LYVE-1 immunohistochemical assessment of lymphangiogenesis in endometrial and lung cancer

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Aims/Methods: Normal and malignant pulmonary and endometrial tissues were analysed for lymphatic vessels to assess the process of lymphangiogenesis and its role at these sites, using specific immunostaining for LYVE-1 and the panendothelial marker CD31.

Results: Lymphatics were clearly demonstrated in some normal tissues (myometrium, bronchial submucosa, and intestinal submucosa), but not in others (endometrium and alveolar tissue). LYVE-1 positive lymphatic vessels were detected at the tumour periphery of endometrial and lung carcinomas, but not within the main tumour mass. Double staining for LYVE-1 and the MIB1 proliferation marker revealed a higher proliferation index in lymphatic endothelial cells at the invading front of endometrial carcinomas, compared with myometrial areas distal to the tumour. Lung and endometrial carcinomas did not have an intratumorous lymphatic network.

Conclusions: Although lymphangiogenesis may occur at the invading tumour front, incorporated lymphatics do not survive. Therefore, the dissemination of cancer cells through the lymphatics may occur by invasion of peripheral cancer cells into the adjacent normal lymphatics, or through shunts eventually produced at the invading tumour front as a consequence of active angiogenesis and lymphangiogenesis.

Materials and Methods

Formalin fixed, paraffin wax embedded tissues from 28 endometrial adenocarcinomas of the endometrioid cell type were retrieved from the archives of the department of pathology, Democritus University of Thrace, Alexandroupolis, Greece. All cases in the series were stage I disease so that lymphangiogenesis could be assessed in the early stages of endometrial tumour development. The patients were treated with total abdominal hysterectomy without lymph node dissection.

Similarly, surgical material from operable (T1,2–N0, stage 1) non-small cell lung carcinomas (19 adenocarcinomas and 9 squamous cell carcinomas) were selected for study.

Abbreviations: APAAP, alkaline phosphatase/antialkaline phosphatase; DAB, diaminobenzidine; MoAb, monoclonal antibody; PBS, phosphate buffered saline; TBS, Tris buffered saline; VEGF, vascular endothelial growth factor; VEGFR, vascular endothelial growth factor receptor.
49 squamous cell carcinomas) was also retrieved from the department of cellular science, University of Oxford, UK. In addition, 15 surgical samples from normal endometrium of various phases of the menstrual cycle and eight samples from normal lungs (from patients who underwent surgery for reasons other than cancer) were retrieved.

**LYVE-1 immunohistochemistry**

Mouse antihuman LYVE-1 clone 8C (IgG1 isotype) was raised against an immunoglobulin fusion protein of human LYVE-1 lymph vessel endothelial hyaluronan receptor. Sections were dewaxed and antigen retrieval was carried out by microwaving in Dako retrieval buffer (pH 6.0; Dako, Glostrup, Denmark), three times for four minutes each. Slides were incubated in phosphate buffered saline (PBS) with 5% human serum for five minutes. Peroxidase was quenched in PBS for 15 minutes. The primary antibody (in PBS plus 5% fetal calf serum) was applied for 45 minutes. After washing with PBS, sections were incubated with a secondary antirabbit horseradish peroxidase conjugated antibody (Kwik Biotinylated Secondary; Shandon-Upshaw, Pittsburgh, Pennsylvania, USA) for 15 minutes and washed in PBS. The colour was developed by a 15 minute incubation with diaminobenzidine (DAB) solution and sections were weakly counterstained with haematoxylin.

Normal tissue sections from the small intestine were used as positive controls. Normal mouse IgG was substituted for primary antibody as the negative control (same concentration as the test antibody).

**CD31 immunostaining**

The JC70 MoAb (Dako), recognising the CD31 panendothelial antigen (platelet/endothelial cell adhesion molecule) was used for microvessel and single endothelial cell staining on 5 μm thick paraffin wax embedded sections. We used the alkaline phosphatase/antialkaline phosphatase (APAAP) procedure. Sections were dewaxed, rehydrated, and predigested with protease type XXIV for 20 minutes at 37 °C. JC70 (1/50 dilution) was applied at room temperature for 30 minutes and washed in Tris buffered saline (TBS). Rabbit antimouse antibody at a dilution of 1/50 (vol/vol) was applied for 30 minutes, followed by application of mouse APAAP complex at a dilution of 1/1 (vol/vol) for 30 minutes. After washing in TBS, the last two steps were repeated for 10 minutes each. The colour was developed by 20 minutes of incubation with new fuchsin solution.

**Microvessel and lymphatic vessel counting**

Microvessel and lymph vessel counting was performed at the tumour invading front and in the inner tumour areas. The areas of highest vascularisation were chosen at low power (×100) and vessel counting followed on three chosen high power (×200) fields of the highest density. The microvessel density was the median of the vessel counts obtained in these three fields. Vessels with a clearly defined lumen or well defined linear vessel shape, but not single endothelial cells, were taken into account for microvessel counting.

