Converging Evidence for Efficacy from Parallel EphB4-Targeted Approaches in Ovarian Carcinoma

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Abstract

EphB4 is a transmembrane receptor tyrosine kinase that plays an important role in neural plasticity and angiogenesis. EphB4 is overexpressed in ovarian cancer and is predictive of poor clinical outcome. However, the biological significance of EphB4 in ovarian cancer is not known and is the focus of the current study. Here, we examined the biological effects of two different methods of EphB4 targeting (a novel monoclonal antibody, EphB4-131 or siRNA) using several ovarian cancer models. EphB4 gene silencing significantly increased tumor cell apoptosis and decreased migration (P < 0.001) and invasion (P < 0.001). Compared with controls, EphB4 siRNA–1,2-dioleoyl-sn-glycero-3-phosphatidylethanolamine alone significantly reduced tumor growth in the A2780-cp20 (48%, P < 0.05) and IGROV-af1 (61%, P < 0.05) models. Combination therapy with EphB4 siRNA–1,2-dioleoyl-sn-glycero-3-phosphatidylethanolamine and docetaxel resulted in the greatest reduction in tumor weight in both A2780-cp20 and IGROV-af1 models (89–95% reduction versus controls; P < 0.05 for both groups). The EphB4-131 antibody, which reduced EphB4 protein levels, decreased tumor growth by 80% to 83% (P < 0.01 for both models) in A2780-cp20 and IGROV-af1 models. The combination of EphB4-131 and docetaxel resulted in the greatest tumor reduction in both A2780-cp20 and IGROV-af1 models (94–98% reduction versus controls; P < 0.05 for both groups). Compared with controls, EphB4 targeting resulted in reduced tumor angiogenesis (P < 0.001), proliferation (P < 0.001), and increased tumor cell apoptosis (P < 0.001), which likely occur through modulation of phosphoinositide 3-kinase signaling. Collectively, these data identify EphB4 as a valuable therapeutic target in ovarian cancer and offer two new strategies for further development. Mol Cancer Ther; 9(8); 2377–88. ©2010 AACR.

Introduction

Ovarian cancer is the fifth leading cause of death among women in the United States and remains the most common cause of death from a gynecologic malignancy (1). There is no effective screening tool; in the majority of cases, diagnosis is not made until the disease is at an advanced stage. Despite the use of primary cytoreductive surgery and combination chemotherapy regimens, most patients eventually develop recurrent cancer and die of their disease. Thus, there is a critical need for novel therapeutic strategies to improve the outcome of this deadly disease.

EphB4 is a protein tyrosine kinase located on chromosome 7q22, a member of the largest family of receptor tyrosine kinases with at least 16 Eph receptors and 9 ephrin ligands (2). Ephin-B2 is the sole known ligand for EphB4, and binding of EphB4 to ephin-B2 requires cell-to-cell contact, which initiates bidirectional signaling in both EphB4- and ephin-B2-expressing cells (3). EphB4 also plays an important role in angiogenesis as well as a variety of processes during embryonic development, including cell aggregation and migration, segmentation, neural development, and vascular remodeling (4, 5). EphB4 is normally expressed on venous endothelial cells whereas its cognate ligand, ephin-B2, is expressed on arterial endothelial cells. Ligand-receptor binding is required to produce vascular hierarchy and vessel maturation (3). Studies have shown that EphB4 and ephin-B2 expression persists in adult vasculature and plays an important role in adult angiogenesis in wound healing and in the female reproductive system (6).

Increased EphB4 expression has been shown in a number of human cancers, including breast (7), prostate (8), bladder (2), and lung carcinoma (9). EphB4 has been...
shown to provide a survival advantage for tumor cells by interfering with apoptotic pathways and by promoting tumor cell migration and invasion (2, 10). We have previously shown that EphB4 overexpression is predictive of poor clinical outcome in ovarian cancer patients (11). In addition, we showed decreased tumor cell viability, migration, and invasion associated with inhibition of EphB4 expression. The purpose of this study is to further elucidate the biological function of EphB4 in ovarian cancer growth and downstream pathway modulation, and to develop therapeutically relevant approaches for EphB4 targeting.

