Temporal pO2 instability was observed in all experiments. Red cell fluxes in microvessels surrounding pO2 measurement locations were measured using fluorescently labeled red cells. Temporal pO2 instability was observed in all experiments. Median pO2 was inversely related to radial distance from microvessels. Transient fluctuations above and below 10 mm Hg were consistently seen, except in one experiment near the microvessels. Median pO2 was <5 mm Hg. Vascular stasis was not seen in these experiments. These results show that fluctuations in red cell flux, as opposed to vascular stasis, can cause temporal variations in pO2 that extend from perivascular regions to the maximum oxygen diffusion distance. (Cancer Res 2006; 66(4): 2219-23)

Introduction

Tumor hypoxia has classically been categorized as developing in two different forms: “chronic” and “acute.” Chronic hypoxia has been attributed to large diffusion distances between tumor microvessels whereas acute hypoxia has been thought to be the result of transient vascular occlusion (1-4). We and others have argued that this categorization misleadingly implies that two independent mechanisms are responsible for hypoxia (5-7). Tumor hypoxia develops because oxygen demand exceeds supply (8-10). Causes of supply deficiency include abnormal vascular geometry (8), relatively low vascular density as compared with most normal tissues (11), and relative paucity of arteriolar supply vessels (12). These features of tumor vasculature contribute to exaggerated longitudinal vascular oxygen gradients, leading to hypoxia within some microvessels (13-15). Previously, we showed that temporal fluctuations in microvessel red cell flux cause variations in perivascular pO2 (16). Using theoretical simulations, we predicted that the temporal flux variations would also cause fluctuations in pO2 throughout the subtended tissue region with a substantial fraction of tissue exhibiting transient episodes of hypoxia. In this study, we conducted experiments to test this prediction.

Materials and Methods

The protocol was approved by the Duke University Animal Care and Use Committee. R3230Ac mammary carcinoma cells were transplanted into dorsal skin-fold window chambers of female Fischer 344 rats as previously described (17). Circular 1-cm diameter segments of epidermis were removed from opposing surfaces of the skin flap, leaving one to two fascial layers, containing s.c. vasculature. The two halves of an anodized aluminum chamber were sutured to this tissue such that the dissected circular fascial layer was contained within the window portion of the chamber. R3230Ac tumors were 1-cm diameter fragments of 0.1-mm diameter, obtained from a donor animal, were placed onto one side of the fascial layer. The two surfaces were then covered with glass coverslips. Tumors were allowed to grow until they reached a diameter of 2 to 3 mm, which typically required 7 to 9 days. During the period of tumor growth, animals were housed individually in an environmental chamber maintained at a temperature of 34°C and 50% humidity. Recessed, gold-plated cathode tip oxygen microelectrodes (tip diameters of 6-10 μm) were manufactured according to the method of Linsenmeier and Yancey (18). Electrodes were calibrated before and after each experiment in saline bubbled with gas mixtures of 0%, 5%, 15%, and 21% O2.

Experiments in which calibrations drifted >20% from pre-to-post measurement time points were not used. All electrodes showed linear relationships between current generated and pO2 with an average sensitivity of 2.99 ± 1.31 pA/mm Hg. Current was measured using a Chemical Microsensor (model 1201, Diamond General Development Corp., Ann Arbor, MI).

Erythrocytes were fluorescently labeled with DiI using the method of Unthank et al. (19). DiI was dissolved in ethanol to a concentration of 0.5 mg/mL. Blood from an anesthetized donor rat was withdrawn via cardiac puncture and the erythrocytes were isolated by centrifugation and washed in PBS. One hundred microliters of packed erythrocytes were suspended in 10 mL of PBS and 100 μL of DiI solution were added. The suspension was incubated at room temperature for 30 minutes. The labeled cells were removed from the DiI-containing solution by washing with PBS and additional centrifugation. Cells were kept in the dark once DiI was added to prevent photobleaching.

 Videomicroscopy of the window chamber was used to measure red cell fluxes and to place microelectrodes to measure interstitial pO2. A Zeiss intravital microscope workstation (Zeiss Axioskop, New York, NY) equipped with optics for transillumination and fluorescence microscopy was used to visualize the microvasculature and labeled erythrocytes during experiments. For transillumination, images were obtained using a CCD video camera (MTI CCD-72, Dage MTI, Michigan City, IN). For epifluorescence, a silicon-intensified tube camera was used (model C2400-08, Hamamatsu Photonics, Hamamatsu City, Japan). Images were recorded onto SVHS videotape (SVO-9500MD Sony Corporation of America, San Jose, CA) for later assessment of red cell fluxes.

Liposomes of 100-nm average diameter were prepared (16) for use as a fluorescent blood pool marker to assist in identifying vascular networks. The liposomes contained polyethylene glycol in the membrane to extend circulation time. The composition was hydrogenated soy...
phosphatidylethanolamine/cholesterol/polyethylene glycol (2000 M)–
Lipids were purchased from Avanti Polar Lipids (Alabaster, AL).

