGFR/ErbB1, EGF has been shown to induce epithelial-mesenchymal transition (EMT), a crucial mechanism for the acquisition of metastatic capabilities. Several mechanisms, such as mitogen-activated protein/extracellular signal-regulated kinase kinase (MEK)-dependent up-regulation of TWIST (8), Snail, or Slug (9) may mediate the effects of EGF on EMT and acquired invasiveness.

Epithelial protein lost in neoplasm (EPLIN) is an actin-binding protein that functions as a key molecule linking the cadherin-catenin complex to the actin cytoskeleton (10, 11). Two EPLIN isoforms, the 600-residue EPLIN-α and 759-residue EPLIN-β, are expressed in epithelial and endothelial cells in a context-dependent manner (11–13, 15). EPLIN was initially thought to be a potential tumor suppressor that is preferentially expressed in human epithelium but frequently lost in cancerous cells (11, 16). Recently, we reported a novel role of EPLIN in the regulation of EMT and invasiveness (17). EPLIN depletion in epithelial-like PCa cells promoted the disassembly of adherens junction, structurally distinct actin remodeling, and activation of β-catenin signaling. We further identified a subset of putative EPLIN target genes associated with EMT, invasion, and metastasis. Significantly, EPLIN down-regulation was associated with clinical lymph node metastases of human solid tumors.

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2 The abbreviations used are: PCa, prostate cancer; EPLIN, Epithelial Protein Lost in Neoplasm; EGFR, EGF receptor; EMT, epithelial-mesenchymal transition; SCCHN, squamous cell carcinoma of the head and neck.
EGF Induces EPLIN Protein Degradation during EMT

TABLE 1
Antibodies used in Western blot analyses, immunoprecipitation, confocal microscopy, and immunohistochemistry

<table>
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<th>Antibody</th>
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<tr>
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- EPLIN (immunoblotting/immunofluorescence) BD Transduction Laboratories 61273
- EPLIN (immunohistochemistry) Santa Cruz Biotechnology Sc-135553
- E-cadherin (immunoblotting/immunofluorescence) BD Transduction Laboratories 610181
- E-cadherin (immunohistochemistry) Santa Cruz Biotechnology Sc-7670
- β-catenin (immunoblotting/immunofluorescence) Santa Cruz Biotechnology sc-7199
- β-catenin (immunohistochemistry) Santa Cruz Biotechnology 96818
- TRP Santa Cruz Biotechnology A5316
- β-actin Cell Signaling Technology 42677
- EGF Cell Signaling Technology 2236
- p-EGFR (Tyr-1068) Cell Signaling Technology 2236
- neu Santa Cruz Biotechnology sc-12351
- p-neu (Tyr-1221/1222) Abcam ab2428
- GFP Santa Cruz Biotechnology 2243
- p-serine Enzo Life Sciences BML-PW8810
- ERK1 Upstate 05-1000
- p-ERK1/2 Santa Cruz Biotechnology sc-9996
- Immunofluorescence and Confocal Imaging—Immunofluorescence was performed as described previously (17). Mouse anti-EPLIN antibody, rabbit anti-E-cadherin antibody, or rabbit anti-β-catenin antibody were used (Table 1). Actin filaments were stained by phalloidin (Invitrogen) at a dilution of 1:500. Cells were imaged on a Zeiss LSM 510 META. In all cases, either a 63× or 100× Zeiss Plan-Apo oil objective was used (numerical aperture of 1.3 and 1.4, respectively). All images had contrast expansion performed in Adobe Photoshop.

EXPERIMENTAL PROCEDURES

Cell Culture—Human PCa cell lines ARCaP-E, ARCaP-M, PC3, and C4-2 were routinely maintained in T-medium (Invitrogen) with 5% fetal bovine serum (FBS). Human SCCHN cell lines 686LN, M4e, 37A, and 37B (21) were maintained in Dulbecco’s modified Eagle’s medium (DMEM)/F12 supplemented with 10% fetal bovine serum. Where specified, cells were starved overnight with serum-free RPMI 1640 and treated with recombinant human EGF (R&D Systems, Minneapolis, MN). For blocking ERK1/2 signaling, U0126 (1 μM; Calbiochem, San Diego, CA) or PD98059 (10 μM; Calbiochem) were preincubated with PCa cells for 1 h prior to EGF treatment. Proteasome inhibitors PS341 (1 μM; LC Laboratories, Woburn, MA) or MG132 (10 μM; LC Laboratories) were preincubated with PCa cells for 2 h prior to EGF treatment for the analysis of protein ubiquitination.

