Polymer dots enable deep \textit{in vivo} multiphoton fluorescence imaging of microvasculature

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Abstract: Deep \textit{in vivo} imaging of vasculature requires small, bright, and photostable fluorophores suitable for multiphoton microscopy (MPM). Although semiconducting polymer dots (pdots) are an emerging class of highly fluorescent contrast agents with favorable advantages for the next generation of \textit{in vivo} imaging, their use for deep MPM has never before been demonstrated. Herein, we characterize the multiphoton properties of three pdot variants and perform deep \textit{in vivo} MPM imaging of cortical rodent microvasculature. We find pdot brightness exceeds conventional fluorophores, including quantum dots, and their broad multiphoton absorption spectrum permits imaging at wavelengths better-suited for biological imaging and confers compatibility with a range of longer excitation wavelengths. This results in substantial improvements in signal-to-background ratio (>3.5-fold) and greater cortical imaging depths ($z = 1,300 \mu m$). Ultimately, pdots are a versatile tool for MPM due to their extraordinary brightness and broad absorption, enabling interrogation of deep structures \textit{in vivo}.

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1. Introduction

\textit{In vivo} imaging with multiphoton fluorescence microscopy (MPM) is widely used due to its ability to provide diffraction-limited, three-dimensional images of intact tissue at depths ranging from a few hundred microns to more than 1 mm [1–5]. Recent reports of deep tissue, \textit{in vivo} imaging with two- and three-photon excitation have demonstrated the potential of imaging structures in the brain beyond the cortex, including the corpus callosum and hippocampus [6,7]. These advances have been driven in part by the availability of new ultrafast laser sources at longer wavelengths ($\lambda_{\text{ex}} = 1,000 – 1,600 \text{ nm}$) than those traditionally used in two-photon fluorescence imaging ($\lambda_{\text{ex}} = 700 – 900 \text{ nm}$). Equally important to the availability of laser sources is the availability of contrast agents with strong multiphoton absorption properties at these longer wavelengths. In this paper, we show that an emerging class of fluorophores termed polymer dots [8] possess strong two- and three-photon excitation across a broad range of wavelengths and enable deep imaging of microvasculature in the brain.

MPM of vasculature requires the intravenous injection of bright, biocompatible contrast agents that preferably exhibit large absorption cross sections under near-infrared (NIR) excitation ($\lambda_{\text{ex}} = 700 – 1,000 \text{ nm}$) and prolonged blood circulation times. Traditional exogenous contrast agents for \textit{in vivo} MPM include organic dyes such as dextran-conjugated fluorescein and indocyanine green or inorganic semiconductor quantum dots [9,10]. Organic dyes, however, suffer from poor photostability and low quantum yields in aqueous biological environments [11–13]. Although quantum dots offer improved brightness and photostability,
they present substantial toxicity concerns and are prone to bioaccumulation in organs and tissues [14–16]. Thus, the biological imaging community is eager for a safer, brighter, and more stable probe for deep, high-resolution *in vivo* imaging.

The highly fluorescent semiconducting polymer dot (pdot) is a promising candidate for *in vivo* MPM with material properties that can potentially overcome many of the limitations faced by other probes [17]. Although pdots are similar to quantum dots with respect to size (~10 – 100 nm, Fig. 6 in Appendix A) and quantum yield, pdots are brighter, more photostable, and present no clear evidence of biotoxicity [17,18]. A useful measure of fluorescence brightness is the action cross section, which is given by the product of the peak absorption cross section and the fluorescence quantum yield [17,19]. Pdots rival both quantum dots and organic dyes in that their two-photon action (2PA) cross-sections are one to two orders of magnitude larger than inorganic quantum dots and three to five orders of magnitude greater than commonly used fluorescent dyes [17,20]. Moreover, polymer dots are readily amenable to functionalization and bioconjugation, enabling their used for tagged localization assays and molecular targeting [18,21]. Herein, for the first time, we present evidence that we can take advantage of pdots’ many favorable properties to produce high-quality *in vivo* multiphoton images of vasculature with excellent signal-to-background ratio (SBR) at depths exceeding 1 mm.

