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Alterations in NO- and PGI₂- dependent function in aorta in the orthotopic murine model of metastatic 4T1 breast cancer: relationship with pulmonary endothelial dysfunction and systemic inflammation

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Abstract

Background: Patients with cancer develop endothelial dysfunction and subsequently display a higher risk of cardiovascular events. The aim of the present work was to examine changes in nitric oxide (NO)- and prostacyclin (PGI₂)-dependent endothelial function in the systemic conduit artery (aorta), in relation to the formation of lung metastases and to local and systemic inflammation in a murine orthotopic model of metastatic breast cancer.

Methods: BALB/c female mice were orthotopically inoculated with 4T1 breast cancer cells. Development of lung metastases, lung inflammation, changes in blood count, systemic inflammatory response (e.g. SAA, SAP and IL-6), as well as changes in NO- and PGI₂-dependent endothelial function in the aorta, were examined 2, 4, 5 and 6 weeks following cancer cell transplantation.

Results: As early as 2 weeks following transplantation of breast cancer cells, in the early metastatic stage, lungs displayed histopathological signs of inflammation, NO production was impaired and nitrosylhemoglobin concentration in plasma was decreased. After 4 to 6 weeks, along with metastatic development, progressive leukocytosis and systemic inflammation (as seen through increased SAA, SAP, haptoglobin and IL-6 plasma concentrations) were observed. Six weeks following cancer cell inoculation, but not earlier, endothelial dysfunction in aorta was detected; this involved a decrease in basal NO production and a decrease in NO-dependent vasodilatation, that was associated with a compensatory increase in cyclooxygenase-2 (COX-2)- derived PGI₂ production.

Conclusions: In 4T1 metastatic breast cancer in mice early pulmonary metastasis was correlated with lung inflammation, with an early decrease in pulmonary as well as systemic NO availability. Late metastasis was associated with robust, cancer-related, systemic inflammation and impairment of NO-dependent endothelial function in the aorta that was associated with compensatory upregulation of the COX-2-derived PGI₂ pathway.

Keywords: Metastasis, Endothelial dysfunction, Inflammation, 4T1

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Background

The vascular endothelium, considered the body's largest autocrine/paracrine/endocrine organ, responds to various physical and chemical stimuli and maintains vascular homeostasis. Endothelial dysfunction, characterized by impaired production of vasoprotective endothelial mediators such as nitrite oxide (NO) and excessive activity of pro-oxidant, pro-thrombotic, pro-inflammatory mediators, plays a key role in the development of atherosclerosis and its cardiovascular complications, as well as in other cardiovascular diseases [1–3]. Cardiovascular complications observed in cancer patients and cancer survivors [4, 5] are also associated with the development of endothelial dysfunction that displays a similar phenotype to that described in patients with *atherothrombosis* [6].

In target organs for metastatic spread, endothelial inflammation may promote cancer metastasis. The endothelium acts as a barrier to cancer cell migration from blood to tissue. It has been proposed that the migration and invasion of circulating cancer cells into distant tissues is supported by the same mechanisms as those involved in leukocyte recruitment [7]. In fact, tumour cells appear to respond to the chemokines and cell adhesion molecules engaged in leukocyte trafficking and endothelial inflammatory response [8, 9]. Interestingly, recent *in vitro* studies have demonstrated that factors released from dysfunctional endothelium activate NF- κ B and STAT3 signalling pathways within cancer cells, and promote their invasiveness *in vitro* [10]. At the same time, vasoprotective endothelial mediators such as for example prostacyclin (PGI₂) display anti-metastatic activity, further supporting a crucial relationship between endothelial function and cancer metastasis [11, 12].

Metastases development is therefore related with changes of the endothelial phenotype in the affected organs (e.g. lung, liver) that contribute to the pathophysiology of metastasis. In addition, development of systemic endothelial dysfunction (e.g. in the conduit arteries) may contribute to cardiovascular complications in cancer patients. Obviously, the mechanisms mediating local changes in endothelial phenotype at metastatic sites and those involved in the development of systemic endothelial dysfunction may be different.

Among various available mouse models of cancer metastasis, the most appropriate for studying local and systemic endothelial response would be models, in which cancer cells are transplanted into the tissue of their origin (e.g. mammary gland, prostate or intestine), so that they form tumours that develop and progress from primary lesions to metastases in their natural microenvironment, i.e. as orthotopically-inoculated cancer cells syngrafted into immune-competent mice.

In the present study, we took advantage of the spontaneously metastasizing 4T1 murine mammary gland

carcinoma model to examine the alterations in endothelial NO- and PGI₂-dependent function in aorta that accompany metastatic expansion. The prolonged disease progression in a model established in our laboratory allowed us to identify a temporal relationships between the development of local, pulmonary inflammation, alterations in pulmonary endothelial function, systemic inflammation and endothelial dysfunction in conduit vessel (aorta) in relation to primary tumour growth and lung metastases. We showed that an early decrease in pulmonary and systemic NO availability coincided with the development of lung inflammation in the early metastatic phase, while late systemic endothelial dysfunction in aorta coincided with robust, systemic, cancer-related inflammation.

Methods

Animals

Seven- to eight-week-old BALB/c female mice were obtained from Charles River Laboratories Polska -Animal Lab (Poznan, Poland). Mice were housed in specific pathogen-free conditions (SPF) and fed a standard laboratory diet and water *ad libitum*.

All experimental procedures used in the present study were followed according to the Guidelines for Animal Care and Treatment of the European Communities and the Guide for the Care and Use of Laboratory Animals published by the US National Institutes of Health (NIH Publication No. 85–23, revised 1996). All procedures were approved by the First Local Ethical Committee on Animal Testing at the Jagiellonian University (Krakow, Poland), permit no: 140/2013.

Tumour cell line

Mouse mammary adenocarcinoma 4 T1 cells were obtained from American Type Culture Collection (ATCC). Cells were cultured in RPMI 1640 (Laboratory of Analytical Chemistry, IJET) with Opti-MEM® media (Life Technologies) (1:1 *v/v*) and 5% fetal bovine serum (HyClone, ThermoFisher), supplemented with 4.5 g/L glucose, 2 mM glutamine, 1.0 mM sodium pyruvate (all from Sigma-Aldrich) and antibiotics (penicillin and streptomycin; Polfa Tarchomin). Cell cultures were maintained at 37 °C in a humidified atmosphere with 5% CO₂.

Murine model of metastatic breast cancer

1 × 10⁴ viable 4 T1 tumour cells suspended in 0.05 ml of Hanks Balanced Salt Solution were orthotopically inoculated into the right mammary fat pad of female BALB/c mice. Analyses were conducted at 2, 4, 5 and 6 weeks after 4 T1 cancer cell transplantation. Prior to each analysis, animals were randomly divided into two experimental groups (one group designated for the analysis and the other for further tumour development), so that the

mean tumour volumes and tumour volume distributions were similar between experimental groups. Healthy BALB/c mice were used as a control group, and were analysed simultaneously with the tumour-bearing mice at 2, 4, 5 and 6 weeks after 4 T1 cancer cell transplantation.

Animals were anesthetized by intraperitoneal (*i.p.*) injection of a mixture of ketamine and xylazine (100 mg ketamine and 10 mg xylazine/kg body weight). Blood samples were collected from the right ventricle of the heart using a syringe containing anticoagulant (EDTA 1.6 mg/ml).

Assessment of the primary tumour and number of metastases in the lungs

Primary tumours were carefully dissected from the surrounding tissues and weighed. Isolated lungs were washed in saline, weighed, and fixed with 4% formalin buffered solution. The number of metastases was macroscopically assessed; metastatic sites visible on the lung surface were visually counted using a magnification glass.

Macroscopic analysis of the lungs was followed by paraffin embedding, histological HE staining and histopathological assessment of the tissue.

Histological and immunohistochemical analysis of the lungs

Lungs were fixed in 4% buffered formalin (for at least 48 h). After macroscopic analysis of the number of metastases, the lungs were prepared using the paraffin method, cut into 6 μ m sections on an Accu-Cut[®] SRM[™] 200 Rotary Microtome and stained with hematoxylin and eosin.

