Arterio-venous gradients of IL-6, plasma and serum VEGF and D-dimers in human cancer

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The circulating angiogenic factors vascular endothelial growth factor-A, interleukin-6 and the fibrin D-dimer fragment were measured in the mesenteric vein, the uterine vein, as well as in peripheral venous and arterial samples in 21 randomly selected patients with operable colorectal, ovarian and cervical carcinoma. In addition, immunohistochemistry for vascular endothelial growth factor-A and interleukin-6 was performed on colorectal tumours of such patients. Serum and plasma vascular endothelial growth factor-A were not significantly elevated in the vein draining the tumours, despite tumour cell expression of vascular endothelial growth factor-A. Serum vascular endothelial growth factor-A is therefore not all tumour-derived. In contrast, serum interleukin-6 was highly elevated in the draining veins in agreement with expression of interleukin-6 in the cytoplasm of tumour cells. In the megakaryoblastic cell line MEG-01, the expression of vascular endothelial growth factor-A was found to be regulated by interleukin-6. Thus, the higher platelet vascular endothelial growth factor-A load resulting in higher serum vascular endothelial growth factor levels in cancer patients may partly result from an interleukin-6 mediated up-regulation of the expression of vascular endothelial growth factor-A in the precursor of the platelet, i.e. the megakaryocyte. We also confirmed by immunohistochemistry that platelets adhere and aggregate on tumour endothelium. We propose that interleukin-6 indirectly promotes tumour angiogenesis through its up-regulation of the vascular endothelial growth factor-A load in platelets. In addition, the correlations found between peripheral venous interleukin-6 and peripheral venous fibrinogen and D-dimers levels, and the high D-dimer levels found in the draining vein of the tumour, in agreement with fibrin deposits found in the tumour stroma, suggest an important role for interleukin-6 in extra-vascular fibrinogen metabolism. Our results support a pivotal role for interleukin-6 in the intrinsic link between haemostasis and angiogenesis. This might be of importance in the development of anti-angiogenic agents based on interference with haemostasis.

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Haemostasis and angiogenesis are tightly regulated physiological processes involved e.g. in wound healing and deregulated in cancer development (Carmeliet and Jain, 2000). Both may be critical in three inter-related features of tumour progression, i.e. growth, invasion and metastasis (Carmeliet and Collen, 1998).

Vascular Endothelial Growth Factor-A (VEGF-A) is one of the most important positive mediators of angiogenesis. Tumour cells as well as stromal cells express VEGF-A (Leung et al, 1989; Veikkola et al, 2000). VEGF-A not only regulates endothelial cell migration, proliferation and survival, but is also known as Vascular Permeability Factor (VPF) because it enhances endothelial cell permeability even more potently than histamine (Dvorak et al, 1995). VEGF-A also induces tissue factor on endothelium and tumour cells, activating coagulation and fibrin formation (Mechtcheriakova et al, 1999). This is of primary importance for angiogenesis and tumour growth (Nagy et al, 1989; 1995). High serum levels of VEGF-A are predictive for tumour doubling time in patients with colorectal cancer. Elevated levels nearly normalise after resection of the primary tumour (Dirix et al, 1997). These observations indicate that circulating levels of VEGF-A mirror the ongoing growth process in tumours.

A nearly 10-fold higher serum than plasma VEGF-A level is found in both cancer patients as in healthy persons (Banks et al, 1998), indicating that circulating VEGF-A is mainly blood cell associated, in particular in platelets (Verheul et al, 1997; Salgado et al, 1999; Gunsilius et al, 2000). High serum VEGF-A levels are frequently encountered in cancer patients, the platelets of whom indeed have a higher VEGF-A content compared with healthy individuals; a prognostic significance is attributed to these levels (Salven et al, 1999; O’Byrne et al, 2000). How VEGF-A is regulated in platelets or in megakaryocytes in cancer patients is unclear, but VEGF-A in epithelial cells is regulated by IL-6 (Cohen et al, 1996). Platelet production is also regulated by IL-6 (Ishibashi et al, 1989).

The interstitial fibrin matrix, formed by VEGF-A-mediated tissue factor expression on tumour and endothelial cells, or by shedding of cancer pro-coagulant proteins (Mielicki et al, 1990), may further serve as a scaffold for endothelial cell migration and

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proliferation (Contrino et al, 1997). In addition, fibrinogen degradation products have pro-angiogenic properties (Thompson et al, 1985; Stirk et al, 2000). Remarkably, high fibrinolytic derivatives are one of the most frequent coagulation test abnormalities encountered in the oncological setting (Edwards et al, 1987).