Animals were anesthetized with pentobarbital sodium at a dose of 40 to
50 mg/kg and a femoral artery and vein were catheterized for monitoring of
blood pressure and i.v. access, respectively. The animals were placed in
lateral recumbency onto a temperature-controlled heating pad that was
mounted on the microscope stage. Fluorescently labeled red cells were
administered i.v. at a concentration to yield ~1% labeled fraction. Blood
pressure and heart rate were measured using digital signals acquired from a
blood pressure transducer and recorded to a PC (CODAS Micro Software,
Akron, OH). A tracing of all blood vessels containing labeled liposomes was
made from the video monitor.

The glass coverslip was removed and a suffusion medium (Earle’s
balanced salt solution) bubbled with 95% air and 5% CO2 at 34°C (normal
skin temperature) was set to flow across the window at a rate of ~1 ml/min.
A silver-chloride anode was sewn into a s.c. pocket into the hind limb of the
rat. A recessed-tip glass microelectrode was placed into a stage-mounted
micromanipulator (Model M0102E, Narishige, Inc., Narishige, Japan) and
inserted into the medium just beneath the surface. Readings represented an
in vivo measurement equivalent to air (21% O2). The suffusion medium was
replaced with the same medium, but in this case bubbled with 95% N2 and
5% CO2. A tent made of Saran Wrap (Dow Brands, Indianapolis, IN) was
placed around the microscope and stage and nitrogen was blown through
the tent to minimize oxygen diffusion to the tissue. The microelectrode was
reinserted into the suffusion medium and stepped through the medium until
it reached the tumor surface.

The microelectrode was placed into an extravascular region, >50 µm
below the tissue surface, to minimize perturbations of oxygen gradients
between the tissue and the overlying suffusion medium (20). For the next
60 to 80 minutes, pO2 values were continuously recorded. Simultaneously,
1-minute videotape segments of the microvessels containing circulating
fluorescently labeled red cells and liposomes were made every 5 minutes.
Light was turned off between measurements to minimize photobleaching.

At the end of each experiment, animals were euthanized, leaving the
oxygen electrode in place. The current dropped to a value equivalent to zero
pO2 by 10 minutes after death. This value was used as the in vivo zero value
as previously described (13). The pO2 values observed during each
experiment were derived from the in vivo calibration data as previously
described (12, 13, 16, 21). After each experiment, the radial distance from the
measurement point to each microvessel was determined from the tracings of the vessels, with calibration from a stage micrometer.

Fluorescent red cell fluxes were determined by counting the number of
fluorescently labeled cells that passed a preset position near the middle
of each vessel segment (midpoint between branch points) over 1-minute
intervals (16). Total red cell flux was determined by dividing the number of
counted fluorescent cells by the labeled fraction as determined by flow
cytometric analysis of a peripheral blood sample obtained after the
fluorescently labeled red cells were administered. Two samples were obtained
in each experiment, one just after the cells were administered and the second
at the end of the experiment. These values did not change appreciably during
the experiments, indicating that systemic levels of circulating fluorescently
labeled red cells were stable over the period of observation (data not shown).

Results

Eight successful experiments were conducted. Another seven
were not completed because of anesthetic deaths (n = 2) or
technical complications (n = 5). Aside from the two anesthetic
deaths, animals maintained a stable level of anesthesia throughout
these experiments. Blood pressures and heart rates averaged 109 ±
8.3 mm Hg and 389 ± 25 bpm, respectively, and were stable over...
the time of measurement (data not shown). These values are well within limits for awake, unanesthetized rats of this strain (22) and are typical for this anesthetic regimen in our prior experience (12, 16, 21).

Examples of results are shown in Figs. 1 and 2. In the first example, a network of three vessels was identified (Fig. 1A). One of the vessels (V3) carried a much larger red cell flux than the other two (Fig. 1B). During the period of observation, this microvessel exhibited two brief periods of flow reversal, at 25 and 35 minutes. Effects on interstitial pO2 value were discernable, particularly at 25 minutes (Fig. 1B). The influence of red cell flux on interstitial pO2 can be examined by removing the time factor and plotting pO2 as a function of red cell flux (Fig. 1C and D). The relationship between the two was linear when comparing data for each vessel alone (Fig. 1C and D) and for total red cell flux (flux added together for all three microvessels; data not shown).

In the second example, a network of four vessels was examined (Fig. 2A). Again, pO2 and red cell flux were temporally unstable (Fig. 2B). The relationship between pO2 and red cell flux was complex, indicating a mixture of relatively oxygenated and deoxygenated blood entering this network. The relationship between red cell flux and pO2 showed a positive slope for vessel 1 (V1, Fig. 2C), a negative slope for vessel 4 (V4, Fig. 2D), and no detectable relationship for vessels 2 and 3 (data not shown). There was no significant relationship between total red cell flux and pO2 in this case (data not shown; \( r^2 = 0.02 \)).

