Abstract. The role of Eps8 in human breast cancer was studied, and we found that Eps8 was overexpressed in >60% of human breast cancer samples compared with adjacent normal breast tissues by immunohistochemical analysis. Eps8 was highly expressed in the highly invasive breast cancer cell line MDA-MB-231 compared with the weakly invasive breast cancer cell lines MCF7 and MDA-MB-468. MCF7 cell line stably expressing Eps8 was established by G418 screening, and the ectopic expression of Eps8 enhanced MCF7 breast cancer cell growth and survival as assessed by MTT analysis, cell viability and liquid colony formation, whereas the lentiviral expression of Eps8 shRNA in MDA-MB-231 cells resulted in a significant reduction in cellular growth and proliferation in vitro and in vivo. Furthermore, Eps8 knockdown inhibited breast cancer cell migration in wound healing assays, decreased the number and size of EGF-induced filopodia and increased the sensitivity of breast cancer cells to cisplatin analyzed by MTT assays. Eps8 knockdown suppressed a partial EMT-like transition and showed a significant increase in E-cadherin and decrease in N-cadherin and vimentin. These results suggest that Eps8 is overexpressed in human breast cancers, possibly by regulating ERK signaling, MMP9, p53 and EMT-like transition to affect breast cancer cell growth, migration and invasion. Therefore, Eps8 might represent a novel potential target in human breast cancer therapy.

Introduction

Breast cancer is the most commonly malignancy of women worldwide (1), the primary cause of the development and progression of breast cancer is multiple genetic changes, whose accumulation results in the metastatic invasion (2). Some critical genes including HER-2/neu, p53, nm23, and cyclin D are studied in clinical diagnosis for prognostic or predictive outcome (3-6). Therefore, further understanding the molecular etiology of breast cancer is helpful for estimating disease prognosis and guiding treatment.

Eps8 extensively functions as an oncogene in various types of human solid tumors, including squamous cell carcinoma, pancreatic cancer and cervical cancer, as well as colon cancer, pituitary tumors and glioma (7-13) and even hematologic malignancies (14). High expression and concomitant tyrosine phosphorylation of Eps8 were detected in a serial of human tumor cell lines (15). Moreover, elevated Eps8 in cancer patients might be a biomarker of poor prognosis for decreased overall survival (8,14,16). Eps8 could increase cell growth and motility, by modulating the downstream pathways including the mTOR/STAT3/FAK, PI3K/AKT/FOXM1/MMP9, p53/p21WAF1/CIPI1-dependent pathway, EGFR signaling via Rac, and Ras/Raf/MEK/ERK pathways in cancer cells, thus affecting EGFR endocytosis, actin dynamics, RNA processing, cell cycle progression, angiogenesis, cell migration and invasion (7-9,11,12). Furthermore, the effects of downregulated expression of Eps8 on chemotherapeutic agents are also studied. Mithramycin downregulates Eps8 expression and inhibits human epithelial carcinoma cell proliferation and migration (17). Additionally, the cytotoxicity of cisplatin is increased in Eps8-attenuated cervical cancer and lung cancer cells (8,18). Moreover, the loss of Eps8 protein in colorectal adenoma and carcinoma plays a role in the development of a subset of colorectal cancers, suggesting the significance of personalized medicine in patient treatment (19).
Although Eps8 plays a critical role in the development of many malignancies (20), the expression and function of Eps8 in breast cancer are not clear. Combined cDNA array comparative genomic hybridization (CGH) and serial analysis of gene expression analysis (SAGE) identified candidate amplicon target Eps8 as novel putative oncogene in breast cancer (21). In this study, we further determine whether Eps8 is involved in breast cancer malignancy, including cell proliferation and migration. Eps8 was found overexpressed in breast cancer samples and highly invasive cell lines. Overexpression of Eps8 promoted MCF7 cell growth, whereas downregulation of Eps8 in MDA-MB-231 cells significantly inhibited cell proliferation, migration and cell motility. These data suggest that Eps8 serves as a novel growth regulator in breast cancer cells and could be a target for the diagnosis and treatment of breast cancer.