In the present work we measured the circulating factors VEGF and IL-6 in veins draining the tumour in order to provide direct evidence of tumour secretion of these factors. We further investigated how platelets of cancer patients acquire a higher VEGF-A content. We then analysed whether circulating fibrin degradation products, D-dimers, are derived from intra-tumoural activation of haemostasis. We more specifically focused on the role IL-6 could have in both increasing the VEGF-content in platelets as well as in mediating fibrin formation in tumours.

MATERIALS AND METHODS:

Patients
Two patients with cervical carcinoma (FIGO Ib), five with ovarian carcinoma (FIGO III), two of them with extensive disease (FIGO IIC), and 14 with colorectal carcinoma (Dukes A – C), of which four had extensive disease at the moment of surgery (Dukes D), were scheduled for elective abdominal surgery. Exclusion criteria were: previous, concurrent or non-recteable cancer, local or systemic inflammatory disease. The following staging procedures were performed: chest X-ray, bone scintigraphy, CT-scan and echography of the liver. All patients received 0.3 mg fraxiparin subcutaneously for prevention of deep venous thrombosis (DVT). All routine precautions for prevention of DVT were taken.

Tissue
As part of the normal surgical resection procedure, of the main tumour mass, a representative, full cross section of the tumour sample surrounded by adjacent mucosa was taken from all patients involved. The tissue was fixed in buffered formalin and paraffin-embedded. Five µm sections were cut and mounted on poly-L-lysine coated slides. Successive sections were used for immuno-staining.

Blood coagulation tests
During the surgical procedure, blood sampling occurred simultaneously in the mesenteric or uterine vein as well as in a platelet pellet. Positive controls included staining of platelets from paraffin-embedded normal endometrium. Positive controls consisted of the omission of the primary antibody, the addition of an irrelevant primary antibody directed against glucose-oxidase from Aspergillus Niger species (this enzyme is not inducible in mammalian cells) and paraffin embedded normal endometrium tissue. Positive controls included staining of platelets in paraffin embedded thrombus and placenta with prominent fibrin deposits as well as in a platelet pellet.

Cell culture experiments
We used the human megakaryoblastic cell line MEG-01 to evaluate whether IL-6 is able to modulate the expression of VEGF in megakaryocytes. Ogura and colleagues established the MEG-01 cell line from a patient in a megakaryoblastic crisis of chronic myelogenous leukaemia. This cell line possesses many megakaryocytic specific markers and does not possess any marker of B-cells, T-cells and myeloid cells (Ogura et al, 1985). The MEG-01 cell line has the receptor for IL-6 and blocking IL-6 with anti-IL-6 antibodies did not have any significant effects on cell growth (Kellar et al, 1995). In order to confirm this we analysed with the Alamar blue assay (Trek Diagnostics, USA) whether alterations in cell proliferation occur after adding IL-6 (10 and 100 ng ml⁻¹) to the culture medium. The cells were cultured in Iscove’s modified Dulbecco’s medium (IMDM), with 10% foetal calf serum and antibiotics. The experiments were initiated by seeding 5 × 10⁴ cells well⁻¹ in a 24-well plate and allowed to proliferate for 24 h in 10% FCS medium before adding 10 or 100 ng ml⁻¹ IL-6 in addition with 1/10 diluted Alamar blue

IL-6 Deparaffinisation and quenching of endogenous peroxidase was performed. The primary 1/10 diluted polyclonal anti-IL-6 antibody (Genzyme Co., Cambridge, MA, USA) was then applied on sections of the colorectal carcinomas followed by a biotinylated secondary antibody. Streptavidin conjugated with peroxidase was subsequently applied on the sections (DAKO, Glostrup, Denmark). Finally, amino-ethylcarbamazime (AEC) was used as a substrate reagent. Negative controls included the omission of the primary antibody and the use of an irrelevant primary antibody.

Fibrin After deparaffinisation and quenching of endogenous peroxidase, protein digestion was performed with protease. Slides were placed on a Ventana NexES (Ventana Medical Systems, Inc.) automated immunostainer and stained with a 1/100 diluted monoclonal T2 G1 antibody (Acc. Chem. Sci. Corp., USA) overnight at 4°C. This antibody is specific for the peptide Bz 15-42 of fibrinogen and for fibrin II. Fibrin II is obtained after cleavage of fibrinogen by thrombin at the bond Az 16-17 and Bf 14-15. Fibrinopeptides A and B are released after cleavage by thrombin. Consequently, intact fibrinogen was not stained (Kudryk and Bini, 2000). The staining was completed with a Vector Basic DAB detection kit (Ventana Medical Systems, Inc.). Negative controls included the omission of the primary antibody. Paraffin embedded placental tissue with peri-villous fibrin deposits was used as a positive control.

