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Effect of bFGF on invasion of ovarian cancer cells through the regulation of Ets-1 and urokinase-type plasminogen activator

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Abstract
The aim of the study was to explore the role of basic fibroblast growth factor (bFGF) in ovarian cancer progression. This was done by investigating the effects of bFGF on both the secretion of urokinase-type plasminogen activator (uPA) and the invasion of tumor cells in SKOV₃ ovarian cancer cells. Human ovarian cancer cell line SKOV₃ was cultured in vitro. The expression of uPA gene and protein was induced in SKOV₃ cells; the impact of bFGF on the expression of uPA gene in SKOV₃ cells was studied by RT-PCR, and the impact of bFGF on the expression of uPA protein was tested by ELISA. Ets-1 antisense oligonucleotides were transfected into SKOV₃ cells by liposome protocol. The effects of bFGF on Ets-1 expression and the invasion ability of SKOV₃ cells were determined both before and after exposure to different concentrations of bFGF for 24 h. The expression of both uPA gene and protein was induced in SKOV₃ cells, \( p < 0.05 \). The expression of uPA was suppressed by Ets-1 antisense oligonucleotides in SKOV₃ cells, \( p < 0.05 \). The invasion ability of SKOV₃ cells was increased by 2.3-fold, and this effect was also suppressed by Ets-1 antisense oligonucleotides. bFGF can enhance the invasion ability of ovarian cancer cells in vitro by inducing the expression of uPA, and this effect is also regulated by the transcription factor Ets-1.

Keywords: Basic fibroblast growth factor; Ets-1; invasion; ovarian carcinoma; uPA

Introduction
Basic fibroblast growth factor (bFGF), a 146 amino-acid polypeptide, is a member of a family of molecules encoded by seven genes; it is involved in embryogenesis, angiogenesis, and wound healing (Pötgens et al., 1995). As a potent angiogenesis factor, bFGF is frequently found in malignant tumor and tumor-adjacent tissue, related to the microvessels density of a tumor.

In ovarian cancer, bFGF expression has been found in tumor tissues, cancer cell lines, serum, and ascetic and malignant effusions, related to tumor grading and staging (Di Blasio et al., 1995; Barton et al., 1997; Yoneda et al., 1998; Ueda et al., 2001; Davidson et al., 2002a). Investigations by Davidson and colleagues (2002a) showed that bFGF was the major angiogenic factor in ovarian carcinoma. Also, bFGF has a prognostic value. In fact, the level of serum bFGF combined with CA125 as an ovarian tumor marker has a higher specificity than CA125 alone (Davidson et al., 2002b; Le Page et al., 2006).

The way in which bFGF promotes the progression and metastasis of malignant tumors through regulating angiogenesis has been well elucidated (Pötgens et al., 1995). Sako et al. (2003) found that bFGF could increase the expression of vascular endothelial growth factor (VEGF) of mesothelial cells and promote peritoneal metastasis. More recently, bFGF was found to be related to paclitaxel resistance (Gan et al., 2006). All these findings have thrown light on the mechanism by which bFGF participates in tumor progression and metastasis, but the underlying mechanisms are not yet well understood. In endothelial cells, bFGF can promote cell invasion and migration by stimulating the expression of proteases such as urokinase-type plasminogen activator
(uPA) and metalloproteinases (MMPs) (Iwasaka et al., 1996). In cervical carcinoma, glioma tumor, and gastric cancer, etc., bFGF has been found to be related to protease expression (Mori et al., 2000; Forbes et al., 2003; Kanda et al., 2005; Zhao et al., 2005). Among the large number of proteases involved in cellular invasion, uPA plays an important role as it initiates the activation of metalloproteinases and the conversion of plasminogen to plasmin (Collen, 1999; Danø et al., 2005). The overexpression of uPA is found in nearly all kinds of malignant tumors, including advanced ovarian cancer. Cai et al. (2007) found that uPA was involved in metastasis and had a prognostic value related to epithelial ovarian cancer (Schmalfeldt et al., 2001). Based on all these findings, we hypothesize here that bFGF can stimulate uPA expression in ovarian cancer cells, and promote the invasion of cancer cells. Also, as uPA is one of the target genes regulated by transcription factor Ets-1 (Watabe et al., 1998), we hypothesize that the effect of bFGF is regulated by Ets-1.