Materials and methods

Immunohistochemistry. Seventy-two breast cancer samples were examined and 3 adjacent normal mammary tissues were used as the control. The study was approved by the Hunan Normal University Human Ethics Committee, and informed consent was obtained from all patients. The immunohistochemical analysis was performed on polyformalin-fixed and paraffin-embedded tissues. Sections (5 µm) were deparaftinized by two 10-min washes in xylene, then rehydrated through successive graded ethanol solutions. Endogenous peroxidase was quenched with 3% H2O2 in methanol for 10 min and washed for 5 min in PBS. Antigen retrieval was achieved by microwaving sections in 0.01 M citrate buffer (pH 6.0) for 10 min at 800 W. The tissues were blocked in 5% bovine serum albumin (BSA) in PBS for 1 h before the addition of the mouse monoclonal anti-Eps8 antibody or mouse IgG control (diluted 1:500, BD Biosciences, CA, USA) at 4˚C for overnight. The sections were incubated with HRP-conjugated goat anti-mouse secondary antibody (diluted 1:100, Sigma-Aldrich, St. Louis, MO, USA) for 45 min and then with 3,3-diaminobenzidine (DAB)/H2O2 for 10 min. Sections were counterstained with hematoxylin, mounted and photographed using an optical microscope (Olympus CX41, Tokyo, Japan). The percentage of tumor cells stained was scored as: 0 (no cell staining), + or 1 (≤30%), ++ or 2 (31-60%) and +++ or 3 (61-100%). The staining between two score values was given 0.5.

Cell culture. Human breast cancer cells MDA-MB-231, MCF7 and T47D were cultured in RPMI-1640 medium (Gibco BRL, Grand Island, NY, USA) containing 100 U/ml penicillin, 100 µg/ml streptomycin and 10% fetal bovine serum (FBS, Hyclone, Australia). MDA-MB-453, MDA-MB-468 cells and normal mammary cells HBL100 were cultured in Dulbecco’s modified Eagle’s medium (DMEM, Gibco) supplemented as described above. All cells were kept at 37˚C in a humidified atmosphere with 5% CO2.

Establishment of MCF7 cell lines overexpressing GFP and GFP-Eps8. MCF7 cells were transiently transfected with constructed pEGFP-C3-Eps8 or pEGFP-C3 control plasmids using Lipofectamine 2000 (Invitrogen, Carlsbad, CA, USA) (13). Stably transfected cells were screened by 400 µg/ml G418 (Sigma) (22,23). The transfection efficiency was detected using an invert fluorescence microscope (Olympus IX71, Tokyo, Japan).

Eps8 RNAi lentivirus generation. Efficient siRNA sequence targeting Eps8 (NM_004447.5) was from the position 93-111 relative to the start codon, stem-loop DNA oligonucleotides were synthesized by Shanghai GeneChem Co. Ltd., China and inserted into the lentivirus-based RNAi vector pGCSIL-GFP (GeneChem) (13). A non-targeting stem-loop DNA was also cloned into pGCSIL-GFP vector as a negative control (NC). Lentiviral particles were prepared as described previously (24). Briefly, the lentivirus expression plasmid and packaging plasmids (pHelper 1.0 and pHelper 2.0) were cotransfected into 293T cells, supernatants were harvested 48 h after transfection and filtered through a 0.45-µm pore size filter (Millipore, Billerica, MA, USA) and concentrated by ultracentrifugation. The infectious titer was determined using hole-by-dilution titer assay. MDA-MB-231 cells at a density of 100,000 cells per well in 6-well plates were infected with Eps8-RNAi-lentivirus or NC-RNAi-lentivirus and 5 µg/ml polybrene (Sigma) at the multiplicity of infection (MOI) 10 and detected on the 4th day by the invert fluorescence microscope.

Cell proliferation assays. For cell viability assay, 3,000 cells were seeded in octuplicate in 96-well plates untreated or treated with 30 µM of cisplatin (DDP, Sigma) for 24 h. On days 1, 3 and 4 or days 1, 3 and 6, cells were analyzed with 4-(3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT, Sigma) at 37˚C for 4 h. Then 100 µl dimethyl sulfoxide (DMSO)/well was added to dissolve the formazan crystals. The absorbance at 490 nm was obtained using a spectrophotometer (UV-2102C, Changsha, China).

