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Near-Infrared Squaraine Dye Encapsulated Micelles for In Vivo Fluorescence and Photoacoustic Bimodal Imaging

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**ABSTRACT:** Combined near-infrared (NIR) fluorescence and photoacoustic imaging techniques present promising capabilities for noninvasive visualization of biological structures. Development of bimodal noninvasive optical imaging approaches by combining NIR fluorescence and photoacoustic tomography demands suitable NIR-active exogenous contrast agents. If the aggregation and photobleaching are prevented, squaraine dyes are ideal candidates for fluorescence and photoacoustic imaging. Herein, we report rational selection, preparation and micelle encapsulation of an NIR absorbing squaraine dye (D1) for *in vivo* fluorescence and photoacoustic bimodal imaging. D1 was encapsulated inside micelles constructed from a biocompatible non-ionic surfactant (Pluronic F-127) to obtain D1-encapsulated micelles (D1_micelle) in aqueous conditions. The micelle encapsulation retains both photophysical features and chemical stability of D1. D1_micelle exhibits high photostability and low cytotoxicity in biological conditions. Unique properties of D1_micelle in the NIR window of 800 - 900 nm enable the development of squaraine-based exogenous contrast agent for fluorescence and photoacoustic bimodal imaging above 820 nm. *In vivo* imaging using D1_micelle, as demonstrated by fluorescence and photoacoustic tomography experiments in live mice, shows contrast-enhanced deep tissue imaging capability. The usage of D1_micelle proven by pre-clinical experiments in rodents reveals its excellent applicability for NIR fluorescence and photoacoustic bimodal imaging.

**KEYWORDS:** bioimaging, fluorescence tomography, micelles, photoacoustic tomography, squaraine dye
Novel optical imaging techniques, such as photoacoustic tomography (PAT), and their capabilities to probe structural, functional and molecular states of biological specimens have been widely investigated during the past decades, and some of them have been accepted for clinic usage.\textsuperscript{1-3} On the other hand, the development of background-free optical imaging techniques that can provide images with rich optical contrast \textit{in vivo} by capitalizing on selective photon absorption is of great significance in the field of biomedical research.\textsuperscript{4} However, each single imaging modality suffers inherent limitations such as limited tissue penetration, low sensitivity and failing to provide anatomy information precisely.\textsuperscript{5} Multimodal combinations of imaging modalities are expected to compensate for inherent limitations of individual imaging modality, and thus emerging as an inevitable trend in the development of new imaging techniques.\textsuperscript{6,7} In optical imaging, a potential imaging approach that could offer tremendous opportunities for future clinic applications is to blend the salient features of fluorescence imaging and PAT at near-infrared (NIR) excitation.\textsuperscript{8,9} Given the unique capabilities of such emerging multimodal imaging combinations, the design and development of new multimodal contrast agents in the NIR region are greatly expected.\textsuperscript{10-12} However, the design of NIR contrast agents that can be used in conjugation with these multimodal noninvasive optical imaging modalities remains challenging.\textsuperscript{13-15} An ideal contrast agent for NIR fluorescence and photoacoustic imaging should typically offer excellent NIR fluorescence emissions and intense absorption along with good photochemical stability and low cytotoxicity in aqueous and biological conditions.\textsuperscript{16,17} As a promising class of NIR dyes, squaraines have proven their versatility in a wide range of imaging applications.\textsuperscript{18-22} In this context, the selection of squaraines as bimodal contrast agents for NIR fluorescence and photoacoustic imaging is justified, since they possess excellent chemical and photophysical properties in the NIR region.
Figure 1. Schematic preparation of D1\textsubscript{micelle} for fluorescence and photoacoustic bimodal imaging. (A) Scheme showing the preparation of D1\textsubscript{micelle} and a single micelle containing D1. (B) Scheme showing intravenous injection of D1\textsubscript{micelle} for fluorescence and photoacoustic bimodal imaging in a mouse.

