Near-infrared optical-resolution photoacoustic microscopy

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Compared with visible light (380–700 nm), near-infrared light (700–1400 nm) undergoes weaker optical attenuation in biological tissue; thus, it can penetrate deeper. Herein, we demonstrate near-infrared optical-resolution photoacoustic microscopy (NIR-OR-PAM) with 1046 nm illumination. A penetration depth of 3.2 mm was achieved in chicken breast tissue ex vivo using optical fluence within the American National Standards Institute (ANSI) limit (100 mJ/cm²). Beyond ~0.6 mm deep in chicken breast tissue, NIR-OR-PAM has shown finer resolution than the visible counterpart with 570 nm illumination. The deep imaging capability of NIR-OR-PAM was validated in both a mouse ear and a mouse brain. NIR-OR-PAM of possible lipid contrast was explored as well. © 2014 Optical Society of America

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With 100% sensitivity to optical absorption, optical-resolution photoacoustic microscopy (OR-PAM) can provide both anatomical and functional information with high spatial resolution [1–5]. In OR-PAM, high lateral resolution is achieved by tightly focusing the laser beam into a diffraction-limited spot [6]. In a highly scattering medium such as biological tissue, the focusing capability of OR-PAM degrades with the imaging depth [7]. In previous OR-PAM studies, ultraviolet (UV) or visible (VIS) light was used for illumination, mainly because of the strong absorption of DNA/RNA, cytochromes, hemoglobin, and melanin in these wavelength regions [8,9]. However, the strong optical scattering limits the penetration depth of OR-PAM in biological tissue.

To improve the imaging depth, we propose near-infrared (NIR) OR-PAM. Compared with VIS illumination, NIR illumination yields several benefits. First, NIR-OR-PAM can penetrate deeper into tissue for a given signal-to-noise ratio (SNR), primarily due to weaker absorption of NIR light by blood [10]. Second, with weaker scattering of NIR light, NIR-OR-PAM can maintain its lateral resolution at greater depths. Third, NIR-OR-PAM can image other tissue components, such as water and lipid [11,12]. The optical absorption spectra of four tissue components from 250 to 1150 nm are shown in Fig. 1(a) [13,14]. Fourth, the American National Standards Institute (ANSI) permits stronger NIR light intensity on the tissue surface [15]. Herein, we apply NIR light in OR-PAM and demonstrate these advantages.

In the NIR-OR-PAM system (Fig. 1(b)), an Nd:YLF laser (INNOSAB, Edgewave, GmbH) generates laser pulses at its fundamental wavelength of 1046 nm. The laser pulses then pass through a second harmonic generator to generate 523 nm laser pulses. After the second harmonic generator, 523 nm and residual 1046 nm laser beams are split by a dichroic mirror. A dye laser (C8R-D, Sirah, GmbH) is pumped by the 523 nm beam and emits laser pulses with tunable wavelengths. We selected 570 nm for photoacoustic (PA) imaging—referred to as VIS-OR-PAM—to compare with NIR-OR-PAM. A flip mirror is used to select either the 570 nm or 1046 nm laser beam. Afterward, the laser beam is reshaped by an iris (ID25SS, Thorlabs) and then attenuated by a neutral density filter (NDC-50C-1400 nm) undergoes weaker optical attenuation.

![Schematic of the NIR-OR-PAM system](image-url)

**Fig. 1.** Absorption spectra of four tissue components and the schematic of near-infrared optical-resolution photoacoustic microscopy (NIR-OR-PAM). (a) Absorption spectra of oxyhemoglobin (HbO₂) (150 g/L or 2.3 mM in blood), deoxyhemoglobin (HbR) (150 g/L or 2.3 mM in blood), water (80% by volume in tissue), and lipid (20% by volume in tissue) from 250 to 1150 nm. Fourth, the American National Standards Institute (ANSI) permits stronger NIR light intensity on the tissue surface. (b) Schematic of the NIR-OR-PAM system. ConL, condenser lens; CorL, correction lens; DC, dye cell; DM, dichroic mirror; FM, flip mirror; M1, M2, and M3, mirrors; ND, neutral density filter; OL, objective lens; PH, pinhole; PL, pump laser; RAP, right-angle prism; RhP, rhomboid prism; SHG, second harmonic generator; SOL, silicone oil layer; UT, ultrasonic transducer; WT, water tank.
the beam further. The laser beam is then focused into the sample by an objective (AC127-050-A, Thorlabs; numerical aperture: 0.1 in air). A high-frequency ultrasonic transducer is placed confocally with the objective to detect the generated PA waves. The acoustic–optical coaxial alignment is achieved by using a beam combiner composed of a thin layer of silicone oil sandwiched between a right-angle prism (NT32-545, Edmund Optics) and a rhomboid prism (NT49-419, Edmund Optics). Each laser pulse generates a one-dimensional PA image (A-line) by recording the time course of the PA signal. Volumetric imaging is provided by two-dimensional motor scanning.