For cell survival assay, 100,000 MCF7 cells stably expressing GFP-Eps8 or GFP and parental cells were plated in triplicate in 6-well plates in complete medium containing 400 µg/ml G418. MDA-MB-231 cells (50,000) infecting GFP-Eps8-RNAi-LV or NC-RNAi-LV were plated in triplicate in 6-well plates. After 3-6 days, viable cell numbers were counted with a hemocytometer after trypan blue staining of dead cells.

Liquid colony formation was performed. 2,000 cells were seeded in triplicate in 6-well plates and grown in complete culture medium for >10 days. Colonies were fixed with methanol, stained with Giemsa (BBI International, Cardiff, UK) and photographed with a digital camera (Canon IXUS 125 HS, Tokyo, Japan). Only colonies containing >30 cells were counted under an inverted microscope (Zeiss Axiovert 25, LLC, USA). All experiments were carried out at least three times.

Cell cycle analysis. Lentivirus-infected and parental MDA-MB-231 cells were plated onto 6-well plates for 24 h. Serum was withdrawn when cells were 70% confluent. After 36 h, 10% FBS was added in the medium for an additional 18 h. Cells were collected gently, fixed in 70% ethanol and stained with propidium iodide (PI, BD Pharmingen, San Diego, CA, USA), the DNA content was analyzed on a BD FACSCalibur cytometer (Becton-Dickinson, San Jose, CA, USA) using CellQuest Pro and ModFit software (BD).

Apoptosis assay. Lentivirus-infected and parental MDA-MB-231 cells were harvested and washed with cold PBS. The pellet
was resuspended in 1X binding buffer and the sample solution was incubated with PI and FITC-conjugated Annexin V (BD Pharmingen) for 15 min at room temperature in the dark. The samples were analyzed by the BD FACSCalibur cytometer using the CellQuest software.

**Tumor formation in nude mice.** The mouse experiments were carried out according to the ethical guidelines for laboratory animal use and approved by the Ethics Committee of Hunan Normal University. Approximately 10³ of lentivirus-infected MDA-MB-231 cells in 0.2 ml of sterile PBS were injected subcutaneously into the left and right dorsal regions of 4- to 5-week-old female nude mice (BALb/c). Mice were checked every 2 days and the formed tumors were measured with a micrometer (25). After 26 days, mice were sacrificed, and tumors were excised, weighed and photographed.

**Wound-healing migration assays.** MDA-MB-231 cells were cultured in 24-well plates until >90% confluence. A 100-µl pipette tip was used to generate wounds. After wound creation, the medium was changed to remove cellular debris. Three wounded areas in each well were marked on the bottom of plates and photographed at 1 and 4 days with an invert microscope.

**Immunofluorescence assays.** Stimulation of membrane ruffling with 60ng/ml of human EGF (Sigma, Deisenhofen, Germany) in RPMI-1640 for 30 min was performed after serum deprivation for 48 h. MDA-MB-231 cells were fixed with 3.7% paraformaldehyde for 10 min, permeabilized with 0.1% Triton X-100 for 10 min, and blocked with 5% BSA in PBS. Alexa fluor 595 conjugated to phalloidin (diluted 1:500, Molecular Probes, Eugene, OR, USA) and 1 µg/ml Hoechst 33258 (Sigma) were incubated for 20 min to stain F-actin and the nucleus in the dark. All specimens were viewed with an upright fluorescence microscope (Zeiss Axioskop 2, LLC, USA). At least 20 cells were measured in three independent experiments.