**MIB1 immunostaining**

In eight selected endometrial carcinoma tissue samples (considered to have the highest LYVE-1 vessel reactivity), simple LYVE-1 and double LYVE-1/MIB1 staining was performed. The LYVE-1 staining procedure was performed as described above, whereas for double staining the MIB1 antibody (clone PRO224; YLEM, Rome, Italy) was applied immediately before the colour development, and incubated overnight. After washing with TBS, sections were incubated with a secondary rabbit antirabbit antibody (Kwik Biotinylated Secondary; Shandon-Upshaw) for 15 minutes and washed in TBS. Kwik streptavidin peroxidase reagent (Shandon-Upshaw) was applied for 15 minutes and the sections were again washed in TBS. The colour was developed by a 15 minute incubation with DAB solution and sections were weakly counterstained with haematoxylin. Because LYVE-1 does not stain nuclei, such a double staining, in direct comparison with parallel sequential sections stained with LYVE1 alone, allowed the detection of proliferating lymphatic endothelia.

The number of lymphatic vessels with and without nuclear reactivity was recorded in three ×200 optical fields in the invading tumour front and in three areas within the myometrium (away from the invading tumour area). The mean value was calculated.

**Statistical analysis**

Statistical analysis was performed using GraphPad Prism® 2.01 software (San Diego, California, USA). The unpaired two tailed t test was used for testing relations between categorical tumour variables (vascular and lymphatic densities compared as continuous variables). All p values are two sided and p values < 0.05 were considered to be significant.

**RESULTS**

**LYVE-1 staining in normal tissues**

The expression of LYVE-1 was examined in normal small intestine (used as staining control), and also in the normal uterus and the normal lung. Clear staining of submucosal lymphatics, but not of the adjacent blood vessels, was noted in the intestine. This exclusive expression of LYVE-1 in the lymphatics was also confirmed in the bronchial submucosa and the myometrium. In contrast, there were no LYVE-1 positive vessels in normal endometrial and alveolar tissues, whereas a dense CD31 positive vascular network was noted in both these tissues.

**LYVE-1 vessel staining**

In malignant endometrium, the mean microvessel density for each ×200 optical field (using the anti-CD31 MoAb) was 29 (range, 4–77) in the tumour invading front, and this was dramatically reduced in inner tumour areas (median, 11; range, 1–42). The mean microvessel density in lung carcinomas (assessed by anti-CD31 staining) was 29 (range, 6–64) in the invading front and 15 (range, 3–50) in inner tumour areas.

The tumorous vessels within the tumour body of endometrial and lung carcinomas were completely negative for LYVE-1. Myometrial lymphatics were clearly noted at the invading tumour front of some tumours (fig 1A), whereas in others the lymphatics were seen at a distance of about one ×200 optical field from the invading tumour front, suggestive of exclusion or even destruction of the lymph vessels by the invading tumour. Lymph vessel density at the invading tumour front ranged from 0 to 7 vessels (median, 2) for each ×200 optical field. In the myometrium adjacent to the tumour invading zone, the lymph vessel density ranged from 10 to 35 (median, 22), whereas in myometrial areas away from the tumour front it ranged from 32 to 41 (median, 35) (p < 0.0001), apparently reflecting variation in the normal regional distribution. In lung cancer, where alveolar tissue was deprived of lymphatics, LYVE-1 positive vessels were only noted in areas adjacent to entrapped bronchi. LYVE-1 positive areas, with a rather degenerating morphology, were occasionally noted among cancerous glandular structures or nests, and may correspond to incorporated lymphatic structures undergoing regression and breakdown.

**Lymphatic proliferation**

In eight cases of endometrial carcinoma, where the presence of lymphatics at the invading tumour front was confirmed,
double staining with MIB1 and LYVE-1 revealed that lymphatic endothelial cells were actively proliferating. In the normal myometrium, the median lymphatic vessel density was 36 (range, 31–40), and the median lymphatic vessel density with MIB1 nuclear staining was 0 (range, 0–2), giving a median percentage of 0% (0 of 36 lymphatics; p < 0.0001). In the myometrium proximal to the tumour edge, the median lymphatic vessel density was 23 (range, 13–35) and the median MIB1 positive lymphatic vessel density was 2 (range, 1–4), which gives a median percentage of 9% (two of 23 lymphatics). At the invading tumour edge, the median lymphatic density was 2 (range, 0–6), and was accompanied by intense proliferation of lymphatics (MIB1 nuclear staining: range, 0–4; median, 1; median percentage, 14%). Figure 1A shows LYVE-1 positive lymphatics and LYVE-1 negative blood vessels in the invading tumour area, whereas fig 1B shows MIB-1 nuclear staining of the vascular and lymphatic endothelium.

DISCUSSION

The production of angiogenic factors by tumour and stromal cells leads to endothelial cell migration, endothelial cell proliferation, and the formation of new tube-like structures, which sprout from preexisting capillaries of the host tissue adjacent to the invading tumour front. Newly formed and parental blood vessels are subsequently incorporated into the growing tumour mass, forming a unique vascular network composed of mature and immature vessels, which have an irregular distribution, discontinuities, shunts, and collapsed ill functioning branches. Arterial branches and disrupted veins communicate with this immature vascular network, and thereafter drain into the veins of the systemic circulation.