Materials and Methods

Cell lines and cultures
A2780-par, A2780-cp20, HeyA8, SKOV3ip1, and IGROV-af1 cell lines were cultured in RPMI 1640 supplemented with 10% to 15% fetal bovine serum and 0.1% gentamicin sulfate (Gemini Bioproducts). The derivation and sources of the cell lines have been reported previously (12, 13). The IGROV-af1 variant was derived from ascites arising in a nude mouse given an intraperitoneal (i.p.) injection of IGROV cells. All experiments were done at 60% to 80% confluence, and cell lines were routinely fixed in formalin for paraffin embedding or snap frozen.

siRNA transfections in vitro
EphB4-targeted siRNA was purchased from Qiagen and used to silence EphB4 expression in ovarian cancer cell lines (target sequence 5'-CCCAGCCAATAGC-CACTCTAA-3'). Control nontargeting siRNA (target sequence 5'-AATTCTCAGGAACGTATCGT-3'; confirmed to have no sequence homology with any known human mRNA by BLAST analysis) was used as control siRNA for all in vitro and in vivo experiments. For in vitro transfections, 2 × 10^6 cells per well were plated in six-well plates. After cells were attached, medium was replaced and cells were incubated with 5 μg siRNA (EphB4 or control) with 30 μL of RNAiFect transfection reagent (Qiagen). The following day, medium was replaced and cells were incubated in 3 μg siRNA (EphB4 or control) with 18 μL of RNAiFect transfection reagent. Medium was again replaced 6 hours after transfection.

Liposomal siRNA preparation for in vivo delivery
For in vitro experiments, siRNA constructs were incorporated into neutral nanoliposomes [1,2-dioleoyl-sn-glycerol-3-phosphatidylcholine (DOPC)], lyophilized, and stored at −20°C as previously described (14). Before in vivo delivery, siRNA-DOPC preparations were rehydrated with PBS to appropriate concentrations.

Anti-EphB4 therapy in orthotopic murine ovarian cancer models
Female athymic nude mice were purchased from the National Cancer Institute-Frederick Cancer Research and Development Center (Frederick, MD) and housed in specific pathogen-free conditions. The animals were cared for in accordance with the guidelines set forth by the American Association for Accreditation for Laboratory Animal Care and the USPHS Policy on Human Care and Use of Laboratory Animals, and all studies were approved and supervised by the M.D. Anderson Cancer Center Institutional Animal Care and Use Committee.

Development and characterization of the orthotopic mouse model of advanced ovarian cancer used in these experiments has been previously described by our laboratory (14, 15). Therapy trials were designed to test the effects of EphB4 siRNA alone and in combination with docetaxel. EphB4 siRNA-DOPC dosing schedule was determined through in vivo downregulation studies (data not shown). Tumor cells were trypsinized, resuspended in PBS, and injected into the peritoneal cavity (1 × 10^6 A2780-cp20 cells or 2 × 10^6 IGROV-af1 cells per mouse). One week after tumor cell injection, mice were randomized into four groups (n = 10 mice/group) and treated with i.p. injections of the following agents: control siRNA-DOPC (5.0 μg/mouse i.p., twice weekly), EphB4 siRNA-DOPC (5.0 μg/mouse i.p., twice weekly), docetaxel only (35 μg/mouse i.p., weekly), or EphB4 siRNA-DOPC plus docetaxel (EphB4 siRNA-DOPC, 5 μg/mouse i.p., twice weekly; docetaxel, 35 μg/mouse i.p., weekly). To show a persistent decrease in EphB4 expression in xenografts at completion of the in vivo experiments, EphB4 mRNA levels were evaluated for the different treatment groups using quantitative reverse transcription-PCR (RT-PCR) for the A2780-cp20 cell line.

Additional therapy trials were designed to test the effects of EphB4-131, a murine IgG1 monoclonal antibody that recognizes the FN-1 domain of EphB4 and induces degradation of the receptor (developed by Vascene Therapeutics, Inc.; ref. 16), alone and in combination with docetaxel. EphB4-131 dosing schedule was determined through in vivo downregulation studies (data not shown). One week after tumor cell injection (A2780-cp20 or IGROV-af1), mice were randomized into four groups (n = 10 mice per group) and treated with i.p. injections of the following agents: control antibody (10 mg/kg/mouse i.p., twice weekly), EphB4-131 (10 mg/kg/mouse i.p., twice weekly), docetaxel alone (35 μg/mouse i.p., weekly), or EphB4-131 plus docetaxel (EphB4-131, 10 mg/kg/mouse i.p., twice weekly; docetaxel, 35 μg/mouse i.p., weekly).

During the therapy experiments, mice were monitored daily for adverse effects and sacrificed when moribund. Mouse and tumor weight, tumor distribution, number of tumor nodules, and amount of ascites were recorded at necropsy. Although longitudinal growth curves are frequently used with subcutaneously injected tumors, it is difficult to perform serial measurements with i.p. tumors. Therefore, final tumor weight determinations and nodule counts were done at the completion of the experiments. Mice that did not develop tumor after i.p. injection were excluded from analysis. Tissue specimens were fixed in formalin for paraffin embedding or snap frozen.
Western blot analysis

