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Anti-Vascular endothelial growth factor therapy impairs endothelial function of retinal microcirculation in colon cancer patients – an observational study

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Abstract

Background: To assess acute effects of bevacizumab (anti-VEGF therapy) on cerebral microvessels and systemic cardiovascular regulation.

Design and subjects: 20 consecutive patients with colorectal cancer (median age: 60.4 years, range 45.5-73.9 years) received bevacizumab intravenously (5 mg/kg) uncoupled of chemotherapy. Prior to and within the first 24 hours after bevacizumab infusion, patients were investigated for retinal endothelial function. A series of a triple 24-hour ambulatory blood pressure measurement was conducted. Retinal endothelial function was determined as flicker light-induced vasodilation. The integrity of baroreflex arc and autonomic cardiovascular control was examined by stimulatory manoeuvres.

Results: Bevacizumab therapy significantly reduced the vasodilatory capacity of retinal arterioles in response to flicker light. A slight decrease in diastolic pressure and heart rate was observed after bevacizumab infusion but this was unrelated to changes in retinal function. The pressure response upon nitroglycerin was largely preserved after bevacizumab infusion. The proportion of patients with abnormal nocturnal blood pressure regulation increased under anti-angiogenic therapy. Autonomic blood pressure control was not affected by bevacizumab treatment.

Conclusions: Bevacizumab acutely impairs microvascular function independent of blood pressure changes. Imaging of the retinal microcirculation seems a valuable tool for monitoring pharmacodynamic effects of bevacizumab.

Trial registration: NCT ID: NCT00740168

Keywords: Angiogenesis, Endothelium, Microcirculation, Cerebral hemodynamics, Nitric oxide

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Background

Vascular endothelial growth factor (VEGF) is a key mediator of physiological and pathological angiogenesis. As all solid tumors are dependent on pathological angiogenesis, anti-VEGF therapy has been approved in various advanced and metastatic cancer types [1,2].

The rationale for treating cancer patients with angiogenesis inhibitors has been on the assumption that unlike in tumor vessels survival and functioning of the normal vasculature is largely independent of VEGF signalling [3]. Although clinical trials have proven the clinical benefit of bevacizumab and other angiogenesis inhibitors for the treatment of advanced cancers [1,2,4,5], their use has been associated with hypertension and thromboembolic events [5-8]. Several mechanisms of the underlying cardiovascular toxicity have been proposed including the inhibition of vasomotor effects by VEGF but also structural and functional microvascular rarefaction [9-11]. Accordingly, changes in endothelial function have been observed in cancer patients treated by angiogenesis inhibitors [10,11] and may serve as an important clinical marker for cardiovascular complications and therapy outcome. This hypothesis is corroborated by multiple clinical trials showing an association between bevacizumab-induced hypertension and response to therapy [12].

Here, we investigated the acute impact of bevacizumab administration on retinal vessel reactivity in relation to acute changes of cardiovascular regulation inclusive blood pressure using the dynamic retinal vessel analysis and evaluation of the autonomic nervous system. We hypothesised that the dynamic retinal vessel analysis represents a useful non-invasive and easy applicable tool to monitor functional microvascular alterations following VEGF inhibition.

Materials and methods

Participants and ethics

Patients with histological confirmed metastatic or locally advanced, inoperable colorectal cancer were enrolled into the study. All patients had an indication for treatment with bevacizumab / fluoropyrimidine containing chemotherapy regimens and had not previously received drugs directly targeting the VEGF pathway. Bevacizumab was administered at a dose of 5 mg/kg one day before the start of chemotherapy (day 0). To separate effects, fluoropyrimidine based chemotherapy was started after completion of all functional vascular measurements on day 1 as described below. Patients were evaluated by dynamic retinal vessel analysis, autonomic function tests and 24-hour ambulatory blood pressure monitoring (ABPM) the day before (day -1) and the day after bevacizumab infusion (day 1). ABPM was also applied on the day of bevacizumab treatment (day 0). The protocol was approved by the Ethics Review Board of the

Medical Faculty of the Dresden University of Technology. All patients signed a written informed consent before inclusion into the study. The procedures conformed to the principles of the Declaration of Helsinki (Clinical Trials Registration NCT ID: NCT00740168)

Evaluation of the autonomic nervous system

All measurements were performed in a temperature and humidity controlled specialized autonomic laboratory. Consumption of caffeine or other xanthine containing foods were not permitted on the examination day. Alcohol and tobacco usage was ceased at the day preceding the measurements. For continuous cardiovascular monitoring using the SUEMPATHY device (Suess Medizin-Technik, Aue, Germany) including the non-invasive blood pressure monitoring CBM3000 device (Nihon Colin Co, Komaki, Japan) the patient was placed in the supine position on a tilt table [13]. After a resting period of 20 minutes the following manoeuvres were performed. First blood pressure and heart rate responses to sublingual administration of 400 µg glyceroltrinitrate (Nitrolingual® akut Spray, Pohl Boskamp, Hohenlockstedt, Germany) were recorded for five minutes followed by deep metronomic breathing at a rate of six breaths per minute for two minutes. For evaluation of orthostatic reaction, patients were tilted to a 60° upright position for 15 minutes. After 10 minutes of head-up tilting, 400 µg glyceroltrinitrate was repeatedly administered while tilting was continued for another five minutes. During head-up tilting the subsequent decrease in ventricular filling and stroke volume evokes a transient fall in arterial pressure which is counter-regulated by the baroreflex arc with resultant tachycardia (vagal withdrawal) and vasoconstriction (sympathetic activation). This reaction is amplified by the application of vasodilatory agents such as glyceroltrinitrate. Orthostatic hypotension may occur as a result of sympathetic dysfunction which is defined as a reduction in systolic blood pressure of at least 20 mmHg and/or diastolic pressure of at least 10 mmHg within three minutes of upright tilt [14].