Light microscopic examination and photographic documentation were performed using an Olympus BX53F microscope equipped with a digital camera. Pictures were taken under the magnification 20 \times and 200 \times .

For immunohistochemical staining of VCAM-1 and vWF in the lung vasculature following deparaffinization, sections were pretreated according to the citrate-based HIER protocol and then preincubated with 5% goat serum (Jackson ImmunoResearch) and 2% dry milk to minimize non-specific binding of antibodies. Primary rat-anti-mouse VCAM-1 (Chemicon) or rabbit-anti-mouse vWF (Abcam) antibodies were used, followed by Cy3-conjugated goat-anti-rat or Cy3-conjugated goat-anti-rabbit secondary antibodies (Jackson ImmunoResearch), respectively. Images were acquired using the AxioObserver D2 inverted fluorescent microscope (Carl Zeiss) and an AxioCam HRm monochromatic digital camera and stored as TIFF files. Fluorescence intensity was analysed automatically by Columbus software (Perkin Elmer).

Assessment of NO production in the isolated lung preparation

Lungs were isolated from anaesthetized (pentobarbitone, 140 mg/kg, *i.p.*) animals and perfused at a

constant flow of about 1.50 ml/min with low glucose DMEM with 4% albumin and 0.3% HEPES, mounted in a water-jacketed artificial thorax and ventilated with negative pressure at a rate of 90 breaths/min. The end-expiratory pressure in the chamber was set to -3 cm H₂O and inspiratory pressure was adjusted between -6 and -10 cm H₂O to yield an initial tidal volume (TV) of about 0.2 ml. Every 5 min during the experiment, a deep breath of -21 cm H₂O end-inspiratory pressure was automatically initiated in order to avoid atelectasis. The PAP was set to around 3 cm H₂O. Venous pressure was set to 2-5 cm H₂O. All lungs preparations were allowed to equilibrate for at least 15 min under perfusion. Nitrate and nitrite concentrations were then measured in the effluent from the isolated lungs perfused with constant, non-recirculating flow. The samples were analysed via sensitive high-pressure liquid chromatography (HPLC) -based techniques (ENO-20 NO_x Analyser; EiCom, Kyoto, Japan).

Assessment of NO-dependent endothelial function in isolated rings of mice aorta

Mice aorta preparations and endothelial function assessment of aortic rings were conducted as previously described [13]. Briefly, NO-dependent endothelial function was measured by response to acetylcholine (Ach; 0.01-10 μ M) in phenylephrine (Phe; 0.1-1 μ M) pre-contracted vessels. Endothelium-independent vasodilatation was determined using sodium nitroprusside (SNP; 0.001-1 μ M). To ensure that entire Ach-evoked response was NO-dependent the response to Ach and SNP was also measured in the presence of L-NAME. Responses were recorded using a data acquisition system and recording software (Power Lab, Lab Chart, AD Instruments, Australia).

Assessment of nitrite production in isolated aorta rings

Basal NO production by the aorta was estimated using measurements of nitrite. Segments from the aortic arch were longitudinally opened, placed in a 96-well plate with endothelium facing up, and incubated for 1 h in 120 μ l K-H buffer at 37 °C using a specially-designed closed chamber (BIO-V (Noxygen)) that was equilibrated with carbogen gas mixture (95% O₂, 5% CO₂). Samples from the incubation buffer were put on ice and used for measurement of nitrite with reductive gas-phase chemiluminescence in 1% wt/vol KI in acetic acid using Sievers[®] Nitric Oxide Analyzer NOA 280i, according to the manufacturer's instructions. The averaged blank signal (without aortic rings) for a given set of experiments was subtracted as a background signal, to account for nitrite contamination in the buffer and/or laboratory atmosphere. Nitrite concentration was expressed as ng/ml/mg of dry weight of aortic rings.

Assessment of PGI₂ production in isolated rings of mice aorta

PGI₂ production by aortic rings was quantified on the basis of the formation of 6-keto PGF_{1α}, a stable metabolite of PGI₂, in the supernatant of the aortic rings.

Aortic rings were incubated on a thermoblock (Liebisch Labortechnik) at 37 °C, in 1 ml of K-H buffer equilibrated with carbogen gas mixture (95% O₂, 5% CO₂), either in the absence or presence of selective COX-2 inhibitor DuP-697 (1 μM) or nonselective COX-1/COX-2 inhibitor indomethacin (5 μM). Both DuP-697 and indomethacin were dissolved in DMSO. Control rings were incubated with DMSO (1 μl/ml).

After equilibration and pre-incubation in the presence or absence of inhibitors, the aortic rings were placed in fresh buffer and incubated with or without inhibitors for 30 min. Effluent samples were taken after 3 (initial) and 30 (final) minutes of incubation. PGI₂ production was calculated as the difference between final and initial 6-keto PGF_{1α} concentrations and was expressed as pg/ml/mg of

dry weight of aortic rings. 6-keto-PGF-1_α concentration was measured using an EIA kit (Enzo, Life Technologies).

Immunohistochemistry of aortic endothelium

Aortic rings were placed perpendicularly in OCT compound (Thermo) and then snap-frozen at - 80 °C. The blocks were mounted on the cryostat holder and cut into 10-μm-thick cross-sectional slides using the Leica CM1950 automatic cryostat. The sections were placed on polylysine-covered (Sigma-Aldrich) microscopic slides (Super Frost, Mentzel Gläser), and then acetone fixed (10 min). Pre-incubation with 2.5% horse serum (Vector Labs) and 2% dry milk was performed to minimize non-specific binding of antibodies. For indirect immunohistochemical detection of von Willebrand factor in endothelium, sections were incubated inside humid chambers with polyclonal rabbit anti-mouse vWF Ig (Abcam) following rinsing in PBS secondary biotinylated horse anti-rabbit Ig (Vector Labs). After another rinse in PBS, sections were incubated with Cy3-conjugated streptavidin

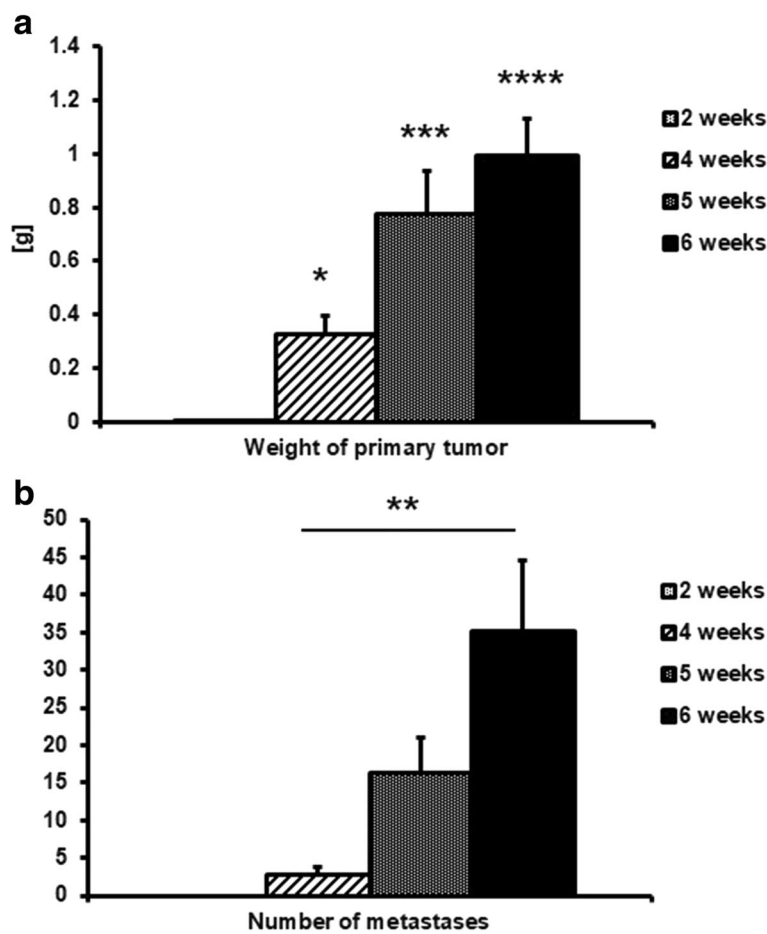


Fig. 1 Weight of primary tumour (a), * indicates $p < 0.05$; *** indicates $p < 0.001$; **** indicates $p < 0.0001$ vs. 2-week group, number of metastases in the lungs (b), ** indicates $p < 0.01$ at 2, 4, 5 and 6 weeks after cancer cell inoculation in mice. Results are expressed as mean \pm SEM, $n = 10-17$

(Jackson ImmunoResearch), and then mounted in glycerol-PBS. Images of the immunostained sections were acquired using the AxioObserver D2 inverted fluorescent microscope (Carl Zeiss) connected to a AxioCam HRm monochromatic digital camera, and stored as TIFF files. Fluorescence parameters were analysed automatically by Columbus software (Perkin Elmer).