Whole-cell lysates were prepared from HeyA8, SKO-V3ip1, A2780-par, A2780-cp20, and IGROV-af1 ovarian cancer cell lines. To assess the effects of EphB4 siRNA on EphB4 silencing, whole-cell lysates were collected from A2780-cp20 and IGROV-af1 cell lines 24 to 72 hours after EphB4 siRNA transfection. To assess the effects of EphB4-131 on EphB4 downregulation, whole-cell lysates were collected from A2780-cp20 and IGROV-af1 cell lines 48 hours following treatment with control antibody or EphB4-131 antibody at varying doses (1, 5, 10, and 20 μg/mL). Cell lysates were also collected from cells treated with 10 μg/mL control antibody or EphB4-131 at varying time points (12, 24, 36, 48, and 72 hours). To investigate the effects of EphB4 stimulation on phospho-AKT levels, A2780-cp20 cells were first starved in serum-free medium. Eight hours after starvation began, 20 nmol/L rapamycin or 3 μmol/L LY294002 [phosphoinositide 3-kinase (PI3K) inhibitor; Calbiochem] were added for 16 hours. Cells were then stimulated with 2.5 μg/mL EphrinB2-Fc (Vasgene Therapeutics) for 30 minutes before lysis. To determine the effects of EphB4 silencing on PI3K signaling, A2780-cp20 cells were transfected with EphB4 siRNA (green fluorescent protein siRNA as control) and then treated 48 hours later with 20 nmol/L rapamycin or 3 μmol/L LY294002 for an additional 16 hours. Cells were then lysed. To prepare whole-cell lysates, cells were washed with PBS and then lysed with radioimmunoprecipitation assay lysis buffer supplemented with 1× protease inhibitor cocktail (Roche) and 1 mmol/L sodium orthovanadate for 20 minutes on ice. Lysates were collected and then centrifuged at 13,000 rpm for 20 minutes at 4°C. The protein concentration of the samples was determined by a bicinchoninic acid Protein Assay Reagent kit. Typically, 50 μg of protein from whole-cell lysate were fractionated by 10% SDS-PAGE and transferred to a nitrocellulose membrane. The membrane was then blocked with 5% nonfat milk for 1 hour at room temperature, and probed with primary antibody against EphB4 (MAb 265, Vasgene Therapeutics) or AKT protein phosphorylated at Ser473 with primary antibody against EphB4 (MAb 265, Vasgene Therapeutics) or AKT protein phosphorylated at Ser473 (Cell Signaling) at 4°C overnight. Blots were then incubated with appropriate horseradish peroxidase–conjugated secondary antibodies (The Jackson Laboratory) and developed with an enhanced chemiluminescence detection kit (ECL; Amersham Pharmacia Biotech). To ensure equal protein loading, a monoclonal β-actin antibody (Chemicon International) as per manufacturer’s instructions. The CDNA was subjected to PCR amplification of EphB4 with β-actin as a housekeeping gene (primers: human EphB4 sense: 5′-GCTGG- ACTACGGTTCAAT-3′, human EphB4 antisense: 5′-TCATAAGTGAGGGTGCGAT-3′; murine EphB4 sense: 5′-CAGCAGTTGCTTTCCTGTA-3′, murine EphB4 antisense: 5′-TTTGGCAATTCCTCAC-3′). The PCR cycling condition was as follows: 94°C for 2 minutes, followed by 25 cycles of 94°C for 30 seconds, 55°C for 45 seconds, and 72°C for 2 minutes. Amplified PCR products were analyzed by electrophoresis on 1% agarose gel with Tris-borate-EDTA buffer and visualized under UV light after staining with ethidium bromide. For real-time RT-PCR, we obtained quantitative values (each sample was normalized based on its 18S content), as previously described (17).

Immunohistochemistry

Tumor tissue was fixed with formalin and embedded in paraffin. Tissue sections (5-μm thick) were deparaffinized and hydrated. For antigen retrieval, sections were brought to a boil in 10 mmol/L sodium citrate buffer (pH 6.0) and then maintained at a sub-boiling temperature for 10 minutes. Endogenous peroxidase activity was blocked by incubation in 3% H2O2, followed by blocking of nonspecific sites with blocking buffer (5% normal goat serum in Tris-buffered saline, 0.1% Tween 20) at room temperature for 1 hour. Sections were then incubated with primary antibody diluted in SignalStain antibody diluent (Cell Signaling) overnight at 4°C. After washing in Tris-buffered saline, 0.1% Tween 20, sections were incubated with biotinylated secondary antibodies for 1 hour at room temperature. ABC reagent (Vector Laboratories) was used to develop signal following the manufacturer’s procedure. Nuclei were counterstained with hematoxylin. Antibodies against AKT, phosphorylated AKT (Ser473), S6, and phosphorylated S6 (Ser235/236) were from Cell Signaling. Antibodies against CD31 and proliferating cell nuclear antigen (PCNA) were from PharMingen and DAKO, respectively. For quantification of PCNA expression, the number of PCNA-positive tumor cells was counted in 10 random fields at ×200 magnification. To quantify microvessel density (MVD), microvessel-like structures consisting of endothelial cells that were stained with the anti-CD31 antibody were counted in similar fields.