Western Blot Analysis—Total cell lysates were prepared using radioimmune precipitation assay buffer (Santa Cruz Biotechnology). Nuclear proteins were extracted using a Novagen kit (EMD Biosciences, San Diego, CA). Immunoblotting analysis followed standard procedure (22). Information for the antibodies used in this study is described in Table 1.

EGF induces EPLIN protein degradation during EMT. These data were further supported by recent reports that EPLIN may be a negative regulator of tumor aggressiveness in PCa, breast cancer, and esophageal cancer (18–20).

Using experimental models of PCa progression, we observed that EGF could induce EMT and promote invasiveness in epithelial-like PCa cells. Intriguingly, the morphological and behavioral changes were associated with a significant decrease in EPLIN expression at protein level. We further elucidated that EGF could activate the phosphorylation, polyubiquitination, and degradation of EPLIN via an extracellular signal-regulated kinase 1/2 (ERK1/2)-dependent signaling cascade. Mutation of certain serine residues of EPLIN rendered resistance to EGF-induced protein degradation. These studies revealed a novel mechanism of EPLIN regulation that could contribute to PCa metastasis.
were serum-starved overnight and further incubated in the absence or presence of EGF at varying concentrations. Luciferase activities were measured 48 h later using a Dual-Luciferase reporter assay system (Promega). Relative luciferase units were expressed as firefly luciferase intensity normalized to Renilla luciferase activity.

Quantitative Reverse Transcription PCR and RT-PCR—Total RNA was prepared with Qiagen RNeasy kit (Valencia, CA). The first-strand cDNA was synthesized using SuperScript® III First-Strand synthesis system (Invitrogen). Quantitative PCR was performed by the LightCycler 480 system (Roche Applied Science) using a Brilliant® SYBR® Green QPCR Master Mix (Stratagene) according to the manufacturer’s instructions. For end point RT-PCR, the SuperScript® III One-Step RT-PCR kit (Invitrogen) was used following the manufacturer’s protocol. The specific primer pairs are described in Table 3. Glyceraldehyde-3-phosphate dehydrogenase (GAPDH) mRNA was amplified with a pair of primers described previously (22) and used to normalize RNA inputs.

Determination of Protein Half-life (T1/2)—ARCaP, or C4-2 cells or PC3 cells transiently transfected (48 h) with individual wild-type and mutated EPLIN-GFP cDNAs, were incubated with cycloheximide (10 μg/ml, Calbiochem) to inhibit further protein synthesis. Following incubation for 0, 1, 2, 4, and 6 h, cells were harvested and lysed, and Western blotting was performed as described above. A rabbit anti-EPLIN (Novus Biologicals, Inc) was used for the detection of endogenous EPLIN protein in ARCaP or C4-2 cells, or an anti-GFP antibody was used to recognize the ectopically expressed EPLIN-GFP proteins in PC3 cells (Table 1). Desired protein bands from the Western blots were quantitated and normalized by the intensity of corresponding β-actin controls using the ImageJ program (National Institutes of Health), and the data were graphed using the SigmaPlot program (Systat Software Inc., San Jose, CA). Protein degradation rate is expressed as half-life (T1/2), the time for degradation of 50% of the protein, which was determined by exponential decay fitting algorithm.

In Vitro Invasion Assay—A modified Boyden chamber with individual 8-μm polyester membrane inserts of a 24-well plate was coated with Matrigel. 1 × 10³ cells in 200 μl of defined medium were seeded into the upper chamber, and 300 μl of serum-free RPMI 1640 medium supplemented with recombinant EGF (50 ng/ml) were added to the lower chamber. Eighteen h later, invasive cells to the lower surface of the membrane were fixed with methanol, stained with 5% Giemsa solution, and counted with a light microscope.