Assessment of nitrite in plasma and NOHb in erythrocytes

Blood samples were centrifuged (1000 x g, 5 min, 4 °C) to isolate plasma and erythrocytes. 50 µl plasma was mixed with 100 µl of cold ethanol and kept on ice for protein precipitation (30 min), centrifuged (14,000 x g, 5 min.) and the resulting supernatant was used immediately after sample preparation to determine nitrite concentration using reductive chemiluminescence analysis (Sievers* Nitric Oxide Analyzer NOA 280i), as described above.

EPR spectra of the isolated erythrocytes were used for the detection of nitrosylhemoglobin (NOHb), as described previously [14, 15]. Briefly, erythrocytes were snap-frozen in insulin syringes and EPR measurements were performed in liquid nitrogen (77 K) using a Bruker EMX Plus spectrometer. Nitrosylhaemoglobin levels were expressed as the EPR amplitude of the second hyperfine line of the NOHb spectra in arbitrary units, and normalized to sample weight.

Blood count and cytokine analysis in plasma

Blood count was performed immediately after blood collection using a blood counter (abc Vet, HORIBA). To obtain blood plasma, samples were centrifuged for 7 min (1000 x g). Acute-phase protein concentration in plasma was measured using magnetic-beads-based immunoassay (Multiplex, Millipore). SAA and IL-6 plasma concentrations were measured with ELISA kits (Invitrogen and R&D Systems, respectively).

Statistical analysis

For statistical analysis, STATISTICA 10 software (StatSoft, Inc.) and OriginPro 9 software (OriginLab, Northampton, MA) were used. The Shapiro-Wilk test was used to verify whether the data were normally distributed. Levene's test was used to determine homogeneity of variances. The significance of between-group differences was evaluated by the Kruskal-Wallis test or ANOVA with post-hoc Tukey multiple comparisons test or by Mann-Whitney U test, depending on the data distribution. Spearman or Pearson's correlation coefficient test was used to assess dependence between two parameters. Results are presented as mean ± SEM. Differences between means were considered significant if $p < 0.05$.

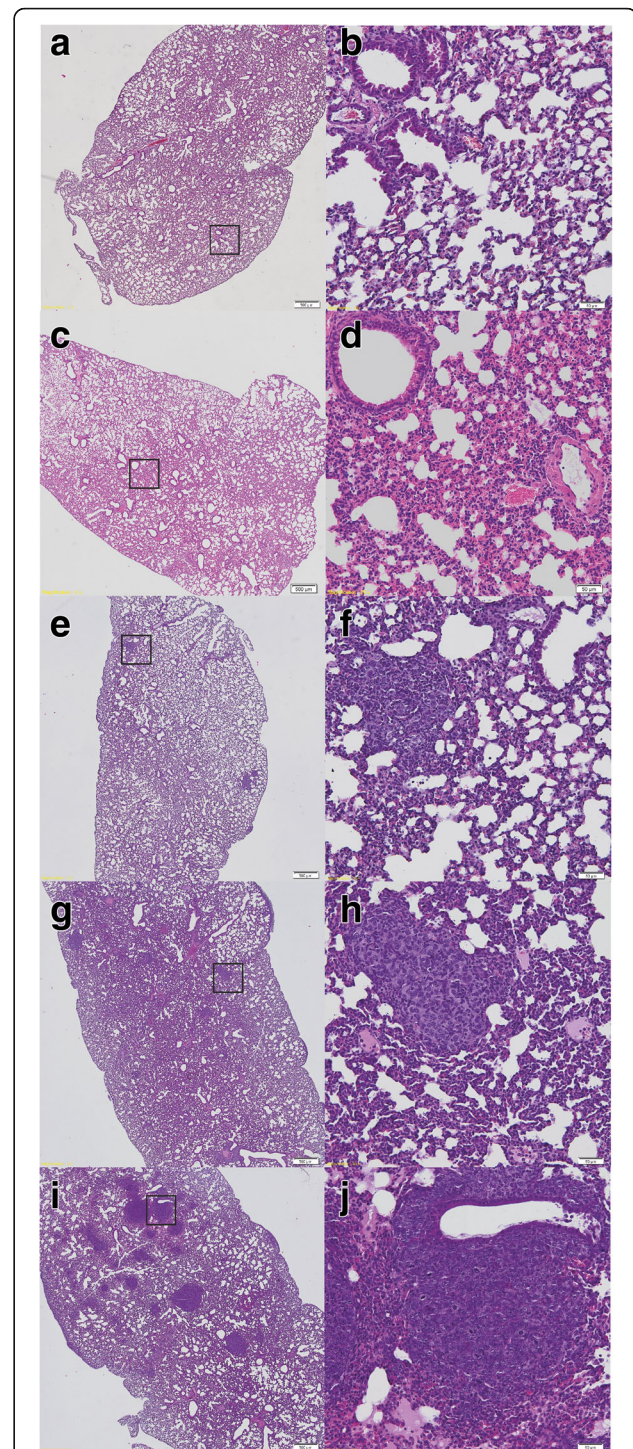


Fig. 2 Histopathological analysis of the mice lung. Control – normal lung tissue (a, b); lungs at 2 weeks after cancer cell injection with visible sites of inflammation, mainly granulocyte infiltration (c, d); 4-week group lungs with few small metastatic foci (e, f); 4-week group lungs with many visible scattered metastatic sites of different sizes (g, h); and lungs at 6 weeks after cancer cell injection, with numerous, extensive, merging metastatic areas (i, j). Magnification 20x: a, c, e, g, i; 200x: b, d, f, h, j (magnification of chosen area)

Results

Primary tumour growth and lung metastases development

Small palpable tumours (0.004 ± 0.001 g) were observed 2 weeks after cancer cell transplantation into mammary glands of BALB/c female mice. As shown in Fig. 1a, mean tumour weight increased progressively over the course of the study to a final weight of 0.99 ± 0.1 g, observed on the 6th week of the experiment.

No metastatic nodules were visible in lungs isolated from mice 2 weeks after cancer cell transplantation. However, the mean number of metastases developed in lungs isolated from tumour-bearing mice 4, 5 or 6 weeks after cancer cell transplantation gradually increased, reaching a mean of 35.2 ± 9.3 metastatic nodules at week 6 of the study (Fig. 1b).

Early pro-inflammatory changes in the lungs and impaired NO production in the pre-metastatic phase

As early as 2 weeks after cancer cell transplantation, local inflammatory cell infiltration into lung parenchyma was detected, identified as a population consisting mainly of granulocytes (Fig. 2d). However, no cancer cell clusters were observed in the lungs at this time point in microscopic cross-section preparations of the lungs (Fig. 2c). Inflammatory cells infiltration was accompanied by enhanced expression of VCAM-1 in the pulmonary vasculature, as evidenced by immunohistochemical staining (Fig. 3a). On the other hand, vWF staining in the lungs showed no significant difference between early pre-metastatic mice and controls (Fig. 3b). Moreover, nitrite and nitrate concentrations in buffer from isolated lungs perfused in a non-recirculated manner were lower in mice 2 weeks after cancer cell inoculation as compared to control mice (nitrite: 0.938 ± 0.301 μ M and 1.749 ± 0.346 μ M in tumour-bearing

and control animals, respectively; nitrate: 3.00 ± 0.013 μ M and 4.714 ± 0.445 μ M in tumour-bearing and control animals, respectively; $n = 3$ in both groups).