In this study, we observed the effects of bFGF on the expression of uPA and Ets-1, and on cell invasion in SKOV3 ovarian cancer cell lines, before and after the inhibition of Ets-1 expression using Ets-1 antisense oligonucleotides.

Materials and methods

Cell culture

SKOV3 cells were cultured in RPMI 1640 medium (Invitrogen) supplemented with 10% heat inactivated fetal bovine serum (FBS), 100 units/mL of penicillin, and 100 µg/mL of streptomycin, and the cells were incubated at 37°C in a humidified 95% air and 5% CO2 atmosphere. When the cells reached subconfluence, basal medium (Invitrogen) containing 0.5% FBS with different concentrations of bFGF (0.1, 1, 10, 100 ng/mL) were incubated for 24 h. Recombinant human bFGF was purchased from Invitrogen, and solutions were prepared in Tris with 0.5% bovine serum albumin (Sigma).

Reverse transcription polymerase chain reaction (RT-PCR) analysis

The cell total RNA was extracted using a Total RNA Isolation Kit (Takara) and treated with RNase-free DNase. The reverse transcription (RT) reaction was carried out using the following: 4 µg of total RNA, 1 µL of oligo primer, 1 µL of 10 mM dNTP, 4 µL of 25 mM MgCl2, 2 µL of 0.1 mM dithiothreitol (DTT), and 2 µL of SuperScript II Reverse Transcriptase (Invitrogen) at 42°C for 50 min.

For the polymerase chain reaction (PCR), 2 µL of cDNA was incubated with 2 µL of 5X buffer, 1.6 µL of dNTP, 0.1 µL of DNA polymerase (Invitrogen), 2 µL of 25 mM MgCl2, 1 µL of primer pairs, and 10.3 µL of water. PCR was performed for 33 cycles of 1 min at 94°C, 1 min at 58°C, 1 min at 72°C, and a final elongation step of 7 min at 72°C.

The human Ets-1 PCR primers were 5’-TCACAGAGCTCTATCGACGC-3’ (sense) and 5’-GTCCTATGAGGTACAGCAG-3’ (antisense); the human uPA primers were 5’-AGAATTCTACCGACTATC-3’ (sense) and 5’-ATTCTCTTCCTTGGTGAC-3’ (antisense); and the human β-actin primers were 5’-GTGGGGCGCCCCAGGCACCA-3’ (sense) and 5’-TCCTTAATGTACGCCAGATTTC-3’ (antisense).

Amplified products were separated on 2% agarose gel and visualized by means of ethidium bromide ultraviolet fluorescence. The expression of a respective target gene was calculated by the density of the corresponding PCR product divided by that of β-actin.

Measurement of uPA levels by ELISA

The supernatant was obtained from cultures of SKOV3 cells treated with different concentrations of bFGF using a uPA enzyme linked immunosorbent assay (ELISA) kit (Adlitteram Diagnostic). Samples were assayed in triplicate. All data were within the linear range of the standard curve generated by use of 0.0, 0.10, 0.25, 0.50, 0.75, and 1.0 ng/mL uPA, which was provided with the kit.

Antisense oligonucleotide treatment of cell monolayers

A 20-mer antisense oligonucleotide was used to knock down Ets-1 expression. The sequences were: antisense 5’-AGATCGACGGCCGCTTCAT-3’, and sense control 5’-ATGAAGGCACGGCTTCAT-3’. The oligonucleotides were purified by high-pressure liquid chromatography. Cells were treated with oligonucleotides by the method described in the manufacturer’s instructions for DOTAP (liposomal transfection reagent; Roche). Briefly, cells were grown to be about 60–70% confluent and rinsed with serum-free RPMI 1640, and then incubated in serum-free RPMI 1640 with 6 µg/mL DOTAP (Roche) and 16 µg/mL oligonucleotides for 6 h.

