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High-speed Label-free Functional Photoacoustic Microscopy of Mouse Brain in Action

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Abstract

We present fast functional photoacoustic microscopy (PAM), which is capable of three-dimensional high-resolution high-speed imaging of the mouse brain, complementary to other imaging modalities. A single-wavelength pulse-width-based method was implemented to image blood oxygenation with capillary-level resolution and a one-dimensional imaging rate of 100 kHz. We applied PAM to image the vascular morphology, blood oxygenation, blood flow, and oxygen metabolism in the brain in both resting and stimulated states.

Many biomedical imaging techniques, especially small-animal functional magnetic resonance imaging (fMRI), two-photon microscopy (TPM), and wide-field optical microscopy, have profoundly impacted hemodynamic studies of the mouse brain by providing structural, blood oxygenation, and flow dynamic information at various length scales. However, small-animal fMRI is insufficient to resolve brain hemodynamic activities at microscopic length scales finer than 50 μm ¹; phosphorescence-lifetime-based TPM suffers from slow measurement of blood oxygenation²; and wide-field optical microscopy lacks depth resolution³.

Given these limitations, photoacoustic (PA) tomography (PAT) can play a complementary role. Previously reported PAT techniques variously lacked capillary-level resolution, wide-field imaging speed, or blood oxygenation imaging capability^{4–8}. Here, we present fast functional photoacoustic microscopy (PAM), which is capable of high-resolution high-speed imaging of the mouse brain through an intact skull *in vivo*. PAM has achieved a lateral spatial resolution of $\sim 3 \mu\text{m}$, which is 5 times finer than that of our previous fast-scanning

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AUTHOR CONTRIBUTIONS

J.Y., J.Z. and L.V.W. conceived and designed the study. J.Y., J.M.Y., K.M., L.W. and C.H. constructed the imaging system. J.Y., T.W., and L.L. performed the experiments and analyzed the data. J.Y., L.W. and L.V.W. wrote the manuscript. L.V.W. supervised the whole study. All authors discussed the conceptual and practical implications of the methods.

COMPETING FINANCIAL INTERESTS

L. V. Wang has a financial interest in Endra, Inc., and Microphotoacoustics, Inc., which, however, did not support this work. K. I. Maslov also has a financial interest in Microphotoacoustics, Inc. The other authors declare no competing financial interests.

system⁷, 25 times finer than that of our previous acoustic-resolution system⁶, and more than 35 times finer than that of ultrasound-array-based photoacoustic computed tomography⁵. By using a new single-wavelength pulse-width-based method, PAM allows three-dimensional (3D) blood oxygenation imaging with capillary-level resolution at a one-dimensional (1D) imaging rate of 100 kHz. PAM's blood oxygenation imaging speed is 50 times higher than that of our fast-scanning PAM⁸, 100 times higher than that of our acoustic-resolution system⁶, and more than 500 times higher than that of phosphorescence-lifetime-based TPM².

In PAM, both the excitation laser beams and the detection acoustic axis are confocally steered by a customized water-immersible MEMS (i.e., microelectromechanical system) scanning mirror (Fig. 1a, Supplementary Fig. 1, Supplementary Note 1). The lateral resolution—in the direction perpendicular to the acoustic axis—is $\sim 3\ \mu\text{m}$ at the optical focus, and the axial resolution—in the direction along the acoustic axis—is $\sim 15\ \mu\text{m}$. The high laser pulse repetition rate of 500 kHz, 5 times greater than that in our previous work⁷, enables dense sampling for morphological capillary-resolution imaging over a large scanning range. With a 1D time-resolved imaging rate of 500 kHz, PAM has achieved a 2D frame rate of 400 Hz over a $\sim 3\ \text{mm}$ scanning range, and a 3D volumetric rate of 1 Hz over a $3\times 2\ \text{mm}^2$ field of view (Fig. 1b).

By using a new single-wavelength pulse-width-based method (PW- sO_2), PAM is capable of high-speed imaging of the oxygen saturation of hemoglobin (sO_2) (Fig. 1c, Online Methods). The two forms of hemoglobin, oxy- and deoxy-hemoglobin (HbO_2 and HbR), have different saturation intensities, defined as the excitation intensity that reduces the absorption coefficient to half its initial value (Supplementary Note 2, Supplementary Fig. 2)⁹. When first excited by a picosecond pulse and subsequently by a nanosecond pulse of the same wavelength and pulse energy, HbO_2 and HbR display different saturation levels (Supplementary Fig. 3). From PA signals acquired with the two laser pulses, the relative concentrations of HbO_2 and HbR are quantified, and thus sO_2 can be computed. PW- sO_2 does not suffer wavelength-dependent optical attenuation as the traditional wavelength-tuning method does. Nevertheless, the maximum PW- sO_2 imaging depth is limited by optical attenuation to the point where saturation becomes insufficient.