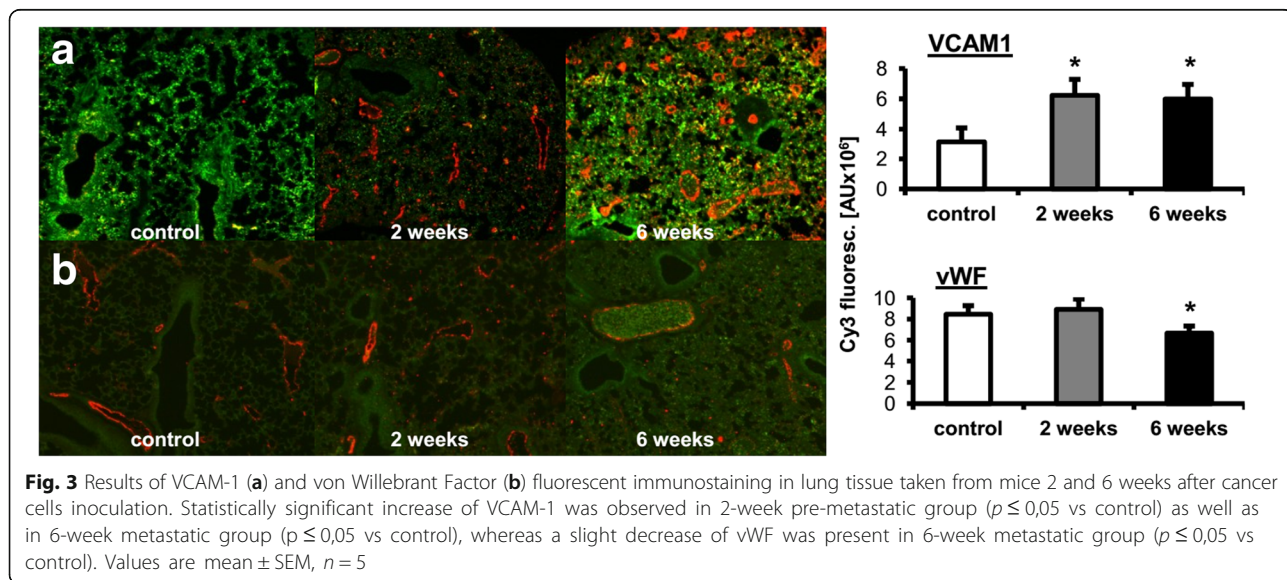
Lung histopathology in the advanced metastatic phase

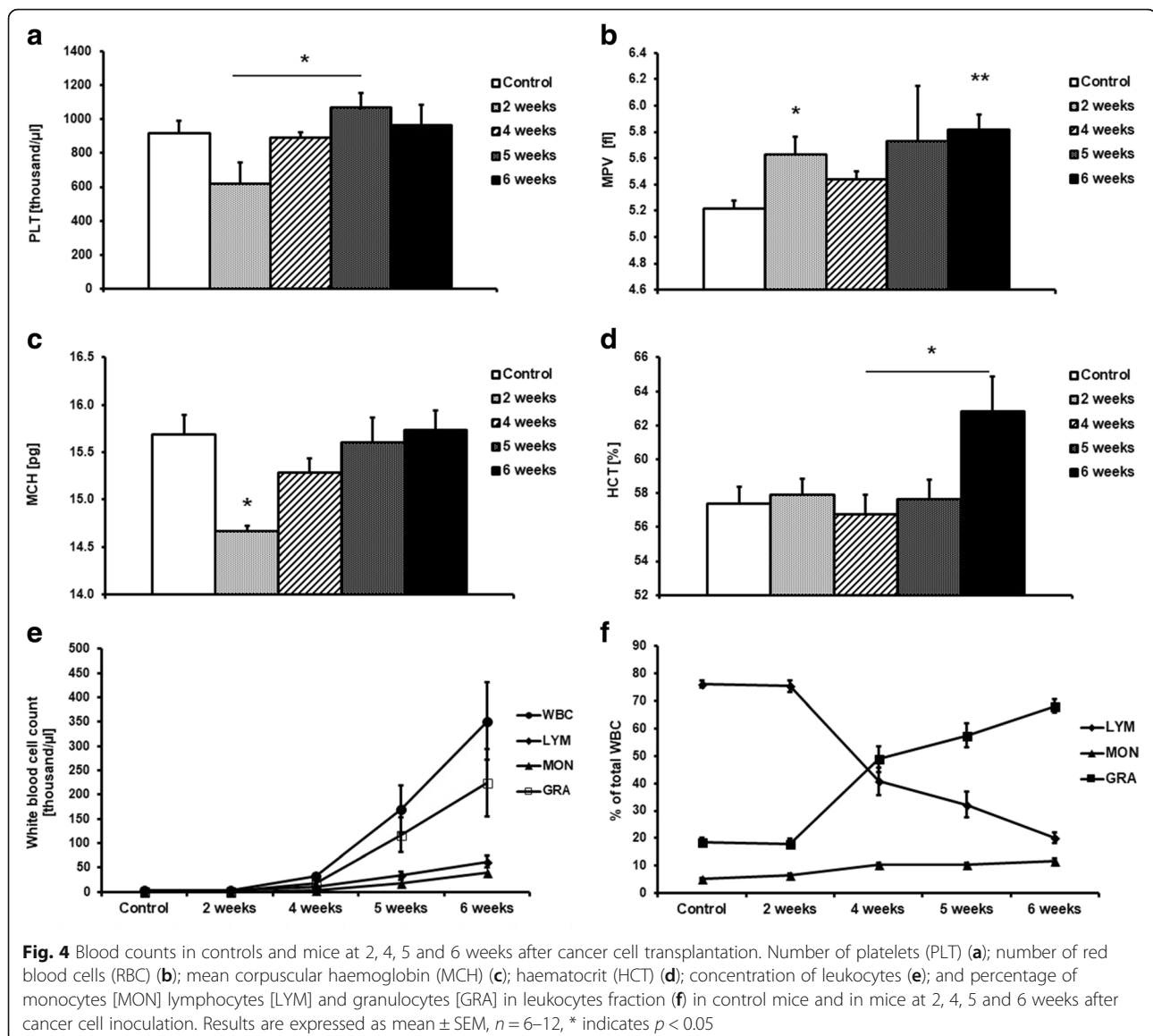
In contrast to the lungs isolated from tumour-bearing mice at 2 weeks following cancer cell transplantation, lungs analysed at 4 weeks displayed small, clear-cut metastatic foci (Fig. 2e, f). 5 weeks after cancer cell transplantation, scattered metastatic sites of different sizes were observed in the lung parenchyma and under the pleura (Fig. 2g, h). In the most advanced stage of metastatic disease progression, 6 weeks after 4 T1 cell engraftment, numerous, extensive, merging metastatic foci were evident (Fig. 2i, j).

Development of cancer-associated systemic inflammation Leukocytosis and other changes in blood count

At 2 weeks after cancer cell transplantation, mild changes in the blood count of tumour-bearing mice were found, including a drop in platelet counts that normalized over the course of the study (Fig. 4a) and an increase in mean platelet volume (MPV) that remained at an elevated level until the late stage of metastatic disease development (Fig. 4b). A significant decrease in mean cell haemoglobin (MCH) was also noted at 2 weeks (Fig. 4c), while in the late stage of disease (6 weeks), haematocrit (HCT) and average red blood cell volume (MCV) were significantly increased (Fig. 4d; Table 1), compensating for the drop in MCH seen in the early stage of cancer progression (2 weeks).

