Promotion of epithelial-mesenchymal transition by Frizzled2 is involved in the metastasis of endometrial cancer

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Received February 3, 2016; Accepted March 15, 2016

DOI: 10.3892/or.2016.4885

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Keywords: endometrial cancer, Frizzled2, epithelial-mesenchymal transition, metastasis, Wnt signaling

Abstract. The Wnt signaling pathway is essential for embryonic development, and genetic alteration in this network is closely correlated with tumorigenesis and progression. Previous research has shown that Wnt receptor Frizzled2 (Fzd2) is elevated in many metastatic cancer cell lines and high grade tumors. Yet, little is known about the Fzd2 expression and activity in human endometrial cancer (EC). In this study, we present evidence of a direct role of Fzd2 in human EC. We found that Fzd2 expression was higher in EC than that in adjacent normal tissues, and was correlated with epithelial-mesenchymal transition markers. Next, it was determined that the stable overexpression of Fzd2 in HEC-1B and Ishikawa cells promoted cell migration and induced an EMT phenotype. Conversely, RNA interference-mediated depletion of Fzd2 inhibited EC cell migration. Additionally, mechanistic investigation revealed that elevated Fzd2 expression activated canonical Wnt signaling and was blocked by canonical Wnt signaling inhibitor XAV939. However, Fzd2 did not influence the proliferation of EC cells. Thus, Fzd2 may be a potential marker for EC metastasis and a target for future therapies for this disease.

Introduction

Endometrial cancer (EC) is one of the most common cancers of the female reproductive system with an estimated 54,870 newly diagnosed cases and 10,170 deaths in the USA alone in 2015 (1). In China, the incidence of EC has increased markedly with a higher prevalence in younger women due in part to factors such as obesity and lifestyle changes (2,3). EC is usually diagnosed in early stages (90%) and is often successfully treated with surgical intervention (4). However, EC may metastasize to the pelvic, para-aortic lymph nodes or to distant sites via different routes, and most deaths from EC are caused by metastases that are resistant to conventional therapies. Therefore, it is significant to elucidate the molecular mechanism underlying EC metastasis so as to gain insight into better diagnostic and prognostic biomarkers, as well as novel therapies.

Tumor metastasis consists of several steps, all of which are required for the spread of tumor cells (5). Notably, during tumor progression, epithelial-mesenchymal transition (EMT) is activated in certain cancer cells and enables them to acquire cellular characteristics associated with high-grade malignancy, including the capacity to complete various steps in the metastatic cascade (6). In addition, several studies have elucidated a link between EMT and stem cell characteristics and drug resistance, reinforcing the opinion that EMT is closely related to tumor progression (7,8). During EMT, epithelial cells undergo extensive alterations in gene expression patterns, resulting in the loss of apico-basal polarity, fracture of intercellular adhesive junctions, and degradation of basement-membrane components (9). In this way, epithelial cells adopt mesenchymal traits by altering their morphology, cellular architecture, adhesion, and migratory capacity (10).
However, the mechanisms and pathways that initiate EMT are not comprehensively clear.

EMT is a dynamic procedure and triggered by interactions between extracellular components from the microenvironment and secreted factors, such as the wingless-type MMTV integration site family members (Wnts), transforming growth factor-β (TGF-β), fibroblast growth factors, and epidermal growth factors (9). These factors participate in multiple signaling pathways and initiate the expression of downstream transcription factors, such as Snail and Twist, as well as cytokines, such as MMP2 and MMP9. In the involved signaling pathways, the Wnt signaling pathway plays a critical role in inducing EMT (11-13). Wnt family proteins bind to and activate one or more of the 10 seven-transmembrane Fzd family receptors, playing roles in proliferation, migration, and invasion (14). Previous studies have shown that during EMT, Wnt5a/b ligand and/or its cognate receptor Fzd2 are generally overexpressed in cell lines derived from late-stage mesenchymal-type cancers, such as melanoma and cancers of the breast, lung, colon, liver, and the gastric tract (15-18). However, whether Wnt5a/b-Fzd2 induces EMT in EC and a mechanistic understanding of signaling pathway regulation have been left unanswered by previous investigations.