The main challenge that prevents the effective use of NIR properties of squaraine dyes relies on their ease formation of aggregates in aqueous/biological conditions.\textsuperscript{23} Thus, researchers have introduced new strategies to control their self-assembly, thereby enabling the efficient use of squaraines for various biological applications.\textsuperscript{24-26} Recently, a new type of modified
dicyanovinyl substituted squaraines was reported and their photophysical properties were studied.\textsuperscript{27-29} The substituted dicyanovinyl squaraine core allows for efficient tuning of NIR properties in the first NIR window. On the other hand, the encapsulation of squaraine dyes within micelles for multiphoton fluorescence bioimaging was established.\textsuperscript{30} Therefore, a rational selection of a squaraine derivative in conjugation with micelle-encapsulated protection strategy en route the development of a new class of bimodal exogenous contrast agents for fluorescence and photoacoustic bimodal imaging in the first NIR window of 650-950 nm is of great importance.

Herein, we report the preparation of a new 4-methylquinolium based dicyanovinyl substituted squaraine with oxy-ethylene side chains (D1) and the encapsulation of D1 inside micelles constructed from a biocompatible non-ionic surfactant (Pluronic F-127) to afford squaraine-encapsulated micelle D$_{1\text{micelle}}$ (Figure 1). It was observed that the as-synthesized D$_{1\text{micelle}}$ exhibits high chemical stability and low cytotoxicity, and could retain photophysical properties of D1 in the NIR region. The promising potential of D$_{1\text{micelle}}$ for \textit{in vivo} NIR fluorescence and photoacoustic bimodal imaging was then demonstrated. Based on a literature survey as shown in Table S1 of the Supporting Information, the present work using squaraine-encapsulated micelle (D$_{1\text{micelle}}$) is the first example of such kinds, demonstrating an excellent bimodal imaging capability at 840 nm.

RESULTS AND DISCUSSION

D1 with oxy-ethylene side chains was synthesized based on previous literature reports with some modifications.\textsuperscript{27,31} Condensation of dicyanovinyl substituted squaric acid with two equivalents of \textit{N}-glycolated 4-methylsquinolium iodide in a 1:1 azeotropic mixture of benzene
and n-butanol afforded D1 in 9-10% yield. As-synthesized dye was then characterized by NMR and high-resolution mass spectrometry (see the Supporting Information for more synthesis and characterization details). Photophysical properties of D1 were investigated in dimethylsulphoxide (DMSO). D1 (1 × 10^{-4} M) in DMSO exhibits intense absorption at 867 nm and emission at 907 nm with a Stoke shift of 40 nm (Figure 2a,b). Inset of Figure 2b shows the false-color pixel intensity map corresponding to the emission of D1 in DMSO at room temperature.

**Figure 2.** Photophysical properties of D1. a) UV/Vis absorption spectrum and b) emission spectrum (excited at 840 nm) of D1 in DMSO (1 × 10^{-4} M). Inset of b) shows fluorescence of D1 in DMSO as false-color pixel intensity. c) Changes in the absorption spectra of D1 in DMSO (1
× 10⁻⁴ M) upon increasing content of water from 0-90%. d) False-color pixel intensity map showing fluorescence images of D1 in DMSO and 90% water/DMSO mixture under 800 nm excitation.

The absorption spectra of D1 in DMSO show a significant blue shift followed by bleaching of optical properties upon increasing the water content from 0 to 90%. Figure 2c reveals the changes in the absorption spectra of D1 in DMSO upon the addition of water, indicating the feasibility of rapid aggregate formation (H-aggregate) of D1 in aqueous conditions.²³ The false-color pixel intensity mapping also exhibits a significant emission quenching in water, which further confirms the formation of aggregates in situ (Figure 2d). In order to preserve the photophysical properties and stability of D1 in aqueous conditions, we adopted the micelle encapsulation strategy for further studies.