Another advantage of pdots is their broad absorption, which enables multiphoton imaging with a variety of ultrafast laser sources, including ytterbium-fiber lasers (yb-fiber, \(\lambda_{ex} = 1,060\) nm) and longer wavelength optical parametric amplifiers (OPA). Such broadband compatibility takes advantage of the favorable photophysical characteristics of these unique laser sources to improve SBR beyond the capabilities of conventional 2P titanium-sapphire (Ti:S) microscopy. Moreover, the ability to excite pdots at longer wavelengths allows us to approach an ideal biological imaging wavelength situated around 1,300 nm where absorption and tissue scattering events are minimized [6,22]. For instance, the photophysical advantages of longer wavelength excitation of poly[9,9-dioctylfluorenyl-2,7-diyl-co-1,4-benzo-{2,1′-3′-thiadiazole}] (PFBT; \(\lambda_{ex} = 1,060\) nm) coupled with the ytterbium-fiber laser’s intrinsic pulse characteristics results in a 3.5-fold improvement in SBR, and an overall 50 µm gain in penetration depth *in vivo*. Multiphoton imaging of poly[2-methoxy-5-(2-ethylhexyloxy)-1,4-(1-cyanovinylene-1,4-phenylene)] (CNPPV) labeled vasculature using an OPA (\(\lambda_{ex} = 1,225\) nm) increases SBR by ~8.2 fold (\(z = 700\) µm), and extends imaging depth 450 µm further into the brain. Notably, PFPV and CNPPV both exhibit a partial three-photon power dependence at these longer wavelengths, which contributes to a larger SBR by the suppression of out-of-focus fluorescence [6,23]. Overall, semiconducting pdots are ideal for MPM due their enhanced brightness over traditional fluorophores, and spectrally wide absorption range. These advantages coupled with their intrinsically low cytotoxicity help overcome the longstanding limitations of quantum dots and establish the immense potential of pdots for *in vivo* biological imaging.

2. Materials and methods

2.1 Power dependence measurements

Adjacent sample wells in a multi-channel slide (µ-Slide VI0.4, Ibidi) were filled with 40 µl of 100 nM CNPPV, poly[9,9-dioctyfluorenyl-2,7-diyl-co-1,4-benzo-{2,1′-3′-thiadiazole}] (PFBT), or PFPV alongside a blank solution (0.9% blood bank saline, Fisher). The sample was then exposed to laser light excitation and the emitted photon counts were monitored by a digital photon counting board (DPC-230, Becker & Hickl GmbH). Three consecutive measurements of photon counts from the sample and blank were monitored using a five second acquisition duration in absolute timing mode. Signal was maximized immediately before each recording by adjusting the z-position of the sample surface relative to the objective via a motorized lab jack (L490MZ, Thorlabs). Measurements were repeated at increasing power
intensities and the logarithmic values of the background-subtracted signal were plotted as a function of the logarithmic excitation value. A linear model was then fit to the data and the slope was extracted to determine the power dependence. A fresh sample was used for each distinct wavelength to avoid the potential introduction of any artifacts related to photostability. Excitation wavelengths of the Ti:S source (Mira 900, Coherent) and the ytterbium-fiber laser (home-built) were measured with a UV-Vis-NIR spectrometer (USB2000, Ocean Optics) whereas the longer optical parametric amplifier wavelengths were measured with an NIR spectrometer (Avaspec-NIR256-1.7, Avantes) [4].

2.2 Two-photon microscopy

In the two-photon laser system, a titanium:sapphire (Ti:S) oscillator (Mira 900, Coherent) beam is steered to a pair of galvanometer scanners (6125HB, Cambridge Technology) driven by servo driver amplifier boards (671215H-1HP, Cambridge Technology). A Keplerian telescope beam expander which consists of a B-coated scan lens (f = 80.0 mm, AC254-080-B, Thorlabs) and tube lens (f = 200.0 mm, LA1979-B-N-BK7, Thorlabs) is used to fill the back aperture of the microscope objective (XLUMPLFLN20XW 0.95 NA or XLPLN25XSVMP2 25X 1.0 NA, Olympus). Excitation and emission paths are separated with a 775 nm cutoff dichroic mirror (FF775-Di01-52x58, Semrock). Fluorescence is epi-collected, transmitted through either a 510/84 bandpass filter (FF01-510/84-25, Semrock) or a 609/181 bandpass filter (FF01-609/181-25, Semrock), and detected by a photomultiplier tube (H10770PB-40, Hamamatsu Photonics). Image acquisition was controlled using custom software (LabVIEW, National Instruments) and image frames were collected at a 512 x 512 pixel size. Image stacks were collected at a z-resolution of 5 µm and three frames were averaged from 0 – 200 µm cortical depths, five frames from 200 – 500 µm, eight frames from 500 – 700 µm, and twelve frames beyond 700 µm. All mice specimens imaged by Ti:S were excited at λ_{ex} = 800 nm.

2.3 Three-photon microscopy

The three-photon laser schematic is provided in Fig. 7 in Appendix B. A 5W laser is used to pump a λ_{ex} = 800 nm mode-locked Ti:S oscillator (Mira 900, Coherent) which is stretched by an external stretcher/compressor to seed a regenerative amplifier (RegA 9000, Coherent). The amplified pulse is compressed and the spectrum of this 800 nm pulse can then be shifted and tuned by a customized optical parametric amplifier at 511 kHz (OPA 9800, Coherent) [3]. Three-photon in vivo imaging was performed at λ = 1,225 nm using a 25x multiphoton objective (XLPLN25XSVMP2 25x 1.0 NA, Olympus). Again, image stacks were collected at a z-increment of 5 µm and all frames were 512 x 512 pixels. Three frame running averages were recorded for the first 500 µm and an additional two frames were averaged each 200 µm interval beyond that.