A $5\times 10\ \text{mm}^2$ region of the mouse brain was imaged by PAM through an intact skull with the scalp removed (Fig. 1d, Supplementary Video 1) (acquisition time: ~ 15 seconds). The optical focal plane was fixed at $\sim 250\ \mu\text{m}$ beneath the skull surface. The imaging parameters for all the key experiments are summarized in Supplementary Table 1. By additional depth-scanning of the optical focal zone with a z -step size of $100\ \mu\text{m}$, PAM provided an imaging depth of $\sim 0.7\ \text{mm}$ (Fig. 1e, Supplementary Fig. 4, Supplementary Video 2), giving an effective pixel count of ~ 47 in focus along the depth direction. The optical scattering of brain tissue degraded the lateral resolution and image contrast of PAM with increasing imaging depth, as in any depth-resolved optical microscopy; thus, deep capillaries cannot be resolved by the current version of PAM. PAM of the brain vasculature was confirmed by TPM (Supplementary Fig. 5). The acoustic sectioning of PAM could not resolve the blood vessels along the z -axis as well as the optical sectioning of TPM. The skull degraded the

image quality of PAM by blurring the optical focusing and attenuating the PA signal (Supplementary Note 3, Figs. S6 and S7).

sO₂ of the mouse brain was mapped vessel-by-vessel by using the pulse-width-based method (Fig. 1f) (acquisition time: ~40 seconds). A pulse energy of 400 nJ was used for all the sO₂ measurements unless otherwise stated. The optical fluence at the optical focus was estimated to be ~0.3 J/cm². The non-saturated PA signal acquired with nanosecond excitation was used to correct for optical attenuation and the laser spot size. We observed that the averaged sO₂ in the skull vessels was lower than that in the cortical vessels, consistent with the low-oxygenation microenvironment in bone marrow¹⁰.

The PW-sO₂ method was validated on blood phantoms, with an average measurement error of ~2.7% (Fig. 1g). The PW-sO₂ method was also compared with the traditional two-wavelength-based method (TW-sO₂) *in vivo*, with an average difference of <5% for superficial vessels (Supplementary Fig. 8)⁴. To quantify the underestimation of PW-sO₂ induced by light attenuation with increasing depth, we measured the sO₂ in blood phantoms and in a mouse ear *in vivo* with pulse energies from 50 nJ to 1000 nJ (Fig. 1h, Figs. S9–S10). When the pulse energy was 300 nJ, the measurement error was ~3% for absolute PW-sO₂.

We carefully investigated the potential for tissue damage induced by PAM. First, bright-field microscopy of a single layer of mouse RBCs, before and after the PAM imaging, confirmed that the PAM-imaged RBCs were intact, with clear donut shapes (Supplementary Fig. 11). Second, TPM of a mouse brain after PAM imaging with the laser pulse energy intentionally increased to 1000 nJ ruled out its potential to induce bleeding (Supplementary Fig. 12a). A few vessels were imaged by TPM but not by PAM, probably due to the lack of RBC perfusion¹¹. Last, standard H&E histology on a mouse brain after PAM imaging (Online Methods) showed no burn damage to the brain tissue (Supplementary Fig. 12b). As a positive control, a part of the brain was intentionally burned and was also studied histologically. Representative histological slices from the inside and outside of the burned area, as well as the imaged area, were compared, revealing no burns in the imaged area (Supplementary Fig. 13).

Directly imaging hyperaemia in the brain can help understand neurovascular coupling. Here, we demonstrate the high-speed functional imaging capability of PAM by studying mouse cortical hemodynamic responses to electrical stimulations to the hindlimbs (Supplementary Fig. 1a). Upon stimulations, the PA amplitude in the contralateral somatosensory region started to increase until the end of the stimulations (Fig. 2a, Supplementary Video 3). Meanwhile, the ipsilateral somatosensory region followed a similar trend but responded more weakly (Figs. S14a–b), suggesting vascular interconnection between the two hemispheres¹². We also observed that the sagittal sinus region responded to both left and right hindlimb stimulations, possibly due to the fact that it drains blood from both hemispheres simultaneously¹². The depth-resolved responses revealed that the responding region covered a depth range of 50–150 μm beneath the cortical surface (Fig. 2b). The deep capillary beds showed stronger amplitude responses than the major arteries and veins (Figs. S15a–b)³.

