Expression of angiogenesis inhibitors in human bladder cancer may explain rapid metastatic progression after radical cystectomy

WOLF-DIETRICH C. BEECKEN1,2, TOBIAS ENGL1,2, DIETGER JONAS2 and ROMAN A. BLAHETA2

1Clinic of Urology, Vitalicum®, 2Clinic of Urology, University of Frankfurt/Main, Frankfurt/Main, Germany

Received September 8, 2008; Accepted October 27, 2008
DOI: 10.3892/ijmm_00000125

Abstract. Angiogenesis is essential for tumor growth and progression. It has been demonstrated that the expression of angiogenesis stimulators (e.g. basic fibroblast growth factor and vascular endothelial growth factor) correlates to tumor progression in various human tumor types. Furthermore, endogenous angiogenesis inhibitors (e.g. angiostatin and endostatin) have been isolated from human tumor models and have been successfully used to treat tumors in mice and humans. In the present study, the expression of angiostatin, endostatin and thrombospondin-1 in four different human bladder carcinoma cell lines with different tumorigenic potential (MGH-U4, RT-4, RT-112 and UMUC-3) were investigated. A subset of bladder carcinoma patients demonstrates rapid metastatic progression after removal of the primary tumor, although no evidence of metastasis is diagnosed before the surgical procedure. A potential mechanism to explain this phenomenon is suggested. Angiostatin, endostatin and thrombospondin-1 was detected in the conditioned media of four human bladder cancer cell lines using Western blotting. Angiostatin was purified and amino acid sequenced via mass spectrometry. The biological activity of angiostatin was determined by proliferation assays using endothelial cells, smooth muscle cells and fibroblasts. Tumor characteristics of the four human bladder carcinoma models were investigated in vitro and in vivo. All the bladder carcinoma cell lines employed in this study produced two biologically active variants of the angiostatin molecule (38 and 49 kDa). Endostatin and thrombospondin-1 were only produced by the low malignancy MGH-U4 and RT-4 bladder carcinoma models. This study identified the expression of different antiangiogenic molecules in human bladder carcinoma. The expression of antiangiogenic molecules seems to be a characteristic of low malignancy bladder carcinomas. The sudden lack of expression of antiangiogenic molecules as a consequence of surgical removal of highly malignant bladder carcinomas may explain the rapid metastatic progression of a subset of bladder carcinomas.

Introduction

Angiogenesis, the formation of new blood vessels from pre-existing capillaries, is essential for tumor growth and progression (1). The formation of new blood vessels supports tumor growth by two different mechanisms: the supply of nutrients and oxygen and the paracrine stimulation of tumor growth through the release of factors produced by the endothelial cells (2). Angiogenesis is modulated by an assortment of stimulators and inhibitors of endothelial cell growth. This angiogenic balance can be shifted by the over-expression of stimulators or the reduction of inhibitors of endothelial cell growth, either of which will cause vessel growth.

Approximately 67,000 new cases of bladder cancer are diagnosed yearly and 14,000 patients die (3) of the disease. Bladder carcinomas (~95%) are transitional cell carcinomas, arising from the urothelium. The majority of bladder carcinomas (70 to 80%) demonstrates superficial growth, while 20 to 30% are invasive at first presentation (3). The gold standard therapy for organ confined, locally invasive bladder carcinomas is radical cystectomy. About 50% of these patients, preoperatively metastases-free, die from distant metastases within one to two years after the surgical procedure (4).

In 1979, Prout and colleagues recognized this clinical phenomenon and stated: ‘...either metastases are a direct cause of the operation or have been present before the procedure’ (5). It is our hypothesis that these metastases have already been seeded prior to surgery and that a factor secreted by the primary tumor is responsible for maintaining a dormant state, as long as the primary tumor is in place.

In 1994, the first tumor-derived angiogenesis inhibitor, angiostatin, was isolated (6). Angiostatin is a 38 kDa C-terminal cleavage product of plasminogen that demonstrates strong antiangiogenic activity (6). Angiostatin was isolated from the serum and urine of mice bearing a Lewis lung carcinoma with a low metastatic phenotype (LLC-LM). This mouse tumor exactly mirrors the clinical course of bladder carcinoma patients with rapid metastatic progression after surgery (6).

The LLC-LM is a mouse tumor variant that, when injected subcutaneously in mice, metastasizes to the lungs. However, as
long as the primary tumor is in place the metastases are dormant and only microscopically visible. Generally, the mice die of tumor burden within 21 to 28 days after tumor cell implantation. If the primary tumor is resected, the animals die of rapidly growing lung metastases within 15 to 21 days after primary tumor resection (6). O’Reilly and coworkers demonstrated that the angiogenesis inhibitor angiostatin, when supplied exogenously, maintained the lung metastases in a dormant state (6). In this study, we provide evidence that angiostatin and other angiogenesis inhibitors are produced by human bladder cancer. Furthermore, we show that angiostatin produced by human bladder cancer is biologically active and specific for the inhibition of endothelial cell proliferation. These findings may explain the rapid growth of dormant metastases after resection of invasive bladder carcinomas in humans.