2.4 Fiber laser microscopy

A commercial fiber oscillator (Origami-10, OneFive GmbH) was used to seed a custom-built ytterbium fiber amplifier [4]. The amplifier consisted of a 6-meter long segment of double-clad ytterbium-doped polarization-maintaining large-mode-area optical fiber with a core diameter of 25 µm and cladding diameter of 250 µm (YB1200-25/250DC-PM, Thorlabs), coiled to a radius of less than 7 cm to suppress higher-order modes and achieve single-mode output. The fiber was pumped by a fiber-coupled laser diode emitting up to 30 W of 915-nm light (K915FA3RN-30.00W, BWT Beijing). With a pump power of 23W, the amplifier produced 6 W of output at 1060 nm with a repetition rate of 80 MHz, and a grating pair was used to compress the pulse width to ~120 fs in the sample plane. The laser power was adjusted using a Glan-type calcite polarizer/half wave-plate combination.
2.5 Animal preparation and vessel line scanning

Eight to ten week old C57 mice were prepared with cranial window implants and administered with retro-orbital injections of fluorescent contrast agents using methods previously described in Perillo et. al 2017 [5]. Line scans were acquired along vessel axes by collecting the fluorescence intensity while the galvo mirrors rapidly scanned a linear, user-defined path along a capillary. Each line consisted of 200 pixels imaged at a rate of 450 kHz, enabling a line rate of 940 Hz. To build a trajectory, the fluorescence intensity is plotted as a function of time (line number) and space (xy-coordinate, or pixel in one line). A sufficient number of lines were scanned along a trajectory to quantify the blood flow from analyzing the dark streaks. A Radon transform was used to determine the inclination angle of the dark streaks which, when combined with the known temporal and length scales, allowed for calculation of the RBC velocity [24–26].

3. Experimental results

3.1 Two- and three-photon excitation power dependence of polymer dots

Two-photon excitation of pdots has been successfully demonstrated by other research groups, although it has been more generally referred to under the larger and less specific umbrella of multiphoton imaging [20]. With mounting evidence of the advantages of three-photon microscopy over 2P imaging for deep in vivo imaging, such as the suppression of out-of-focus fluorescence, reduced scattering, and improved SBR, the need to identify specific excitation wavelengths that produce two- versus three-photon absorption becomes imperative [6,27]. Thus, we tested the excitation power dependence of various pdots including a CNPPV and two fluorene-based copolymers, PFBT and PFPV, at wavelengths ranging from 790 - 850 nm, 1,060 nm, and 1,200 – 1,350 nm (Appendix A). We find that all three semiconducting polymers exhibit a strong two-photon dependence at 800 nm (Fig. 1). In the case of CNPPV and PFBT, this effect persists out to 1,060 nm whereas PFPV begins to demonstrate a partial three-photon power dependence at this wavelength (Fig. 1). Pure three-photon fluorescence is most strongly demonstrated at 1,300 nm for CNPPV, 1,350 nm for PFBT, and 1,325 nm for PFPV (Fig. 1). Within the 1,060 – 1,300 nm excitation range, there is a clear rising transition from two-photon to three-photon excitation. This data is especially instructive in that it serves as a reference for the selection of appropriate excitation sources and wavelengths for MPM experiments with pdots.