Animal Study—Six-week-old athymic nude mice (BALB/c, National Cancer Institute) were used. Animal protocols were approved by Emory University Institutional Animal Care and Use Committee. 2 × 10⁶ ARCaP cells per 100 μl per site were injected subcutaneously using a previously established proce-
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with decreased E-cadherin and increased nuclear β-catenin (Fig. 1, C and D). Confocal microscopy further revealed the disassembly of actin stress fibers with a radial structure to short, disordered actin cores, formation of peripheral and dorsal ruffles, and a lower degree of co-localization of EPLIN and actin filaments (Fig. 1E). In a modified Boyden chamber assay, invasion of ARCaPε cells through Matrigel was markedly increased (by 5-fold) in the presence of EGF (Fig. 1F). Consistently, EGF treatment promoted the appearance of a mesenchymal-like morphology in 686LN SCCHN cells (21) (Fig. 1G). These data indicated that EGF could induce EMT in certain human cancer cells, which may involve active remodeling of the actin cytoskeleton and the disruption of epithelial adhesions.

EGF Promotes EPLIN Protein Turnover—Following EGF treatment, a progressive reduction of EPLIN proteins was observed in ARCaPε, C4-2, and PC3 cells, which was accompanied by a decrease in E-cadherin expression (Fig. 2A, left and middle panels). EGF treatment also resulted in a reduction of EPLIN protein in epithelial-like SCCHN cells (Fig. 2A, right panel). On the contrary, EGF did not significantly affect mRNA expression of either EPLIN isoforms in PCa cells, as demonstrated by RT-PCR, quantitative RT-PCR, and EPLIN-β promoter activity assay (Fig. 2B). These results suggested that EGF could promote EPLIN down-regulation primarily at post-transcriptional levels.

We investigated whether EGF induced EPLIN down-regulation by promoting protein turnover. The half-life ($T_{1/2}$) of EPLIN protein in ARCaPε cells was determined in the presence of cycloheximide, a de novo protein synthesis inhibitor (Fig. 2C). In the absence of EGF, EPLIN-ε and -β isoforms have a half-life of $25.3$ and $27.7$ h, respectively. EGF treatment significantly reduced their $T_{1/2}$ to $14.2$ and $6.0$ h, respectively. Addition of a proteasome inhibitor MG132 effectively antagonized EGF-induced EPLIN degradation in ARCaPε cells ($T_{1/2}$ for EPLIN-ε, and $21.2$ h for EPLIN-β). These results indicated that EGF may facilitate protein turnover of both EPLIN isoforms through a proteasome-dependent mechanism.

Polyubiquitination serves as a triggering signal leading to protein degradation in the proteasome (28). To examine the mechanism responsible for EGF-induced EPLIN degradation in ARCaPε cells, EGF treatment (50 ng/ml) induced EMT-like morphological changes in 686LN cells.

*FIGURE 1. EGF induces EMT in PCa and SCCHN cells. A, a functional EGFR signaling system in ARCaP cells. Left panel, expression and phosphorylation of EGFR and neu (ErbB2) in ARCaP cells. A pan-p-EGFR antibody that detects phosphorylation at tyrosine residues 992, 1045, 1068, 1148, and 1173 was used. Right panel, EGF treatment (50 ng/ml, 72 h) induces EMT in ARCaP cells. B, Western blot analyses of E-cadherin expression and nuclear translocation of β-catenin in ARCaPε cells in response to EGF treatment (50 ng/ml). D, confocal microscopy showed that EGF (50 ng/ml) induces down-regulation and dissociation of EPLIN and E-cadherin, and nuclear translocation of β-catenin in ARCaPε cells. E, EGF treatment (50 ng/ml) induces re-organization of the actin cytoskeleton in ARCaPε cells. F, EGF (50 ng/ml, 18 h) treatment increases invasion of ARCaPε cells through Matrigel in modified Boyden chamber. Error bars denote S.E. ($n = 3$). G, EGF treatment (100 ng/ml, 72 h) induces EMT-like morphological changes in 686LN cells.*
FIGURE 2. EGF induces EPLIN protein degradation in PCa and SCCHN cells. A, left panel: EGF treatment (50 ng/ml) induces progressive decrease in protein expression of both EPLIN isoforms in ARCaPE cells. The total lysates were the same as used in Fig. 1C. Middle panel: EGF treatment (50 ng/ml) reduces protein expression of EPLIN and E-cadherin in C4-2 and PC3 cells. Right panel: EPLIN is the predominant EPLIN isoform in several human SCCHN cell lines, which is reduced in the more aggressive cell lines (37B, M4e) when compared with that in the low invasive cell lines (37A, 686LN) (top). EPLIN expression is reduced upon EGF treatment in 37A (50 ng/ml; middle) and 686LN (100 ng/ml; bottom) cells. B, the effects of EGF on the mRNA expression of EPLIN isoforms in ARCaPE cells were assessed by regular (left panel) and real-time quantitative (middle panel) PCR and EPLIN reporter assay (right panel). Expression of GAPDH was used as control. Bars denote S.E. (n = 3). C, EGF (50 ng/ml) promotes protein turnover of both EPLIN isoforms, whereas treatment with a proteasome inhibitor MG132 partially rescues EPLIN in ARCaPE cells. D, EGF induces polyubiquitination of EPLIN in a time-dependent manner. ARCaPE cells were treated with 50 ng/ml EGF. Immunoprecipitation was performed at the indicated times with an anti-EPLIN antibody, and Western blot analyses were performed with an anti-polyubiquitin antibody. Input, total cell lysates; DMSO, dimethyl sulfoxide; CHX, cycloheximide.
EGF Induces Serine Phosphorylation and EPLIN Turnover through ERK1/2 Pathway—Phosphorylation in response to extracellular signals or stress is a common mechanism “priming” certain proteins for subsequent recruitment of degradation machinery (29, 30). We investigated whether EGF could induce phosphorylation of EPLIN in ARCaPE cells, which may be a prerequisite for ubiquitination and degradation of EPLIN. Because current antibodies against phosphorylated mouse EPLIN (31) were not suitable for detecting phosphorylation of human EPLIN, an immunoprecipitation protocol was used to precipitate EPLIN and determine its phosphorylation status using a pan-phosphoserine (p-Ser) antibody in an immunoblotting assay. Serine phosphorylation of EPLIN was found to be rapidly increased, with a peak at 15–30 min following EGF stimulation (Fig. 3A).