Terminal deoxynucleotidyl transferase–mediated dUTP nick end labeling assay

After deparaffinization, samples were treated with proteinase K (1:500 dilution). One set of slides was treated with DNase (1:50 dilution) as a positive control. Samples were incubated with terminal dUTP (1:400) and biotin-16-dUTP (1:200) in terminal deoxynucleotidyl transferase buffer at 37°C for 1 hour and then incubated with 2% bovine serum albumin/normal horse serum in water. Samples were then incubated with peroxidase streptavidin 1:400 in house detection diluent at 37°C for...
40 minutes. For visualization, 3,3′-diaminobenzidine and counterstaining with Gill’s hematoxylin was used. For quantification of terminal deoxynucleotidyl transferase-mediated dUTP nick end labeling (TUNEL) signal, the number TUNEL-positive tumor cells were counted in 10 random fields at ×200 magnification.

**Staining of EphB4- or EphB2-expressing cells with EphB4-131**

Full-length EphB4 or EphB2 was transiently expressed in 293T cells and stained with EphB4 or EphB2 antibodies. EphB4-131–specific staining was examined in EphB4- and EphB2-expressing cells, whereas EphB2–specific MAb 110 was used as a positive control. Nuclei were counterstained with 4′,6-diamidino-2-phenylindole (blue).

**Endocytosis of EphB4-131**

A2780-cp20 ovarian cancer cells were incubated with 10 μg/mL of biotinylated EphB4-131 for 1 hour at 4°C or 37°C. Cells were then fixed with 4% paraformaldehyde, and EphB4-131 was localized using streptavidin–FITC. Cells from 10 random fields (final magnification ×400) were counted at ×100 with a Zeiss confocal microscope.

**Migration assay**

To determine the effects of EphB4 downregulation on tumor cell migration, 1 × 10⁵ ovarian cancer cells (A2780-cp20), pretreated with EphB4 siRNA for 48 hours or EphB4-131, were resuspended in serum-free medium (SFM) and plated onto a 0.1% gelatin-coated membrane matrix using the membrane invasion culture system (18). Bottom wells were filled with SFM, and chambers were incubated for 6 hours at 37°C. At completion, cells in the bottom chambers were removed with 0.1% EDTA, loaded onto a 3.0-μm polycarbonate filter (Osmonics) using an S&H Minifold I Dot-Blot System (Schleicher & Schuell), fixed, stained, and counted by light microscopy (18). Cells from 10 random fields (final magnification ×400) were counted by two investigators (W.A.S. and A.K.S.). Experiments were done in duplicate and repeated once.

**Invasion assay**

The effect of EphB4 downregulation on tumor cell invasion was determined using a membrane invasion culture system as previously described (18, 19). In brief, A2780-cp20 cells were pretreated with EphB4 siRNA for 48 hours or EphB4-131. The following day, 1 × 10⁵ viable cells (resuspended in SFM) were placed onto the top wells of a defined basement membrane matrix consisting of human laminin (Sigma), type IV collagen (Sigma), and gelatin (ICN Biomedical), used as the intervening barrier to invasion. SFM or 5% fetal bovine serum–containing medium was placed into bottom wells as chemotactant, and chambers were then incubated for 24 hours at 37°C. Analysis of cell invasion was done as previously described for migration assay (18). Invasion assays were done in duplicate and repeated once.

**Cell viability assay (MTT assay)**

To determine the in vitro effects of EphB4 downregulation on ovarian cancer cell viability, 2,000 cells per well were plated in a 96-well plate after EphB4 siRNA transfection, with experimental conditions set in triplicate. Cells were then incubated for 72 hours at 37°C. To assess cell viability, 50 μL of 0.15% MTT (Sigma) was added to each well and samples were incubated for 2 hours at 37°C. The medium was then removed from each well and replaced with 100 μL DMSO. Absorbance at 470 nm was analyzed with Ceres UV 900C (Bio-Tek Instrument) within 30 minutes.

**Apoptosis assay**

Apoptosis was studied using Annexin V–phycoerythrin (PE) apoptosis detection kit I (BD Biosciences). Twenty-four, 48, and 72 hours following EphB4 siRNA transfection, A2780-cp20 cells were washed twice with cold PBS and then resuspended in 1× binding buffer at a concentration of 1 × 10⁵ cells/mL. Cells (1 × 10⁵) were then incubated with 5 μL Annexin V–PE and 5 μL of 7-AAD. Cells were gently vortexed and then incubated for 15 minutes at room temperature (25°C) in the dark. After adding 400 μL of 1× binding buffer, samples were analyzed by flow cytometry.