Retinal vessel analysis

The Dynamic Retinal Vessel Analyzer (DVA, IMEDOS GmbH, Jena, Germany) was used for digital fundus imaging to perform static and dynamic retinal vessel analysis. Thirty minutes after initiation of mydriasis of the right pupil using 1% tropicamide eye drops, each participant underwent seven-field fundus photography using the system Visualis (IMEDOS, FF450plus, 535–561 nm, 30° image, 1840×1360 pixel) as previously described [15]. The arteriolar-to-venular ratio (AVR) was calculated from the central retinal arterial and venous equivalents to quantify generalized arterial narrowing. Dynamic vessel analysis was performed allowing non-invasive diagnosis of microvascular function by measuring the diameter of retinal arteries and veins continuously

[16,17]. An optoelectronic shutter was inserted in the retinal camera which interrupted the observation light with a frequency of 12.5 Hz. The baseline diameter was measured for 30 seconds in continuous light followed by three cycles of 20-second flicker provocation and 50-second steady fundus illumination. An arterial and venous segment of approximately 1.5 mm length was subsequently evaluated. An interval of 30 seconds before each flicker stimulation was considered as baseline to which the subsequent diameter response was normalised. The peak dilation was the largest vessel diameter at the end of each flicker stimulation averaged across three flicker periods. Changes in ocular hemodynamic parameters were expressed as percent change over baseline values.

24-h Ambulatory blood pressure monitoring

Validated automated 24-hour blood pressure devices (Boso GmbH, Jungingen, Germany) fulfilling the accuracy criteria of the Association for the Advancement of Medical Instrumentation were used to measure blood pressure and heart rate at intervals of 15 minutes at daytime (6–22 hours) and of 30 minutes at night (22–6 hours) at three different occasions. For each 24-hour measurement, blood pressures were evaluated separately for daytime and nighttime phases. Arterial daytime hypertension was diagnosed according to the criteria of the European Society of Hypertension (systolic blood pressure ≥ 140 mmHg and/or diastolic blood pressure ≥ 90 mmHg). Nocturnal hypertension was defined according to the guidelines of the American Hypertension Association (systolic blood pressure > 120 mmHg and diastolic blood pressure > 75 mmHg). A non-dipping blood pressure profile was existent when the nocturnal blood pressure fall was less than 10% of daytime values [18,19].

Multiple trigonometric regressive spectral analysis

Frequency analysis allows for quantification of cardiovascular regulation by assessing spontaneous oscillations in systemic arterial pressure and heart rate. Multiple trigonometric regressive spectral analysis (MTRS, ANS Consult, Freital, Germany) was applied to corresponding values of systolic blood pressure and of R-R interval derived from electrocardiogram. A non-artifactual global data segment of 2 min was analyzed. The local time window was set at 30 seconds and was shifted beat by beat for temporal determination of frequency and amplitude [20,21]. Two main spectral bands are usually considered: High frequency (HF) oscillations (spectral band between 0.15 and 0.4 Hz) of heart rate relate to respiratory sinus arrhythmia and, therefore, to parasympathetic cardiovagal tone. The other oscillation of interest is in the low-frequency range (spectral band between 0.04 and 0.15 Hz), usually centred around 0.1 Hz (six cycles min^{-1}). Low-frequency oscillations of heart rate (LF R-R) are thought to reflect the

baroreflex-mediated adjustments to the sinus node while LF variations of systolic blood pressure are primarily the result of sympathetically mediated fluctuations in peripheral vasomotor tone. The LF/HF ratio allows for quantification of the relation between the two branches of the autonomic nervous system. The total spectral power describes the amount of variability of a signal or stochastic process at a specific frequency. Baroreflex sensitivity was calculated as the slope of the linear regression line of coherent pairs of the detected oscillations of R-R interval and systolic blood pressure (cross correlation coefficient > 0.7) [20,22,23]. The estimation of a global BRS index was based on the weighted mean of all 300–500 individual BRS values derived from all local data segments analysed according to a variance ratio: (variance reduction of RRI/ variance reduction of SBP)².