Materials and methods

Cell lines. Four human bladder carcinoma cell lines were used to investigate the expression of angiostatin, endostatin and thrombospondin-1 in human bladder cancer. MGH-U4, a kind gift of Dr W.S. McDougal (Massachusetts General Hospital, Boston, MA) represents an atypical dysplasia of the urothelium (7). The RT-4 cell line (ATCC, Manassas, VA) is a model for a human papillary, well-differentiated bladder cancer (8). RT-112 (ATCC) is an invasive (pathological stage T2) moderately differentiated (grade 2/3) model of human bladder cancer (9) and UMUC-3 (ATCC) is derived from a deeply invasive (pathological stage T3) poorly differentiated (grade 3) human bladder tumor (10). All cell lines were grown in roller bottles (Costar Corning, Corning, NJ) maintained with trypsin and the amino acid sequences were analyzed by mass spectrometry as described by Shevchenko et al (11).

Proliferation assays. The purified angiostatin from the tumor cell conditioned media were tested in proliferation assays with bovine capillary endothelial cells, human foreskin fibroblasts and bovine vascular smooth muscle cells. Cells were trypsinized and suspended in 10 ml of DMEM containing 5% BCS and 1% GPS and 1 ng/ml of basic fibroblast growth factor (bFGF, Scios Nova, Mountain View, CA). Various concentrations of the purified angiostatin were added to the assay wells. After 72 h incubation, cells were trypsinized, resuspended in Hank’s balanced salt solution and counted with a Coulter particle counter (Coulter, Miami, FL). The results were compared to an angiostatin-free control in the presence of bFGF.
Growth of the human bladder carcinoma cell lines in immunodeficient mice. Tumor cells (1x10^6 cells in 0.1 ml of saline) of UMUC 3, RT-112, RT-4 and MGH-U4 were implanted under the dorsal skin of severe combined immunodeficient (SCID) mice (Mass. General Hospital, Boston, MA) (n=4/group). Tumor growth was monitored by repeated measurements in two dimensions. Tumor volume was calculated using the formula: Tumor volume = a x b^2 x 0.52, where 'a' is the longest diameter and 'b' is the shortest diameter. Tumor growth was monitored until the mean tumor volume of a group was 1000 mm^3.

Microvascular density of human bladder tumors in mice. To determine the microvascular density of tumors derived from the four different human bladder carcinoma cell lines, tumors were harvested at the end of the tumor growth experiment, fixed in cold-buffered formalin (Buffered Formalde-Fresh, Fisher Scientific, Fair Lawn, NJ) and embedded in paraffin. For immunohistochemistry, 5-μm-thick sections were placed on glass slides and incubated overnight with anti-mouse CD31 antibody (PharMingen, San Diego, CA) at a concentration of 1:250 at 4°C. Hereafter, the sections were incubated with an anti-rat secondary antibody (Vector, Burlingame, CA) at a concentration of 1:200 for 1 h at room temperature. Staining was enhanced by tyramide amplification (New England Nuclear, Boston, MA). 3-amino-9-ethylcarbazole (ACE, BioGenex, San Ramon, CA) was used as a chromogen and Gill's hematoxylin (Fisher Scientific) was used as counterstain (12). Vessel counts were performed in three of the human bladder carcinoma models, excluding the MGH-U4 cell line, which does not grow in mice. Multiple areas of highest vascular density (hot spots) were counted in three tumors of each cell line, as previously described (12). After the hot spots were identified using low magnification (x100), all vessels inside a grid (area 0.19 mm^2) were counted at a high magnification (x400). The mean value of all counts was taken

Figure 1. Western blot analysis of the angiogenesis inhibitors (A) angiostatin, (B) endostatin and (C) thrombospondin-1. Recombinant human angiostatin or endostatin proteins served as the controls.

Figure 2. Amino acid sequences of the angiostatin fragments identified by mass spectrometry after tryptic digestion. Amino acid sequences of the identified fragments (bold and underlined) match those of the 38 kDa (A) or 49 kDa (B) angiostatin molecules.
as the vascular density of tumors derived from the particular cell line.

Results

Identification of angiostatin, endostatin and thrombospondin-1. Two different forms of angiostatin were purified from conditioned media of the human bladder carcinoma cell lines. As determined by Western blot analysis with polyclonal antibodies to plasminogen, all of the investigated cell lines produced these two different angiostatin molecules (Fig. 1A). Amino acid sequencing by mass spectrometry revealed that the smaller angiostatin molecule (38 kDa) comprised the kringle domain 1 to 3 of the plasminogen molecule (Fig. 2A) while the larger angiostatin molecule (49 kDa) comprised the kringle domain 1 to 4 of plasminogen (Fig. 2B).