Platelets Immunohistochemistry for platelet glycoplasticin was performed to demonstrate adherence and aggregation of platelets on endothelium in colorectal tumours. Glycoplasticin is a fragment of the platelet membrane glycoprotein Ib (Steinberg et al, 1987). Deparaffinisation and quenching of endogenous peroxidase was performed. Normal rabbit serum (NRS), diluted 1/5, was applied, followed by 1/20000 diluted primary mouse anti-human glycoplasticin (clone: G28E5) antibody (kind gift from Dr Hoylaerts). Subsequently, the 1/400 diluted, secondary biotinylated rabbit anti-mouse antibody (DAKO, Glostrup, Denmark) was applied. The immune reaction was completed with ABC-streptavidin detection system and DAB (diaminobenzidine) as chromogen. The slides were counterstained with haematoxylin and mounted. Negative controls included the omission of the primary antibody, the addition of an irrelevant primary antibody directed against glucose-oxidase from Aspergillus Niger species (this enzyme is not inducible in mammalian cells) and paraffin embedded normal endometrium tissue. Positive controls included staining of platelets in paraffin embedded thrombus and placenta with prominent fibrin deposits as well as in a platelet pellet.

Angiogenic cytokines Blood sampling was performed as mentioned above. Serum collection and VEGF-A₁₆₅ and IL-6 measurements were performed as mentioned previously (Salgado et al, 1999). Plasma VEGF-A₁₆₅ was measured with an ELISA-kit of R&D systems (R&D systems, Minneapolis, MN, USA) on trisodium citrate-anticoagulated collected blood. For the ELISA-assay of VEGF-A₁₆₅ no cross-reactivity with other members of the VEGF-A family is documented. Samples were assayed in duplicate. Within assay variability has been tested before (Salgado et al, 1999).

Immunohistochemistry

VEGF Immunohistochemistry was performed on colorectal cancer tissue as described previously with the anti-VEGF-A VG-1 antibody that stains VEGF-A efficiently on paraffin sections (Turley et al, 1998). This antibody recognises the 121, 165 and 189 isoforms of the VEGF-A protein.
medium. Fluorescence intensities, as a measure of reduction of the Alamar blue medium by proliferating cells, was measured every 6 h with a Cytofluor (Cytofluor2™ 2300, Millipore).

In subsequent experiments the IL-6 receptor (IL-6R) was blocked for 48 h with a molar excess of monoclonal anti-IL-6R antibody (20 µg ml⁻¹)(R&D Systems, Minneapolis, MN, USA). Cells without antibody served as negative controls.

Cell viability was assessed using Trypan blue exclusion dye. Conditioned medium was collected and ELISA for VEGF-A was performed. Experiments were performed in triplicate.

Ethical committee

The local Ethical Committee approved this study and informed consent was obtained from all patients involved.

Statistical analysis

Statistical analysis was performed with Graphpad Prism, version 2.0 (Graphpad Software, Inc.). Half the detection limit value of the patient samples was used for statistical analysis in case the measured values did not reach the detection limit of the assay. Detection limits were 0.7 pg ml⁻¹ and 9 pg ml⁻¹ for the IL-6 and VEGF-A assays, respectively. Comparisons of continuous variables were performed with the Mann–Whitney U-test. A paired students t-test was used to evaluate observed differences in the cell culture experiments. The relation between continuous variables was analysed with a Spearman rank correlation analysis. Means± standard deviations are given. A P-value <0.05 was considered to be significant.

RESULTS

Analysis of VEGF, IL-6 and D-dimers in patients without distant metastasis

In order to verify whether there was a gradient for VEGF, IL-6 and D-dimers between the afferent and efferent blood flow in a tumour we explicitly used samples from patients without distant metastasis (n=15) as VEGF, IL-6 and D-dimers produced by metasteses could mask observed gradients. We also assumed that the values in the brachial artery samples are comparable with mesenteric/uterine vein compared with arterial samples (17.66±7 pg ml⁻¹ and 354.8±99.8 pg ml⁻¹ for the venous peripheral and tumoural draining vessels respectively (n=15). No statistical differences were observed between VEGF-A in the arterial and efferent draining vein respectively (P=0.79; Table 1).