Statistical methods

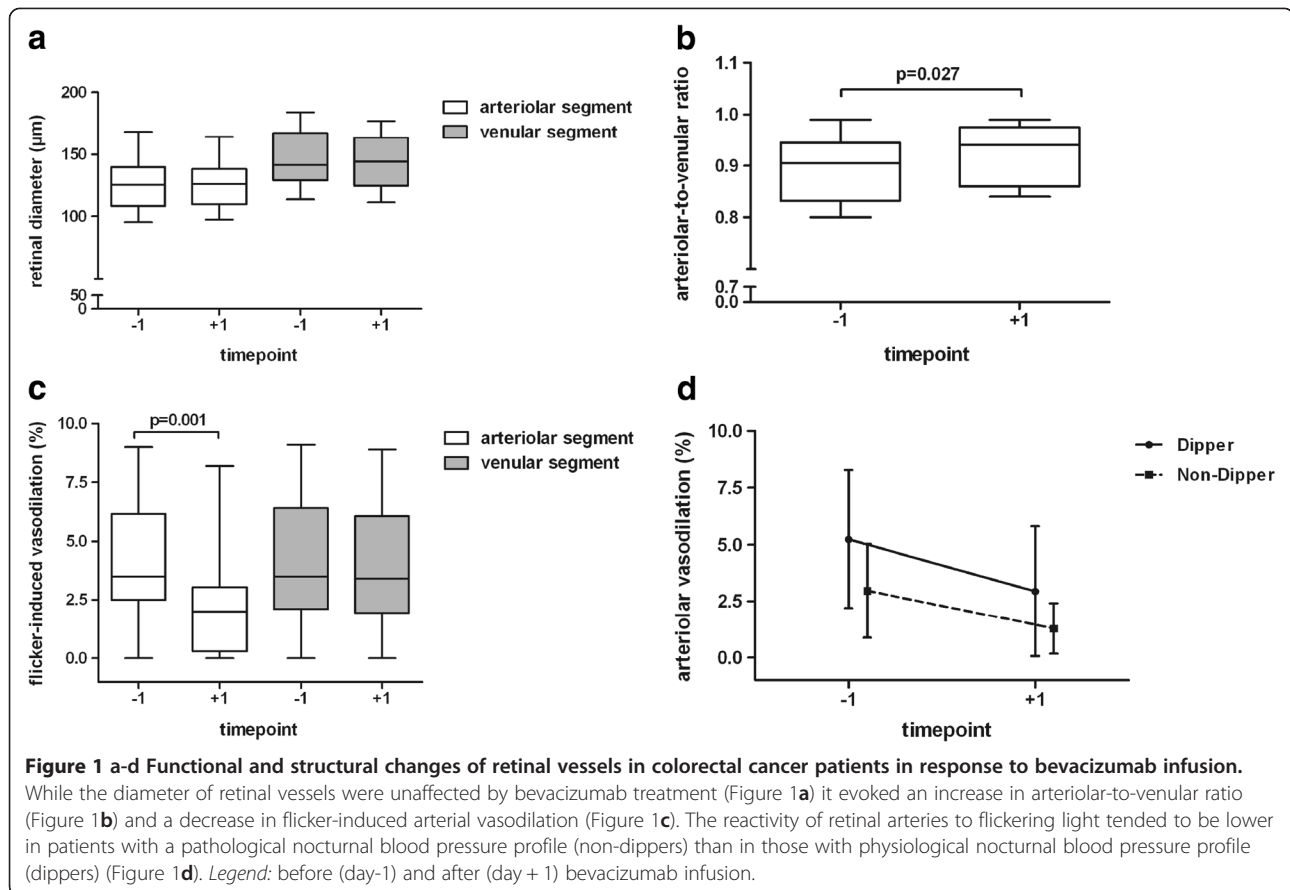
The SPSS package version 17.0 (SPSS Inc., Chicago, IL, USA) was used for all statistical computations. Data are expressed as mean and standard deviation or quartiles. Assuming non-Gaussian distribution, the acute effects of bevacizumab treatment on hemodynamic parameters were analyzed using Wilcoxon and Friedman Test. Spearman correlation coefficients were calculated between changes of continuous variables. Nominal data were compared by Fisher's exact test using cross-tabulation. The overall survival of patients was calculated by the methods of Kaplan-Meier. Patients alive were censored at the day of last contact. A two-tailed *p*-value of < 0.05 was used to indicate significance.

Results

Twenty patients (15 men and 5 women) were enrolled into the study. Eleven patients had rectal and nine colon cancer. The median age was 60.4 [range 45.5–73.9] years. Overall survival of all patients was 23.5 [15.0 – 31.8] months (median [95% CI]). Patients receiving bevacizumab as first line treatment ($n = 9$) had an average survival of 27.1 [21.8 – 32.4] months and those receiving the antibody as second line treatment ($n = 11$) survived on average 21.2 [14.0 – 28.5] months.

Retinal vessel analysis

Static retinal vessel analysis revealed that there was no arteriolar narrowing based on age-normalised reference values of the ARIC (Atherosclerosis Risk in Communities) population. With bevacizumab, no change was observed for arterial diameter, but there was a trend towards a smaller venous diameter (day-1: 146 ± 22 μm vs. day+1: 143 ± 22 μm , $p = 0.062$, Figure 1a) resulting in a significantly increased AVR after bevacizumab administration (day-1: 0.89 ± 0.07 vs. day+1: 0.92 ± 0.06 , $p = 0.027$, Figure 1b). The arterial response to flicker provocation



deteriorated from $4.2 \pm 2.7\%$ to $2.2 \pm 2.2\%$ within 24 hours of VEGF inhibition while maximal venous dilation remained unaffected (Figure 1c).