Figure 3. a) TEM image of D₁₉micelle in water and b) high magnification TEM image of a single D₁₉micelle particle. c) Dynamic light scattering data showing the hydrodynamic size of D₁₉micelle in PBS buffer at pH 7.4.
The micelle encapsulation of D1 was carried out in dichoromethane / phosphate buffered saline (PBS) mixture by following previously reported procedures (see the Methods section for experimental details).32,33 In our study, we selected poloxamer 407, a biocompatible polymer commonly known as Pluoron F-127 (PF-127),34 for the micelle fabrication. PF-127 has been approved by US Food and Drug Administration (FDA) and studied extensively for sustained delivery of pharmaceutical active ingredients.35,36 After the preparation of D1micelle, it was characterized by transmission electron microscope (TEM) and dynamic light scattering (DLS) techniques. Figure 3a shows the TEM image of D1micelle in PBS buffer. The TEM analysis reveals uniform spherical micelle formation with a good contrast. Apparently, the presence of organic dye inside the micelle enhances the visibility of spherical morphology during TEM analysis. A high-resolution TEM image (Figure 3b) of single D1micelle particle clearly indicates the formation of micelles with an average diameter of 100±30 nm. Furthermore, DLS measurement (Figure 3c) shows a hydrodynamic diameter of 184±30 nm with a narrow size distribution for D1micelle, which is consistent with the observation from TEM images.

We then investigated the optical properties, the effect of nucleophiles and the stability of D1micelle in biological fluids. The effect of pH on the photophysical properties of D1micelle in comparison with D1 alone was also studied. UV/Vis absorption spectrum of D1micelle in PBS at pH 7.4 exhibits broad absorption ranging from 800 to 900 nm with the maximum absorbance at 840 nm corresponding to the encapsulated D1 (Figure 4a). This monomeric existence of D1 inside micelles demonstrates successful preservation of its optical characteristics in completely aqueous conditions. The absorption band below 700 nm may be due to the scattering from D1micelle. Figure 4b shows the fluorescence spectrum of D1micelle in PBS at pH 7.4. A blue shift in the emission maximum was observed in aqueous solution, which was a common phenomenon
attributed to the existence of D1 in non-polar microenvironment inside the micelle.\textsuperscript{30,32} A fluorescence false-color pixel intensity mapping of D1\textsubscript{micelle} shows intense emission at 840 nm in PBS buffer at pH 7.4 (Figure 4b, inset).

\textbf{Figure 4.} a) UV/Vis absorption and b) fluorescence emission spectra of D1\textsubscript{micelle} in PBS buffer at pH 7.4. Inset of Figure b shows false-color pixel intensity image corresponding to fluorescence of D1\textsubscript{micelle} at 840 nm ($\lambda_{ex}$ 745/30 nm). c,d) False-color pixel intensity images showing the stability of D1 and D1\textsubscript{micelle}: (c1) HBP alone, (c2) D1 (1 $\times$ 10$^{-4}$ M) in HBP, and (c3) D1\textsubscript{micelle} in HBP; (d1) BSA alone, (d2) D1 (1 $\times$ 10$^{-4}$ M) in BSA, and (d3) D1\textsubscript{micelle} in BSA. e) Fluorescence mapping image of a portion of 96 well plate showing pH effect to D1 (A1-A5), micelle alone
Indocyanine green (ICG) is a commonly used biocompatible dye as a contrast agent in various imaging modalities. We compared the photophysical properties of D1micelle with ICG in water in order to prove the significance of micelle encapsulation strategy. ICG exhibits broader absorption spectrum with a maximum peak at 810 nm in PBS at pH 7.4. The spectral signature of D1micelle shows absorption in the visible and NIR regions with a maximum intensity at 840 nm (Figure S1a in the Supporting Information). Compared to ICG, D1micelle exhibits a 30 nm red shift in the absorption maximum, which indicates its increase in the effective range of analysis. Figure S1b in the Supporting Information shows a comparison of concentration dependent fluorescence mapping of a micro-well plate with D1micelle (A1-A5) and ICG (B1-B5) in PBS at pH 7.4. It is evident that D1micelle exhibits intense fluorescence intensities in buffered aqueous conditions at concentrations ranging from 10 to 0.6 µM. Thus, the potential capability of D1micelle for fluorescence imaging was proven as compared to a known standard dye.