Mean D-dimer levels in the mesenteric/uterine vein were 1706±260 ng ml⁻¹, whereas in the brachial artery a mean value of 798.1±638.1 ng ml⁻¹ was found (P=0.08; Mann–Whitney U-test; Figure 2D; Table 1; n=15). As already mentioned, both aPTT and PT values were in the normal range indicating that the high D-dimer levels encountered in the draining veins are not due to artifactual activation of coagulation during sampling.

The major substrate of D-dimers, fibrin, was encountered both in the stromal as well as on the luminal and/or abluminal side of endothelial cells in the majority of cases examined (Figure 1D; E; n=14). No spatial co-localisation between platelet and fibrin staining was evident.

Platelets (n=15) Mean platelet count in the brachial vein, brachial artery, and mesenteric/uterine vein were 254.4±60.1×10³ µl⁻¹, 253±68.2×10³ µl⁻¹ and 241.9±60×10³ µl⁻¹. No statistical differences were observed between platelet counts in the artery branchialis and efferent draining vein (Table 1; n=15). Activated Partial Thromboplastin Time (aPTT) en PT were in all samples in the normal range indicating that no artifactual coagulation occurred in the samples taken. To merely verify whether platelet adherence and aggregation occurs in tumours we performed immunohistochemistry using an antibody directed at the platelet membrane protein glycoplatin. Positive staining indicates that these platelets are activated since glycoplatin is translocated to the platelet membrane upon activation.

The specificity of the glycoplatin platelet antibody was demonstrated in the positive controls (platelets in a platelet pellet, in a thrombus and in placental fibrin deposits), where clear staining of platelets was shown. Tumour cells, the fibrin matrix in the thrombus, white blood cells as well as in the placental fibrin deposits stained negative. Platelet adherence and aggregation on endothelial cells and extra-vasation of platelets was observed in all tumours (Figure 1C; n=14).

Analysis of VEGF, IL-6 and D-dimers in patients with distant metastasis

In order to find evidence for IL-6 and D-dimer secretion by metastatic cells we compared arterio-venous differences in the primary tumour of patients with metastasis (n=6). If metastatic cells would secrete both IL-6 as well as D-dimers, arterial levels of these would increase and would consequently mask or lessen the arterio-venous gradients as were observed in non-metastatic patients. Mean age of all patients with metastasis involved is 62.8±9.2 years.

Interleukin-6

All patients with colorectal cancer had extensive IL-6 labelling in the tumour cell compartment (Figure 1B; n=14). A 2.5-fold higher serum level of IL-6 was observed in the mesenteric/uterine vein compared with arterial samples (17.66±8.3 vs 7.06±9.8 pg ml⁻¹; P<0.001, Mann–Whitney U-test; Figure 2C; n=15; Table 1). Only one patient had an IL-6 value below the detection limit of the assay.
patients without metastatic disease (7.06±9.8 pg ml\(^{-1}\)). None of the patients had IL-6 values below the detection limit of the assay. Analogous results were obtained when comparing peripheral venous samples (12.95±15.5 pg ml\(^{-1}\)). When analysing arteriovenous differences in these patients with a mean level of IL-6 of 80.20±129.9 pg ml\(^{-1}\) in the mesenteric/uterine vein, a \(P\)-value of 0.09 compared with \(P\leq0.0001\) in the non-metastatic group was found.

D-dimers

Arterial levels of D-dimer concentrations of patients with advanced disease (\(n=6\)) were compared with non-metastatic patients (\(n=15\)). A three-fold higher arterial D-Dimer level (2372±1741 ng ml\(^{-1}\)) was found in advanced metastatic patients compared with those patients without metastatic disease (798.1±638.1 ng ml\(^{-1}\)). Analogous results are obtained when comparing peripheral venous samples (2575±1870 ng ml\(^{-1}\)). When analysing these arterial levels with the D-dimers measured in the veins draining the primary tumour in these patients (\(n=6\); 3066±1741 ng ml\(^{-1}\)), a \(P\)-value of 0.69 compared with \(P=0.08\) in patients without metastasis was yielded.