3.2 Two-photon signal-to-background ratio is improved by the brightness of polymer dots

The primary goal of this study is to evaluate the use of pdots for deep in vivo vascular imaging. C57 mice with intravenous injections of fluorescent dye, quantum dots, or pdots were imaged through an optical cranial window [28] using 800 nm excitation. All imaging experiments were performed in age-matched mice from the same litter to minimize any sources of variability in tissue properties that could potentially obscure depth comparisons. Dextran-conjugated fluorescein (λ<sub>em</sub> = 524 nm; FD2000S, Sigma-Aldrich) served as the organic dye of interest, chosen because of its prevalent usage as a contrast agent in biological experiments [19,29–31]. QD605 (λ<sub>em</sub> = 605 nm; Q10001MP, ThermoFisher Scientific) was the selected semiconductor quantum dot, a probe known for its large quantum yield, atypically high brightness, and photostability [30,32]. Lastly, three pdot variants, PFBT (λ<sub>em</sub> = 538 nm), PFPV (λ<sub>em</sub> = 520 nm), and CNPPV (λ<sub>em</sub> = 590 nm), were selected as they represent different classes of polymers, including poly(phenylene vinylene) and fluorene-based copolymers [17]. Comparing 100 µm thick maximum intensity projections at the same cortical depths, the pdot data visually appears much brighter than either the fluorescein or quantum dot images, demonstrating an enhanced SBR given by pdots (Fig. 2). A quantitative comparison of the contrast agents reveals that PFPV produces the largest SBR, followed by
PFBT then CNPPV as shown in Fig. 8 in Appendix C. Relative to QD605 and fluorescein, pdot SBR is larger throughout the entire depth range. To ensure fair comparisons, the average laser power at each depth was maintained at consistent levels across the separate imaging experiments. Furthermore, appropriate filters were selected for all experiments to maximize the collection efficiency of each fluorophores and ensure fair comparisons of the separate image stacks. Vasculature in mice labeled by QD605 (Fig. 2, Right) appeared noticeably distorted and discontinuous, and imaging was limited to a 550 µm depth relative to the cortical surface. This is explained by the fact that the mouse did not survive the retro-orbital injection of inorganic quantum dots due to toxic effects, resulting in an absence of active circulation to evenly distribute the contrast agent in plasma. Meanwhile, chronic in vivo imaging experiments with repeated intravenous injections of pdots in the same mice over the course of several months showed no signs of cytotoxicity or deleterious effects on the animals, supporting several published claims of pdot biocompatibility [18,21,33,34]. An example of two-photon data collected from a mouse injected with 50 nM of PFPV showcases excellent SBR up to 650 µm deep and appreciable signal levels highlighting clearly delineated blood vessels from 750 – 850 µm (Fig. 3). The enhanced SBR diminishes the requirement for increased frame averaging, enabling rapid data acquisition of dynamic processes.

Fig. 1. Polymer dot power dependence. (A) Logarithmic plots of the dependence of two- and three-photon induced fluorescence on excitation power. The excitation wavelength and fitted slope is indicated in the legend of each graph. The estimated uncertainty of each slope is reported as a standard deviation. Each plot corresponds to a distinct polymer dot species. (B) CNPPV (left), PFBT (middle), and PFPV (right) power dependence versus wavelength. The dashed and dash-dotted lines correspond to pure two- and three-photon power dependence, respectively. The blue shaded region represents the titanium-sapphire tuning range ($\lambda_{\text{ex}} = 700 – 1,000$ nm), the yellow shaded region represents the ytterbium-fiber laser’s bandwidth ($\lambda_{\text{ex}} = 1,060$ nm; $\Delta \lambda = 40$ nm), and the green shaded region represents the optical parametric amplifier tuning range ($\lambda_{\text{ex}} = 1,100 – 1,400$ nm).
Fig. 2. Titanium-sapphire in vivo vascular imaging of C57 mice. Reto-orbital injection labeling with polymer dots (Columns 1–3), dextran-conjugated fluorescein (Column 4), and semiconductor quantum dots (QD605; Column 5); λex = 800 nm; scale bars = 100 µm. Maximum intensity projections spanning 100 µm ranges were taken at various depth intervals in the cortex, with distance relative to the pial surface indicated across the left margin. Polymer dots yield appreciably higher signal-to-background ratio at all depths relative to fluorescein and QD605. Imaging of QD605-labeled vasculature beyond 550 µm was restricted due to the toxicity of the injection. All depth comparisons stem from age-matched litter mates; CNPPV, PFBT, and fluorescein imaging was performed in the same specimen.