24 h ABPM profile

The profiles of 24-h ABPM over three recording days are depicted in Figures 2a and 2b. There was a significant reduction of diastolic pressure after bevacizumab administration (Figure 2a). Twenty-four hour heart rate steadily decreased over time (day -1: 80 ± 12 bpm; day 0: 76 ± 11 bpm; day +1: 74 ± 12 bpm; Friedman $p = 0.047$). All other parameters were not affected by bevacizumab. There was a trend towards a higher proportion of patients with a hypertensive blood pressure profile under VEGF inhibition (Figure 2c). The proportion of patients with non-dipping blood pressure at night increased continuously over the three recording days (Figure 2d). Although not statistically significant, there was a trend towards a lower retinal arterial reactivity in non-dippers versus dippers irrespective of bevacizumab infusion ($p > 0.05$, Figure 1d). Decrements of diastolic blood pressure from day 0 to day +1 were neither associated with changes in retinal vascular function ($r = -0.250$, $p = 0.332$) nor with AVR ($r = -0.422$, $p = 0.298$).

Autonomic testing

Sublingual nitroglycerin (Nitro) evoked similar decreases in systolic blood pressure before and after bevacizumab infusion during supine (rest vs. nitro: $p = 0.001$ at day-1, $p = 0.001$ at day+1) and upright position (rest vs. +nitro: $p = 0.001$ at day-1, $p = 0.002$ at day+1) (Figure 3a). The nitroglycerin-mediated fall in diastolic blood pressure during head-up tilt (+nitro) was diminished after bevacizumab treatment (rest vs. +nitro, $p = 0.002$ at day-1; $p > 0.05$ at day+1). Controlled breathing at six cycles per min resulted in shifts from high to low frequency oscillations of R-R interval (LF/HF ratio of R-R interval: $p = 0.010$ at day -1; $p = 0.006$ at day+1) which was not accompanied by adequate adjustments in cardiovagal baroreflex sensitivity (Figure 3b) as observed in healthy individuals [23]. Orthostatic challenge induced significant increases in low frequency oscillations of R-R interval ($p = 0.003$ at day-1; $p = 0.004$ at day+1) which were paralleled by an expected decrease in baroreflex sensitivity ($p = 0.001$ at day-1; $p = 0.003$ at day+1). Cardiovascular outflow as indexed by high frequency oscillations of R-R interval remained unchanged during orthostasis. Bevacizumab treatment did not modify autonomic adjustments of blood pressure control during cardiovascular perturbation.

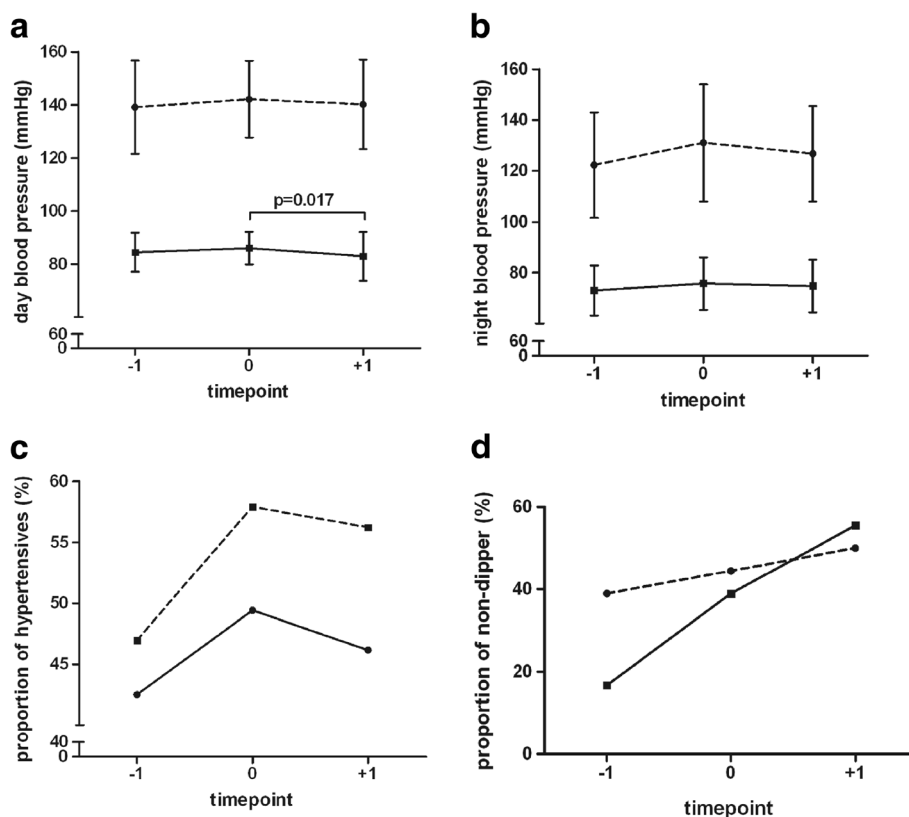


Figure 2 a-d 24-h ambulatory blood pressure profiles in colorectal cancer patients in response to bevacizumab infusion. Diastolic blood pressure during daytime but not during nighttime decreased after bevacizumab infusion (Figure 2a,b). The proportion of individuals with pathological blood pressure profile increased after bevacizumab (Figure 2c,d). Legend: systolic blood pressure (broken lines), diastolic blood pressure (smooth lines); proportions are based on the diagnostic criteria of the European Society of Hypertension and the American Hypertension Association before (day-1), during (day 0) and after (day+1) bevacizumab infusion.