Endostatin was detected in three of the four human bladder carcinoma models (MGH-U4, RT-4 and RT-112). However, endostatin detected in the conditioned media of the RT-112 cell line ran a little longer on SDS gel which might indicate that this cell line produces a variant of endostatin (Fig. 1B).

Thrombospondin-1 was only detected in the MGH-U4 and RT-4 cell lines and ran at ~180 kDa. RT-112 and UMUC-3 showed no detectable thrombospondin-1 (Fig. 1C).

Biological activity and specificity of purified angiostatin from human bladder cancer models. Using different concentrations of the purified angiostatin, a dose response for the inhibitory activity on endothelial cells was determined. Compared to controls, where no angiostatin was added to the assay, 2 μg of the purified angiostatin inhibited endothelial cell proliferation by ~50% (Fig. 3A). This inhibition appeared to be endothelial cell-specific in that neither fibroblasts (Fig. 3B) nor vascular smooth muscle cells (Fig. 3C) were inhibited by the purified angiostatin when used at the same concentration.

Growth of human bladder tumors in mice and vascular density. Tumor growth was not detected in four mice 80 days after MGH-U4 cells were implanted subcutaneously. Since this cell line was derived from a urothelial dysplasia, it might lack certain characteristics for the development of malignant tumors in mice. However, the RT-4, RT-112 and UMUC-3 cell lines did produce tumors that grew to 1000 mm³. The mean tumor volume of 1000 mm³ was reached 80, 44 and 23 days after tumor cell injection, respectively (Fig. 4). Vascular density inversely correlates with the tumorigenic potential of the bladder tumors with the exception of MGH-U4.
In conclusion, the results of this study show that four different angiogenesis inhibitors, angiostatin, endostatin and thrombospondin-1, in a model of bladder carcinoma progression, consisting of four different bladder carcinoma cell lines demonstrating different levels of aggressiveness.

Angiostatin was expressed by all four models of bladder cancer (Fig. 1A). Two different variants of angiostatin were expressed, one comprises the kringle domain 1 to 3 of the plasminogen molecule and has a molecular weight of 38 kDa. The other molecule comprises the kringle domain 1 to 4 of plasminogen and runs at ~49 kDa on SDS-PAGE. Both angiostatin molecules have been described in the medical literature (25,26).

Endostatin was identified in three of the four human bladder carcinoma models. The most aggressive cell line, UMUC-3, did not exhibit any endostatin expression. However, the RT-112 cell line seemed to express a different variant of the endostatin molecule, that to our knowledge has so far not been described (Fig. 1B).

Thrombospondin-1 was expressed only in the lower aggressive MGH-U4 and RT-4 cell lines. The more aggressive RT-112 and UMUC-3 cell lines did not express any thrombospondin-1 (Fig. 1C).

It is well known, that the expression of angiogenesis stimulators correlates to tumor vascular density and tumor aggressiveness, also in bladder cancer (27). We hypothesize that the expression of angiogenesis inhibitors inversely correlates to the malignant potential and vascular density of tumor growth. By injecting mice following the different bladder carcinoma models we produced tumor progression curves and after harvesting the tumors determined the vascular density of the tumors using immunohistochemistry (Figs. 4 and 5).

The expression results of the three different antiangiogenic molecules and the total antiangiogenic activity of these tumors compared with tumor progression and vessel count support our hypothesis. This phenomenon might be explained by the concept that the net angiogenic output of a certain tumor, which is mirrored by its vascular density, relates more reliably to the sum of angiogenesis stimulators and inhibitors produced by the tumor cells than to the expression of a single angiogenic mediator.

Both angiostatin molecules which we detected in human bladder cancer have been described as biologically active (25,26). We show here that the total angiostatin expressed by human bladder carcinoma cells is inhibitory to endothelial cell proliferation (Fig. 3A) and does not show any inhibitory effect on the proliferation of human fibroblasts and bovine smooth muscle cells (Fig. 3B and C).

In conclusion, the results of this study show that four human bladder carcinoma cell lines, independent of differentiation, aggressiveness, rate of growth and vascular
density show different expression of the angiogenesis inhibitors angiostatin, endostatin and thrombospondin-1. The growth of primary tumors of these tumor models in mice appeared to correlate with the expression of the three different angiogenesis inhibitors. In two previous studies, we demonstrated that different models of human bladder cancer in mice, independent of vascular density and growth rate, were sensitive to treatments with angiogenesis inhibitors (12,28).

How the expression of angiogenesis inhibitors relates to the clinical course of bladder carcinoma patients who demonstrate rapid metastatic progression after removal of the primary tumor by cystectomy warrants further investigation. However, this laboratory study, for the first time, begins to explain this phenomenon by a molecular mechanism and could open an avenue for clinical application.

Acknowledgements

This study was supported by grants from the Ernst & Berta Grimmke Stiftung.

References