This study explored the association between the expression of Wnt receptor Fzd2 and EMT markers in EC tissues and investigated the role of Fzd2 in the regulation of EMT in EC cell lines. The findings shed light on the correlation between extracellular components from the microenvironment and secreted factors, such as the wingless-type MMTV integration site family members (Wnts), transforming growth factor-β (TGF-β), fibroblast growth factors, and epidermal growth factors (9).

Materials and methods

Patients and tissues. Thirteen cases of fresh EC and para-tumor normal endometrial tissues were obtained from Chinese female patients who underwent surgical treatment during 2014 and 2015 at the Shanghai First Maternity and Infant Hospital (Shanghai, China). No patient had undergone endocrine therapy, radiotherapy, or chemotherapy before surgery. This study was approved by the Human Investigation Ethics Committee of the Shanghai First Maternity and Infant Hospital. The samples were collected after written informed consent was obtained from the patients.

Cell culture. The human EC cell lines HEC-1B and Ishikawa were obtained from the American Type Culture Collection (ATCC; Manassas, VA, USA). The cells were cultured in Dulbecco's modified Eagle's medium (DMEM)/F12 (Gibco, Auckland, New Zealand), supplemented with 10% fetal bovine serum (FBS) (Gibco Life Technologies, Carlsbad, CA, USA) and 100 U/ml penicillin/streptomycin, and maintained in a 5% CO₂ humidified incubator at 37°C.

Transient and stable transfection. For stable overexpression of human Fzd2 in EC cells, Fzd2 coding sequences were cloned into lentiviral vectors with Ubi-MCS-3FLAG-SV40-puromycin using Gateway technology (Invitrogen Life Technologies, Carlsbad CA, USA) by GeneChem Biotech Co., Ltd. (Shanghai, China). HEC-1B and Ishikawa cells were infected with nontarget or Fzd2-specific lentiviral particles in 6-well plates in the presence of Polybrene (5 mg/ml). The cells were treated with puromycin (1 µg/ml) to generate stable Fzd2-overexpressing clones. The siRNA targeting Frizzled2 (si-Fzd2) and the negative control (si-Ctl) were purchased from Hanyin Biotech (Shanghai, China). The cells were transfected with the siRNA in Opti-MEM using Lipofectamine 2000 (11668-019; both from Invitrogen Life Technologies) according to the manufacturer's instructions.

RNA extraction and qRT-PCR. Total RNA was extracted from the cultured cells using Trizol reagent (Invitrogen Life Technologies) and converted into cDNA with the One Step PrimeScript RT reagent kit (Takara, Dalian, China). The gene expression was detected by real-time polymerase chain reaction (PCR) using SYBR Green Master Mix (Takara) on an ABI Prism 7000 thermal cycler (Applied Biosystems, Foster City, CA, USA). The gene expression was calculated using the 2-ΔΔCt formula and normalized against glyceraldehyde 3-phosphate dehydrogenase (GAPDH). The oligonucleotide primers used for quantitative reverse transcription (qRT)-PCR are listed in Table I. The data were obtained in triplicate from three independent experiments.

Protein extraction and western blot analysis. Total protein was extracted with lysis buffer (Beyotime Biotech) containing a 1% dilution of the protease inhibitor and incubated with antibodies against Fzd2 (Shanghai, China). The gene expression was calculated using the 2-ΔΔCt formula and normalized against glyceraldehyde 3-phosphate dehydrogenase (GAPDH). The oligonucleotide primers used for quantitative reverse transcription (qRT)-PCR are listed in Table I. The data were obtained in triplicate from three independent experiments.

Table I. Primer sequences for real-time PCR analysis.