Discussion

Many efforts have been made to target the VEGF pathway in tackling cancer growth and tumor metastasis. Anti-angiogenic therapies using anti-VEGF antibodies or inhibitors of VEGF receptor 2 are increasingly employed in combination with standard therapy for the treatment of advanced cancer. However, predictive markers of efficacy as well as of toxicity have not yet been established. Imaging studies using diffusion contrast enhanced MRI (DCE-MRI) have documented very early changes in response to treatment with VEGF inhibitors [24,25]. However, DCE-MRI technique currently seems too complex and expensive to be incorporated into large multicenter trials [26].

Arterial hypertension is a candidate prognostic biomarker of response to VEGF therapy [27]. One hypothesis is that VEGF inhibitors induce arterial hypertension by increasing vasoconstriction through alteration of nitric oxide (NO) bioavailability. Our method of dynamic analysis of flicker-light induced vasodilation, originally established for application in ophthalmological and cardiovascular research, was therefore used to describe the

acute impact of bevacizumab administration on non-invasively measured retinal endothelial function in relation to blood pressure. We clearly demonstrate that anti-angiogenic treatment of colorectal cancer acutely reduces flicker-light induced vasodilation of retinal arteries independent of blood pressure changes. Furthermore, blood pressure responses to a nitrovasodilator were not altered by bevacizumab.

The vasodilatory response of retinal vessels to diffuse luminance flicker stimulation is considered to be endothelium dependent through release of NO [28]. Hence, a diminished vasodilatory capacity in response to flicker-light stimulation may indicate endothelial dysfunction. In line with our results, endothelial dysfunction was also reported by two other trials in cancer patients receiving either bevacizumab or telatinib - a small molecule angiogenesis inhibitor [10,11]. In these trials endothelial function was determined by laser Doppler fluxmetry using pilocarpine [10] and flow-mediated vasodilation of the brachial artery [11]. Although these methods measure endothelial function in different vascular beds it seems that the detrimental effect of angiogenesis inhibitors on