The most pronounced alterations in the blood count profile were observed in the late stages of metastatic disease and involved profound and progressive leukocytosis observed at 4–6 weeks after cancer cell transplantation





(Fig. 4e; Table 1). Although the absolute numbers of granulocytes [GRA], monocytes [MON], and lymphocytes [LYM] increased, the percentage changes indicated a substantial increase in granulocyte count, a mild increase in monocyte count, and a relative decrease in lymphocyte count (Fig. 4f). In fact, in control animals as well as in the early stage of disease progression in tumour-bearing mice (2 weeks), the most abundant WBCs were lymphocytes, while in the advanced stages of cancer progression (4–6 weeks) the most abundant WBCs were granulocytes (Fig. 4f; Table 1).

There was a high correlation between number of metastases and leukocytes, irrespective of whether the change in total white blood cells, or in granulocytes, lymphocytes, and monocytes, were analysed (Table 2).

Changes in plasma concentration of acute phase proteins and IL-6

Along with tumour progression, there was a gradual increase in plasma IL-6 concentration in tumour-bearing mice. In the early stage, 2 weeks and 4 weeks after cancer cell transplantation, plasma IL-6 concentration was comparable to that of control mice, while at 5 and 6 weeks after cancer cell transplantation, there was a significant, 3- to 4-fold elevation in plasma IL-6 concentration compared to control (14.5 ± 4.5 pg/ml and 19.8 ± 2 pg/ml vs. 5.8 ± 1.4 pg/ml, respectively; Fig. 5a). Quite a similar response pattern was observed for acute phase proteins in plasma. The concentrations of serum amyloid A (SAA), haptoglobin and serum amyloid protein (SAP) (Fig. 5b-d) were

Table 1 Blood counts in control mice and mice at 2, 4, 5 and 6 weeks after cancer cells inoculation

	Control	2nd week	4th week	5th week	6th week
PLT [K. μl^{-1}]	916.30 \pm 47.39	618.75 \pm 128.13#	892.81 \pm 30.04	1067.25 \pm 87.86	964.58 \pm 119.81
MPV [fl]	5.21 \pm 0.06	5.63 \pm 0.13*	5.44 \pm 0.06	5.73 \pm 0.42	5.82 \pm 0.12**
RBC [M. μl^{-1}]	10.37 \pm 0.17	10.50 \pm 0.15	10.16 \pm 0.22	10.27 \pm 0.23	10.72 \pm 0.30
MCH [pg]	15.69 \pm 0.20	14.67 \pm 0.06*	15.28 \pm 0.15	15.60 \pm 0.27	15.74 \pm 0.21
HCT [%]	57.40 \pm 0.96	57.92 \pm 0.93	56.78 \pm 1.12\$	57.63 \pm 1.15	62.84 \pm 2.01
MCV [fl]	55.33 \pm 0.18	55.15 \pm 0.31	55.92 \pm 0.32	55.10 \pm 0.32	58.75 \pm 0.70*
WBC [K. μl^{-1}]	3.87 \pm 0.21	2.94 \pm 0.26	31.80 \pm 6.14	169.35 \pm 49.59**	351.50 \pm 79.54****
LYM [K. μl^{-1}]	2.88 \pm 0.15	2.65 \pm 0.60	9.96 \pm 1.26	33.62 \pm 8.15**	62.19 \pm 11.43****
MON [K. μl^{-1}]	0.15 \pm 0.02	0.15 \pm 0.02	3.59 \pm 0.76	18.46 \pm 5.67**	40.53 \pm 8.15****
GRA [K. μl^{-1}]	0.84 \pm 0.08	0.62 \pm 0.09	18.46 \pm 4.54	117.27 \pm 36.28*	224.36 \pm 69.17****
LYM [%]	76.14 \pm 1.39	75.35 \pm 2.00	40.83 \pm 5.03****	32.19 \pm 4.69****	20.09 \pm 1.86****
MON [%]	5.25 \pm 0.26	6.69 \pm 0.45	10.27 \pm 0.83****	10.34 \pm 0.64****	11.74 \pm 0.98****
GRA [%]	18.61 \pm 1.39	17.97 \pm 1.71	48.91 \pm 4.70*	57.47 \pm 4.43***	68.17 \pm 2.48****

Data are expressed as mean \pm SEM

PLT platelets, MPV mean platelet volume, RBC red blood cells, MCH mean cell haemoglobin, HCT haematocrit, MCV mean corpuscular volume, WBC white blood cells, LYM lymphocytes, MON monocytes, GRA granulocytes

* $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$, and **** $p < 0.0001$ vs. control group; # $p < 0.05$ vs. 5-week group; \$ $p < 0.05$ vs. 6-week group, ($n = 8-12$)

significantly increased in tumour-bearing mice at 5 and 6 weeks following cancer cell inoculation. Plasma levels of haptoglobin, IL-6 and SAA correlated with the number of lung metastases in tumour-bearing mice (Table 3). On the other hand, metastases development did not significantly influence plasma concentrations of $\alpha 1$ acid glycoprotein (Fig. 5e) or $\alpha 2$ macroglobulin (Fig. 5f).

Changes in NO-dependent function in aorta

In all experimental groups, acetylcholine (Ach) induced a dose-dependent vasodilation of phenylephrine-precontracted aortic rings that was entirely inhibited by L-NAME. The Ach-induced response in aorta was not changed at 2, 4 and 5 weeks after cancer cell inoculation, but was impaired in the latest stage of metastatic cancer progression (6 weeks) as compared to control mice (Fig. 6a). In contrast, the endothelium-independent response induced by SNP remained unaffected for all experimental groups at all time points (Fig. 6b).

Table 2 The correlation between the number of metastases in the lungs of tumour-bearing mice and blood parameters assessed via the Spearman correlation coefficient test

The pair of parameters	R Spearman	p
Number of metastases & WBC	0.795611	0.000000
Number of metastases & LYM	0.785315	0.000000
Number of metastases & MON	0.788080	0.000000
Number of metastases & GRA	0.755175	0.000001
Number of metastases & MCV	0.632756	0.000102
Number of metastases & MCH	0.515611	0.002525

Parallel to the impairment of the NO-dependent Ach-induced response, a deterioration of nitrite production by the endothelium was observed in the isolated aortic rings. As shown in Fig. 7a, basal nitrite production was significantly lower in aortas isolated from tumour-bearing mice at 6 weeks after cancer cell inoculation, supporting an impairment of NO-dependent function.

Changes in systemic NO bioavailability

Mean plasma nitrite concentration in control mice was 487 ± 45 nM. Interestingly, in the early stage of cancer development (2 and 4 weeks after cancer cell transplantation) plasma nitrite concentration and NOHb concentration were lower (significant for NOHb, $p < 0.05$) compared to control mice (Fig. 7b, c).

In contrast to the lower concentration of NO_x metabolites in the early phase of metastasis, the concentrations of both NOHb and nitrite were significantly increased at 6 weeks after cancer cell transplantation (Figs. 6c and 7b).

Changes in PGI₂-dependent function in the aorta

Impairment of NO-dependent function in the aorta was preceded by persistent compensatory up-regulation of PGI₂ production, as shown through an increased concentration of the stable PGI₂ metabolite 6-keto-PGF_{1 α} in the effluent of aortic rings isolated from tumour-bearing mice at 4–6 weeks following cancer cell inoculation. Both DuP-697 (1 μM) and indomethacin (5 μM) profoundly inhibited PGI₂ release from aortic rings, suggesting that COX-2 was the main contributor to the upregulated PGI₂ production in the vascular wall (Fig. 8).