3.3 Polymer dot excitability at longer wavelengths extends vascular imaging depth and signal-to-background ratio

A distinct characteristic of pdots is their wide absorption range [17]. The broad excitability of pdots enables their compatibility with a number of excitation sources including long wavelength, tunable laser sources between λex = 1,100 and 1,400 nm. The maximum imaging depth of MPM is ultimately determined by SBR, which is influenced by the scattering and absorption events that occur in biological tissue as well as the power dependence of an emitter. Excitation light that is attenuated by scattering or absorption before reaching a contrast agent fails to produce any emitted fluorescence, and the fraction of attenuated
photon is heavily dependent on wavelength. The fraction of excitation light reaching the focal volume at a depth, \( z \), can be approximated as \( \exp[-(\mu_a(\lambda) + \mu_s(\lambda))z] \), where \( \mu_a(\lambda) \) and \( \mu_s(\lambda) \) are wavelength-dependent absorption and scattering coefficients [22,35]. Experimental data of effective attenuation lengths in biological tissue between 1,300 nm and 1,700 nm supports the accuracy of this theoretical model [36]. Modeling of this function provided in Appendix D reveals that there is an ideal biological imaging wavelength situated around 1,300 nm [22] where photo-attenuation is minimized in brain tissue. In addition, 1,300 nm light can be used to achieve three-photon excitation of many fluorophores, which reduces out-of-focus excitation and thereby decreases background [7,37,38] compared with two-photon excitation. Therefore, it is readily understood that brighter fluorophores such as pdots that exhibit a strong 3P power dependence in response to 1,300 nm excitation should yield a markedly improved SBR and maximize imaging depth. Unfortunately, the 3PA cross section of a pdot such as CNPPV is quite low at 1,300 nm. However, a suitable compromise for an optimal imaging wavelength is at 1,225 nm where attenuation length remains relatively close to its maximum (Appendix D) and the slope of CNPPV’s power dependence is \( \sim 2.5 \) (Fig. 1), representing a combination of two- and three-photon excitation processes. Indeed, longer wavelength excitation of CNPPV-labeled vasculature (\( \lambda_{ex} = 1,225 \) nm) allows us to exploit these principles for optimized deep imaging and achieve a tissue penetration depth up to 1.3 mm beneath the pial surface (Fig. 4(A)). In contrast, two-photon imaging (\( \lambda_{ex} = 800 \) nm) of the same region of the same mouse only reached a maximum imaging depth of 850 \( \mu \)m (Fig. 4(B)). A comparison of the CNPPV-labeled vascular networks imaged at the different wavelengths shows that SBR is vastly improved by 1,225 nm excitation and a higher-order power dependence (Figs. 4(C) and 4(D)). The effect is pronounced enough that the SBR of the 1,225 nm image stack at \( z = 900 \) \( \mu \)m (SBR \( \sim 7.8 \)) exceeds the SBR of the 800 nm image stack near the cortical surface (SBR \( \sim 7.2 \); \( z = 350 \) \( \mu \)m). A plot of background signal versus depth shows that the difference in SBR is directly owed to the rapid increase of background introduced by 800 nm excitation relative to the modest rise of background seen from 1,225 nm excitation (Fig. 4(D)). Quantification of SBR and background throughout the superficial dura (\( z \sim 0 - 150 \) \( \mu \)m) is omitted from these plots due to high background caused by second harmonic generation of collagen.

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Fig. 4. Longer wavelength excitation \( \lambda_{ex} = 1,225 \) nm versus shorter wavelength excitation \( \lambda_{ex} = 800 \) nm of CNPPV-labeled C57 vasculature. (A) Sagittal (xz) projection of a 365 \( \times \) 365 \( \times \) 1,300 \( \mu \)m\(^3\) image stack at \( \lambda_{ex} = 1,225 \) nm using an optical parametric amplifier source. (B) Sagittal (xz) projection of a 356 \( \times \) 365 \( \times \) 850 \( \mu \)m\(^3\) image stack of the same region show in A at \( \lambda_{ex} = 800 \) nm using a titanium-sapphire source. (C) 2D tangential (xy) projections over shorter depth ranges of the stacks shown in A and B. (D) Comparison plots of signal-to-background ratio (SBR) versus depth (Left) and background intensity versus depth (Right). Scale bars = 100 \( \mu \)m.
3.4 High power fiber laser imaging of polymer dots significantly improves signal-to-background ratio beyond a ~600 µm depth

Pdots’ broad light absorption further permits excitation by a high average power ytterbium-fiber laser ($\lambda_{\text{ex}} = 1,060$ nm). Fiber lasers are a lower-cost alternative to OPA imaging, and offer excellent pulse characteristics as well as ease-of-operability to the user [39]. Our lab has recently demonstrated the use of such a system for deep in vivo imaging [4]. Although the fiber laser’s fixed output wavelength ($\lambda_{\text{ex}} = 1,060$ nm) does not coincide with the ideal biological imaging wavelength ($\lambda_{\text{ex}} = 1,300$ nm), a comparison of in vivo image stacks of PFPV-labeled C57 vasculature collected at $\lambda_{\text{ex}} = 1,060$ vs 800 nm clearly illustrates the advantages of the pdots’ excitability by a fiber laser (Fig. 5). Although the Ti:S penetration limit ($z = 850$ µm) is not significantly extended, the image quality (contrast and SBR) of the yb-fiber images is substantially improved under 1,060 nm excitation, an effect which is most prominent beyond ~600 µm (Fig. 5(A)). This corresponds to the z-position at which maximum Ti:S output power was reached whereas the upper limit of fiber laser output power was not met before all signal was lost. In addition to the higher average power of the yb-fiber relative to the Ti:S source, two primary advantages can be attributed for the improvement in SBR. First, the number of photons lost to scattering and absorption events is reduced by longer wavelength excitation ($\lambda_{\text{ex}} = 1,060$ nm vs. 800 nm) (Fig. 9 in Appendix D). At an 850 µm depth, the remaining photon fraction at $\lambda_{\text{ex}} = 1,060$ nm is approximately five times greater than at $\lambda_{\text{ex}} = 800$ nm. Second, PFPV exhibits a partial three-photon power dependence at 1,060 nm ($n \sim 2.38 \pm 0.08$) versus a two-photon excitation signature at 800 nm ($n \sim 1.93 \pm 0.14$) (Fig. 1). This partial three-photon dependence at $\lambda_{\text{ex}} = 1,060$ nm further improves SBR through suppression of background fluorescence due to a higher-order nonlinear dependence on excitation intensity. Thus, longer wavelength excitation of PFPV is expected to improve both the signal and background of images at all depths. This expectation is empirically demonstrated by line profiles drawn across identical in-plane blood vessels located 700 µm beneath the surface imaged separately at $\lambda_{\text{ex}} = 1,060$ and 800 nm (Figs. 5(B)-5(C)). The data reveals that the SBR of the $\lambda_{\text{ex}} = 1,060$ nm image is ~3.5 times greater than the $\lambda_{\text{ex}} = 800$ nm SBR. SBR improvement at z-positions beyond the depth at which Ti:S excitation becomes power-limited demonstrates that the higher output power of the yb-fiber laser coupled with longer wavelength excitation is a critical advantage for deep in vivo imaging. We emphasize that similar improvements made by longer wavelength excitation cannot be achieved with conventional fluorophores such as fluorescein ($\sigma_2 = 0.31$ at $\lambda_{\text{ex}} = 1,050$ nm) or Texas Red, which exhibit low action cross sections beyond the Ti:S range (>1,000 nm) and are unlikely to undergo a three-photon transition at 1,060 nm [3,40]. A major practical benefit of improved SBR is the ability to collect high-quality vessel line scans at greater depths to quantify blood velocity. Vessel line scanning quantifies flow velocity along the central axis of blood vessels at high frequency, relying on the contrast between injected plasma fluorophores and red blood cells (RBCs) which remain dark [25]. Therefore, high SBR is essential to accurately differentiate RBCs and fluorescent plasma and obtain precise flow velocity measurements. Here, we show that vessel line scans of PFPV-labeled mice form an image containing distinct streaks that represent RBCs traveling at a mean velocity of $1.28 \pm 0.20$ mm/sec at a depth of 750 µm beneath the pial surface (Fig. 5(D)).
**4. Discussion**