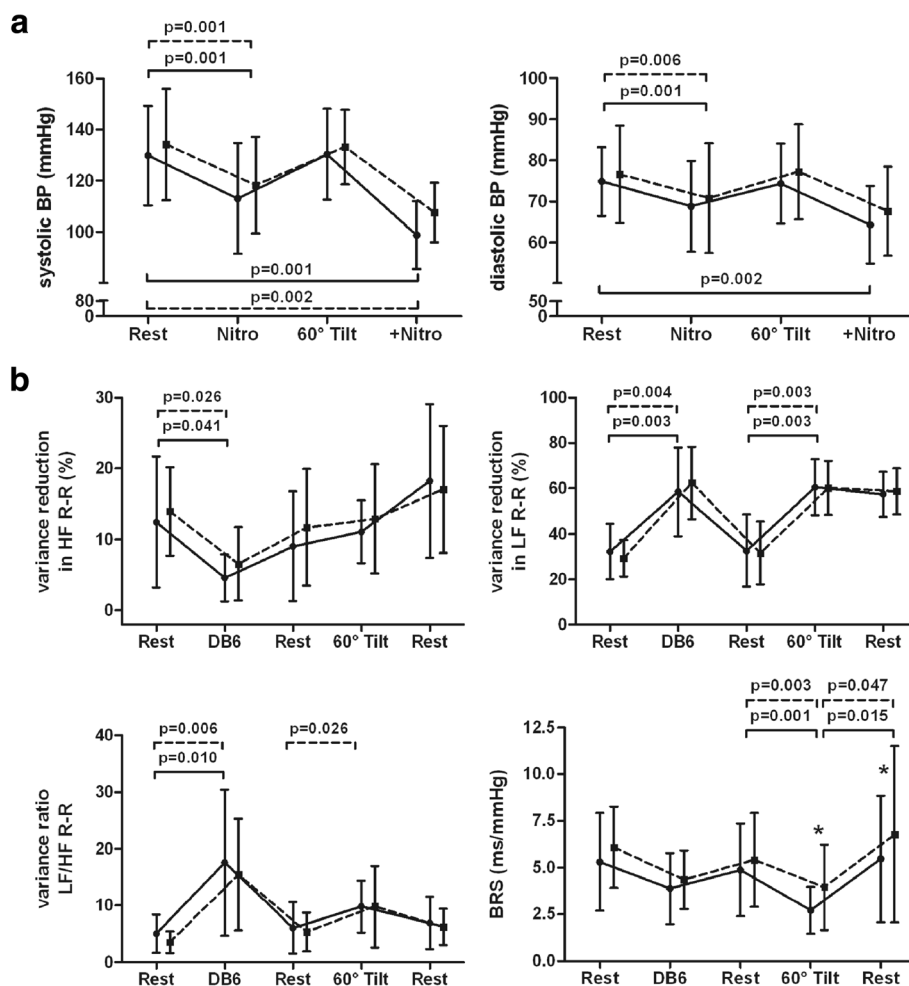


Figure 3 a-b Blood pressures and frequency bands of R-R interval during autonomic testing in response to bevacizumab infusion. The blood pressure response to the vasodilator nitroglycerin was largely preserved except for diastolic blood pressure during passive standing (Figure 3a). Cardiovascular perturbation by deep breathing and head-up tilt evoked a physiological shift from high to low frequency oscillations in R-R interval which was unaffected by bevacizumab treatment (Figure 3b). Adequate adjustments of baroreflex sensitivity were only observed in response to postural change but not to deep breathing irrespective of bevacizumab treatment. Legend: BP, blood pressure; BRS, baroreflex sensitivity; DB6, deep breathing at six cycles per minute; HF, high frequency oscillations; LF, low frequency oscillations; Nitro, nitroglycerin; +Nitro, Nitroglycerin application during head-up tilt before (smooth lines) and after (Broken lines) bevacizumab infusion.

endothelial function is not confined to a certain vascular bed and that arterial endothelial dysfunction is a general phenomenon of pharmacological angiogenesis inhibition. This is not surprising considering that VEGF receptor 2, which confers the hypotensive actions of VEGF [29], is virtually expressed by all endothelial cells.

It is however intriguing that we did not observe any functional change in retinal veins after bevacizumab administration. The lack of effect may be due to a marked regional and segmental heterogeneity in vascular endothelial function [30]. Accordingly, it has been reported that shear stress elicits a smaller NO-dependent vasodilation in veins and venules than in their arterial counterparts even on the same segment [31]. Therefore, the extent of possible changes in endothelium-dependent

vasodilation induced by any stimulus may be inherently less pronounced in the venous system. Furthermore, it is not fully clear whether the endothelial NO-synthase is also modulated by VEGF signaling in the venous circulation and whether VEGF binding to its respective receptors activates the same molecular machinery than in the arterial system. Although VEGF receptors have been identified on endothelial cells of veins a differential pattern of signal transduction has been suggested along the vascular tree [32].

Although elevated blood pressure or hypertension has been the most frequently documented side effect of anti-angiogenic treatment [33] we did not observe meaningful changes in blood pressure nor in short-term blood pressure regulation by anti-angiogenic therapy. The most

probable explanation for this observation is that the frequency of blood pressure elevation is a function of treatment duration [34]. From a range of phase II clinical trials the median interval to incident hypertension was estimated at 4.5 - 6 months from initiation of bevacizumab therapy [35,36]. Under the assumption that there is a progressive evolution of hypertension during chronic VEGF inhibition our findings may imply that endothelial dysfunction is the first step in a sequence of deleterious events evoked by VEGF blockade subsequently evolving into clinical disease [10,11]. *In vitro* and *in vivo* studies clearly demonstrated the role of VEGF in maintaining baseline vascular tone by regulating NO synthesis and prostacyclin release [37]. Loss of action by blockade of VEGF signaling renders endothelial dysfunction highly possible. Consistent with our above postulation, we observed a higher frequency of an abnormal blood pressure profile at night after bevacizumab administration. Furthermore, non-dipping status after anti-angiogenic therapy tended to be related to a stronger impairment of microvascular function. Although the latter data were not statistically significant it still may support our hypothesis of a central role of endothelial dysfunction in the development of VEGF-induced hypertension.

We must acknowledge limitations of our pilot study. The small number of patients precluded us from performing multivariate modeling for analysis of possible interactions, especially co-morbidities, nor from doing correlation analysis of results of functional vascular analysis with outcome data of anti-angiogenic treatment. However, it is important to note that the study design allowed for separating the effect of anti-angiogenic treatment from that of chemotherapy.

Conclusions

We demonstrate that the anti-VEGF antibody bevacizumab acutely impairs endothelial function of retinal arteries irrespective of blood pressure changes. Hence, functional imaging of the retinal microcirculation seems a valuable tool to monitor early effects of anti-VEGF treatment. Considering the close relationship between therapeutic and side effects of VEGF blockade, flicker provocation of retinal vessels may be a valid predictor of therapy success and outcome.

Abbreviations

ABPM: 24-hour ambulatory blood pressure monitoring; DCE-MRI: Diffusion contrast enhanced MRI; HF: High-frequency; LF: Low-frequency; MTRS: Multiple trigonometric regressive spectral analysis; NO: Nitric oxide; VEGF: Vascular endothelial growth factor.

Competing interests

The study was supported by the University Hospital Carl Gustav Carus. The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript. Gunnar Folprecht received honoraries for lectures and advisory boards from Roche / Genentech. All other authors have declared that no competing interests exist. Additionally,

Focke Ziemssen has received speaker's fees or consultancy honoraria from Alcon, Alimera, Allergan, Biogen, Bayer, Pfizer and Novartis.

Authors' contribution

MR performed the statistical analysis and drafted the manuscript. RH assisted in the statistical analyses. TZ, FZ and GF designed the trial. TZ, KT, GF participated in the coordination of the trial and revised the manuscript for intellectual content. GE and HR revised the manuscript for intellectual content. All authors revised and approved the final manuscript.