Meanwhile, the artery dilated substantially in the contralateral hemisphere during the stimulations (Supplementary Fig. 14c, Supplementary Fig. 15a). In the ipsilateral somatosensory region, arterial dilation was also observed but with a much weaker magnitude (Supplementary Fig. 14d). Veins did not show dilations (Supplementary Fig. 15c, Supplementary Fig. 15a)³. Deep capillary beds are reported to dilate less than 0.5 μm in diameter¹³, which is not resolvable by the current version of PAM. Fast line scanning along the vessel axis was repeated to measure the blood flow speed (Supplementary Fig. 16, Online Methods)^{8, 14}. Stimulations induced a substantial increase in blood flow speed in both arteries and veins (Supplementary Fig. 14e and Supplementary Fig. 15d)¹⁴. However, PAM could not detect the flow speed changes in deep capillaries.

Upon stimulations, $s\text{O}_2$ increased substantially in veins and deep capillary beds (Fig. 2c, Supplementary Video 4, Supplementary Video 5). The fractional change in $s\text{O}_2$ diminished with increasing distance from the core responding region (Figs. S17a–b), which was ~ 100 μm below the cortical surface (Supplementary Fig. 17c)³. The $s\text{O}_2$ increase was greater in deep capillary beds than in veins and was insignificant in arteries (Supplementary Fig. 15e). The lack of arterial $s\text{O}_2$ response is consistent with the fact that arterial blood has not yet reached capillaries for oxygen consumption and thus maintains a high oxygenation level³.

In the core responding region, the increase in $s\text{O}_2$ in veins also manifested as a decrease in the oxygen extraction fraction (OEF) (Fig. 2d)¹⁵. The fractional change in the cerebral metabolic rate of oxygen (CMRO_2) can be estimated from the above hemodynamic measurements (Online Methods). A moderate fractional increase in CMRO_2 , peaking at $\sim 15\%$, was observed (Fig. 2d). The ratio between the fractional changes in cerebral blood flow (CBF) and CMRO_2 (i.e., the flow-consumption ratio) was ~ 2.0 , consistent with the literature¹⁶.

In summary, using endogenous contrast, PAM has achieved high-speed high-resolution imaging of the vascular morphology, blood oxygenation, blood flow dynamics, and oxygen metabolism of the mouse brain. In particular, PAM has achieved a 1D time-resolved imaging rate of 500 kHz for morphological imaging and 100 kHz for blood oxygenation imaging. In the future, the axial resolution of PAM can be improved by using an ultrasonic transducer with a wider bandwidth or by using nonlinear photoacoustic mechanisms¹⁷. Near-infrared wavelengths can increase the imaging depth of PAM¹⁷. The potential for optical breakdown in RBCs *in vivo*, which is relevant to excitation pulsewidth, can be further investigated by using photothermal microscopy or more other methods¹⁸.

We would like to end by briefly discussing the advantages and limitations of our PAM for mouse brain hemodynamic imaging. (1) PAM provides better depth resolution and greater absorption-based image contrast than wide-field optical microscopy, but is slower and more expensive. (2) In comparison to TPM, PAM does not need exogenous contrast agents and point-by-point depth scanning, but PAM has worse axial resolution, and at the currently available wavelengths less penetration. (3) In comparison to fMRI, which is sensitive only to HbR, PAM is sensitive to both HbR and HbO_2 , and has higher spatial and temporal resolutions. However, fMRI can provide deeper penetration. Therefore, PAM is highly

complementary to other brain imaging modalities in its contrast mechanism, spatial-temporal resolutions, and functional imaging capability.

ONLINE METHODS

Photoacoustic tomography (PAT)

In PAT, as photons travel in tissue, some are absorbed by biomolecules, and their energy is partially or completely converted into heat. The heat-induced pressure wave propagates in tissue, and is detected outside the tissue by an ultrasonic transducer or transducer array to form an image that maps the original optical energy deposition in the tissue. PAT has a 100% relative sensitivity to optical absorption, which means a given percentage change in the optical absorption coefficient yields the same percentage change in the PA signal amplitude. In PAT operating at 532 nm, the high contrast of cortical microvasculature comes from the absorption of hemoglobin in red blood cells (RBCs), which overpowers the absorption of other absorbers such as water and lipids by three orders of magnitude. As the optical absorption of blood is highly oxygenation dependent, PAT can measure blood oxygenation with high sensitivity. Photoacoustic microscopy (PAM) is a microscopic focused-scanning embodiment of PAT.