**Clonogenic assays**

A2780-cp20 cells (1 × 10⁶) were plated in 10-cm plates. After cells were attached, medium was replaced and cells were incubated with 10 μg siRNA (EphB4 or control) with 60 μL of RNAiFect transfection reagent (Qiagen). The following day, medium was replaced and cells were incubated in 8 μg siRNA (EphB4 or control) with 48 μL of RNAiFect transfection reagent. Medium was again replaced 6 hours after transfection. Twenty-four hours after transfection, cells were trypsinized and replated in six-well plates (5 × 10² cells per well). After approximately 1 week, when control dishes had formed sufficiently large clones, colonies were fixed and stained. After washing with PBS, 1 mL of a mixture of 4% formaldehyde and 1% crystal violet was added to each well for 5 minutes. Cells were then rinsed with PBS and allowed to dry at room temperature. Colonies of at least 50 cells were then counted in 10 random fields (final magnification ×40).

**Cell proliferation assay**

A2780-cp20 cells (3 × 10⁵) were plated in 96-well plates and then treated with EphB4 siRNA for 24, 48, and 72 hours. To measure the effect of EphB4 siRNA on DNA synthesis, cells were pulse labeled with [³H]thymidine (MP Biomedical) for 2 hours followed by lysis in 100 μL of 0.1 mol/L KOH. Cells were harvested onto fiberglass filters, and incorporated tritium was quantified by β-counter.

**Statistics**

All results were expressed as mean ± SD unless indicated otherwise. Continuous variables were compared...
using Student’s t test or ANOVA. For nonnormally distributed data sets, the Mann-Whitney rank sum test was used. Survival curves were plotted by the Kaplan-Meier method, and differences were determined by log-rank test. \( P < 0.05 \) was considered statistically significant.

Results

EphB4 downregulation with EphB4-131 monoclonal antibody

We first characterized EphB4 expression in ovarian cancer cell lines. Western blot analysis showed EphB4 expression in SKOV3ip1, A2780-par, A2780-cp20, and IGROV-af1 cell lines, with highest expression noted in A2780-cp20 and IGROV-af1 cells (Fig. 1A). Compared with control antibody, treatment with EphB4-131 antibody significantly reduced EphB4 protein levels in A2780-cp20 and IGROV-af1 cells at 48 hours (Fig. 1B). EphB4-131 (10 \( \mu \)g/mL) led to peak downregulation of EphB4 in both cell lines, and further increases in antibody concentration (20 \( \mu \)g/mL) did not result in more downregulation (Fig. 1B). To see the time response with EphB4-131 treatment, A2780-cp20 and IGROV-af1 cells were then treated with 10 \( \mu \)g/mL EphB4-131 antibody at varying time points. The greatest EphB4 downregulation was noted at 72 hours (Supplementary Fig. S1A). Antibody-triggered endocytosis of the cell surface receptor is a well-known mechanism for receptor degradation; thus, we sought to determine if EphB4-131 induces EphB4 endocytosis. A2780-cp20 ovarian cancer cells were incubated with biotinylated EphB4-131, which was localized using streptavidin-FITC. Confocal images show that EphB4-131 bound to cell surface at 4°C and was internalized at 37°C (Fig. 1C).

Specificity of EphB4-131 binding

Full-length EphB4 or EphB2 was transiently expressed in 293T cells and stained with appropriate antibodies. EphB4-131 specifically stained EphB4-expressing cells but not EphB2-expressing cells (Fig. 1D). EphB4-131 also did not recognize other EphB receptors (data not shown).

EphB4 downregulation with siRNA

Before performing in vivo experiments, efficacy of EphB4 siRNA was also tested in vitro with A2780-cp20 and IGROV-af1 cancer cell lines. Compared with control siRNA, treatment with EphB4 siRNA resulted in significantly reduced EphB4 mRNA (Supplementary Fig. S1B) and protein (Fig. 2A) levels in A2780-cp20 cells 72 hours after transfection. Similar results were noted in IGROV-af1 cells, with maximum (>80%) downregulation of EphB4 protein expression at 72 hours (data not shown).