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References

1. Miller K, Wang M, Gralow J, Dickler M, Cobleigh M, Perez EA, Shenker T, Cella D, Davidson NE: **Paclitaxel plus bevacizumab versus paclitaxel alone for metastatic breast cancer.** *N Engl J Med* 2007, **357**:2666-2676.
2. Sandler A, Gray R, Perry MC, Brahmer J, Schiller JH, Dowlati A, Lilienbaum R, Johnson DH: **Paclitaxel-carboplatin alone or with bevacizumab for non-small-cell lung cancer.** *N Engl J Med* 2006, **355**:2542-2550.
3. Longo R, Sarmiento R, Fanelli M, Capacetti B, Gattuso D, Gasparini G: **Anti-angiogenic therapy: rationale, challenges and clinical studies.** *Angiogenesis* 2002, **5**:237-256.
4. Escudier B, Pluzanska A, Koralewski P, Ravaud A, Bracarda S, Szczyluk C, Chevreau C, Filipek M, Melichar B, Bajetta E, et al: **Bevacizumab plus interferon alfa-2a for treatment of metastatic renal cell carcinoma: a randomised, double-blind phase III trial.** *Lancet* 2007, **370**:2103-2111.
5. Hurwitz H, Fehrenbacher L, Novotny W, Cartwright T, Hainsworth J, Heim W, Berlin J, Baron A, Griffing S, Holmgren E, et al: **Bevacizumab plus irinotecan, fluorouracil, and leucovorin for metastatic colorectal cancer.** *N Engl J Med* 2004, **350**:2335-2342.
6. Hurwitz H, Saini S: **Bevacizumab in the treatment of metastatic colorectal cancer: safety profile and management of adverse events.** *Semin Oncol* 2006, **33**:S26-S34.
7. Nalluri SR, Chu D, Keresztes R, Zhu X, Wu S: **Risk of venous thromboembolism with the angiogenesis inhibitor bevacizumab in cancer patients: a meta-analysis.** *JAMA* 2008, **300**:2277-2285.
8. Scappaticci FA, Skillings JR, Holden SN, Gerber HP, Miller K, Kabbinnar F, Bergsland E, Ngai J, Holmgren E, Wang J, Hurwitz H: **Arterial thromboembolic events in patients with metastatic carcinoma treated with chemotherapy and bevacizumab.** *J Natl Cancer Inst* 2007, **99**:1232-1239.
9. Facemire CS, Nixon AB, Griffiths R, Hurwitz H, Coffman TM: **Vascular endothelial growth factor receptor 2 controls blood pressure by regulating nitric oxide synthase expression.** *Hypertension* 2009, **54**:652-658.
10. Mourad JJ, des Guetz G, Debbabi H, Levy BI: **Blood pressure rise following angiogenesis inhibition by bevacizumab. A crucial role for microcirculation.** *Ann Oncol* 2008, **19**:927-934.
11. Steeghs N, Gelderblom H, Roodt JO, Christensen O, Rajagopalan P, Hovens M, Putter H, Rabelink TJ, De Koning E: **Hypertension and rarefaction during treatment with telatinib, a small molecule angiogenesis inhibitor.** *Clin Cancer Res* 2008, **14**:3470-3476.
12. Scartozzi M, Galizia E, Chiellini S, Giampieri R, Berardi R, Pierantoni C, Cascinu S: **Arterial hypertension correlates with clinical outcome in**