Fast functional photoacoustic microscopy

To induce photoacoustic signals, a 3-ns pulsed laser beam at 532 nm (AOT-YVO-100Q, AOT Inc.; pulse repetition rate: 100 kHz) is combined with a 3-ps pulsed laser beam at 532 nm (APL-4000-1064, RPMC Lasers, Inc.; pulse repetition rate: 500 kHz) via a polarizing beam splitter (PBS251, Thorlabs, Inc.). The polarizations of the two laser beams are adjusted by wave plates to maximize the combining efficiency. Laser energy fluctuations are monitored by a fast photodiode that samples a small portion of the laser beams. The laser beams are focused by a plano-convex lens (LA1131, Thorlabs, Inc.), then spatially filtered by a 50- μm -diameter pinhole (P50C, Thorlabs, Inc.). The filtered laser beams are focused to ~ 3 μm spots by an objective lens (AC127-050-A, Thorlabs, Inc.; NA: 0.1 in air). The optical focal zone (also known as the depth of focus), defined as the full width at half maximum of the peak intensity, is ~ 83 μm , within which the lateral resolution degrades up to $\sqrt{2}$; correspondingly, the depth range within which the lateral resolution degrades up to a factor of 2 is ~ 144 μm . An optical-acoustic beam combiner, composed of an aluminum-coated prism (NT32-331, Edmund, Inc.) and an uncoated prism (NT32-330, Edmund, Inc.), provides optical-acoustic coaxial alignment. Here, the thin aluminum coating reflects light but transmits sound. An optical correction lens attached to the top surface of the combiner corrects the optical aberration due to the prism. The focused laser beams and the generated photoacoustic waves are both directed by a MEMS scanning mirror in a coaxial configuration. The photoacoustic waves are focused by an acoustic lens and detected by an ultrasonic transducer (V214-BB-RM, Olympus-NDT, Inc.; central frequency: 50 MHz; one-way -6 dB bandwidth: 100%).

Driven by sufficiently strong electromagnetic force, the whole MEMS scanning mirror can operate under de-ionized water in a water tank, which is required to provide acoustic coupling from the sample surface to the acoustic lens. The bottom of the water tank is sealed

with a piece of membrane that is both optically and acoustically transparent. In PAM, because the acoustic flight time provides depth information along the acoustic axis, each excitation laser pulse generates a 1D depth-resolved image. Volumetric imaging is provided by fast angular scanning of the MEMS mirror along the x -axis and slow linear motor-stage scanning of the sample along the y -axis at a speed of 2–4 mm/s. In PAM, a 3 μm lateral resolution at the optical focus and a 15 μm axial resolution have been achieved in clear media. The axial resolution of PAM is jointly determined by the laser pulse width, the frequency-dependent acoustic attenuation in tissue, and the frequency response of the ultrasonic transducer. The maximum in-focus scanning range is ~ 3.0 mm along the x -axis, with a cross-sectional frame rate of 400 Hz. When necessary, additional depth-scanning of the optical focal zone extends the focal range of PAM at the expense of imaging speed. By steering both the optical and acoustic axes simultaneously, PAM maintains confocal alignment and high detection sensitivity over the field of view. The two lasers are triggered with a time interval of 500 ns. The 500 ns delay allows the first PA signal to travel 0.75 mm, which is approximately the maximum penetration depth of PAM in the brain.

PAM of oxygen saturation (sO₂)

In PAM, the two lasers emit the same pulse energy at 532 nm. Because the picosecond pulse has a higher peak intensity than the nanosecond pulse, it results in more saturation (Figs. S3a–b). We define a saturation factor as the ratio of the PA amplitudes under picosecond and nanosecond excitations (Supplementary Fig. 3c).

The relative concentrations of HbR and HbO₂ can be estimated by solving the following linear equations:

$$P_{ns} = \ln(10)kF(\varepsilon_{HbR}C_{HbR} + \varepsilon_{HbO_2}C_{HbO_2}), \quad (1)$$

$$P_{ps} = \ln(10)kF[r_{HbR}(F)\varepsilon_{HbR}C_{HbR} + r_{HbO_2}(F)\varepsilon_{HbO_2}C_{HbO_2}], \quad (2)$$

where P_{ns} and P_{ps} are the PA amplitudes under nanosecond and picosecond excitations, respectively; k is a proportionality coefficient related to the detection system, the are the molar Grüneisen parameter, and the nonradiative quantum yield; ε_{HbR} and ε_{HbO_2} extinction coefficients of HbR and HbO₂, respectively; C_{HbR} and C_{HbO_2} are the molar concentrations of HbR and HbO₂, respectively; and r_{HbR} and r_{HbO_2} are the saturation factors of HbR and HbO₂, respectively, which are functions of the local fluence F (i.e., the photon energy imposed over a unit area).

Note that $\varepsilon_{HbR} \approx \varepsilon_{HbO_2}$ at 532 nm and $r_{HbR} \approx 1$, Eqs. (1)–(2) are reduced to

$$P_{ns} \approx k_1(C_{HbR} + C_{HbO_2}), \quad (3)$$

$$P_{ps} \approx k_1[C_{HbR} + r_{HbO_2}(F)C_{HbO_2}], \quad (4)$$

where $k_1 = \ln(10)k\epsilon_{HbR}F$. Once Eqs. (3)–(4) are solved, the total hemoglobin concentration C_{HbT} is computed by $C_{HbT} = C_{HbR} + C_{HbO_2}$.