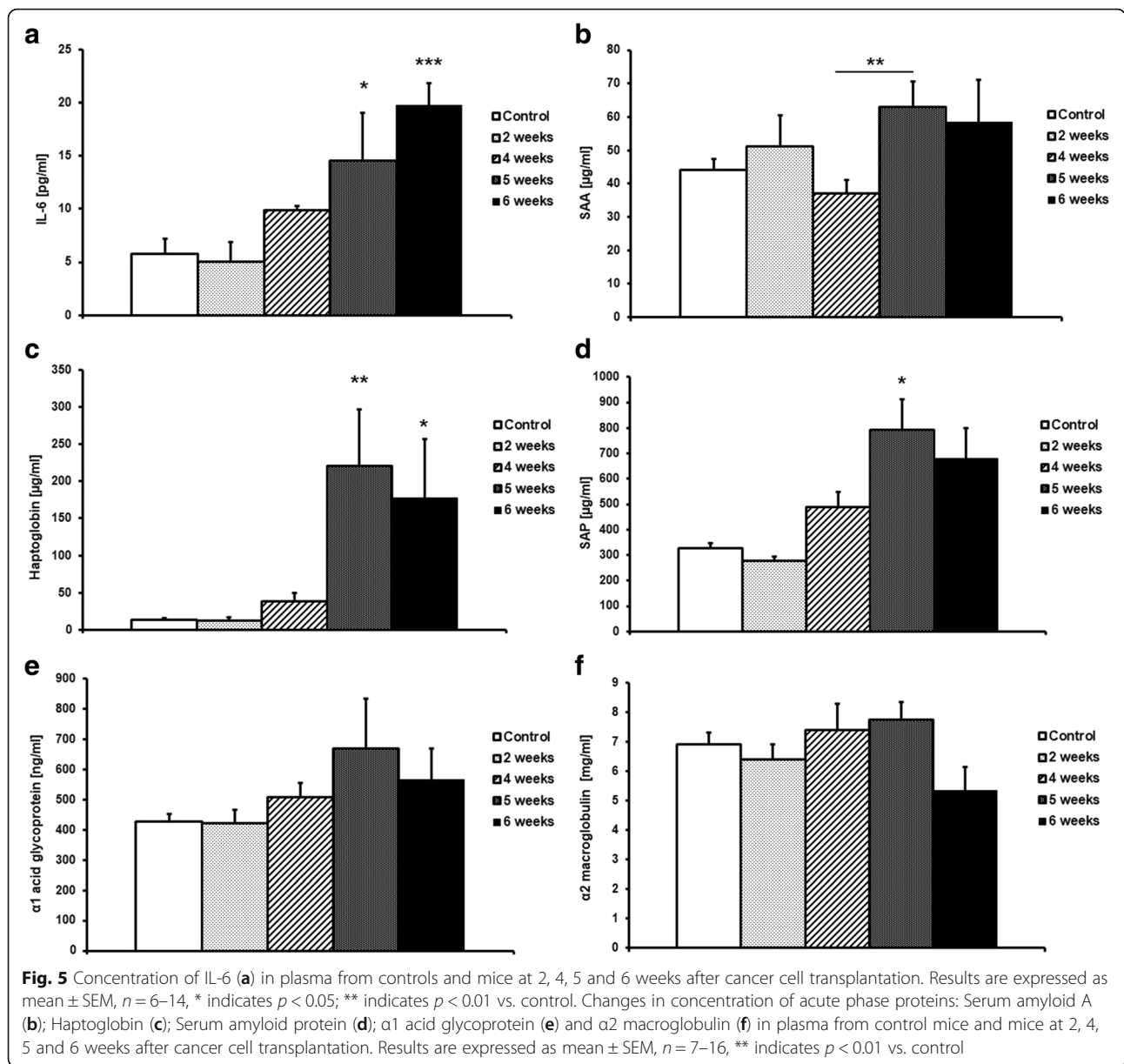


Table 3 The correlation between number of metastases in the lungs of tumour-bearing mice and Interleukin-6 (IL-6), Serum Amyloid A (SAA) and haptoglobin, as assessed via the Spearman correlation coefficient test

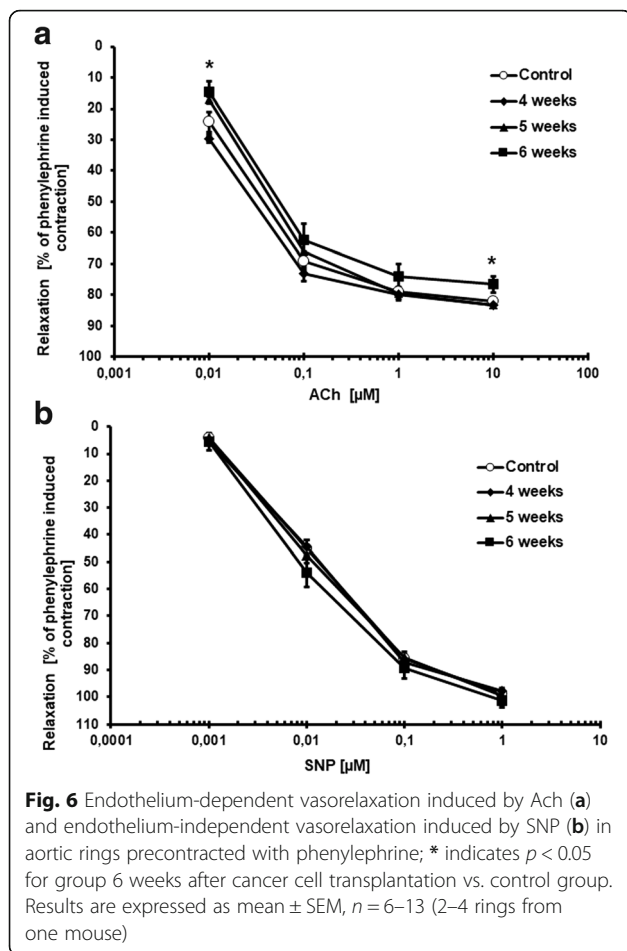
The pair of parameters	R Spearman	p
Number of metastases & IL-6	0.655558	0.000014
Number of metastases & SAA	0.554662	0.000447
Number of metastases & Haptoglobin	0.814386	0.013844

Changes in vWF expression in aorta

As shown in Fig. 9, in the aortic endothelium of tumour-bearing mice, vWF expression was increased compared to control animals. Up-regulated expression of vWF was observed as early as 2 weeks after cancer cell inoculation and progressively increased through the later stages of metastases development (Fig. 9a, b).

Discussion

Healthy endothelium inhibits formation of metastasis, while dysfunctional endothelium has an opposite effect [1, 7]. On the other hand, systemic endothelial dysfunction developing along with tumour progression and metastasis contributes to cardiovascular complications in



cancer patients [6]. The present work has taken advantage of the prolonged time course of the disease progression in the spontaneously metastasizing, orthotopic, 4 T1 murine breast cancer model in order to characterize the temporal relationships between the development of local, pulmonary, and systemic endothelial dysfunction in relation to primary tumour growth and lung metastases formation. We showed that an early decrease in NO bioavailability was temporarily associated with the development of early metastasis and lung inflammation, while late systemic impairment of NO-dependent function in the aorta coincided with robust, systemic, cancer-related inflammation. Our results show that the local pulmonary endothelial response to murine metastatic breast cancer development involves pulmonary inflammation, granulocyte infiltration and decreased NO bioavailability in the lung. On the other hand, impairment of NO-dependent function in the aorta occurred only in the late phase of metastasis formation, most likely as a consequence of robust, cancer-related systemic inflammation mediated by IL-6, TNF α [16] or other mechanisms [17]. Interestingly, impairment of NO-dependent function was associated with the

upregulation of COX-2-derived PGI $_2$ production, which could represent a compensatory mechanism. Indeed, PGI $_2$, the most potent endogenous anti-platelet agent, has been shown to prevent cancer metastasis [11, 12]. The same activity has also been reported for PGI $_2$ analogues [16, 17] and PGI $_2$ -releasing compounds [20].

Although it was recently reported that the development of metastatic breast carcinoma in mice is associated with changes in vascular reactivity of the aorta [16], to our knowledge the present work reports for the first time the temporal relationships between the development of the local, pulmonary endothelial dysfunction that is a pre-requisite for development of metastasis in the lung and the systemic endothelial dysfunction and vascular inflammation that affect general cardiovascular health.