Effect of IL-6 on VEGF-A secretion in the MEG-01 cell line

Cell viability exceeded 98% in all circumstances. The mean IL-6 and VEGF-concentrations (pg ml \(^{-1}\)×10\(^3\) cells) in the conditioned medium of the control cells were 2.84±2.084 pg ml \(^{-1}\)×10\(^3\) and 6.093±2.651 pg ml \(^{-1}\)×10\(^3\) cells, respectively.
Incubating the MEG-01 cells for 48 h with 10 or 100 ng ml\(^{-1}\) IL-6 yielded mean VEGF-values of 5.34 ± 1.6 pg ml\(^{-1}\) and 4.72 ± 0.6 pg ml\(^{-1}\) \(\times\) 10\(^3\) cells, which were not significantly higher than the VEGF concentration in the conditioned medium of control wells.

We then analysed with the Alamar blue assay whether blocking of the IL-6 receptor resulted in a significant inhibition of cell proliferation. No significant changes in cell proliferation were observed (data not shown).

The median ± s.e. (standard error) VEGF concentration after 48 h incubation with an IL-6R blocking antibody was 3.44 ± 1.44 pg ml\(^{-1}\) \(\times\) 10\(^3\) compared with 6.850 ± 1.082 pg ml\(^{-1}\) \(\times\) 10\(^3\) cells for the control cells (\(P=0.03\)) (Figure 4).

**DISCUSSION**

Interleukin-6 is a cytokine involved in haematopoietic cell differentiation, in the regulation of the acute-phase response, in immune

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**Table 1** Comparison between coagulation and angiogenesis factors in arteria brachialis, peripheral vein and mesenteric/uterine vein in patients with non-metastatic cancer (n=15).

| Factor            | Arterial       | Peripheral venous | Efferent        | \(P\)-value* |
|-------------------|----------------|-------------------|-----------------|--------------|
| Platelets         | 253.0 ± 68.2   | 254.4 ± 60.1      | 241.9 ± 60.0    | 0.71         |
| Fibrinogen        | 357.0 ± 76     | 354.1 ± 87        | 354.8 ± 99.8    | 0.79         |
| D-dimers          | 798.1 ± 638.1  | 737.3 ± 448.3     | 1706 ± 2603     | 0.08         |
| Plasma VEGF       | 30.0 ± 15.27   | 24.6 ± 24.6       | 19.92 ± 9.22    | 0.07         |
| Serum VEGF        | 345.2 ± 269.5  | 373.6 ± 403.1     | 482.7 ± 371.1   | 0.77         |
| IL-6              | 7.06 ± 9.8     | 6.936 ± 9.7       | 17.66 ± 18.3    | <0.0001      |

Platelets: \(\times\) 10\(^3\) ml\(^{-1}\); Fibrinogen: mg dl\(^{-1}\); D-dimers: ng ml\(^{-1}\); VEGF and IL-6, pg ml\(^{-1}\). Efferent: mesenteric or uterine vein values. *\(P\)-value: differences between arterial and efferent values.

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**Figure 2** Observed gradients between arterial samples (art.) and tumour efferent blood (EV: efferent venous: mesenteric and uterine venous) for (A) serum VEGF-A, (B) plasma VEGF-A, (C) Interleukin-6 and (D) D-dimers in patients with localised cervical, ovarian and colorectal cancer (n=15).
regulation and is produced by epithelial, bone marrow and blood borne cells (Borden and Chin, 1994; Simpson et al, 1997; Hirano, 1999). Several studies have elaborated on the prognostic impact of circulating IL-6. A poor prognosis has been consistently associated with high levels of serum IL-6 in patients with renal cell, ovarian, and prostate cancer (Ljungberg et al, 1997; Tempfer et al, 1997; Nakashima et al, 2000). We have shown that high serum levels of IL-6 have independent prognostic significance in a series of 96 patients with metastatic breast cancer (data not shown). De Vita et al (2001) have recently demonstrated, in univariate analysis, that high circulating IL-6 levels are associated with reduced overall survival and reduced time to disease progression in patients with gastrointestinal cancer.

Our results demonstrate that circulating IL-6 is mainly derived from the tumour in patients with colorectal, cervical and ovarian cancer (Figure 2C). Moreover, arterial and peripheral venous circulating levels of IL-6 are about two-fold higher in patients with disseminated disease compared with arterial and peripheral venous IL-6 levels of patients with localised disease. These higher levels diminish the arterio-venous differences in the primary tumour of the patients with metastatic disease. This suggests that metastasised cells also produce and secrete IL-6. Nevertheless, to definitively prove that metastases secrete IL-6, gradient studies need to be performed. This study, however, will be difficult to perform due to technical and ethical reasons.