<table>
<thead>
<tr>
<th>Gene</th>
<th>Primer sequence</th>
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<tbody>
<tr>
<td>GAPDH</td>
<td>F: 5'-AGGCGCTGTTTTAACTCTGTT-3'</td>
</tr>
<tr>
<td>N-cadherin</td>
<td>R: 5'-CCCCACTTGATTGGGGAAGGA-3'</td>
</tr>
<tr>
<td>Vimentin</td>
<td>F: 5'-GACGCCATCAACACGGAGTT-3'</td>
</tr>
<tr>
<td>SPP1</td>
<td>R: 5'-CTTTCGTCGGTGTTAGCTGGT-3'</td>
</tr>
<tr>
<td>Cytokeratin7</td>
<td>F: 5'-TCCGCGAGTTCCACCATTAC-3'</td>
</tr>
<tr>
<td>Cytokeratin19</td>
<td>R: 5'-GCTCTGTCAACTCCGTCAT-3'</td>
</tr>
</tbody>
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F, forward; R, reverse.
and GAPDH (1:5,000; Abcam, Cambridge, MA, USA) at 4°C overnight. Peroxidase-linked secondary anti-rabbit (1:2,000) or anti-mouse antibodies (1:2,000; both from Cell signaling Technology) were used to detect the bound primary antibodies, and the blotted proteins were visualized using an enhanced chemiluminescence kit (Pierce Biotechnology, Inc., Rockford, IL, USA). The intensity of protein bands was quantified using ImageJ software (National Institutes of Health, Bethesda, MD, USA). The relative expression of target proteins was described as a ratio relative to the expression of GAPDH, and statistical data from at least three experiments were graphed.

Cell proliferation assay. HEC-1B and Ishikawa cells were seeded into a 96-well plate (3,000 cells/well). Then, 20 µl of 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) (5 mg/ml; Sigma-Aldrich, St. Louis, MO, USA) was added to each well and subsequently incubated at 37°C for 1 h. The absorbance was measured at 490 nm on a plate reader (Bio-Rad, USA). The intensity of protein bands was quantified using ImageJ software (National Institutes of Health, Bethesda, MD, USA). The relative expression of target proteins was described as a ratio relative to the expression of GAPDH, and statistical data from at least three experiments were graphed.

Migration assay. The cells were seeded in 6-well plates and allowed to adhere for 24 h. Confluent monolayer cells were scratched using a 200-µl pipette tip and then washed three times with 1X phosphate-buffered saline to clear cell debris and suspension cells. Fresh serum-free medium was added, and images were captured at 0 and 24 h at the same position of the wound. For the Transwell assay, a total of 4x10^4 cells were resuspended in 200 µl of the serum-free medium and seeded on the top chamber of the Transwell cell culture chambers (8 µm pore size; Corning Costar, no. 3422). The complete medium (800 µl) was added to the bottom chamber as a chemoattractant. After 16 h, the cells that had migrated to the basal side of the membranes were stained with calcein-AM (0.2 µg/ml; Invitrogen Life Technologies, no. C3100MP) for 30 min and counted at a x200 magnification. Recombinant human Wnt5a and Wnt5b (250 ng/ml; R&D Systems, Inc.) were added to both top and bottom chambers. All experiments were repeated at least three times. The number of cells that had migrated was estimated using MetaMorph image analysis software (Molecular Devices, LLC, Sunnyvale, CA, USA), and the data are expressed as mean average ± standard deviation (SD) (n=3).

Construction of reporter plasmids and luciferase assays. T-cell factor/lymphoid enhancer factor (TCF/LEF) reporter M50 Super 8x TOPFlash and M51 Super 8x FOPFlash (TOP Flash mutant) (plasmids #12456 and #12457 respectively; Addgene Cambridge, MA, USA) driving the expression of green fluorescent protein (GFP) (TOP/FOP-GFP) were
gifts from Randall Moon (Cambridge, MA, USA). HEC-1B and Ishikawa cells (2x10^5) were plated in 24-well plates 24 h before transfection. The cells were co-transfected with 500 ng FOP/TOP reporter plasmid and Renilla luciferase plasmid. The luciferase activity was assayed 24 h after transfection and measured using Dual-Glo Luciferase reagents (E1531;
Promega Corp., Madison, WI, USA). The results were normalized against Renilla activity. All experiments were performed in triplicate.

Statistical analysis. Statistical analyses were performed using the Statistical Package for the Social Sciences software version 17.0 (SPSS, Inc., Chicago, IL, USA). All data are represented as the mean ± SD. Measurement data were analyzed using unpaired Student’s t test or one-way analysis of variance for multiple comparisons. $P<0.05$ was considered to indicate a statistically significant result.

Results

Fzd2 expression in EC tissues. A previous study showed Fzd2 overexpression to be associated with many types of human cancers; the present study assessed whether this was also true for EC. Western blot analysis was used to evaluate the expression of Fzd2 in EC tissues and paired adjacent normal tissues. In a panel of 13 patient tissues, Fzd2 was overexpressed in EC tissues relative to the level in normal tissues (Fig. 1A and B). Moreover, the expression of Fzd2 was positively correlated with markers of mesenchymal cells, such as vimentin (VIM) and N-cadherin (CDH2) (Fig. 1A, C and D).