To evaluate pdots’ multiphoton properties, we prepared multiple polymer variants via nanoprecipitation, a simple and rapid procedure. The batch size could be easily varied to prepare ample amounts of the nanopolymers for vascular imaging of tens of mice per preparation. The polymer dots had the additional advantage of a long shelf life (upwards of 5 months) and easy storage, characteristics that allow them to be easily disseminated for broad use [41]. Through our imaging experiments, we demonstrate that pdots offer a wealth of optical properties that make them very well suited for deep, *in vivo* multiphoton fluorescence microscopy. However, the lack of knowledge surrounding their nonlinear excitation properties has prevented their widespread adoption as popular contrast agents in multiphoton fluorescence imaging. To remedy this, we have characterized the two- versus three-photon power dependence of three pdot variants (CNPPV, PFBT, and PFPV) representing different polymer classes including poly(phenylene vinylene) and fluorene-based copolymers (Fig. 1).

We found that all three pdot species demonstrate a 2P power dependence across the conventional Ti:S tuning range ($\lambda_{ex} = 780 – 850$ nm). Meanwhile, researchers who rely on the photophysical advantages that accompany 3PM, such as reduced out-of-focus fluorescence and diminished scattering, in order to improve the signal-to-background ratio in their imaging experiments can do so via excitation of CNPPV at 1,300 nm, PFBT at 1,350 nm, and PFPV at 1,325 nm. In contrast, two of the conventional organic dyes evaluated, Texas Red and ICG, remain rooted in the two-photon regime at comparable wavelengths ($\lambda_{ex} = 1,350$ nm and 1,280 nm, respectively) whereas only fluorescein possessed a 3P power dependence near this spectrum [3]. Furthermore, it is interesting to note that CNPPV undergoes a three-photon transition at a higher energy wavelength than either PFBT or PFPV, despite being the most
red-shifted of the three. This can potentially be explained by the unique chemical structure of poly(phenylene vinylene) polymers relative to fluorene-based co-polymers. The distinct molecular structures of the conjugated polymers affect the final energy levels of the pdot species in their excited states. Specifically, CNPPV has a donor-π-acceptor arrangement whereas PFBT and PFPV have donor-acceptor and donor-π-donor configurations, respectively, which leads to differences in the energy levels of their three-photon transitions.