In all experiments, pO2 was temporally unstable (Fig. 3). However, the degree of fluctuation varied from one experiment to another. To examine whether the magnitude of the fluctuations was influenced by proximity to the vasculature, we plotted the median pO2 and range as a function of distance from the microvessel in the network with the highest red cell flux (Fig. 4). The vessel with the highest red cell flux would be expected to have the greatest influence on tissue pO2, in such circumstances, as this vessel would have the greatest oxygen carrying capacity. Figure 4 includes variations in perivascular pO2 (measurements made on the outer wall of microvessels) obtained from a prior publication using the same tumor model (16). Neither the median nor the range of the fluctuations seemed to be predictably influenced by radial distance. However, it is noteworthy that in experiment 7, the distance from the vessel was maximal (140 μm) and pO2 variations in the range below 5 mm Hg were observed (Fig. 3).

**Discussion**

This study confirms our previous theoretical prediction (16) that fluctuations in red cell flux lead to temporal variations in extravascular pO2 ranging up to the diffusion distance of oxygen. These variations in pO2 are superimposed on the decline in pO2 with distance from the nearest microvessel (Fig. 4). The results are consistent with prior work indicating that fluctuation in red cell flux in tumor microvessels is common and that this leads to changes in extravascular pO2 that are of potential radiobiological importance. In other studies using tumors growing in animal flanks, we observed similar kinetics of pO2 fluctuation (23).
In these experiments, steady-state chronic hypoxia was not observed even at the limit of the oxygen diffusion distance. Experiment 7 showed a condition of chronic hypoxia at a point 140 μm from the vessel with highest red cell flux but fluctuations of ±2 mm Hg were observed in that case as well. In one case (Fig. 2D), pO2 showed a negative correlation with red cell flux in a particular vessel, suggesting that the vessel was acting as an oxygen sink by supplying deoxygenated blood. We have recently identified anastomoses between hypoxic and more aerobic vessels in a mouse mammary carcinoma line, grown in skin-fold window chambers, using hyperspectral imaging to measure hemoglobin saturation in vascular networks (24). These hypoxic vessels were emerging onto the tumor surface from the underlying tumor and presumably were deoxygenated because they had traversed through the tumor before reaching the surface that we were visualizing. We observed these vessels to be consistently hypoxic over several days of observation. It is likely that the diametrically opposed relationship between pO2 and red cell flux in two different vessels of the same network, as depicted in Fig. 2, was due to a similar vascular arrangement.

The mechanisms underlying the observed variations in red cell flux remain to be determined. Potential mechanisms include arterial vasomotion (25) or vascular remodeling occurring concomitantly with angiogenesis (26). Studies using the dye mismatch method, which reflects variations in perfusion (but does not directly indicate oxygenation), suggest that the variations typically involve groups of vessels as opposed to single isolated vessels (7, 27), consistent with our previous findings (16).

The presence of intermittent hypoxia has several implications with respect to tumor biology and therapy. Genetic instability may arise as a result of free radicals generated during hypoxia-reoxygenation injury (28) and/or as a result of reduced capacity to repair DNA damage (29). Intermittent blood flow and oxygenation may contribute to drug resistance (7). Hypoxia-mediated cytotoxins are being developed to

**Figure 3.** Temporal measurement of pO2 in all experiments. Dashed line, 10 mm Hg threshold. With one exception (experiment 7), pO2 values fluctuated across the 10 mm Hg boundary at least once during the period of observation.
target cells that escape radiation treatment in hypoxic regions (30). Fluctuations in tissue oxygenation may compromise the effectiveness of this approach if regions of tumor are not hypoxic during the time of drug exposure. Recently, our group has shown that reoxygenation postirradiation can cause up-regulation of hypoxia inducible factor 1-mediated gene transcription (31).

The prevalence of fluctuating hypoxia in human tumors is unknown. Its characteristics in preclinical models may depend on tumor type (32). Recently, we compared the kinetics of pO2 fluctuations in three rat tumor lines: the R3230Ac mammary tumor, the 9L glioma, and a fibrosarcoma (33, 34). The dominant frequencies were low, <0.3 cycles per minute, for all three tumor types but amplitudes varied widely. Similar results have been reported for early passage human melanoma xenografts (35, 36). Fluctuations in red cell flux have been measured using laser Doppler flowmetry in several human tumor xenograft lines and in human tumors (37–39). It remains to be determined whether fluctuations in pO2 are present and lead to radiobiologically significant fluctuations in tumor hypoxia in human subjects.

References


Direct Demonstration of Instabilities in Oxygen Concentrations within the Extravascular Compartment of an Experimental Tumor

Jennifer Lanzen, Rod D. Braun, Bruce Klitzman, et al.


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