**Western blot analysis.** Cells were lysed in RIPA buffer [50 mM Tris-HCl (pH 7.2), 150 mM NaCl, 1% Triton X-100, 1% sodium deoxycholate, 0.1% SDS and cocktail protease inhibitors]. The lysates (50 µg) were denatured in sample buffer and heated to 105°C for 5 min. Samples were then separated on 10-15% SDS-PAGE gels and transferred to polyvinylidene fluoride (PVDF) membranes (Millipore). The membranes were blocked with 1% BSA in TBS-T [10 mM Tris-HCl (pH 7.5), 150 mM NaCl, 0.1% Tween-20] and incubated overnight with rabbit polyclonal antibodies against GFP and MMP9, mouse monoclonal antibodies against ERK and phosphorylated ERK, cyclin D1 (CCND1), c-Myc, p53, N-cadherin, E-cadherin, vimentin, β-actin and GAPDH (diluted 1:1,000, Santa Cruz Biotechnology, Santa Cruz, CA, USA) in TBS-T containing 1% BSA with gentle shaking at 4°C. HRP-conjugated goat anti-rabbit and goat anti-mouse secondary antibodies (diluted 1:5,000, Sigma) were used. The signals were detected with SuperSignal West Pico chemiluminescent Substrate (Thermo Scientific Pierce, Rockford, IL, USA) and visualized with tanon-5200 system (Bio-tanon, Shanghai, China).

**Statistical analysis.** All statistical analyses were performed using the SPSS 11.0 software (SPSS Inc., Chicago, IL, USA).

### Table I. Patient characteristics of immunohistochemistry.

<table>
<thead>
<tr>
<th>Variable</th>
<th>No. of patients (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Total number</strong></td>
<td>72 (100)</td>
</tr>
<tr>
<td><strong>Age (median, 48 years)</strong></td>
<td></td>
</tr>
<tr>
<td>&lt;48 years</td>
<td>33 (45.8)</td>
</tr>
<tr>
<td>≥48 years</td>
<td>39 (54.2)</td>
</tr>
<tr>
<td><strong>Histological diagnosis</strong></td>
<td></td>
</tr>
<tr>
<td>(invasive carcinoma)</td>
<td></td>
</tr>
<tr>
<td>Ductal</td>
<td>61 (84.7)</td>
</tr>
<tr>
<td>Medullary</td>
<td>11 (15.3)</td>
</tr>
<tr>
<td><strong>Histological grade</strong></td>
<td></td>
</tr>
<tr>
<td>(invasive carcinoma)</td>
<td></td>
</tr>
<tr>
<td>Grade 1</td>
<td>14 (19.4)</td>
</tr>
<tr>
<td>Grade 2</td>
<td>40 (55.6)</td>
</tr>
<tr>
<td>Grade 3</td>
<td>18 (25)</td>
</tr>
<tr>
<td><strong>TNM staging</strong></td>
<td></td>
</tr>
<tr>
<td>Stage IV</td>
<td>4 (5.6)</td>
</tr>
<tr>
<td>Stage III</td>
<td>22 (30.6)</td>
</tr>
<tr>
<td>Stage II</td>
<td>39 (54.2)</td>
</tr>
<tr>
<td>Stage I</td>
<td>7 (9.6)</td>
</tr>
<tr>
<td><strong>Normal tissue</strong></td>
<td>3</td>
</tr>
</tbody>
</table>

Data are shown as mean ± SD from at least 3 independent experiments. Statistical significance was determined using Student’s t-test at P-values <0.05.

**Results**

**Eps8 was overexpressed in breast cancer tissues and high-invasive cell lines.** The expression level of Eps8 was examined in 4 stage IV, 22 stage III, 39 stage II, 7 stage I human breast cancers and 3 adjacent normal mammary tissues by the immunohistochemistry analysis using mouse monoclonal anti-Eps8 antibody. We found that Eps8 was completely localized in the cytoplasm (Fig. 1A). Eps8 expression was detected in 46 (63.9%) of the 72 breast cancers with strong staining (3+), 21 (29%) of the 72 breast cancers were moderately stained (2+), and 5 (6.9%) were weakly stained or negative for Eps8 expression (+/0), which indicated Eps8 was highly expressed in the cytoplasm of breast cancers and 3 adjacent normal mammary tissues by the immunohistochemistry analysis using mouse monoclonal anti-Eps8 antibody. We found that Eps8 was completely localized in the cytoplasm (Fig. 1A). Eps8 expression was detected in 46 (63.9%) of the 72 breast cancers with strong staining (3+), 21 (29%) of the 72 breast cancers were moderately stained (2+), and 5 (6.9%) were weakly stained or negative for Eps8 expression (+/0), which indicated Eps8 was highly expressed in the cytoplasm of breast cancers (P<0.001) according to Student’s t-test. A complete loss of Eps8 expression was observed in normal mammary tissues (Fig. 1B). Therefore, Eps8 expression was significantly increased in human breast cancer samples. Patient characteristics are summarized in Table I.

