BASIC RESEARCH

Expression of vascular endothelial growth factors A and C in human pancreatic cancer

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AIM: To study the expression of vascular endothelial growth factor A (VEGF-A) and VEGF-C and to determine whether the presence of VEGF-A and VEGF-C was associated with the clinicopathologic characteristics of pancreatic cancer.

METHODS: VEGF-A and VEGF-C mRNA transcripts were examined by Northern blot in 6 human pancreatic cancer cell lines and 8 normal pancreatic tissues and 8 pancreatic carcinoma specimens. The expression of VEGF-A and VEGF-C proteins was examined by Western blot in the tested cell lines and by immunohistochemical stain in 50 pancreatic carcinoma samples.

RESULTS: VEGF-A and VEGF-C mRNA transcripts were present in all the 6 human pancreatic cancer cell lines. Immunoblotting revealed the presence of VEGF-A and VEGF-C proteins in all the cell lines. Northern blot analysis of total RNA revealed 3.0-fold and 3.6-fold increase in VEGF-A and VEGF-C mRNA transcript in the cancer samples, respectively. Immunohistochemical analysis confirmed the expression of VEGF-A and VEGF-C in cancer cells within the tumor mass. Immunohistochemical analysis of 50 pancreatic cancer tissue samples revealed the presence of VEGF-A and VEGF-C immunoreactivity in 50% and 80% of the cancer tissue samples, respectively. The presence of VEGF-A in these cells was associated with larger tumor size and enhanced local spread ($\chi^2 = 6.690, P = 0.035<0.05$) but was not associated with decreased patient survival. However, the presence of VEGF-C in the cancer cells was associated with increased lymph node metastasis ($\chi^2 = 5.710, P = 0.017<0.05$), but was not associated with decreased patient survival. There was no correlation between the expression of VEGF-A and VEGF-C in the same cancer cells.

CONCLUSION: VEGF-A and VEGF-C are commonly overexpressed in human pancreatic cancer and may contribute to tumor growth and lymph node metastasis. There is no relationship between the expression of VEGF-A and VEGF-C in pancreatic cancer.

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Key words: Pancreatic cancer; Vascular endothelial growth factor-A; Vascular endothelial growth factor-C; Survival

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INTRODUCTION

Though the mortality rate of pancreatic cancer has decreased and the survival rate of pancreatic cancer patients is improved after surgery, the 1-year survival rate after diagnosis of pancreatic cancer remains below 20% and the 5-year survival rate is only 3%[1, 2]. One reason for this poor prognosis is the propensity of pancreatic cancer to metastasize. Pancreatic carcinoma is characterized by its aggressive local invasion of adjacent structures, perineural invasion and early lymph node and liver metastasis[3-5]. Lymph node metastasis and blood vessel invasion are the definitive factors of the prognosis after resection[6, 7]. Angiogenesis plays an important role in the development, growth and metastasis of carcinoma in mammals[8]. It is also important to consider the possible impact of lymphangiogenesis (development of lymphatic vessels) on cancer growth and metastasis[9]. In fact, tumor spread is dependent on both the angiogenic and lymphangiogenic systems. Many carcinomas metastasize through the lymphatic system, whereas others spread mainly hematogenously.

VEGF-A has a potent mitotic activity specific to vascular endothelial cells[9, 10], suggesting that VEGF-A
contributes to the pathologic angiogenesis associated with solid tumors. Recently VEGF-B to -E showing homology to VEGF-A have been identified\cite{11-13}. Among these, VEGF-C possessing approximately 30% identity with VEGF, has been shown to induce specific lymphatic endothelial proliferation and hyperplasia of the lymphatic vasculature by binding to its specific receptor flt-4 \cite{11, 12, 14-16}. VEGF-C may also be an important factor regulating the mutual paracrine relationship between tumor cells and lymphatic endothelial cells not only in primary tumor but also in adjacent normal tissue.

We have previously reported that VEGF-C and its receptor are overexpressed in pancreatic cancer\cite{17}. We now report the coexpression of VEGF-A and VEGF-C in human pancreatic cancer.