Taking advantage of the slowly progressing 4 T1 breast cancer-based model established by our group [14, 17], which relies on transplantation of a low number of 4 T1 cancer cells and is similar to the recently-reported optimized 4 T1 triple negative syngeneic micro- and macro-metastasis model by Bailey-Downs [21], we clearly differentiated between the occurrence of early endothelial dysfunction in lungs at the pre-metastatic/early metastatic stage and development of peripheral endothelial dysfunction in the conduit artery at the late metastatic stage. These results suggest different mechanisms involved in these responses. The early pre-metastatic phase was characterized by clear-cut evidence of pulmonary inflammation and granulocytic cellular infiltration. The presence of leukocyte infiltration is consistent with previous work [22-24] demonstrating that 4 T1 tumour-bearing mice display granulocytic infiltrates in metastatic sites [25] and the result supports the concept of a pre-metastatic niche formed in the pulmonary target tissue before it is invaded by migrating cancer cells [26]. This hypothesis is also supported by the VCAM-1 immunofluorescent staining results showing a pro-inflammatory phenotype in the pulmonary endothelium as early as 2 weeks following cancer cell transplantation, at the early stage of the metastatic process.

Based on the analysis in the isolated perfused lung preparation, we found that nitric oxide production was decreased in the early phase of metastasis development. Impairment of NO bioavailability may contribute to the adhesion of cancer cells and neutrophils to the endothelium and may facilitate cancer cells extravasation [25-28]. Given that NO displays anti-platelet activity and that platelets are involved in arresting tumour cells in the vasculature at metastatic extravasation sites [29-32], a low-NO microenvironment could also affect the platelet-cancer cell interaction, facilitating metastasis. Interestingly, we detected a decrease in platelet number in the early phase of metastasis, which could reflect an increased

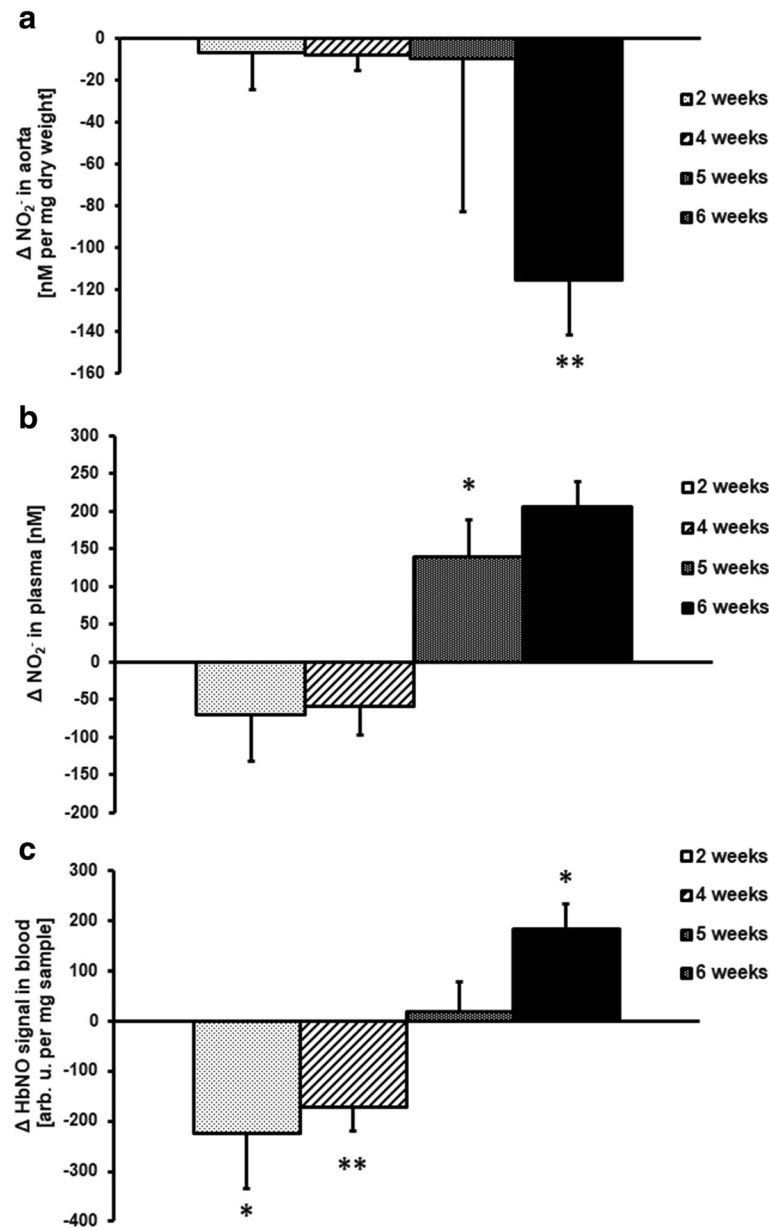
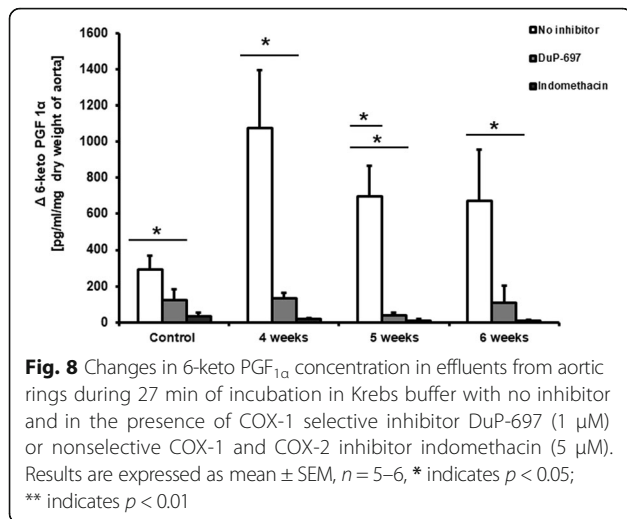


Fig. 7 Basal nitrite production in incubated aorta rings (a), plasma nitrite concentration (b) and nitrosylhemoglobin level (c) in mice at 2, 4, 5 and 6 weeks after cancer cell transplantation (Results presented as Δ metastasis – control, showing [%] difference from appropriate control groups at corresponding age (2, 4, 5 and 6 weeks after cancer cell inoculation). * indicates $p < 0.05$ and ** indicates $p < 0.01$. Results are expressed as mean \pm SEM, $n = 3$ (control groups) and $n = 6$ (metastasis groups)

platelet-cancer cell interaction. Altogether, our findings of early NO deficiency seem to be important in the context of pathogenic endothelial-dependent mechanisms of early metastasis. NO-deficient pre-metastatic niche formation may promote early metastasis in the lung involving cancer cells, leucocytes and platelets.

Interestingly, the pre-metastatic/early metastatic phase was associated with a transient decrease in systemic NO availability, as evidenced by a drop in NOHb content, as