Several EGFR downstream pathways were tested for their potential role in mediating EGF-induced EPLIN phosphorylation. Among them, ERK1/2 were found to be significantly activated by EGF treatment (Fig. 3B). Blockade of the ERK1/2 pathway with specific MEK inhibitors U0126 or PD98059 abrogated EGF-induced EPLIN phosphorylation at serine sites (Fig. 3C). Importantly, inhibition of ERK1/2 activity effectively antagonized EGF-induced EPLIN turnover. As shown in Fig. 3D, EGF treatment significantly accelerated EPLIN degradation in the presence of cycloheximide ($T_{1/2}$ for EPLIN-β and 5.2 h for EPLIN-α, respectively), whereas preincubation with U0126 resulted in longer $T_{1/2}$ (>15.0 h) for both EPLIN isoforms. Similarly, EGF reduced the $T_{1/2}$ of EPLIN proteins in C4-2 cells, an effect that was attenuated by the pretreatment with U0126 (Fig. 3E). These data suggested a pivotal role of ERK1/2 pathway in mediating EGF regulation of EPLIN phosphorylation and degradation in PCa cells.

Ser-362 and Ser-604 Are Required for EGF-induced Phosphorylation and Degradation of EPLIN—Two web-based programs, PhosphoSitePlus™ and the Eukaryotic Linear Motif (ELM) server, were used to predict putative phosphorylation sites