Invasion assay

The invasion ability of cells treated with or without Ets-1 was assessed by the following method. Cells were washed, and resuspended at 105 cells/200 µL in serum-free medium with 10 ng/mL bFGF and Ets-1 oligonucleotides or bFGF only. The cells were distributed to upper transwells coated with Matrigel (BD Bioscience). The bottom chambers were filled with 10 µg/mL fibronectin. The cells were then incubated at 37°C in a humidified 95% air and 5% CO2 atmosphere for 72 h. After that, the
lower surface of the filter was fixed and stained. The number of invading cells was calculated as the sum of cells counted in five fields at a ×40 magnification.

**Statistical analysis**

Results were assessed by Student’s t-test. The SPSS computer package was used for all analyses, with \( p < 0.05 \) considered as significant.

**Results**

**Effects of bFGF on mRNA expression of uPA and Ets-1 in SKOV3 cells**

The expressions of uPA and Ets-1 after incubation in serum-free medium for 24 h were used as basal levels. After incubation with different concentrations of bFGF (0.1, 1, 10, 100 ng/mL) for 24 h, we observed a significant increase of both uPA and Ets-1 expression. The mRNA levels of uPA and Ets-1 were increased by 3.2- and 2.5-fold, respectively, when treated with bFGF (10 ng/mL), compared with control. The up-regulation of these two genes showed a dose-dependent response (Figure 1).

**Effect of bFGF on protein secretion of uPA**

Using ELISA, we found that uPA protein expression was increased with bFGF treatment and was highest with the treatment of 100 ng/mL bFGF. This effect showed a dose-dependent response (Figure 2).

**Effects of antisense Ets-1 oligonucleotides on Ets-1 and uPA expression**

The use of antisense oligonucleotides blocked the induction effect of bFGF on Ets-1 and uPA expression. Sense oligonucleotides, as control, showed no change of effect. uPA mRNA expression was increased by 3.2-fold with 10 ng/mL bFGF treatment and decreased to 1.4-fold when cells were transfected with antisense Ets-1 oligonucleotides. Similar results were observed with uPA protein expression: a 2.5-fold increase reduced to 1.4-fold upon transfection with antisense oligonucleotides. These results indicate that Ets-1, a bFGF-regulated transcription factor, plays an important role in regulation of uPA expression in SKOV3 cells.

**Effect of bFGF on cell invasion of SKOV3 cells**

Finally, as uPA plays key roles in tumor invasion, we wanted to know whether knock-down of Ets-1 expression altered tumor invasion induced by bFGF. After induction by bFGF, we observed a 2.3- and 2.4-fold increase of cell invasion in the control and sense Ets-1 oligonucleotide groups, respectively, but only a 1.5-fold increase in the antisense oligonucleotide group (Table 1). This shows that Ets-1 plays an essential role in tumor invasion induced by bFGF.

![Figure 1. RT-PCR analysis of uPA and Ets-1 genes. After SKOV3 cells were cultured in basal medium and different concentrations of bFGF (0.1, 1, 10, 100 ng/mL) for 24 h, a significant change of expression of uPA and Ets-1 mRNA of SKOV3 cells was observed. The increase of both genes presented a dose-dependent response. β-actin was functioning here as a housekeeping gene.]

![Figure 2. Effect of bFGF on secretion of uPA protein. Each bar represents mean ± SEM. After treatment with different concentrations of bFGF (0.1, 1, 10, 100 ng/mL), supernatant was obtained from cultures of SKOV3 cells, and tested by ELISA. The expression of uPA protein increased due to stimulation with bFGF in a dose-dependent manner.]