EphB4-131 antibody reduced ovarian cancer growth in orthotopic murine model

To assess potential efficacy of targeting EphB4, we first carried out experiments with the EphB4-131 antibody in the A2780-cp20 and IGROV-af1 models of ovarian carcinoma. Seven days after tumor cell injection, treatment was started according to the following groups (\( n = 10 \) mice per group): (a) control antibody, (b) EphB4-131 antibody, (c) docetaxel alone, or (d) EphB4-131 antibody plus docetaxel. After 5 weeks of therapy, mice were sacrificed and necropsies were done. Compared with controls, EphB4-131 alone decreased tumor growth by 83% in A2780-cp20 (\( P < 0.01 \); Fig. 1E) and 80% in IGROV-af1 (\( P < 0.001 \); Fig. 1E) cell lines. Combination therapy with EphB4-131 and docetaxel resulted in the greatest tumor reduction in both A2780-cp20 and IGROV-af1 models (94–98% reduction versus controls; \( P < 0.05 \) for both groups). Compared with docetaxel alone, combination treatment with EphB4-131 and docetaxel significantly decreased tumor growth in both models (87% versus 94%, \( P < 0.05 \); 85% versus 98%, \( P < 0.001 \), respectively). There was no significant difference between treatment with single-agent EphB4-131 or single-agent docetaxel in either cell line. In both experiments, there was no significant difference in feeding behavior or average mouse weights among the various treatment groups (data not shown).
Figure 1. Targeting EphB4 with EphB4-131 monoclonal antibody in ovarian cancer cell lines. A, Western blot analysis of EphB4 expression in ovarian cancer cell lines. B, A2780-cp20 and IGROV-af1 cells were treated with the indicated amounts of EphB4-131 antibody for 48 h. Cells were then lysed and subjected to Western blot analysis to assess EphB4 expression. Corresponding densitometry graphs are included. C, biotinylated EphB4-131 was localized in A2780-cp20 cells using streptavidin-FITC. Confocal images were taken at ×100. EphB4-bound MAb 131 (green) was internalized at 37°C only. D, EphB4-131-specific staining was seen in cells expressing EphB4 (left) but not in cells expressing EphB2 only (middle and right). EphB2-specific antibody MAb 110 was used as a positive control. Nuclei were counterstained with 4′,6-diamidino-2-phenylindole (blue). E, EphB4-131 alone and in combination with docetaxel significantly decreased tumor growth in A2780-cp20 and IGROV-af1 tumor models (*, P < 0.05, compared with control antibody group). An unrelated control antibody was used as negative control.
Biological effects of EphB4 gene silencing

Based on the role of EphB4 in angiogenesis (4, 5), we assessed MVD in A2780-cp20 tumors by CD31 staining (Fig. 3A). Compared with controls, EphB4 siRNA-DOPC treatment significantly decreased MVD by 49% (P < 0.001). The greatest decrease in MVD was seen in the group with combination of EphB4 siRNA-DOPC and docetaxel (66% decrease; P < 0.001).

To determine whether EphB4 affects tumor cell viability, we examined both in vitro and in vivo effects of EphB4 silencing. A2780-cp20 cells were treated in vitro with EphB4 siRNA or control siRNA, in combination with docetaxel, and the number of viable cells was assessed by the MTT assay. EphB4 siRNA treatment led to a 37% reduction in A2780-cp20 cell number compared with untreated cells at 72 hours (Fig. 3B). To determine the cause for reduced cell numbers following EphB4 knockdown as observed by the MTT assay, we examined the in vitro effects of EphB4 silencing on tumor cell apoptosis using Annexin V–PE labeling. Compared with control, EphB4 siRNA treatment increased apoptosis (9% versus 25%) at 72 hours, a 2.5-fold increase (Fig. 3C).

We next examined the effects of EphB4 siRNA-DOPC, docetaxel, and combination therapy on apoptosis in vivo by immunohistochemical analysis of TUNEL assay in tumor tissues (Fig. 3A). Compared with treatment with control siRNA-DOPC, EphB4 siRNA-DOPC alone led to a 153% increase in tumor cell apoptosis (P < 0.001). A similar increase in apoptosis (169%; P < 0.001) was observed in the group treated with docetaxel alone. The greatest increase in tumor cell apoptosis was observed in the group with combination of EphB4 siRNA-DOPC and docetaxel (415%; P < 0.001).

To determine if EphB4 siRNA has direct antiproliferative effects on tumor cells, A2780-cp20 cells were treated with either control or EphB4 siRNA for 48 hours, and cell proliferation was determined by pulse labeling of cells with [3H]thymidine. EphB4 downregulation did not affect in vitro tumor cell proliferation (Fig. 3D). However, EphB4 silencing decreased tumor cell proliferation in vivo, as evidenced by PCNA staining (Fig. 3A). Both EphB4 siRNA-DOPC and docetaxel alone significantly reduced tumor cell proliferation compared with controls (40% and 41%; P < 0.001; Fig. 3A). The greatest effect was observed with combination treatment (69%; P < 0.001). Given the absence of direct in vitro effects, these results suggest an indirect effect, possibly due to reduced angiogenesis.

EphB4 has been shown to regulate the long-term clonogenic potential of colorectal tumor cells (21). To determine if EphB4 silencing affects the capacity of cells to produce colonies in ovarian cancer, A2780-cp20 cells were treated with either control or EphB4 siRNA and then plated in six-well plates for 1 week. Compared with control, EphB4 siRNA treatment decreased colony formation by 63% (P < 0.001; Fig. 3E).
EphB4 regulates cell migration and invasion

Tumor growth and metastasis is regulated, in part, by the ability of tumor cells to degrade the surrounding matrix and migrate. EphB4 has been shown to promote the migration of various cells, including endothelial cells (22). Therefore, we next determined whether EphB4 siRNA could decrease tumor cell migration and invasion using a membrane invasion culture system (18, 19). Compared with control, tumor cell migration was reduced by 60% following treatment with EphB4 siRNA and 58% following treatment with EphB4-131 (*, P < 0.001; Fig. 4A).