To measure the penetration depth of NIR-OR-PAM, we imaged a 50-μm-diameter black human hair obliquely inserted into a piece of fresh chicken breast tissue. To conform to the ANSI safety limit [15], the light fluences on the tissue surface at 570 and 1046 nm were adjusted to 20 and 100 mJ/cm², respectively. NIR-OR-PAM was able to image the black hair 3.2 mm beneath the skin surface with an SNR of 6 dB compared to only 2.3 mm by VIS-OR-PAM with the same SNR (Figs. 2(a) and 2(b)). NIR-OR-PAM can penetrate deeper than VIS-OR-PAM. First, NIR light undergoes weaker optical attenuation than VIS light, which allows it to penetrate deeper. Second, ANSI permits stronger NIR light fluence on tissue surface, which also gives us larger penetration depth. The same time-gain compensation (TGC) (0.75 dB/MHz/cm) for acoustic attenuation was applied to improve the visibility in both cases.

To measure the lateral resolution of the NIR-OR-PAM system in clear media, a sharp metal edge was imaged with a step size of 0.625 μm and a scanning range of 100 μm along the x axis. By fitting the measured data to an edge spread function, the full width at half-maximum (FWHM) of the corresponding line spread function was estimated as the lateral resolution. The lateral resolution of NIR-OR-PAM in a clear medium was estimated to be 6.2 μm (Fig. 3(b)), and that of VIS-OR-PAM was estimated to be 2.9 μm (Fig. 3(a)), both of which are close to the theoretical lateral resolutions of OR-PAM, which are determined by the focal diameters of the laser beams. The axial resolution of NIR-OR-PAM was estimated to be ~27 μm based on the transducer bandwidth and the speed of sound in tissue [16]. In scattering media such as biological tissue, both the lateral and axial resolutions decrease with imaging depth due to optical scattering and frequency-dependent acoustic attenuation in tissue.

The lateral resolutions of NIR-OR-PAM and VIS-OR-PAM at different imaging depths were measured in scattering media. A sharp metal edge obliquely inserted into a piece of fresh chicken breast tissue was imaged at different depths while the optical focus was maintained on the metal edge. As optical focusing deteriorated in tissue, the line spread functions were no longer Gaussian functions [7]; thus, the lateral resolution could no longer be estimated by fitting the edge spread functions using error functions, as we did for the clear medium. Therefore, we estimated the lateral resolution as the FWHM of the numerical first derivative of the measured edge spread function. Figure 3(c) shows the lateral resolutions of both NIR-OR-PAM and VIS-OR-PAM as functions of imaging depth. Due to optical scattering in the chicken tissue, the optical focusing degrades with depth, deteriorating the lateral resolution of OR-PAM. At 1046 nm, scattering in chicken breast tissue is weaker than that at 570 nm. With weaker optical scattering, NIR light can focus better than VIS light after traveling through a certain depth, which results in finer lateral resolution in NIR-OR-PAM over VIS-OR-PAM. For VIS-OR-PAM, the lateral resolution is finer than 40 μm up to ~1.6 mm depth, while for NIR-OR-PAM, the lateral resolution is finer than 40 μm up to ~2 mm depth. Close to the tissue surface (~<0.6 mm), NIR-OR-PAM has a worse lateral resolution than VIS-OR-PAM due to the longer wavelength; at ~0.6 mm depth, NIR-OR-PAM and VIS-OR-PAM show similar resolutions. As the imaging depth goes beyond 0.6 mm, NIR-OR-PAM has finer lateral resolutions than VIS-OR-PAM until the
optical focusing becomes so ineffective that OR-PAM turns into acoustic-resolution PAM, where the acoustic focusing of the 50 MHz ultrasonic transducer (∼43 μm) determines the lateral resolution.