**FIGURE 3. EGF induces serine phosphorylation and degradation of EPLIN through the ERK1/2 pathway.** A, EGF (50 ng/ml) induces EPLIN phosphorylation at serine residues in ARCaPE cells. B, EGF (50 ng/ml) activates ERK1/2 signaling in ARCaPE cells. C, blockade of ERK1/2 signaling with PD98059 or U0126 inhibits EGF-induced EPLIN phosphorylation in ARCaPE cells. Input: total cell lysates. D, blockade of ERK1/2 signaling inhibits EGF-induced EPLIN degradation in ARCaPE cells. E, blockade of ERK1/2 signaling with U0126 inhibits EGF-induced EPLIN degradation in C4-2 cells. CHX, cycloheximide; DMSO, dimethyl sulfoxide.
within EPLIN-β. Ten serine-containing sequences were identified as potential ERK1/2 substrates, including these spanning amino acids 359–365 and 601–607. A recent study searching for candidate substrates for ERK1/2 by a proteomic approach found that ERK1/2 phosphorylates mouse EPLIN at several serine residues including Ser-360 and Ser-602 (31). Their counter-part residues in human EPLIN protein are Ser-362 and Ser-604, respectively. Of particular interest, Ser-362 appears to be a consensus ERK phosphorylation site (Pro-Leu-Ser-Pro) (Fig. 4A, left panel) (32). To investigate whether these serine residues are required for EGF-induced phosphorylation of EPLIN, we characterized two single-point mutants (S362A, S604A) and a double-point mutant (S362A/S604A) using GFP-tagged wild-type EPLIN-β cDNA as the template (Fig. 4A, right panel). Wild-type (WT) and mutated EPLIN-GFP cDNAs were transiently expressed in the PC3 cell line for its high transfection efficiency and relatively low endogenous EPLIN expression (17). Immunoprecipitation with anti-GFP antibody and subsequent blotting with p-Ser antibody were performed on cells treated with EGF. Consistent with our previous observation (Fig. 3A), EGF treatment significantly induced serine phosphorylation of wild-type EPLIN-GFP within 30 min, whereas point mutations at Ser-362 or Ser-604 inhibited EPLIN phosphorylation by EGF (Fig. 4B, left panel). The double-point mutation appeared to be more effective in antagonizing EGF-induced EPLIN phosphorylation at serine residues (Fig. 4B, right panel). Furthermore, immunoblotting with anti-ubiquitin antibody showed that mutations of S362A, S604A, and S362A/S604A attenuated polyubiquitination of EPLIN in response to EGF treatment (Fig. 4C). The data indicated that these serine residues are required for EGF-mediated EPLIN phosphorylation and ubiquitination.

To examine whether Ser-362 and Ser-604 residues are involved in EGF-induced EPLIN degradation, the stability of wild-type and mutated EPLIN-GFP proteins in the presence of EGF were compared in PC3 cells (Fig. 4D). Following addition of cycloheximide, expression of wild-type EPLIN-GFP was rapidly reduced with a half-life of ~1.6 h. S362A and S604A mutants exhibited longer half-lives of ~15.9 and 5.9 h, respectively, and the double mutation at Ser-362 and Ser-604 significantly increased the stability of EPLIN-GFP protein in PC3 cells, with a predicted $T_{1/2}$ of 29.2 h. These results indicated that the presence of these two serine residues is required for EGF-mediated EPLIN turnover in PCa cells.

Mutations at Ser-362 and Ser-604 Inhibit EGF-induced Down-regulation of E-cadherin in PCa Cells—Using E-cadherin down-regulation as a molecular indicator of EMT, we examined whether a stabilized EPLIN could inhibit EGF-induced EMT process in PCa cells. Consistent with Fig. 2A, EGF induced a decrease in protein expression of both EPLIN and E-cadherin in PCa cells transfected with wild-type EPLIN-GFP. On the contrary, transient expression of mutated EPLIN-GFP (S362A, S604A, and S362A/S604A) effectively blocked EGF-induced E-cadherin down-regulation (Fig. 4E). These data indicated that these “degradation-resistant” EPLIN proteins could render PCa cells less sensitive to EGF-induced EMT.

EGF Induces in Vivo Down-regulation of EPLIN and E-cadherin in ARCaP_e Xenograft Tumors—To investigate the physiological significance of EGF-EPLIN signaling in PCa progression, we determined whether activation of EGFR signaling is associated with EPLIN down-regulation and EMT in an ARCaP_e xenograft model. As shown in Fig. 5A, a 6-week treatment with recombinant EGF did not significantly affect ARCaP_e tumor growth (1.07 ± 0.49 cm³ versus 0.80 ± 0.14 cm³ of control group). However, immunohistochemistry analyses showed that tissue expression of EPLIN and E-cadherin was markedly reduced following EGF treatment. Increased p-EGFR in EGF-treated tissues suggested an activation of the EGFR signaling, which was associated with increased β-catenin at the tissue level (Fig. 5B). The data indicated that activation of EGF signaling could promote in vivo EMT and EPLIN down-regulation in PCa.

**DISCUSSION**

Our recent studies demonstrated that EPLIN could function as a suppressor of tumor metastasis by negatively regulating EMT in PCa cells (17). Down-regulation of EPLIN, therefore, could significantly contribute to the progression of epithelial cancer toward metastatic status. Yet, no previous studies have attempted to elucidate the regulatory mechanism of EPLIN in cancer cells. In this study, we provide biochemical evidence supporting a mechanism by which EGF negatively regulates EPLIN expression through ERK1/2-dependent protein turnover. Gene mutation experiments identified two amino acid residues that are required for the phosphorylation, polyubiquitination, and protein degradation of EPLIN. Animal studies further supported a physiological function of EGF in promoting EPLIN down-regulation and in vivo EMT in an experimental model of PCa. Taken together, these results elucidated an important mechanism for EPLIN down-regulation in epithelial cancer cells.