Therefore, sO_2 can be computed as

$$sO_2 = \frac{C_{HbO_2}}{C_{HbT}} \approx \frac{1 - \frac{P_{ps}}{P_{ns}}}{1 - r_{HbO_2}(F)} = k_2 \left(1 - \frac{P_{ps}}{P_{ns}}\right), \quad (5)$$

where $k_2 = 1/[1 - r_{HbO_2}(F)]$.

From Eq. (5), we can see that r_{HbO_2} is needed for absolute sO_2 measurement. r_{HbO_2} is calibrated for using the local fluence F (Supplementary Fig. 3c), which is proportional to the unsaturated PA signal P_{ns} (Supplementary Fig. 3b). Note that the local fluence change due to the varied laser spot size at different depth is also accounted for by P_{ns} . r_{HbO_2} can also be calibrated for according to the neighboring arteries.

In studies where only fractional changes in sO_2 are of interest, we have

$$\frac{\Delta sO_2}{sO_2} \approx \frac{\Delta \left(\frac{P_{ps}}{P_{ns}}\right)}{1 - \frac{P_{ps}}{P_{ns}}}. \quad (6)$$

Eq. (6) shows that local fluence is not needed for measuring fractional changes in sO_2 , as long as the saturation is sufficient.

PAM of cerebral metabolic rate of oxygen (CMRO₂)

If the cortical region of interest has well-defined feeding arteries and draining veins, and the volumetric blood flow rates in the feeding and draining vessels are conserved, CMRO₂ can be calculated as

$$CMRO_2 = \xi \times (sO_{2-artery} - sO_{2-vein}) \times C_{HbT} \times CBF/W, \quad (7)$$

where ξ is the oxygen binding capacity of hemoglobin (1.36 mL O₂/gram hemoglobin or 87.7 L O₂/mol hemoglobin); $sO_{2-artery}$ and sO_{2-vein} are the oxygen saturation averaged in the artery and vein, respectively; CBF is the volumetric blood flow rate (L/s); and W is the weight of the region of interest (grams).

Oxygen extraction fraction (OEF) is defined as the fractional difference between the arterial and venous oxygen saturation:

$$OEF = \frac{sO_{2-artery} - sO_{2-vein}}{sO_{2-artery}}. \quad (8)$$

Under normal conditions, arterial blood is close to fully oxygenated (i.e., $sO_{2-artery} \approx 1$). We can rewrite Eq. (7) as

$$CMRO_2 = \xi \times OEF \times sO_{2-artery} \times C_{HbT} \times CBF/W \quad (9)$$

$$\approx \xi \times OEF \times C_{HbT} \times CBF/W.$$

Since the volumetric blood flow rates are conserved, we have

$$CBF = \frac{\pi}{4} (d_{vein})^2 v_{vein}, \quad (10)$$

where d_{vein} is the diameter of the vein, and v_{vein} is the average blood flow speed in the vein.

Assuming W does not change during the brain activity, we have

$$1 + \frac{\Delta CMRO_2}{CMRO_2} \approx \left(1 + \frac{\Delta OEF}{OEF}\right) \left(1 + \frac{\Delta CBF}{CBF}\right) \left(1 + \frac{\Delta C_{HbT}}{C_{HbT}}\right) \quad (11)$$

$$\approx \left(1 - \frac{\Delta sO_{2-vein}}{sO_{2-vein}}\right) \left(1 + \frac{\Delta d_{vein}}{d_{vein}}\right)^2 \left(1 + \frac{\Delta v_{vein}}{v_{vein}}\right) \left(1 + \frac{\Delta C_{HbT}}{C_{HbT}}\right).$$

From Eq. (11), we can calculate the fractional change in $CMRO_2$ from the fractional changes in other parameters measured individually.

Experimental animals

Six female ND4 Swiss Webster mice (Harlan Laboratory, Inc.; 16–30 g, 3–10 weeks old) were used for the studies. The laboratory animal protocols were approved by the Animal Studies Committee of Washington University in St. Louis. During the experiment, the mouse's temperature was kept at 37 °C by a water-circulating heating pad. An intraperitoneal dose of 100 mg/kg α -Chloralose was used for anesthesia, which had a relatively mild effect on the brain functions¹⁴. The mouse was then taped to a lab-made animal holder, which was mounted to the PAM system. The head of the mouse was fixed in a stereotaxic frame. Before imaging, the scalp was surgically removed, while the skull was left intact. Bloodstains on the skull surface—even when invisible to naked eyes—could generate strong photoacoustic signals; thus, the exposed skull surface was carefully cleaned with phosphate buffered saline (PBS) solution. Ultrasound gel was then applied on the skull surface to retain moisture and couple the acoustic signals. A water tank filled with de-ionized water was then placed on top of the mouse head. The membrane at the bottom of the water tank was in gentle contact with the ultrasound gel. The translation of the animal holder by the motor-stage at a speed of 2–4 mm/s did not induce significant disturbance to the animal and the water in the tank.