- colorectal cancer patients treated with first-line bevacizumab. *Ann Oncol* 2009, **20**:227–230.
13. Ziemssen T, Reichmann H: **Cardiovascular autonomic testing in extrapyramidal disorders.** *J Neurol Sci* 2011, **310**:129–132.
 14. Freeman R, Wieling W, Axelrod FB, Benditt DG, Benarroch E, Biaggioni I, Cheshire WP, Chelimsky T, Cortelli P, Gibbons CH, et al: **Consensus statement on the definition of orthostatic hypotension, neutrally mediated syncope and the postural tachycardia syndrome.** *Clin Auton Res* 2011 Apr, **21**(2):69–72.
 15. Vilser W, Nagel E, Lanzl I: **Retinal Vessel Analysis—new possibilities.** *Biomed Tech (Berl)* 2002, **47**(Suppl 1 Pt 2):682–685.
 16. Reimann M, Prieur S, Lippold B, Bornstein SR, Reichmann H, Julius U, Ziemssen T: **Retinal vessel analysis in hypercholesterolemic patients before and after LDL apheresis.** *Atheroscler Suppl* 2009, **10**:39–43.
 17. Reimann M, Julius U, Bornstein SR, Fischer S, Reichmann H, Rudiger H, Ziemssen T: **Regular lipoprotein apheresis maintains residual cardiovascular and microvascular function in patients with advanced atherosclerotic disease.** *Atherosclerosis Suppl* 2013, **14**:135–141.
 18. Ohkubo T, Hozawa A, Yamaguchi J, Kikuya M, Ohmori K, Michimata M, Matsubara M, Hashimoto J, Hoshi H, Araki T, et al: **Prognostic significance of the nocturnal decline in blood pressure in individuals with and without high 24-h blood pressure: the Ohasama study.** *J Hypertens* 2002, **20**:2183–2189.
 19. Schmidt C, Berg D, Prieur S, Junghanns S, Schweitzer K, Globas C, Schols L, Reichmann H, Ziemssen T: **Loss of nocturnal blood pressure fall in various extrapyramidal syndromes.** *Movement disorder* 2009, **24**:2136–2142.
 20. Reimann M, Friedrich C, Gasch J, Reichmann H, Rudiger H, Ziemssen T: **Trigonometric Regressive Spectral Analysis reliably Maps Dynamic Changes in Baroreflex Sensitivity and Autonomic Tone: The Effect of Gender and Age.** *PLoS One* 2010, **5**:e12187. doi:12110.11371/journal.pone.0012187.
 21. Ziemssen T, Reimann M, Gasch J, Ruediger H: **Trigonometric Regressive Spectral Analysis An innovative tool for Evaluating the Autonomic Nervous System.** *J Neural Transm* 2013, in press.
 22. Gasch J, Reimann M, Reichmann H, Rudiger H, Ziemssen T: **Determination of baroreflex sensitivity during the modified Oxford maneuver by trigonometric regressive spectral analysis.** *PLoS One* 2011, **6**:e18061.
 23. Friedrich C, Rüdiger H, Schmidt C, Herting B, Prieur S, Junghanns S, Schweitzer K, Globas C, Schöls L, Berg D, Reichmann H, Ziemssen T: **Baroreflex sensitivity and power spectral analysis during autonomic testing in different extrapyramidal syndromes.** *Mov Disord* 2010, **25**(3):315–24.
 24. Lockhart AC, Rothenberg ML, Dupont J, Cooper W, Chevalier P, Sternas L, Buzenet G, Koehler E, Sosman JA, Schwartz LH, et al: **Phase I study of intravenous vascular endothelial growth factor trap, aflibercept, in patients with advanced solid tumors.** *J Clin Oncol* 2010, **28**:207–214.
 25. Morgan B, Thomas AL, Drevs J, Hennig J, Buchert M, Jivan A, Horsfield MA, Mross K, Ball HA, Lee L, et al: **Dynamic contrast-enhanced magnetic resonance imaging as a biomarker for the pharmacological response of PTK787/ZK 222584, an inhibitor of the vascular endothelial growth factor receptor tyrosine kinases, in patients with advanced colorectal cancer and liver metastases: results from two phase I studies.** *J Clin Oncol* 2003, **21**:3955–3964.
 26. Hamstra DA, Rehemtulla A, Ross BD: **Diffusion magnetic resonance imaging: a biomarker for treatment response in oncology.** *J Clin Oncol* 2007, **25**:4104–4109.
 27. Murukesh N, Dive C, Jayson GC: **Biomarkers of angiogenesis and their role in the development of VEGF inhibitors.** *Br J Cancer* 2010, **102**:8–18.
 28. Dornier GT, Garhofer G, Kiss B, Polska E, Polak K, Riva CE, Schmetterer L: **Nitric oxide regulates retinal vascular tone in humans.** *Am J Physiol Heart Circ Physiol* 2003, **285**:H631–H636.
 29. Li B, Ogasawara AK, Yang R, Wei W, He GW, Zioncheck TF, Bunting S, De Vos AM, Jin H: **KDR (VEGF receptor 2) is the major mediator for the hypotensive effect of VEGF.** *Hypertension* 2002, **39**:1095–1100.
 30. Boegehold MA: **Heterogeneity of endothelial function within the circulation.** *Curr Opin Nephrol Hypertens* 1998, **7**:71–78.
 31. Kuo L, Arko F, Chilian WM, Davis MJ: **Coronary venular responses to flow and pressure.** *Circ Res* 1993, **72**:607–615.
 32. Laham RJ, Li J, Tofukuji M, Post M, Simons M, Sellke FW: **Spatial heterogeneity in VEGF-induced vasodilation: VEGF dilates microvessels but not epicardial and systemic arteries and veins.** *Ann Vasc Surg* 2003, **17**:245–252.
 33. Sica DA: **Angiogenesis inhibitors and hypertension: an emerging issue.** *J Clin Oncol* 2006, **24**:1329–1331.
 34. Grothey A, Sugrue MM, Purdie DM, Dong W, Sargent D, Hedrick E, Kozloff M: **Bevacizumab beyond first progression is associated with prolonged overall survival in metastatic colorectal cancer: results from a large observational cohort study (BRiTE).** *J Clin Oncol* 2008, **26**:5326–5334.
 35. Vaklavas C, Lenihan D, Kurzrock R, Tsimberidou AM: **Anti-vascular endothelial growth factor therapies and cardiovascular toxicity: what are the important clinical markers to target?** *Oncologist* 2010, **15**:130–141.
 36. Yang JC, Haworth L, Sherry RM, Hwu P, Schwartzentruber DJ, Topalian SL, Steinberg SM, Chen HX, Rosenberg SA: **A randomized trial of bevacizumab, an anti-vascular endothelial growth factor antibody, for metastatic renal cancer.** *N Engl J Med* 2003, **349**:427–434.
 37. Horowitz JR, Rivard A, van der Zee R, Hariawala M, Sheriff DD, Esakof DD, Chaudhry GM, Symes JF, Isner JM: **Vascular endothelial growth factor/vascular permeability factor produces nitric oxide-dependent hypotension. Evidence for a maintenance role in quiescent adult endothelium.** *Arterioscler Thromb Vasc Biol* 1997, **17**:2793–2800.

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