**MATERIALS AND METHODS**

**Cell culture**

ASPC-1, CAPAN-1, MIA-PaCa-2 and PANC-1 human pancreatic cancer cells were obtained from the American Type Culture (Rockville, MD, USA). COLO-357 and T3M4 human pancreatic cancer cells were a gift from R. S. Metzger at Duke University. Cells were grown in monolayer culture containing humidified 50% L. CO2 and 95% air at 37°C. ASPC-1, CAPAN-1 and T3M4 cells were grown in RPMI-1640 medium. COLO-357, MIA-PaCa-2 and PANC-1 cells were grown in Dulbecco’s modified Eagle’s medium (DMEM). Madia contained 10% fetal bovine serum, 100 U/mL penicillin and 100 mg/mL streptomycin.

**Tissues for Northern blot analysis**

Normal pancreatic tissue and pancreatic cancer tissue samples were frozen in liquid nitrogen and stored at -80°C for RNA extraction.

**Tissues for immunohistochemistry**

Pancreatic cancer tissue samples were obtained from 50 patients (35 men and 15 women; mean age, 64.3 ± 10.8 years; range, 48-81 years) undergoing surgery for pancreatic cancer. The tumors were classified as previously described\cite{18}. Histologically, there were 15 well, 31 moderately and 4 poorly differentiated adenocarcinomas. There were 0 stage I, 3 stage II, 9 stage III and 38 stage IV tumors. Tissue samples were fixed in 10% paraformaldehyde solution for 18-24 h and embedded in paraffin.

**Cloning and labeling of VEGF-A and VEGF-C probes**

Reverse-transcription polymerase chain reaction (RT-PCR) was carried out using RT-PCR kit according to the manufacturer’s instructions. Briefly, total RNA was extracted from human placenta using the acid guanidinium thiocyanate method. The prepared RNA (1 µg) was mixed with reverse transcription reagents in a total volume of 20 µL, incubated for 30 min at 42°C into first-strand cDNA. One µL cDNA was used for PCR amplification. The upstream primer for VEGF-A was 5’-CGGAATTCAACACAGTT TTCATAATAGA-3’ with an expected size of 1167 bp. Amplification conditions were denaturation at 94°C for 5 min, followed by 32 cycles of denaturation at 95°C for 1 min, annealing at 55°C for 1 min, extension at 72°C for 1 min. The PCR products were electrophoresed on 1.5% agarose gels and the target bands containing the specific cDNA were cut down and purified using an ultraclean DNA purification kit (Mo Bio Laboratories, Solana Beach, CA, USA) according to the manufacturer’s instructions. The accuracy of the RT-PCR products was confirmed by sequencing of cloned cDNA. The prepared cDNA probes were labeled by PCR-DIG labeling mix according to the manufacturer’s instructions.

**Northern blot analysis**

Total RNA was extracted from human tissue samples and human pancreatic cancer cell lines using the acid guanidinium thiocyanate method. RNA was fractionated on 0.8% agarose/2.2% formaldehyde gels, electrotransferred onto nylon membranes and cross-linked by ultraviolet irradiation\cite{19}. The blots were prehybridized in hybridization buffer for 1 h at 42°C, hybridized with the indicated DIG-labeled cDNA probes overnight at 42°C, and washed twice for 5 min with 2×SSC/SDS (0.1%) at room temperature, twice for 15 min with 0.1×SSC/SDS (0.1%) at 68°C. DIG-labeled nucleic acids were detected with CSPD according to the manufacturer’s instructions at room temperature. Briefly, after rinsed in washing buffer, the membrane was incubated for 30 min in blocking solution and anti-DIG-AP conjugate, respectively. Then the membrane was washed twice in washing buffer for 15 min and in detection buffer for 5 min and incubated for 5 min in CSPD solution. The damp membrane was sealed in a hybridization bag, incubated for 15 min at 37°C and exposed to radiographic film for 15 min at room temperature. Equivalent loading of RNA in each lane was confirmed by hybridizing the total RNA with a β-actin cDNA. All cDNAs were labeled with DIG as above. Densitometric analysis of the signals were performed with a gel imaging system.