Electrical stimulations to hindlimbs

Electrical stimulations were introduced by two pairs of needle electrodes inserted under the skin of the right and left hindlimbs, respectively. The electrodes were connected to a function generator (DS345, Stanford Research Systems, Inc.) through a manual switch. The whole procedure consisted of five periods and lasted for five minutes. The first, third and fifth periods were resting states, while the second period, from 60 seconds to 80 seconds, was left hindlimb stimulation, and the fourth period, from 200 seconds to 220 seconds, was right hindlimb stimulation. Each stimulation period consisted of a train of electrical pulses

with an amplitude of 2 mA, a pulse width of 0.25 millisecond and a repetition rate of 2 Hz. The stimulation period and intensity were controlled without inducing any paw motions. Five trials were performed on each mouse.

Automatic vessel segmentation

The PA signal amplitude was extracted through the Hilbert transformation of each 1D depth-resolved signal. The data was then processed using a customized vessel-segmentation algorithm. All the data analysis was performed in 3D. Briefly, a cross-sectional image was first converted to a binary image, where the cross-section of each vessel was identified and labeled. By tracking the cross-sections of each vessel throughout all the cross-sectional images, the vessels were individually labeled and thus segmented. All the segmented vessels were visually evaluated and corrected if necessary. The final segmentation information was then stored for future use. The vessel segmentation algorithm can be performed along different orientations.

The vessel segmentation was used to measure blood vessel diameters and correct the sO_2 calculation. To measure the vessel diameter, we measured the length of a line across the vessel at different angles relative to the vessel's axis, and chose the shortest path length as the vessel diameter. To correct the sO_2 calculation, any overlapping vessels were separated at the junction through segmentation, and the sO_2 of each vessel was calculated individually.

PAM of blood flow speed

Line scanning along the axis of a vessel can be used to measure the blood flow speed. Moving RBCs imaged at a sufficient rate appear as bright–dark streaks in the resulting space–time map. The slope of the bright–dark streaks, measured from the vertical direction, is proportional to the centerline (axial) flow speed. The slope is measured by using a 2D Fourier transformation of the space–time map. The flow direction can be determined from the sign of the slope and the direction of the line scan sweep. To measure high flow speeds more accurately, we imaged the same cells with at least five line scans. Therefore, the maximum measurable flow speed corresponds to a travel distance of 3 mm within 12.5 ms, which translates to a flow speed of ~24 cm/s.

H&E histology

The right hemisphere of a mouse was imaged by PAM with a picosecond pulse energy of 1 μ J and a pulse repetition rate of 500 kHz. Immediately after the imaging, the mouse was transcardially perfused with 0.9% saline followed by 4% paraformaldehyde in PBS. The brain was removed and postfixed in 4% paraformaldehyde for 24 h. Coronal sections (5 μ m thick) were cut with paraffin embedding. Standard H&E staining was performed on the sections, which were examined using bright-field microscopy (NanoZoomer, Hamamatsu) with a 20 \times objective (NA = 0.67). In the positive control experiment, the left hemisphere of a mouse was illuminated by a continuous-wave laser (GM-CF02-100, Information Unlimited, Inc.; wavelength: 532 nm; power: 100 mW; spot size: ~0.25 mm²) for one minute to induce thermal coagulation (i.e., burn). The right hemisphere was imaged by PAM for one minute with a pulse energy of 1 μ J and a pulse repetition rate of 500 kHz. H&E histology was then performed on the brain with the same procedure as above.

Two-photon microscopy imaging

To rule out the potential for causing vessel leakage by PAM, a mouse was imaged by two-photon microscopy (Fluoview 1000, Olympus, Inc.) after the PAM imaging (pulse energy: 1 μ J; pulse repetition rate: 500 kHz). For two-photon microscopy, the skull was thinned to \sim 30 μ m using a dental drill and a microsurgical blade as previously described¹⁴. FITC-dextran solution in PBS (150 μ L, 2.5% w/v) was injected via a tail vein before the two-photon imaging. A 4 \times objective (NA = 0.10) was used to find the same imaging area as that in PAM, and then a 20 \times objective (NA = 0.70) was used to acquire high-resolution images (excitation wavelength: 800 nm; emission filter wavelength: 495–540 nm). Depth-scanning was performed from the skull surface to a depth of 600 μ m into the cortex, with a step size of 5 μ m. The same procedure was also used for validating PAM by using two-photon microscopy (Supplementary Fig. 5).