We next analyzed the expression of Eps8 proteins in five human breast cancer cell lines. Higher expression of Eps8 proteins was evident in MDA-MB-231, MDA-MB-453 and T47D cells than in MDA-MB-468, MCF7 cells and normal breast epithelial cells HBL100 (Fig. 1C). Thus, the level of Eps8 expression was high in the highly invasive breast cancer cell line MDA-MB-231 and low in the weakly invasive breast.
cancer cell line MCF7. Both cell lines were further selected to knockdown and overexpress Eps8 proteins, respectively.

**Eps8 overexpression enhances the proliferation of MCF7 cells.** To further investigate the role of Eps8 in human breast cancer cells, we overexpressed Eps8 and asked whether Eps8 upregulation enhances the proliferation of breast cancer cells. The pEGFP-C3/Eps8 and pEGFP-C3 plasmids were transfected to established MCF7 cell lines stably overexpressing GFP/Eps8 and GFP by G418 screening (Fig. 2A). As shown in Fig. 2B, GFP and GFP/Eps8 target proteins were expressed at high levels in MCF7 cells.

We then investigated whether Eps8 overexpression in the breast cancer MCF7 cells leads to cell proliferation. The same amount of cells was plated in triplicate in 6-well plates and cell number was counted on days 1, 3 and 4. We found that the overexpression of Eps8 in MCF7 cells shows an increased cellular growth compared with controls (Fig. 2D). Further, the liquid colony formation assays indicated a great increase in colony number and size (Fig. 2E). Therefore, these results suggested that Eps8 could contribute to breast cancer cell survival and proliferation *in vitro*.

**Eps8 knockdown inhibits the proliferation and survival of MDA-MB-231 cells in vitro and in vivo.** The above results indicated that Eps8 overexpression significantly promotes MDA-MB-231 cellular proliferation. To gain further supporting data, we attempted to knock down Eps8 expression and asked whether shRNA-mediated Eps8 knockdown inhibits the proliferation of breast cancer cells. Then, lentivirus-based RNAi vector pGCSIL-Eps8 containing the efficient
Eps8 shRNA1 and packaging plasmids were cotransfected to 293T cells (13). Lentiviral particles were prepared to infect MDA-MB-231 cells. The fluorescence intensity was markedly increased 4 days after infection and the infection efficiency was close to 90% in MDA-MB-231 cells (Fig. 3A). Additionally, western blot analysis showed that Eps8 shRNA1 could significantly suppress the expression of endogenous Eps8 proteins when compared with negative control shRNA (Fig. 6A).

Next, we examined whether Eps8 is a critical regulator of breast cancer cell proliferation and investigated the effect of Eps8 knockdown on MDA-MB-231 breast cancer cell growth. From Fig. 3B and C, we observed that Eps8 deletion significantly inhibits the proliferation of MDA-MB-231 cells, whereas control shRNA has no effect. The liquid colony formation assays showed that Eps8 knockdown displays much fewer and smaller colonies in MDA-MB-231 cells, while control shRNA has no effect compared with uninfected parental cells (Fig. 3D). We further explored the molecular mechanism involved in Eps8-mediated cancer cell survival, the effects of Eps8 knockdown on cell cycle and apoptosis were investigated. The flow cytometry analysis showed that...
Eps8 knockdown results in a significant decrease from mean 91.9 to 74.66% in living MDA-MB-231 cells and exhibits an increase in early and late apoptotic cells compared to control shRNA-infected and parental cells (Fig. 3E). Moreover, we compared the DNA content between Eps8 repressed and control cells and found that the proportion of S-phase cells was significantly decreased in Eps8 knockdown cells (31.33%) compared with MDA-MB-231 cells (39.05%), and the propor-
tion of S-phase cells were even lower in Eps8 repressed cells compared with NC control cells (44.83%) (Fig. 3F). These observations suggested that Eps8 knockdown arrested cells at the G1/S checkpoint. Therefore, the above data showed that Eps8 knockdown suppresses proliferation, and cell cycle progression enhancing cell apoptosis of MDA-MB-231 cells.