**Immunoblotting**

Human pancreatic cancer cells were solubilized in lysis buffer containing 50 mmol Tris, 150 mmol NaCl, 1 mmol ethyleneglycoltetraacetic acid, 1% NP40, 1% sodium deoxycholate, 1 mmol sodium benadate, 50 mmol sodium fluoride, 2 mmol EDTA (pH 8.0), 100 µg/mL benzamidine, 50 µg/mL aprotinin, 10 µg/mL leupeptin, 10 µg/mL pepstatin A, and 1 mmol phenylmethylsulfonyl fluoride. Proteins were subjected to sodium dodecyl sulfatepolyacrylamide gel electrophoresis and transferred to Immobilon P membrane. Membranes were incubated for 90 min with highly specific anti-VEGF-A or anti-VEGF-C polyclonal antibody, then washed and incubated with a horseradish peroxidase-coupled secondary goat anti-rabbit antibody for 60 min. After washing, antibodies were visualized by enhanced chemiluminescence. Equivalent loading of protein in each lane was confirmed by hybridizing the total protein with β-actin.
Immunohistochemistry

The same anti-VEGF-A and anti-VEGF-C polyclonal antibodies were used for immunohistochemical analysis. Paraffin-embedded sections (4 µm) of pancreatic cancer and noncancerous pancreatic tissues were subjected to immunostaining using the streptavidin-peroxidase technique. After deparaffinization, antigen retrieval was performed by immersing the sections in 10 mmol citrate buffer (pH 6.0) and heated twice in a microwave oven (95 °C) for 5 min each time. Endogenous peroxidase activity was blocked by incubation with 1% hydrogen peroxide in distilled water for 10 min. Tissue sections were incubated with normal donkey serum for 20 min at room temperature and then incubated with polyclonal anti-VEGF-A antibody (2 µg/mL) or anti-VEGF-C antibody (2 µg/mL) in normal donkey serum for 2 h at room temperature. Bound antibodies were detected with biotinylated goat anti-rabbit IgG secondary antibody and streptavidin-peroxidase complex, using diaminobenzidine tetrahydrochloride as the substrate. The sections were counterstained with Mayer’s hematoxylin and incubated with nonimmunized rabbit IgG or without primary antibodies, which did not yield positive immunoreactivity.

Scoring was carried out by two independent observers blinded to the patient’s status. Positive staining was defined as the presence of VEGF-A and VEGF-C immunoreactivity in at least 10% of the cancer cells.

Statistical analysis

Differences in distribution of VEGF-A or VEGF-C were determined using the χ² test. Kaplan-Meier survival analysis was used to estimate survival time and log-rank test was used to compare differences in survival time between VEGF-A positive and negative groups or between VEGF-C positive and negative groups. P<0.05 was considered statistically significant.

RESULTS

Expression of VEGF-A and VEGF-C in human pancreatic cancer cell lines

Northern blot analysis of total RNA isolated from pancreatic cancer cell lines indicated that the 6 cell lines expressed the 4.1 kb VEGF-A transcript and 2.4 kb VEGF-C transcript (Figures 1A and 1B). Immunoblotting with a highly specific anti-VEGF-A or anti-VEGF-C antibody revealed an approximately Mr 43000 band (occasionally also a Mr 41000 band) of VEGF-A protein or Mr 55000 band (occasionally proteolytic forms of Mr 41000 molecules) of VEGF-C protein in all the cell lines (Figures 1 A and 1B). Both Mr 43000 band and Mr 41000 band corresponded to the VEGF165 isoform homodimer.