A mouse ear was imaged in vivo with NIR-OR-PAM. All experimental animal procedures were carried out in conformity with the laboratory animal protocol approved by the Animal Studies Committee at Washington University in St. Louis. A 2 × 2 mm² area of the mouse ear was imaged by both VIS-OR-PAM and NIR-OR-PAM (Figs. 4(a) and 4(b)). VIS-OR-PAM is able to image only the blood vessels (Fig. 4(a)). NIR-OR-PAM, however, is able to image more contrasts (Fig. 4(b)). In addition to blood vessels, the circular structures imaged by NIR-OR-PAM are suspected to be fat cells surrounding sebaceous glands [17,18]. Close-up images and B-scan images (Figs. 4(c) and 4(d)) further demonstrate the improvement of NIR-OR-PAM in lipid contrast and imaging depth in vivo.

We also imaged a mouse brain in vivo with NIR-OR-PAM. The same area of the mouse brain was imaged by VIS-OR-PAM and NIR-OR-PAM. The scalp was removed, but the skull was intact. Due to the strong blood absorption and brain tissue scattering, few blood vessels deeper than 0.5 mm can be imaged by VIS-OR-PAM (Figs. 5(a) and 5(b)). However, NIR-OR-PAM can detect signals from blood vessels deeper than 0.5 mm (Figs. 5(c) and 5(d)).

NIR-OR-PAM can not only penetrate deeper than VIS-OR-PAM but can also image structures in deep regions with finer resolution. As shown in Figs. 6(a) and 6(b), at both illumination wavelengths, OR-PAM was able to clearly image the cortical vascular network. However, close-ups of the vessels more than 200 μm deep in Figs. 6(a) and 6(b) show that the vessels are less blurred in the NIR-OR-PAM images than in the VIS-OR-PAM images, as shown in Fig. 6(c). The lateral resolution improvement for in vivo imaging by NIR-OR-PAM was estimated by comparing the apparent diameters of vessels at the same depths. The apparent diameters of two representative vessels were 120 and 160 μm in VIS-OR-PAM, while they were 50 and 80 μm in NIR-OR-PAM, respectively (Fig. 6(d)). The apparent diameters of blood vessels were approximately two times smaller in NIR-OR-PAM.

Due to strong absorption, an overlying blood vessel may shadow vessels underneath in VIS-OR-PAM. At
1046 nm, the optical absorption of hemoglobin is approximately 40 times less than that at 570 nm (Fig. 1(a)). Therefore, 1046 nm NIR light has a 1/e penetration depth of 1.2 mm in blood and is able to image multiple layers of blood vessels. This advantage was demonstrated by imaging a bleeding area in a mouse brain. The bleeding layer appears weaker in NIR-OR-PAM than in VIS-OR-PAM (Figs. 7(a) and 7(b)). By peeling off the bleeding layer in data processing, we can clearly see that the optical shadowing from the bleeding layer is weaker in NIR-OR-PAM, and the vascular network underneath can thus be better resolved with greater SNR. However, due to the strong absorption, nearly all the VIS light is absorbed by the blood pool; this generated strong PA signals that were reverberated by other underlying brain structures, showing as a "ghost" image of the blood pool.

Lower optical absorption in the NIR region allows deeper penetration but results in weaker PA signals. However, with nanosecond pulse excitation, the tissue damage mechanism is mainly photothermal [19]. Therefore, lower absorption allows for stronger excitation fluence, which can partially compensate for the reduced signal strength.

In conclusion, we have demonstrated that NIR-OR-PAM offers new imaging contrast and better resolution at deeper imaging depths than VIS-OR-PAM. By employing NIR light with our fast-scanning OR-PAM [20] and incorporating multiple wavelengths [21], we should be able to measure hemodynamic parameters, including blood flow speed, the oxygen saturation of hemoglobin, and the metabolic rate of oxygen [22].

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