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgments

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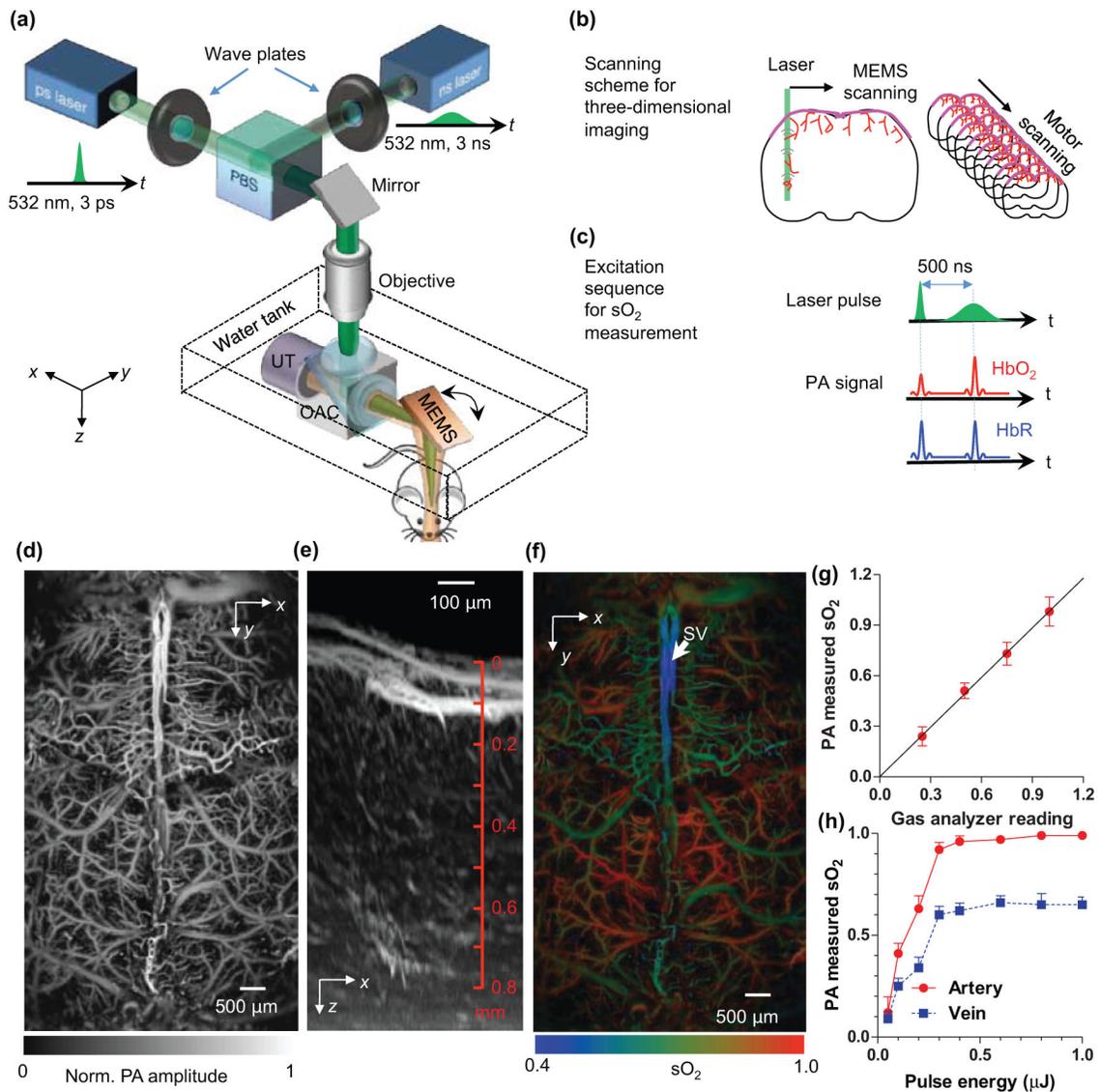