We have also demonstrated in this study that neither serum or plasma VEGF-A levels are significantly elevated in the vein draining the tumour, despite a high tumour cell expression of VEGF-A (Figure 2A,B). This contradicts the common notion that serum VEGF-A is mostly derived from spill over of tumour cell produced VEGF-A in patients with cancer. Landriscina et al (1998) also did not find significantly higher serum levels of VEGF-A in mesenteric blood compared with peripheral blood in patients with colorectal cancer. This finding was recently confirmed in patients with rectal cancer (Werther et al, 2001). These observations suggest that serum VEGF-A might be derived from other sources. However, it might be possible that continuous spilling of tumour-derived VEGF in the circulation might lead, especially considering the half-life of VEGF-A of about 30 min (Eppler et al, 2002), to equivalent levels of venous and arterial VEGF-A.

Several groups have confirmed the role of platelets in the storage of circulating VEGF-A (Verheul et al, 1997; Banks et al, 1998; Salgado et al, 1999; Gunsilius et al, 2000; Salgado et al, 2001). In a previous study we have shown that serum IL-6 correlates not only with the platelet count (illustrating its thrombopoietic property), with serum VEGF-A (illustrating the importance of IL-6 in the regulation of VEGF-A), but also with the VEGF-A load in platelets (Salgado et al, 1999). Since thrombopoietic cytokines, such as thrombopoietin (TPO) and interleukin-3 (IL-3), enhance VEGF-A expression in megakaryocytes, and, as is the case for TPO, in parallel with megakaryocytic differentiation, we hypothesised that IL-6 derived from primary tumour cells as well as
from disseminated tumour cells might enhance VEGF-A expression in platelet progenitors much in the same manner (Imai et al., 1991; Mohle et al., 1997; Bobik et al., 1998; Salgado et al., 1999).

Our in vitro data demonstrate that the endogenous production of IL-6 in a megakaryocytic cell line, MEG-01, is substantially high. The observation that adding IL-6 did not have any effect on VEGF-A production, might suggest that the IL-6 receptors are already maximally stimulated and saturated. We therefore opted for blocking this autocrine pathway with a molar excess of IL-6 receptor blocking antibody. Our results indicate that VEGF-A production is, at least partly, regulated by IL-6 (Figure 4).

We elaborate on an important role of tumour cell produced IL-6 in the enhanced platelet VEGF-A load of cancer patients explaining herewith the previously found correlation between serum IL-6, VEGF-A and the VEGF-A load in platelets. Moreover, as we did not find higher serum or plasma VEGF-A levels in the efferent blood vessel of tumours, the role of bone marrow, thus megakaryocyte-derived platelet VEGF-A, may contribute more to the total blood vessel of tumours, the role of bone marrow, thus megakaryocyte-derived platelet VEGF-A, may contribute more to the total

veins exhibit an enhanced, VEGF-A induced, vascular permeability and kinetics of extravasation of several circulating proteins are enhanced in cancer patients, e.g. fibrinogen (Dvorak et al., 1984). Recent evidence, however, demonstrates that tumour cells themselves produce fibrinogen, which is consequently deposited in the extra-cellular matrix (Rybarczyk and Simpson-Haidaris, 2000).

Fibrinogen is, in the pro-coagulant environment of the tumour stroma, rapidly converted to fibrin. Interleukin-6 is able to induce tissue factor expression on e.g. monocytes and may therefore contribute to the intra-tumoural activation of coagulation (Neumann et al., 1997). Measuring the fibrin degradation product D-dimer in the efferent vessel draining the tumour indicates that circulating D-dimer levels are also derived from intra-tumoural fibrinogen metabolism (Figure 2D). The borderline significance found (P=0.08) may probably gain significant values if a higher number of patients are involved and may also partly be explained by the involvement of patients with different histology in our study. However, our results confirm previous findings of intra-tumoural activation indicated by measuring D-dimers in veins draining lung tumours (Kalweit et al., 2000). The higher arterial and peripheral venous D-dimer levels encountered in patients with extensive disease compared with non-metastatic patients, masking observed arterio-venous gradients compared with non-metastatic patients, suggest that fibrinogen turnover also occurs in metastatic deposits. We have detected fibrin in metastatic skin deposits from patients with breast cancer (unpublished data). The importance of plasma D-dimers is illustrated by our previous finding that high circulating D-dimer levels correlate with progression kinetics (P<0.0001), with tumour volume (P<0.0001) and with reduced overall survival in patients with metastatic breast cancer (Dirix et al., 2002).

In conclusion, this study provides further evidence that IL-6 may be pivotal in haemostasis and angiogenesis associated with cancer.

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