**Table 1.** Effect of bFGF on cell invasion of SKOV3 cells and the effect of antisense oligonucleotides on that procedure.

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Migrated cells (vs. control)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Basal medium (control)</td>
<td>1.9±0.7</td>
</tr>
<tr>
<td>10 ng/mL bFGF</td>
<td>4.4±1.0</td>
</tr>
<tr>
<td>10 ng/mL bFGF + sense Ets-1</td>
<td>4.5±1.0</td>
</tr>
<tr>
<td>oligonucleotides</td>
<td></td>
</tr>
<tr>
<td>10 ng/mL bFGF + antisense Ets-1</td>
<td>2.9±0.9</td>
</tr>
<tr>
<td>oligonucleotides</td>
<td></td>
</tr>
</tbody>
</table>
Discussion

Among the angiogenic factors, bFGF is best characterized. It stimulates endothelial cells to secrete proteases such as MMPs and uPA, which results in degradation of the vessel basement membrane, allows cells to invade the surrounding matrix, and contributes to the neovascularization of tumors (Mandriota & Pepper, 1997). However, little was known about the mechanism of the effect of bFGF on protease secretion or cell invasion in cancer cells.

In this study, we tested the effect of bFGF on the expression of uPA in ovarian cancer cell line SKOV3. As we hypothesized, bFGF stimulated the expression of uPA in ovarian cancer cells in a dose-dependent manner. A similar finding was reported for glioma cells in a previous study (Mori et al., 2000).

Urokinase-type plasminogen activator is one of the most important proteases in tumor progression, not only because it participates in angiogenesis through degradation of the vessel basement membrane, but also because it contributes to tumor invasion and metastasis by degradation of the extracellular matrix, thus allowing cells to overcome the constraints of cell–cell and cell–matrix interactions (Collen, 1999; Danø et al., 2005). In this study, we found that bFGF increased the invasion ability of SKOV3 cells and this effect was accompanied by increased uPA expression. This finding showed a clear correlation of uPA expression and tumor cell invasion. This possibility was also supported by the fact that Ets-1-inhibited expression significantly diminished the invasion ability of SKOV3 cells induced by bFGF, and decreased uPA expression. Together with previous findings in glioma, our studies support a novel mechanism of the effect of bFGF on tumor progression: in addition to contributing to angiogenesis, bFGF promotes tumor invasion through the up-regulation of uPA expression (Kitange et al., 1999).

Finally, we tested the possible pathway of bFGF in uPA regulation. As uPA is one of the target genes of Ets-1 in breast cancer cells (Watabe et al., 1998), we hypothesized that Ets-1 participates in this regulation. Using an antisense oligonucleotide of Ets-1, we found that the expression of uPA gene and protein was significantly decreased by suppressing the Ets-1 gene, and so was the cell invasion ability. Our studies indicated that bFGF regulation of uPA expression and cell invasion is mediated by Ets-1. Kitange et al. (1999) described a similar phenomenon in a study of glioma cells. A recent study by Fujimoto et al. (2004) also showed a significant relationship between Ets-1 and invasive ovarian cancer (Davidson et al., 2003).

The serum concentration of bFGF was reported to range from 4 to 33 ng/mL in ovarian cancer patients (Di Blasio et al., 1995). However, the concentration of bFGF in the peritoneal cavity is much higher, since cellular injury or death may release more bFGF. Hence, we chose the following concentration gradient of bFGF: 0.1, 1, 10, 100 ng/mL. In this range, we found a dose-dependent response in the regulation of uPA expression.

Our experiments were performed in vitro, and the effects may not be the same in vivo, but the results enable us to reach a better understanding of the various mechanisms that regulate the metastatic process; furthermore, they can help to provide reasonable targets for the development of new antimitastatic therapies.

Declaration of interest: The authors report no conflicts of interest.

References