The inhibitory effect of Eps8 knockdown on breast cancer cell proliferation in vitro suggested that Eps8 depletion might suppress tumor growth in vivo. Tumorigenicity assays were performed by subcutaneous injection of ~10^7 of NC shRNA-infected cells (right dorsal flank) and Eps8 shRNA1-infected cells (left dorsal flank). The measurement of volume of the excised tumors. These data are mean ± SD in 3 independent experiments. *p<0.05 compared with controls.

**Figure 4. Eps8 knockdown results in decreased cell proliferation of MDA-MB-231 cells in vivo.** (A and B) Representative images of nude mice and the tumors following injection of ~10^7 of NC shRNA-infected cells (right dorsal flank) and Eps8 shRNA1-infected cells (left dorsal flank). (C) The measurement of volume of the excised tumors. These data are mean ± SD in 3 independent experiments. *p<0.05 compared with controls.

Eps8 knockdown suppresses breast cancer cell migration, decreases the number and size of EGF-induced filopodia and increases the sensitivity of breast cancer cells to cisplatin. We next assessed the effect of Eps8 on breast cancer cell migration by wound-healing assays. As shown in Fig. 5A, Eps8 depletion produced 62% inhibition of cell migration in MDA-MB-231 cells. In contrast, control groups dramatically promoted cell migration. Thus, Eps8 knockdown was able to significantly inhibit cell motility, which confirmed the role of Eps8 in breast cancer cell migration.

As Eps8 was previously described to be essential for actin dynamics and cytoskeletal organization in pancreatic cancer (12), we examined the effects of Eps8 knockdown on filopodia in MDA-MB-231 cells. Cells were cultivated for 48 h in serum-free medium and then EGF was added for 30 min. Eps8-deletion cells exhibited reduction of filopodial density and length and inhibited the filopodial growth (Fig. 5B), indicating that Eps8 is involved in a rearrangement of F-actin in breast cancer cells.

We further investigated the implication of Eps8 in chemosensitivity of breast cancer cells. MDA-MB-231 cells were treated with cisplatin for 24 h. Sixty-two percent reduction of MTT absorbance was detected in Eps8 shRNA cells compared with 48% inhibition in their control cells (Fig. 5C). Thus, Eps8 knockdown was able to sensitize MDA-MB-231 cells to the cytotoxicity of cisplatin as reported in cervical cancer cells (8).

Eps8 affects the expression of ERK signaling, MMP9, p53 and EMT markers. Because Eps8 influences many genes involved in the development and progression of human cancers, we decided to examine whether Eps8 regulates these target genes in breast cancer cells. As shown in Fig. 6A, Eps8 knockdown had no effect on ERK protein, but it decreased the phosphorylated level of ERK and downregulated the expression of e-Myc and cyclin D1 (CCND1) as downstream mitogenic targets of ERK signaling (26). Moreover, Eps8 knockdown exhibited inhibited MMP9 and enhanced the tumor suppressor p53. The level of EMT epithelial marker E-cadherin was concomitantly increased in Eps8-depleted cells. In contrast, Eps8 knockdown decreased the levels of mesenchymal markers N-cadherin and vimentin. Taken together, these data showed that Eps8 regulates breast cancer cell proliferation and migration, at least in part, by affecting ERK signaling, MMP9, p53 and EMT markers.