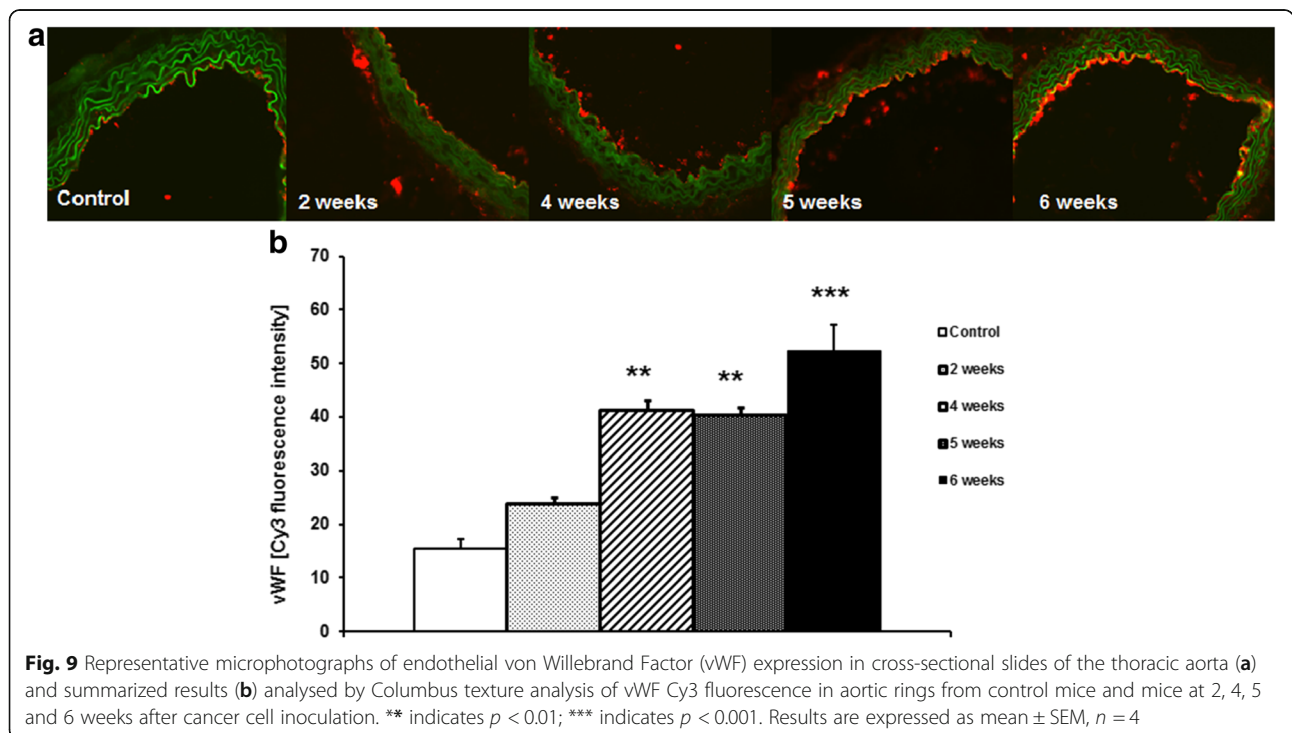
well as decreased nitrite concentration in plasma. These results suggest that the magnitude of the impaired NO production in the lung was high enough to impose an effect on systemic NO bioavailability. Given that the pulmonary endothelial surface represents approximately 30% of the total endothelial surface of the cardiovascular system, this finding is not surprising. Later in the course of metastatic development, nitrite and NOHb increased, an effect most likely linked to systemic inflammation and NOS-2 induction [33].



In contrast to the pre-metastatic and early metastatic phases, the late metastatic phase of disease (5–6 weeks after cancer cell transplantation) was characterized by a robust systemic inflammation, including substantial leucocytosis, increases in RBC number, MCV and HCT, elevated platelet number, mean platelet volume, as well as increased plasma concentration of acute phase proteins (haptoglobin, serum amyloid A) and IL-6. Leukocytosis seems to be compatible with the production of colony-stimulating factors released by tumour cells in 4 T1 tumour-bearing mice, enhanced myelopoiesis in bone marrow and extramedullary myelopoiesis occurring in the

spleen [25], and may support metastasis by creating an environment that facilitates the recruitment and growth of circulating tumour cells. Indeed, G-CSF neutralization was shown to diminish tumour cell infiltration into lungs, and consequently to reduce metastasis [24].

In our study, progression of metastasis was also associated with an increased concentration of IL-6, a major proinflammatory cytokine that affects cell proliferation, survival, and metabolism and is known to be associated with the progression of various types of cancer [34–36]. IL-6 stimulates the expression of adhesion molecules on the endothelial surface that could favour the adhesion of tumour cells [37]. Interestingly, since IL-6 has been found to be elevated in cancer patients with distal metastases, as compared with non-metastatic patients, an increased IL-6 concentration has been proposed as a prognostic marker in some types of metastatic cancer [37, 38]. In the present work, metastasis was linked with the activation of the acute phase response, as evidenced by the elevation of serum amyloid A, serum amyloid P and haptoglobin, a response that could be triggered by an increased IL-6 concentration. It has previously been reported that haptoglobin is elevated in breast cancer and may be indicative of metastases [40, 41]. Interestingly, altered glycan content in some acute phase proteins, for example, haptoglobin, interact with a number of receptors on macrophages in the tumour micro-environment, potentially modulating macrophage activity and thereby contributing to tumour cell survival, growth, and metastasis [41, 42]. In turn, SAA stimulates its own transcription as well as that of the proinflammatory



S100A8 and S100A9 proteins, strongly enhancing adhesion, migration and invasion of human and mouse tumour cells [43]. Thus, the acute phase response may contribute to the metastasis formation via various mechanisms [41]. In our study, metastasis in 4 T1 tumour-bearing mice was associated with substantial leukocytosis, activation of IL-6-dependent mechanisms and an acute phase response, all of which could contribute to the further acceleration of metastasis formation [23, 39, 40]. In fact, there was a high correlation between leukocyte number, IL-6, haptoglobin, SAA plasma concentration and a number of lung metastases in tumour-bearing mice. At the stage of advanced metastasis associated with robust systemic inflammation, the NO-dependent response in aorta was impaired, while the endothelium-independent response was preserved. Interestingly, downregulation of a NO-dependent endothelial activity was associated with a compensatory increase in COX-2-mediated synthesis of PGI₂, a vasoprotective molecule with known anti-metastatic potential [11, 12, 18–20, 44].

In our hands, impaired NO-dependent function in the aorta was associated with up-regulation of vWF expression, suggesting a pro-thrombotic phenotype of dysfunctional endothelium in aorta, compatible with cancer-related activation of pro-thrombotic endothelial mechanisms [6]. Interestingly, vWF did not increase in the pulmonary endothelium, but was even lower in the late stage of metastasis formation compatible with a possible involvement of vWF in cancer cell metastasis to the lungs [45, 46].

Conclusion

In the present work in orthotopic murine model of metastatic 4 T1 breast cancer we have characterized the temporal relationships between the development of endothelial dysfunction in the pulmonary and systemic circulation in relation to primary tumour growth and lung metastases. We have demonstrated that the local pulmonary endothelium response to cancer cells, that is the pre-requisite response for the development of metastases in the lung, involves decreased NO bioavailability in the lung, that is also reflected by a decrease in systemic NO bioavailability and is associated with pulmonary inflammation with granulocyte infiltration. All these early changes precedes the systemic inflammation and systemic endothelial dysfunction induced by cancer development. Late impairment of endothelial NO-dependent function in the aorta was associated with compensatory upregulation of the COX-2-derived PGI₂ pathway and coincided with robust, systemic, cancer-related inflammation. Our results obtained in an murine, orthotopic model of breast cancer metastasis, relevant to human pathology, suggest a heterogeneity of mechanisms of the early endothelial dysfunction in metastatic

organ paving the way for metastases formation and the late systemic endothelial dysfunction contributing to systemic vascular inflammation.

Abbreviations

Ach: Acetylcholine; COX-2: Cyclooxygenase-2; CTCs: Circulating tumour cells; DMSO: Dimethyl sulfoxide; EMT: Epithelial mesenchymal-like-transition; HPLC: High-pressure liquid chromatography; IL-6: Interleukin 6; K-H buffer: Krebs-Hanseleit buffer; L-NAME: N- ω -Nitro-L-Arginine Methyl Ester; NO: Nitric oxide; NOHb: Nitrosylhaemoglobin; NOS: Nitric oxide synthase; PDGF: Platelet-derived growth factor; PGI₂: Prostacyclin; Phe: Phenylephrine; SAA: Serum amyloid A; SNP: Sodium nitroprusside; TGF- β : Transforming growth factor β ; VCAM-1: Vascular cell adhesion molecule 1; VEGF: Vascular endothelial growth factor; vWF: von Willebrand factor

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Availability of data and materials

All data generated or analyzed during this study are included in this published article.

Authors' contributions

EB and SC conceived and design research; EB, AD, LM, BP, TW, BS, AF, AJ, EK, AC-S carried out experiments; RG contributed with analytic tools; EB, BP, LM, AJ, performed data analysis; EB, SC drafted the manuscript. AD, BP, JW revised the manuscript; EB, SC wrote the final version of the manuscript. All authors read and approved the final manuscript.

Ethics approval and consent to participate

All experimental procedures used in the present study were followed according to the Guidelines for Animal Care and Treatment of the European Communities and the Guide for the Care and Use of Laboratory Animals published by the US National Institutes of Health (NIH Publication No. 85-23, revised 1996). All procedures were approved by the First Local Ethical Committee on Animal Testing at the Jagiellonian University (Krakow, Poland), permit no: 140/2013.

Competing interests

The authors declare that they have no competing interests.

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