Similarly, compared with controls, tumor cell invasion was decreased by 51% with EphB4 siRNA and 67% with EphB4-131 (*, P < 0.001; Fig. 4B).

Figure 3. Biological effects of EphB4 silencing. A, effects of EphB4 siRNA-DOPC with or without docetaxel on MVD (CD31 staining), proliferation (PCNA staining), and apoptosis (TUNEL assay). Bar graphs correspond to the figures on the left. All photographs were taken at an original magnification of ×200 (error bars, SEM; *, P < 0.01, compared with the control siRNA-DOPC group). B, effect of EphB4 si RNA on A2780-cp20 cell viability in vitro at 72 h. C, effect of EphB4 siRNA on A2780-cp20 apoptosis analyzed by Annexin V–PE staining. D, effect of EphB4 siRNA in vitro on A2780-cp20 tumor cell proliferation ([3H]thymidine uptake). E, effect of EphB4 siRNA on A2780-cp20 colony formation (*, P < 0.05, compared with the control siRNA group). Representative pictures are shown below the quantitation bar graphs.
**EphB4 stimulates PI3K activation**

The PI3K pathway plays a pivotal role in ovarian cancer growth and progression. Therefore, we examined the effects of EphB4 activation on PI3K signaling in ovarian cancer cells. Before stimulation, A2780-CP20 cells had been starved for 24 hours to reduce background pAKT (AKT phosphorylated at Ser473, an important marker of PI3K pathway activation) signal. In some settings, cells were also treated by LY294002 (PI3K inhibitor) or rapamycin (mTOR inhibitor) for 16 hours. Cells were then stimulated with 2.5 \( \mu \text{g/mL} \) EphrinB2-Fc for 30 minutes. This stimulation resulted in increased pAKT level (Fig. 5A). Treatment with LY294002 (PI3K/AKT inhibitor) effectively blocked this increase in pAKT level, whereas rapamycin had no effect. This lack of effect by rapamycin is likely due to the interrupted feedback mechanisms in serum-starved cells (23).

Basal AKT activation was observed in A2780-cp20 cells cultured in fetal bovine serum containing complete growth medium, indicating constitutive activation (Fig. 5B). This basal AKT (PI3K) activation was significantly decreased by LY294002, as well as EphB4 silencing by siRNA (Fig. 5B). Thus, AKT activation in these cells is for the most part mediated by EphB4. In addition, LY294002 also reduced EphB4 level (Fig. 5B). We also examined the effect of EphB4-131 antibody on PI3K signaling in vivo. Immunohistochemical staining was done to check the level of total AKT and pAKT in ovarian xenograft tumor tissues treated by control antibody or EphB4-131. The level of pAKT was significantly reduced in EphB4-131 antibody–treated tumors, whereas no obvious difference in total AKT was observed between the control group and EphB4-131 antibody–treated tumors (Fig. 5C).

**Discussion**

In this study, we found that EphB4 siRNA-DOPC and EphB4-131 therapy alone and in combination with chemotherapy substantially reduced tumor growth in ovarian cancer models. These effects seem to be due to a multitude of biological effects on ovarian cancer cells, including decreased tumor cell proliferation and invasion, induction of apoptosis, and decreased tumor angiogenesis.

EphB4 plays an important role in a number of diverse cellular functions, including cell migration, axon guidance, angiogenesis, and vascular remodeling. Aberrant EphB4 expression has been reported in multiple malignancies, including breast (7), colon (21), prostate (8), bladder (2), and lung carcinoma (9). In a previous study, we showed EphB4 overexpression in a substantial proportion of ovarian cancers (11). Although we were able to silences EphB4 expression using antisense oligonucleotides, we wanted to develop more clinically relevant therapeutic approaches against EphB4. Therefore, in the current study, we used two parallel approaches that have potential for clinical development.