EGF-induced EMT has been demonstrated as a crucial mechanism for the acquisition of invasiveness in a variety of solid tumors, including breast, ovarian, and head and neck cancer (8, 33). Mechanistically, EGF could activate distinct signaling pathways, including ERK1/2, Akt, Wnt-β-catenin, and signal transducer and activator of transcription 3, to suppress the expression of epithelial proteins (e.g. E-cadherin) and to increase the expression of mesenchymal proteins (e.g. vimentin). Although activation of EGF-EGFR pathway has been associated with aggressiveness and progression-free interval in PCa patients, it remains largely unknown on the mechanisms by which EGF-EGFR signaling promotes metastasis (1–6, 34–36). Earlier studies in DU145 PCa cells found that EGF induced the disruption of epithelial cell adhesion to the extracellular membrane through dephosphorylation and inactivation of the focal adhesion kinase signaling, resulting in enhanced motility and invasion (37). EGF also caused the disruption of cell-cell adherens junctions by caveolin-1-mediated E-cadherin endocytosis, followed by nuclear translocation of β-catenin and activation of T-cell-specific factor-dependent transcription (38). Recently, Gan et al. (39) demonstrated that EGF treatment led to up-regulation of Snail and down-regulation of E-cadherin through an Akt-dependent mechanism, which eventually promoted EMT and enhanced invasiveness in DU145 and PC3 cells. In the present study, we elucidated a novel signaling mechanism by which PCa cells undergo EMT in response to EGF stimulation.

**EGF Induces EPLIN Protein Degradation during EMT**
FIGURE 4. Ser-362 and Ser-604 are critical to EGF-induced phosphorylation and degradation of EPLIN. A, left panel: Ser-362 is an ERK substrate and adjacent to a putative PEST sequence in human EPLIN protein. Right panel: schematic diagram of the EPLIN-GFP construct and its point mutants at Ser-362, Ser-604, or Ser-362/Ser-604. B, mutation at Ser-362, Ser-604, or Ser-362/Ser-604 abrogates EGF-induced EPLIN phosphorylation in PC3 cells. Immunoprecipitation with anti-GFP antibody and subsequent blotting with p-Ser antibody were performed. Input: total cell lysates. C, mutation at Ser-362, Ser-604, or Ser-362/Ser-604 inhibits EGF-induced ubiquitination of EPLIN in PC3 cells. Immunoprecipitation with anti-GFP antibody and subsequent blotting with poliubiquitin antibody were performed. Input: total cell lysates. D, mutation at Ser-362, Ser-604, or Ser-362/Ser-604 inhibits EGF-induced EPLIN degradation in PC3 cells. Protein T1/2 assays were performed in PC3 cells transiently expressing wild type (WT) EPLIN or the point mutants. An anti-GFP antibody was used to detect the presence of EPLIN-GFP proteins. E, mutation at Ser-362, Ser-604, or Ser-362/Ser-604 inhibits EGF-induced down-regulation of EPLIN and E-cadherin in C4-2 and PC3 cells.
demonstrated that EPLIN, a key component of the cell-cell adhesion complex, is a downstream target of EGF-EGF signaling in PCa cells. Down-regulation of EPLIN via accelerated protein turnover could result in the disruption of the adherens junctions and active reorganization of the actin cytoskeleton, both characteristics of EMT and indicators of invasive cancer cells. In addition to these structural alterations, EPLIN down-regulation may also affect the expression of multiple target genes, as we described previously (17). We acknowledged that in parallel to EPLIN degradation, other signaling pathways (such as E-cadherin down-regulation and Wnt/β-catenin activation) could simultaneously be activated by EGF treatment, which together contribute to EGF-induced EMT in PCa cells.