Expression of VEGF-A and VEGF-C mRNA in human pancreas

Northern blot analysis of total RNA isolated from human pancreatic tissue revealed the 4.1 kb VEGF-A transcript and 2.4 kb VEGF-C transcript (Figures 2A and 1B). Immunoblotting with a highly specific anti-VEGF-A or anti-VEGF-C antibody revealed an approximately Mr 43000 band (occasionally also a Mr 41000 band) of VEGF-A protein or Mr 55000 band (occasionally proteolytic forms of Mr 41000 molecules) of VEGF-C protein in all the cell lines (Figures 1 A and 1B). Both Mr 43000 band and Mr 41000 band corresponded to the VEGF165 isoform homodimer.
Expression of VEGF-A and VEGF-C protein in human pancreatic tissue

The same anti-VEGF-A or anti-VEGF-C antibody was used to localize VEGF-A or VEGF-C immunohistochemically. In the normal pancreatic tissue, moderate VEGF-A and VEGF-C immunoreactivity was present in the cytoplasm of endocrine islet cells, a weak immunoreactivity in some ductal cells within the small ductules and in a few acinar cells (data not shown). In the pancreatic cancer tissue, moderate to strong VEGF-A and VEGF-C immunoreactivity was present in the cytoplasm of many cancer cells (Figures 3A and 3B). Among the 50 pancreatic cancer cases, 25 (50%) were positive for VEGF-A and 40 (80%) for VEGF-C immunoreactivity in the cytoplasm of cancer cells.

Correlation between VEGF-A and VEGF-C expression and patient survival

χ² analysis indicated that the presence of VEGF-A in cancer cells was associated with a statistically significant increase in tumor size and local extension (T category). However, there was no correlation between the presence of VEGF-A in cancer cells and lymph node metastasis or distant metastasis or tumor stage or histological grade of cancers (Table 1). The presence of VEGF-C in cancer cells was associated with a statistically significant increase in the presence of lymph node metastasis (N category), but there was no correlation between the presence of VEGF-C in cancer cells and other factors (Table 1).

Table 1 VEGF-A and VEGF-C expression in relation to the clinicopathologic characteristics of 50 patients with pancreatic cancer

| Age (yr) | VEGF-A | VEGF-C |
|---------|--------|--------|
| +       | 64±10  | 65±10  |
| -       | 62±9   | 62±8   |
| Gender  |        |        |
| Male    | 18     | 28     |
| Female  | 7      | 12     |
| TNM     |        |        |
| T1      | 1      | 8      |
| T2      | 9      | 13     |
| T3      | 15     | 19     |
| N0      | 6      | 6      |
| N1      | 19     | 34     |
| M0      | 20     | 35     |
| M1      | 5      | 5      |
| Stage   |        |        |
| I       | 0      | 0      |
| II      | 1      | 2      |
| III     | 5      | 6      |
| IV      | 19     | 32     |
| Differentiated degree | | |
| well    | 10     | 14     |
| moderate| 13     | 23     |
| poor    | 2      | 3      |
| TNM: tumor node metastasis.

Table 2 Correlation between VEGF-A and VEGF-C expression in 50 patients with pancreatic cancer

| VEGF(-A) | VEGF(-C) | P  |
|----------|----------|----|
| +        | 21       | 19 |
| -        | 4        | 6  |

mean survival time of the VEGF-A negative and positive groups was 20.80 ± 15.00 months and 16.04 ± 9.00 months respectively. Thus, the survival time of patients with VEGF-A positive tumors was shorter. However, Kaplan-Meier analysis and log-rank test indicated that there was no significant difference in survival between these two groups (Figure 4A). The mean survival time of VEGF-C negative and positive groups was 30.7 ± 14.00 months and 15.4 ± 10.00 months respectively. The survival time of patients with VEGF-C positive tumors was shorter. But Kaplan-Meier analysis and log-rank test indicated that there was no significant difference in survival between these two groups (Figure 4B).

Correlation between VEGF-A and VEGF-C expression in pancreatic cancer

There was no correlation between VEGF-A and VEGF-C expression in pancreatic cancer (Table 2).