Wnt5-Fzd2 regulates cell migration. To determine whether Fzd2 is required for cell growth or migration, EC cell lines HEC-1B and Ishikawa were stably transfected with lentiviral vectors encoding human Fzd2 (HEC-1B/LV-Fzd2, Ishikawa/LV-Fzd2) and an empty vector as a control (HEC-1B/LV-Ctl, Ishikawa/LV-Ctl). To examine the efficiency of Fzd2 overexpression, the levels of mRNA and protein expression were detected before cellular assays (Fig. 2A). No differences were observed in cell viability between the control and Fzd2-overexpressing cells (Fig. 2B). However, wound-healing and Transwell migration assays both demonstrated that the migration ability of the HEC-1B and Ishikawa cells was markedly increased by 2- to 3-fold after Fzd2 overexpression (Fig. 2C and D). Additionally, exposure of HEC-1B and Ishikawa cells to human recombinant protein Wnt5a or Wnt5b also appropriately increased cell migration potential (Fig. 2E). Furthermore, when Fzd2 was depleted in HEC-1B and Ishikawa cells using siRNA (Fig. 3A), a significant reduction in migration (Fig. 3B) was found. Overall, these data showed that Fzd2 plays a causal role in EC cell motility.

Fzd2 overexpression promotes the EMT phenotype in EC cells. Because Fzd2 levels are correlated with mesenchymal markers in EC tissues, and the processes involved in EMT are closely correlated with cell motility and cancer metastasis, it was hypothesized that Fzd2 overexpression may drive EMT. To test this, the cellular morphology of the Fzd2-overexpressing EC cells was microscopically examined. HEC-1B/LV-Fzd2 and Ishikawa/LV-Fzd2 cells gained a spindle-shaped morphology and lost cell-cell contacts compared with their control cells, suggesting an EMT phenotype (Fig. 4A). To identify whether this transformation represented EMT, the levels of EMT-associated genes were detected by qRT-PCR and western blot analysis (Fig. 4B and C). Relative to the controls, the levels of epithelial marker E-cadherin in the HEC-1B/LV-Fzd2 and Ishikawa/LV-Fzd2 cells were decreased, whereas the levels of mesenchymal markers CDH2 and VIM were increased. Thus, it was concluded that Fzd2 was involved in the EMT of EC cells.

Fzd2-mediated cell migration is dependent on the canonical Wnt pathway. The activation of the Wnt pathway plays a vital role in EMT during cancer progression. Previous data showed that Fzd2 could activate β-catenin-dependent (canonical) signaling by activating the transcription factor TCF, whose activity can be monitored using well-characterized TOP/FOP-GFP reporter plasmids. Overexpression of Fzd2 in the HEC-1B and Ishikawa cells induced a 2-fold increase in luciferase activity compared with that noted in the vector-only cells (Fig. 5A). Consistently, the levels of β-catenin protein expression in these cells were also elevated (Fig. 5B). To investigate whether Fzd2-mediated migration depended on the β-catenin-TCF pathway, HEC-1B/LV-Fzd2 and Ishikawa/LV-Fzd2 cells were treated with an inhibitor of β-catenin stabilization (XAV939). XAV939 abolished cell migration compared with dimethyl sulfoxide (DMSO) (Fig. 5C). All in all, all these data suggested that Fzd2-mediated cell migration depended on the β-catenin-TCF pathway in EC cell lines HEC-1B and Ishikawa.
Figure 4. Fzd2 overexpression induces the EMT phenotype in EC cells. (A) Fzd2 overexpression induced a mesenchymal morphology in the HEC-1B and Ishikawa cells (magnification, x200). (B) mRNA levels of EMT markers as analyzed by qRT-PCR in the HEC-1B and Ishikawa cells. GAPDH was included as an internal control. (C) Western blot analysis of EMT-related markers in the HEC-1B and Ishikawa cells (left), and blots were further quantified by densitometry of triplicate blots (right). GAPDH was used as an internal control. *P<0.05, **P<0.01, ***P<0.001.