Furthermore, we justify the use of pdots over conventional fluorophores for multiphoton imaging experiments in the 800 – 900 nm excitation range due to their increased brightness relative to quantum dots, organic dyes, and fluorescent proteins. However, we were unable to measure absolute cross-sections at longer wavelengths since excitation becomes a mixture of two- and three-photon absorption processes, which makes defining a cross-section ambiguous. Nevertheless, previously reported cross-section measurements demonstrate that pdots [20] are brighter than quantum dots, organic dyes, and fluorescent proteins [10, 40] within a 770-870 nm excitation range with peak cross-sections around ~2.02 to 5.50 × 10^5 GM, as illustrated in Fig. 10 in Appendix E. When compared to fluorescent dyes and proteins, pdot cross sections are three to five orders of magnitude larger. Most notably, this comparison includes organic molecules specifically engineered for enhanced two-photon absorption [30, 42]. In particular, many fluorescent dyes can be problematic within in vivo biological settings due to a reduced quantum yield and poor photostability in aqueous environments [11–13]. In contrast, pdots composed of hydrophobic conjugated polymers are strongly resistant to this effect [17]. Of course, highly emissive quantum dots such as QD605 [30] and QD535 [40] do come close to matching a dimmer pdot such as MEH-PPV in brightness. Therefore, one might be tempted to employ quantum dots as their preferred class of contrast agents due to their well-regarded photostability and spectrally broad light absorption, an essential feature for multicolor imaging experiments. However, pdots share these same properties [18] as well as the critical advantage of exhibiting no evidence of cytotoxicity. There are conflicting reports regarding quantum dot biocompatibility which can most likely be attributed to physiochemical and environmental factors [16]; yet in our hands, intravenous injection of QD605 was acutely toxic and fatal to mice whereas identical chronic studies with pdots did not produce any observed health concerns. Nevertheless, a biological mechanism of pdot blood clearance has yet to be thoroughly assessed and future research efforts should be directed to determine whether bioaccumulation in the spleen, liver, kidney or other major organs is a valid concern.

Next, we were able to show that pdots’ brightness and excitability in the NIR regime improves SBR at considerable depths in mouse cortex at an 800 nm excitation wavelength relative to identical experiments performed with an exceptionally bright quantum dot species, QD605, and a commonly used organic dye, dextran-conjugated fluorescein (Fig. 2). As noted above, QD605 is cytotoxic when administered intravenously, particularly at the high concentrations necessitated by deep imaging experiments. Ex vivo animal studies may not be hindered by such a consequence; however, in the case of vascular imaging, continuous circulation and blood flow is essential to avoid distorted vessel appearance and photobleaching of the now stagnant probes (Fig. 2, right). The overall imaging depth of the fluorescein-labeled vasculature matched that of the pdot-labeled mice; however, the SBR of the fluorescein images were less satisfactory resulting in reduced vascular clarity (~1.8-fold at \(z = 550 \mu m\); Appendix C). The implications of poor SBR include obfuscated analysis and degradation of automated image segmentation schemes. In this case, SBR improvement from brighter contrast agents implies that depth penetration and corresponding SBR was signal-limited. However, in contexts where background fluorescence becomes the limiting factor, such as in regions with large overlying blood vessels or high vascular density, background increases commensurately with signal, and SBR is predicted only by numerical aperture, depth, wavelength, and attenuation length, rather than by the brightness of the contrast agent [6].
Another substantial limitation of fluorescein and other conventional organic dyes is their narrow absorption range. At $\lambda_{ex} = 800$ nm, fluorescein’s 2PA action cross section is ~36 GM and at $\lambda_{ex} = 1,050$ nm its cross section is a barely detectable 0.31 GM [40]. In contrast, pdots exhibit broad multiphoton absorption, meaning that they can be efficiently excited with a variety of laser sources all the way out to 1,400 nm. We were able to demonstrate that the use of longer excitation wavelengths and higher-order nonlinear excitation allows us to improve SBR and overcome the tissue scattering limits of imaging depth observed with conventional 2PM. Specifically, excitation at $\lambda_{ex} = 1,225$ nm attains a 1,300 µm imaging depth whereas $\lambda_{ex} = 800$ nm results in an 850 µm cortical depth (Fig. 4). Moreover, the SBR of the images collected at the longer wavelength excitation greatly exceed those recorded at an 800 nm excitation at all depths, producing a much higher quality 3D volume. Through modeling and our power dependence characterization (Fig. 1) we are able to attribute this gain in penetration depth and SBR to the reduced scattering and photo-attenuation of longer excitation wavelengths, and the ancillary reduction of background signal due to CNPPV’s partial 3P power dependence at 1,225 nm. In regard to the neuroscience community, the extended imaging depth with longer wavelength excitation enables researchers to investigate neural layers of anatomy beyond the cortex and corpus callosum, most notably the hippocampus [7]. In addition, the biological imaging community can take advantage of pdots’ overlapping broad absorption spectra, which permits several pdot species emitting at discrete bands to be excited simultaneously at a single wavelength. This characteristic coupled with the fact that pdots can be readily functionalized and used to tag unique cellular structures [21] allows researchers to design simple and effective multicolor imaging experiments. An important caveat to consider, however, is that pdots were delivered intravenously in our studies, and labeling neural structures located in high-density extravascular brain tissue could pose a challenge due to the relatively large diameters of pdots (~20-30 nm). Recent efforts have produced pdot nanoparticles with sub-5 nm diameters, yet the yield from these preparations is still quite low [43].