In contrast to the process of haemangioiogenesis, less information is at present available with regard to lymphangiogenesis. The growth factor VEGF-C, produced by several different tumours, was thought to induce proliferation of lymph vessels after binding to the VEGFR-3 receptor. However, because KDR, the receptor that binds to VEGF-A, VEGF-B, and VEGF-C, is also expressed by the lymphatic endothelium, these VEGFs could be lymphangiogenic as well. The formation of a new lymphatic network within tumours would greatly facilitate tumour growth by draining waste products of metabolism and promoting lymphatic dissemination of tumour cells.

“A striking absence of LYVE-1 positive lymphatics was noted within the tumour body of all cases studied, even though a CD31 positive vascular network was consistently noted.”

Using the lymph vessel specific LYVE-1 MoAb, which recognises an endocytic receptor for hyaluronan, we assessed whether lymphangiogenesis does indeed occur in certain human malignancies, namely endometrioid adenocarcinoma of the uterus, lung adenocarcinoma, and lung squamous cell carcinoma. Staining of normal lung, myometrium, and small intestine confirmed the specificity of LYVE-1 immunostaining of lymphatics. LYVE-1 positive lymphatics were also identified within the normal myometrium and the normal bronchial submucosa. LYVE-1 positive lymphatic vessels were not seen in the normal alveolar tissue or within the normal endometrium. These findings demonstrate the variable nature of the lymphatic network among different tissues.
A striking absence of LYVE-1 positive lymphatics was noted within the tumour body of all cases studied, even though a CD31 positive vascular network was consistently noted. In the invading edge of endometrial cancer two different patterns could be identified, namely: (1) LYVE-1 positive vessels entrapped by the spreading tumour, and (2) absence of LYVE-1 positive vessels. In lung cancer, such a feature was noted only at invading tumour edges adjacent to bronchi and not adjacent to alveolar tissue, where lymphatics were absent. The lack of immunohistochemically identifiable lymphatics within the main tumour mass suggests that even though lymphatics may be incorporated into the tumour mass, these cannot survive, but rather regress and vanish. Indeed, we showed previously that the survival of vessels in the inner tumour areas is problematic, and that the vessel density rapidly decreases in tumour areas 4–6 mm distant from the invading tumour front. Cancer sections stained for lymphatics did occasionally show LYVE-1 positive stromal areas, squeezed among malignant tissue, that may represent degenerative lymphatic structures incorporated into the tumour.

Assessment of the proliferation status of endothelial cells at the invading front of this type of endometrial carcinoma exhibiting a growth pattern with incorporating lymphatics showed that lymphatic endothelium (similar to non-lymphatic vascular endothelium) had a higher proliferation rate than the endothelium located at distal areas. This shows that lymphangiogenesis may occur in the invading tumour edge, where tumour interacts with the lymphatic rich normal tissue. However, this was not a constant feature and it may differ among tumours. The lack of intratumorous lymphangiogenesis in human melanomas suggested in a study by Fallowfield and Cook, where double staining for UEA1 and collagen IV was used to discriminate between blood and lymphatic vessels, argues in part with our findings. In contrast, two recent studies in melanoma (using the anti-LYVE-1 MoAb) confirmed the presence of focal areas exhibiting intratumorous lymphatic proliferation. Importantly, increased lymphatic density was linked with lymphatic proliferation even in areas of intense haemangiogenesis. This shows that there may be a growth pattern with incorporating lymphatics and intercommunicating non-lymphatic and lymphatic vessels, agrees in part with our findings. In melanoma, these cannot survive, but rather regress and vanish. Indeed, we showed previously that the survival of vessels in the inner tumour areas is problematic, and that the vessel density rapidly decreases in tumour areas 4–6 mm distant from the invading tumour front. Cancer sections stained for lymphatics did occasionally show LYVE-1 positive stromal areas, squeezed among malignant tissue, that may represent degenerative lymphatic structures incorporated into the tumour.

In conclusion, using specific immunostaining for LYVE-1 and the proliferation marker MIB1, lymphangiogenesis was detected in a subset of endometrial adenocarcinomas, and was located exclusively at the invading tumour edge. Such an event is difficult to confirm in lung carcinomas because alveolar tissue is deprived of vessels and such a process should be sought only in the proximity of tumour to bronchial structures. The lack of lymphatics within the tumour body, regardless of the presence or absence of peripheral lymphangiogenesis, suggests that tumour lymph drains directly into the immature intratumorous vascular network, and only in peripheral tumour areas is the lymph drained into the lymphatics of the surrounding normal tissue. Lymphatic dissemination of cancer cells may occur either by active invasion of cancer cells into host lymphatics adjacent to tumour tissue or by invasion of actively dividing and intercommunicating non-lymphatic and lymphatic vessels generated at the invading tumour edge.

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