Figure 5. Fzd2-mediated cell migration is dependent on the canonical Wnt pathway. (A) Overexpression of Fzd2 in the HEC-1B and Ishikawa cells induced an increase in luciferase activity compared with the vector-only cells. (B) Western blot analysis of β-catenin in the HEC-1B and Ishikawa cells. (C) Canonical pathway inhibitor XAV939 (10 µM) abolished cell migration after pretreatment for 72 h. Data are expressed as the mean ± SD from one representative experiment of three independent experiments. *P<0.05, **P<0.01, ***P<0.001. Fzd2, Frizzled2.
Tumor metastasis and dissemination are the leading causes of death in EC and as much as 90% of cancer-associated mortality in general (19). Regrettably, the progress in developing efficient strategies specifically targeting tumor metastasis or cells with metastatic potential has been limited (20). Enhancing the understanding of the molecular mechanisms of the metastatic process might improve the clinical management and outcomes of patients with the disease.

Malignant epithelial tumor cells can disseminate from the primary tumor and invade distant organs through a variety of mechanisms. EMT is considered to be an important means of tumor metastasis in many common cancers, including EC (21). Previous studies have indicated that the activation of Fzd2-mediated signaling might be important in metastatic and late-stage cancers, and a high expression of Fzd2 and its ligand Wnt5a could be a potential marker of poor outcome of patients with hepatocellular carcinoma and prostate cancer, respectively (17,18). The present results are consistent with previous findings and prove for the first time that Fzd2 plays an important role in tumorigenesis and acquisition of the metastatic phenotype in EC. This study found that overexpression of Fzd2 in EC cell lines HEK-1B and Ishikawa could promote cell migration potential and an EMT phenotype with an increase in E-cadherin and concomitant reduction in CDH2 and VIM. The loss of E-cadherin protein appears to be a crucial step, reducing cell-to-cell adhesion and destabilizes the epithelial architecture (22). Additionally, even with limited patient numbers, Fzd2 was overexpressed and correlated with EMT markers in EC tissues relative to paired normal tumor-adjacent tissues. Although these studies are not sufficient to confirm Fzd2 as a prognostic factor for EC, the expression data combined with functional data indicate that Fzd2 may serve as a target for anticancer therapy.

However, how Fzd2 regulates EMT transition in EC development remains unknown and must be further investigated. The Wnt signaling pathway is highly conserved with metastatic potential has been limited (20). Enhancing the understanding of the molecular mechanisms of the metastatic process might improve the clinical management and outcomes of patients with the disease.

EMT induction. Furthermore, a series of additional factors have been found capable of inducing EMT in various types of epithelial cells, including TGF-β, Notch, Sonic hedgehog, and multiple growth factors secreted into the microenvironment of tumor cells. All things considered, it is also possible to reveal an alternative mechanism in the tumor microenvironment for inducing and maintaining the mesenchymal state in EC.

A delicate balance between estrogen and progesterone signaling underlies the normal functioning of the female reproductive tract and menstrual cycle. The development of EC, especially Type I, is correlated with estrogen excess (29). Previous studies have shown that estrogen enhances Wnt/β-catenin signaling in the proliferative phase during the menstrual cycle, while progesterone inhibits Wnt/β-catenin signaling, restraining proliferative actions of estrogens in the secretory phase (30,31). Thus, when exposed to enhanced or unopposed estrogen signaling, the constitutive activation of Wnt/β-catenin signaling in endometrium would trigger endometrial hyperplasia, which may develop further into EC.

Although surgery is the standard treatment for early-stage EC patients, patients with lymph node or distant-organ metastases also require chemoradiotherapy and often have poor clinical outcomes. Therefore, identifying biomarkers correlated with cell metastatic potential might help to optimize treatment strategies. This study found that Fzd2 levels in EC tissues are positively correlated with the markers of EMT. Furthermore, our findings concerning EC cells showed that Fzd2 overexpression promoted the EMT phenotype, and these effects involved the activation of the Wnt/β-catenin pathway. Thus, Fzd2 might be a potential marker for EC metastasis and a target for future therapies for this disease.

Acknowledgements

This study was supported by the National Natural Science Foundation of China (no. 81172476, 81272885 and 1472427), the Science and Technology Commission of Shanghai Municipality (no. 13JC1404501), the Doctoral Fund of Ministry of Education of China (no. 212007310090), the Program for Young Excellent Talents in Tongji University (no. 1400813).

References