We also take advantage of pdots’ broad light absorption to image PFPV-labeled mice and resolve vasculature using a custom, home-built ytterbium-fiber laser [4]. Again, we observe that reduced scattering of longer wavelength excitation light and a partial 3P power dependence improves imaging depth slightly ($z_{800 \text{ nm}} = 850$ µm; $z_{1,060 \text{ nm}} = 900$ µm) (Fig. 5). We note that the SBR of the 1,060 nm excitation images is improved most substantially beyond ~600 µm, an imaging depth which corresponds to the $z$-position at which the Ti:S output power was saturated.

Future advances in polymer engineering, structure based-design, and mutagenesis can vastly improve pdot performance and enhance multiphoton absorption at 1,300 nm for optimized deep imaging. Redesign and optimization of pdots’ molecular structure for improved three-photon imaging will involve an exploration of the effects of donor-$\pi$-donor, donor-$\pi$-acceptor-$\pi$-donor, and acceptor-$\pi$-donor-$\pi$-acceptor conformations on absorption cross sections and quantum yield. Overall, pdots present an exciting new approach to multiphoton in vivo imaging due to their enhanced brightness, broad excitability, and nontoxic features. With brighter, biocompatible probes, researchers will be able to resolve vascular architecture in living organisms with improved clarity and depth, enabling critical insights into fundamental biological problems.

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Disclosures

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Appendix A: Polymer dot structures, size distributions, and spectra

![Polymer dot structures, size distributions, and spectra](image)

Fig. 6. (A) Molecular structures of CNPPV (Left), PFBT (Center), and PFPV (Right). (B) Size distributions of CNPPV (21.89 nm), PFBT (31.47 nm), and PFPV (22.74 nm) as measured by dynamic light scattering. (C) Absorption (Left) and emission (Right) spectra of CNPPV, PFBT, and PFPV [17].

Appendix B: Schematic of the optical parametric amplifier imaging system

![Schematic of the optical parametric amplifier imaging system](image)

Fig. 7. A 5W pump (Verdi, Coherent) is used to seed a Ti:Sapphire (Ti:S) oscillator (Mira 900, Coherent), which is then stretched by a modified external stretcher/compressor and amplified by a Ti:S regenerative amplifier (RegA, Coherent) seeded by a 18W pump (Verdi G18, Coherent). The amplified pulse is then converted to a longer wavelength by an optical parametric amplifier tunable over a 1,100 – 1,400 nm range. The OPA output is compressed by a set of prisms before entering the scanning optics resulting in an objective-focused beam with a pulse width of ~45 fs.
Appendix C: *In vivo* imaging signal-to-background ratio as a function of depth

![Graph](image)

Fig. 8. Normalized signal-to-background (SBR) ratio versus depth comparison of polymer dots relative to fluorescein and QD605; \(\lambda_{ex} = 800\) nm. SBR is diminished at the surface due to high background signal from the dura resulting from second harmonic generation of collagen. At \(\sim 150\) \(\mu\)m SBR is at a maximum for all contrast agents and gradually decreases with depth. Relative to QD605 and fluorescein, all three polymer dots retain a higher SBR beyond \(\sim 150\) \(\mu\)m. To ensure fair comparisons, the average laser power at each depth was maintained at consistent levels across the separate imaging experiments.

Appendix D: Modeling of light attenuation due to scattering and absorption

![Graph](image)

Fig. 9. An ideal biological imaging wavelength is situated at 1,300 nm [22,35]. The blue line indicates the photon fraction at a 1 mm depth in brain tissue versus wavelength. The red line indicates the percent of photon absorbed versus wavelength. The wavelength excitation regions of the titanium-sapphire (beige), ytterbium-fiber laser (gray), and optical parametric amplifier (blue) laser systems are delineated by color.
Appendix E: Action cross-sections of polymer dots relative to quantum dot, organic dye, and fluorescent protein species

![Graph showing action cross-sections of polymer dots relative to quantum dot, organic dye, and fluorescent protein species.]

Fig. 10. Semilog plot of polymer dot brightness relative to quantum dots (QDs), organic dyes (ODs), and fluorescent proteins (FPs). The two-photon action cross-sections of polymer dots are plotted as circles connected by a solid line (PPFV in green, and MEH-PPV in red-orange) [20]. The blue shaded region corresponds to the expected range of quantum dots’ two-photon action (2PA) cross sections [10,30]. The wavelength-dependent 2PA cross section of an atypically bright quantum dot species, QD605, is represented by black triangles with a dashed and dotted line [30]. The red shaded region corresponds to the expected range of 2PA cross sections of fluorescent dyes and proteins [10]. The 2PA cross sections of two typical fluorescent dyes are represented by diamonds connected by dashed lines (rhodamine B in magenta, fluorescein in dark green) and the 2PA cross sections of two typical fluorescent proteins are represented by squares connected by dotted lines (eGFP in blue and YFP in black) [40].

References