EphB4 overexpression has been shown to provide a survival advantage for cancer cells and promote tumor cell invasion, metastasis, and angiogenesis by ligand-dependent and ligand-independent mechanisms. To
Figure 5. EphB4 signals through PI3K/AKT in ovarian cancer cells. A, A2780-cp20 cells were starved overnight and then stimulated with 2.5 μg/mL ephrin-B2-Fc for 30 min. In some settings, cells were also treated with LY294002 (PI3K inhibitor) or rapamycin (mTOR inhibitor) for 16 h. B, cells were transfected with EphB4 siRNA or GRP siRNA as control. Forty-eight hours posttransfection, cells were treated with LY294002 or rapamycin for another 16 h. In both A and B, whole-cell lysates were subjected to Western blot analysis with EphB4, pAKT, and β-actin antibodies. Na3VO4, an inhibitor of phosphatase, was used as a positive control to show phosphorylation of AKT. Quantitation was done by Image J (NIH), and the bar graphs are shown below the picture. The number labels in the bar graph correspond to the number labels in the picture. C, immunohistochemical staining of tumor tissues treated with control or EphB4-131 antibody. Pictures of AKT, pAKT, S6, and phospho-S6 (Ser235/236) were taken at an original magnification of ×200.
evaluate the effects of EphB4 targeting in ovarian cancer, we used a novel monoclonal antibody as well as a highly effective method of systemic siRNA delivery to target EphB4 in orthotopic ovarian cancer models. We showed that targeting EphB4 using EphB4 siRNA-DOPC or EphB4-131 significantly decreased tumor growth. Combination therapy with EphB4 siRNA-DOPC or EphB4-131 and taxane-based chemotherapy had the greatest effect, with no observed side effects. Our results are consistent with other tumor xenograft studies that have shown a marked reduction in tumor growth associated with EphB4 silencing (8, 10).

Angiogenesis is a complex and highly regulated process that is critical for tumor growth and metastasis. EphB4 expression on tumor cells has been shown to promote tumor vascularization through binding with its ligand, ephrin-B2, on endothelial cells (22). Targeting EphB4 on tumor cells is thought to decrease interactions with ephrin-B2–positive endothelial cells, resulting in decreased tumor angiogenesis (22). Our experiments showed a significant decrease in vessel density following EphB4 gene silencing. These results are supported by previous studies showing that blocking EphB4 signaling interferes with tumor angiogenesis and vessel organization (24). In a murine breast cancer xenograft model, EphB4 knockdown significantly decreased tumor growth, with a 44% reduction in tumor angiogenesis (7). Therefore, targeting EphB4 may offer a dual benefit in ovarian cancer by directly targeting tumor cells and indirectly targeting the tumor vasculature.

In addition to decreased angiogenesis, EphB4 targeting led to a significant decrease in tumor cell proliferation and viability in our in vivo experiments. These findings are also supported by studies that have shown EphB4 to function as a survival factor. For example, Xia and colleagues (2) showed a decrease in tumor cell proliferation and increase in apoptosis following EphB4 knockdown in a murine bladder cancer model. Downregulation of EphB4 expression in breast cancer cell lines led to decreased cell viability and activation of caspase-8–mediated apoptosis (7). Consistent with our in vivo results, EphB4 silencing led to a significant reduction in A2780-cp20 ovarian cancer cell viability compared with untreated cells, accompanied by induction of apoptosis.

Given the prosurvival role of EphB4 in ovarian cancer, we suspected that EphB4 may promote other features of malignant disease, such as tumor cell migration and invasion. Munarini and colleagues (25) have shown that transgenic mice with mammary tumors resulting from targeted overexpression of neuT have localized tumors, whereas EphB4/neuT double-transgenic mice frequently develop lung metastasis. Yang and colleagues (26) showed the role of EphB4 in melanoma cell invasion and migration through its influence on RhoA-mediated actin cytoskeleton reorganization. EphB4 silencing decreased RhoA activity and cell migration. Our in vitro studies confirm that EphB4 silencing leads to a significant decrease in ovarian cancer cell migration and invasion. Although EphB4 silencing also contributed to loss of tumor cell viability, the effects on invasion and migration are likely to be independent because they were observed at shorter time points before the observed effects on apoptosis.

Little is known about the downstream signaling of EphB4 in ovarian cancer cells, although activation of the PI3K pathway, including AKT phosphorylation, has been observed in endothelial cells (27). Here, we show increased levels of AKT phosphorylation following activation of EphB4 in ovarian cancer cells. In addition, ovarian cancer cells showed basal levels of AKT phosphorylation, and EphB4 silencing with siRNA effectively blocked this phosphorylation. Moreover, ovarian cancer xenograft tumors treated with the EphB4-131 antibody had decreased levels of phosphorylated AKT and downstream S6 protein. These findings suggest that PI3K signaling in ovarian cancer is modulated by targeting EphB4 in vivo. Thus, the role of EphB4 in ovarian cancer cell migration, proliferation, and survival is likely through activation of the PI3K pathway.

In summary, targeting EphB4 with siRNA-DOPC or EphB4-131 was highly effective in reducing tumor growth in orthotopic ovarian cancer models. Thus, EphB4 may represent a novel target for biological therapy in ovarian cancer.

Disclosure of Potential Conflicts of Interest

No potential conflicts of interest were disclosed.

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References


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