DISCUSSION

Recently, VEGF-B to -E showing homology to VEGF-A have been identified[11-13]. Among these, VEGF-C has
Expression of VEGF-A and VEGF-C was detected in normal and cancer tissues in the present study. Overexpression of VEGF-A has been reported in mammary, colorectal, liver, gastric and pancreatic malignancies. Expression of VEGF-A and its receptor KDR correlates with vascularity, proliferation and metastasis of colonic cancer, and with vessel count and tumor stage of gastric cancer as well as with growth of pancreatic cancer. Recent studies demonstrated that expression of VEGF-C in various human tumors, including lymphoma, melanoma, fibrosarcoma, squamous cell carcinoma, breast cancer, gastric cancer, colorectal cancer, pancreatic cancer and hepatocellular carcinoma. VEGF-C is expressed only in lymph node-positive breast cancer. Gastric cancer patients with a high expression of VEGF-C protein have a much poorer prognosis than those with a low VEGF-C expression. VEGF-C promotes lymph node metastasis of rectal cancer. Expression of VEGF-C and its receptor flt-4 correlates with lymphangiogenic process and metastasis of pancreatic cancer.

The present study showed that VEGF-A and VEGF-C expression was abnormal in pancreatic cancer in vivo. Northern blot analysis revealed a single 4.1 kb VEGF-A mRNA transcript and a single 2.4 kb VEGF-C mRNA transcript in all normal and pancreatic cancer tissue specimens, as well as 3.0-fold and 3.6-fold increase of VEGF-A and VEGF-C mRNA transcripts in pancreatic cancer tissue specimens, indicating that both VEGF-A and VEGF-C are overexpressed in pancreatic cancer. The distribution of VEGF-A and VEGF-C in normal and cancerous tissue specimens was different. In the normal pancreas, VEGF-A and VEGF-C were relatively abundant in endocrine islets, less frequently present in ductal cells and only occasionally present in acinar cells. In contrast, VEGF-A and VEGF-C were abundant in ductal-like cancer cells of pancreas. VEGF-A is a specific mitogen toward endothelial cells and VEGF-C is a specific mitogen to lymphatic endothelial cells, suggesting that VEGF-A has a potential to act in a paracrine manner on endothelial cells within the pancreatic tumor mass, leading to enhanced angiogenesis and tumor growth, while VEGF-C has a potential to act in a paracrine manner on lymphatic endothelial cells within the pancreatic tumor mass, leading to enhanced invasion to lymphatic vessels and lymph node metastasis. Cancer with active lymphangiogenesis could be predisposed to metastatic spread via the lymphatic system, resulting in a poor survival rate. It appears that tumor cells need to penetrate the lymphatic endothelium twice to translocate into the interstitium of a lymph node. Because lymphatic vessels have no tight junctions or continuous basal laminae, their penetration may only require tumor cell adhesion to the endothelium and transmigration through intracellular gaps between lymphatic endothelial cells. VEGF-A can increase microvascular permeability by enhancing the transmigration activity of vesicular-vacuolar organelles. VEGF-C and flt-4 may be components of a paracrine signaling network between cancer cells and the endothelium and may be involved in modifying the permeabilities of both blood and lymphatic vessels and metastasis.

Univariate analysis of the immunohistochemical data demonstrated that the presence of VEGF-A and VEGF-C in pancreatic cancer cells was associated with tumor size and lymph node metastasis respectively, indicating that VEGF-A and VEGF-C contribute to pancreatic tumor growth and lymphatic metastasis in vivo. Though the survival time was shorter in patients with VEGF-A and VEGF-C positive tumors, this correlation was not statisti-
cally significant. Niedergethmann et al. reported that the survival time of patients with pancreatic ductal carcinoma showing high expression of VEGF-A is significantly shorter than that of those with low or no expression of VEGF-A. Kurahara et al. reported that VEGF-C and VEGF-D expression in tumor cells is significantly associated with the 5-year survival rate of patients with pancreatic head cancer. In our study, VEGF-A or VEGF-C did not emerge as an independent prognostic parameter. Moreover, our previous study showed that the survival time of patients with VEGF-C positive tumors is shorter. Because the majority of patients in the present study had no detectable metastatic disease at the time of surgery, whether VEGF-A or VEGF-C expression correlates with enhanced propensity of pancreatic cancer to metastasize remains unknown. Further studies are needed.

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