So far, information on the regulation of EPLIN expression is very limited (12). Although the two EPLIN isoforms appear to exert similar functions in the maintenance of epithelial structures (10, 40), their varied expression in different cancer types suggested that EPLIN may be delicately regulated in a context-dependent manner (11, 12, 17, 18). One can envision that extracellular signals may play a major role in the regulation of EPLIN expression. Indeed, mRNA level of EPLIN-α, but not of EPLIN-β, could be readily induced in NIH3T3 fibroblasts upon serum stimulation. A transcription complex composed of serum response factor and megakaryoblastic acute leukemia was formed and subsequently bound a region containing a conserved CARG consensus site within EPLIN-α promoter (41). In PCa cells, however, it appeared that EGF did not have a significant effect on mRNA expression of either EPLIN isoforms, which led us to postulate that EGF-induced EPLIN down-regulation may primarily occur at protein level by affecting protein stability. In fact, as a major EGFR downstream signaling pathway, ERK1/2 has been shown to be capable of directly binding and phosphorylating certain serine residues on mouse EPLIN (31). Intrigued by these observations, we investigated whether phosphorylation of EPLIN could result in the down-regulation of EPLIN. In the present study, we identified at least two serine residues (Ser-362 and Ser-604) of human EPLINprotein are involved in the regulation of EPLIN degradation. It is interesting that the double-point mutation of Ser-362 and Ser-604 appears to be more effective in attenuating EGF-induced phosphorylation and polyubiquitination of EPLIN-GFP, suggesting a compensatory and synergistic effect between Ser-362 and Ser-604 residues. Nonetheless, it is potentially possible that other residues/sequences may also be involved in the regulation of EPLIN stability. For example, it has been demonstrated that the presence of one or more PEST motifs, i.e. the primary sequence rich in Pro, Glu, Ser, and Thr flanked by positively charged amino acids, may serve as a signal to direct ubiquitination for proteolytic degradation. Using a web-based algorithm PESTfind (42), we identified a PEST sequence (RASSLSESSPPK) with a PEST score of +5.89 (PEST scores greater than +5 are considered significant) (Fig. 4A, left panel). Interestingly, this PEST sequence is adjacent to one of the putative ERK1/2 phosphorylation site Ser-362 and contains five serine residues and one lysine residue that may be phosphorylation and ubiquitination sites. It is also worth of noting that the two putative ERK1/2 phosphorylation sites are present in both EPLIN isofoms, which may explain the observed effect of EGF on EPLIN protein expression regardless of which isoform is more prevalent in various cancer cells (Fig. 2A).

Interaction between epithelial cancer cells and adjacent tumor-associated “reactive” stroma is essential to the acquisition of invasive phenotypes (43, 44). Clinical data have indicated that reactive stroma could be a predictor of biochemical-freeness recurrence in PCa patients independent of Gleason score and other pathologic variables (45, 46). A recent microarray study found that EGF transcripts increased significantly in laser-captured reactive prostatic stroma compared with that in matched normal stroma (47). It is possible that EGF abundantly expressed by reactive prostatic stroma could provide a niche to promote EMT and convert certain PCa cells into a highly invasive form (48). Supporting this concept, our animal studies showed that short-term administration of recombinant EGF could significantly reduce EPLIN expression in ARCaPelectrode graft tumors, which was associated with increased p-EGFR and β-catenin at tissue level. Importantly, the expression of E-cadherin in ARCaPelectrode tumors was markedly reduced following EGF treatment, indicating the occurrence of in vivo EMT. It is equally important to note that the in vivo administration of recombinant EGF did not significantly affect tumor growth in athymic nude mice. Consistently, EGF did not exhibit any mitogenic effect in PCa cells (data not shown).

These findings could have important clinical implications. In fact, although EGFR overexpression has been observed in ~30% of PCa patients (49), it remains controversial on the contribution of EGFR signaling to PCa progression. Most of the speculations were based on the failure of a clinical trial with gefitinib (Iressa), a small molecule EGFR inhibitor, in showing significant benefits in PCa patients (1, 50). Our present study
highlighted the importance of EGF-EPLIN signaling in EMT, suggesting a primary role of EGFR signaling in the regulation of PCa invasiveness, but not of PCa growth. Therefore, these observations could provide a molecular explanation for the ineffectiveness of gefitinib in suppressing the growth of primary PCa in clinical setting. On the other hand, the blockade of EGFR signaling could be more effective in preventing and retarding PCa progression toward metastasis. Supporting this notion, gefitinib treatment significantly reduced bone metastatic incidence of the experimental model of PCa (14).

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