**Discussion**

Eps8 is important in regulating the development and progression of many human cancers. However, the role of Eps8 in human breast cancer has not been reported. In the present study, we examined the expression level of Eps8 in histological samples from a large number of patients with previously untreated breast cancer. As detected in the immunohistochemistry analysis, Eps8 was overexpressed in most of breast cancer tissues. Moreover, Eps8 protein was significantly higher in 55.6% of stages II to III breast cancer samples compared to normal mammary tissues. Furthermore, 5 breast cancer cell lines were detected to confirm that the protein level of Eps8 is high in highly invasive breast cancer cells, and low or lost in weakly invasive breast cancer cells. Thus, Eps8 expression may predict cancer recurrence and outcome in stages II-III breast cancer patients.

Evidence presented here demonstrated the effect of Eps8 on breast cancer cell proliferation, we overexpressed...
Eps8 expression is critical for cell growth and proliferation of breast cancer cells in vitro and in vivo and possibly involved in the development and progression in human breast cancer.

Actin polymerization and formation of lamellipodia are believed to play important roles in cell migration during metastasis (27). Eps8, Abi-1 and Sos1 form a tricomplex, induce Rac-specific guanine nucleotide exchange factor (GEF)
activity and transduce signals from Ras to Rac leading to actin remodeling (28-30). Moreover, Eps8 binds with F-actin and colocalizes with actin-based membrane protrusions such as lamellipodia, filopodia and membrane ruffles (28,29). Likely, Eps8 knockdown reduces filopodial density and length and inhibits cell migration in breast cancer cells, suggesting the important role of Eps8 in cellular movement and migration of tumor cells.

Eps8 is an important signal molecule and integrates multiple pathways. For example, Eps8 could control the Ras-Raf-MEK-ERK signaling cascade (31), which plays a crucial role in regulating cellular processes including differentiation, proliferation, survival and apoptosis (32). We demonstrated that Eps8 affects the expression of phosphorylated ERK and downstream genes c-Myc and CCND1 in cellular growth of breast cancer. Apart from regulating cell survival, the ERK pathway promoted invasiveness in tumor cells by upregulation of MMPs such as MMP9 for extracellular matrix remodeling (33). We provided evidence that the levels of MMP9 are decreased in Eps8 shRNA infected cells. In addition, the tumor suppressor gene p53 inhibits the development and growth of the majority of human tumors (34). Eps8 knockdown also enhanced p53 upregulation, resulting in growth arrest or apoptosis of breast cancer cells. The results supported the conclusion that Eps8 regulates the expression of c-Myc, CCND1 and MMP9 through the ERK pathway as well as p53.

Cancer progression toward malignancy is mostly associated with the loss of epithelial differentiation and a switch toward a mesenchymal phenotype. Epithelial-mesenchymal transition (EMT) is considered a critical process for tumor invasion and metastasis (35). Many epithelial proteins including E-cadherin, α-catenin and β-catenin were downregulated, whereas mesenchymal proteins were upregulated, such as fibronectin, N-cadherin, α-SMA and vimentin (36). Loss of E-cadherin promotes metastasis by disrupting intercellular contacts in metastatic dissemination (37). N-cadherin promotes proliferation, adhesion and invasion of prostate cancer cells (38). Moreover, vimentin is associated with a highly invasive cellular phenotype (39). In the present study...
we found that Eps8 knockdown has a significant impact on EMT by increasing E-cadherin expression and decreasing N-cadherin and vimentin, and might achieve lower motility and invasiveness (Fig. 6B). Therefore, these data suggested that Eps8 might partly mediate EMT-like transition and invasion in breast cancer.

In conclusion, our results revealed the pivotal role of Eps8 in breast cancer progression. Eps8 was highly expressed in breast cancer tissues and highly-invasive cell lines. Eps8 regulated the proliferation, migration and invasion of breast cancer cells in vitro and in vivo. Eps8 knockdown inhibited cell migration, decreased the number and size of EGF-induced filopodin and increased the cytotoxicity of cisplatin in breast cancer cells, at least in part, by affecting the expression of ERK signaling, MMP9, p53 and EMT markers. The oncoprotein Eps8 might have an important role as a molecular therapeutic target and even prognostic marker in human breast cancer.

Acknowledgements

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