Fig. 1. Fast functional photoacoustic microscopy (PAM) of the mouse brain

(a) Schematic of the PAM system. OAC, optical-acoustic combiner; PBS, polarizing beam splitter; UT, ultrasonic transducer. (b) Scheme of PAM scanning. 3D imaging is achieved by fast MEMS mirror scanning along the x -axis and slow motor-stage scanning along the y -axis. (c) Sequence of PAM excitation and detection. The picosecond pulse incident on oxy-hemoglobin (HbO_2) results in more saturation and thus a weaker PA signal than the following nanosecond pulse, whereas the difference for deoxy-hemoglobin (HbR) is negligible. (d) A representative x - y projected brain vasculature image through an intact skull ($n = 6$). (e) A representative enhanced x - z projected brain vasculature image acquired over a 0.6×0.6 mm² region with depth scanning, where the signal amplitude was normalized depth-wise ($n = 6$). (f) PAM of oxygen saturation of hemoglobin (sO_2) in the same mouse brain as in (d), acquired by using the single-wavelength pulse-width-based method (PW- sO_2) with two lasers. The averaged sO_2 in the skull vessels was lower than that in the cortical vessels. SV, skull vessel. (g) Comparison of the PW- sO_2 measurements in four blood phantoms and the

gas analyzer readings. **(h)** *In vivo* PW-sO₂ measurements in an artery-vein pair in a mouse ear with varied excitation pulse energies. The data in **(g)** and **(h)** are averaged within the samples, and the error bars are standard deviations.

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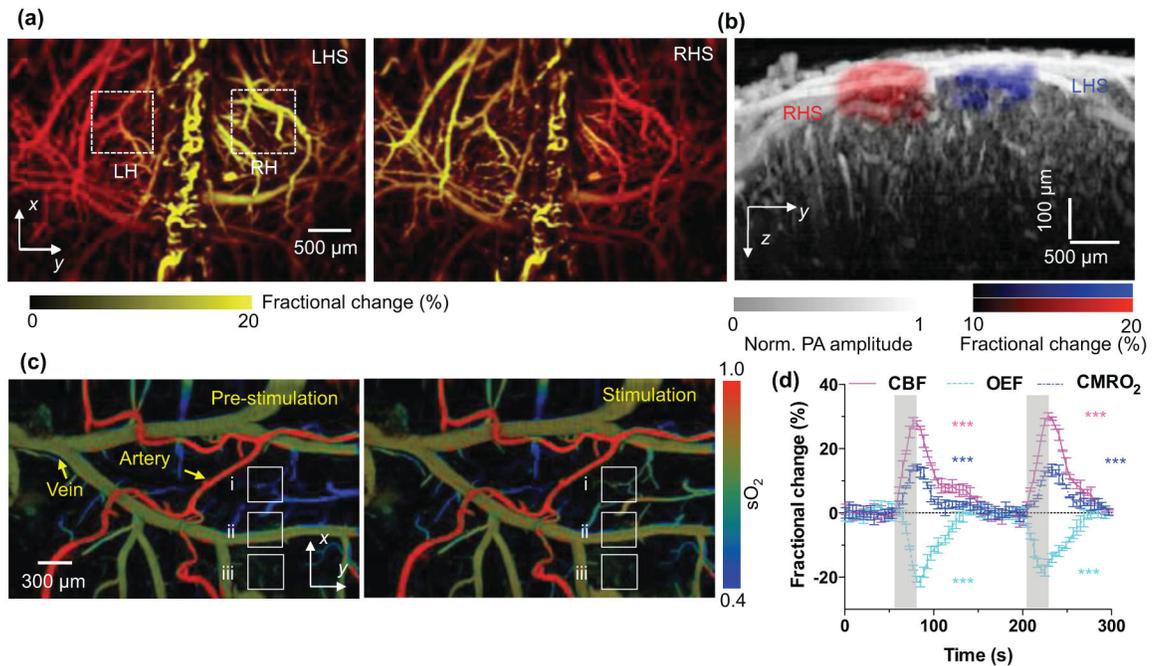


Fig. 2. PAM of brain responses to electrical stimulations to the hindlimbs of mice ($n = 6$)
(a) Fractional PA amplitude changes (shown in yellow) in response to left hindlimb stimulation (LHS) and right hindlimb stimulation (RHS), superimposed on the vascular image (shown in red). LH/RH, left/right hemisphere. **(b)** Depth-resolved PA amplitude responses. The responding areas in the LH and RH are shown in red and blue, respectively, and superimposed on the gray-scale y - z projection image. The signal amplitude in the y - z projection image was normalized depth-wise. **(c)** Fast sO_2 imaging before (left panel) and during (right panel) stimulations to the left hindlimb. Three $0.3 \times 0.3 \text{ mm}^2$ subregions (i, ii and iii) are further analyzed (Supplementary Fig. 17). **(d)** Time courses of the fractional changes in the cerebral blood flow (CBF), oxygen extraction fraction (OEF), and cerebral metabolic rate of oxygen ($CMRO_2$) in the core responding region. All the sO_2 measurements were acquired with two lasers. The data in **(d)** are averaged over five trials on each of the six mice, and the error bars are standard errors. Statistics: paired student's t -test. P values: *** < 0.001 .