D1 in the presence of a nucleophile such as cysteine undergoes the addition reaction, thereby bleaching all its photophysical properties (Figure S2a in the Supporting Information). However, D1micelle shows no changes in the absorption maximum at 840 nm, which indicates the immunity of D1micelle towards cysteine addition even at higher concentrations (Figure S2b in the Supporting Information). Photophysical stability of D1 (1 × 10^{-4} M) and D1micelle in biological fluids was then investigated by incubating them in human blood plasma (HBP) and bovine serum albumin (BSA) at 37°C for 1h. Fluorescence mapping (\(\lambda_{em} \ 840\ nm, \ \lambda_{ex} \ 745/30\ nm\)) was then carried out. Figure 4c shows the fluorescence false-color pixel intensity mapping of HBP after 1h.
incubation with D1 and D1\textsubscript{micelle} at 37°C, respectively. D1 underwent the aggregation in HBP, which was indicated by visible precipitation in HBP. On the other hand, D1\textsubscript{micelle} was found to be stable, and its fluorescence was retained in HBP, which was obviously evident from the fluorescence intensity map. The stability of D1 and D1\textsubscript{micelle} was also investigated in BSA solution, and similar phenomena to the case in HBP were observed. Figure 4d shows the corresponding fluorescence mapping images. Furthermore, the pH effect on the stability and photophysical properties of D1, micelle alone, and D1\textsubscript{micelle} was studied in the pH range of 6.6-7.4. Figure 4e shows the false-color fluorescence mapping image ($\lambda_{em}$ 840 nm) of a 96-well plate section filled with D1 (A1-5), micelle alone (B1-5), and D1\textsubscript{micelle} (C1-5) in the pH range of 6.6-7.4 (from no. 1 to no. 5 plate). Intense fluorescence at 840 nm was observed from D1\textsubscript{micelle} in comparison with D1 and micelle without the dye. Figure 4f shows the summary of fluorescence intensity variation based on Figure 4e. Thus, the high stability of D1 inside the micelle (D1\textsubscript{micelle}) in different pH is evident from the comparison plot of fluorescence intensity at 840 nm.

Inherent cytotoxicity of D1\textsubscript{micelle} was evaluated using the MTT (3-(4,5-dimethylthiazolyl-2)-2,5-diphenyltetrazolium bromide) viability assay with human hepatoma cell line (Huh-7) incubated for 24 h. The cell viability was measured by varying the concentrations of D1\textsubscript{micelle} in micro molar range. Summary of the viability assay is shown in Figure S3 of the Supporting Information, which clearly indicates that D1\textsubscript{micelle} has no considerable cytotoxicity at low to moderate concentrations. The low cytotoxicity and good solubility in aqueous conditions as well as retained NIR photophysical properties encouraged us to further investigate the imaging capability of D1\textsubscript{micelle} both in vitro and in vivo.
Figure 5. Fluorescence microscope images of Huh-7 cell line incubated with (a-c) D1 and (d-f) D1\textsubscript{micelle} (10 µM) for 12h. Fluorescence images of cells incubated with D1: a) the cell nuclei were stained with DAPI (excitation at 405 nm, DAPI channel), b) emission at 820-850 nm obtained by exciting at 808 nm, and c) overlay image of a and b. Fluorescence images of cells incubated with D1\textsubscript{micelle}: d) DAPI channel (excitation at 405 nm), e) emission at 820-850 nm obtained by exciting at 808 nm, and f) overlay image of d and e. Scale bar: 20 µM.

The \textit{in vitro} fluorescence imaging capability of D1\textsubscript{micelle} was assessed in comparison with D1 alone by NIR fluorescence imaging in Huh-7 cell line. Aqueous solutions of D1 alone and D1\textsubscript{micelle} were incubated separately with Huh-7 cells for 12 h, and were then imaged under 808 nm excitation. Figure 5 shows the fluorescence microscopy images of Huh-7 cells (40x objective) treated with D1 alone and D1\textsubscript{micelle}, respectively. The blue fluorescence from the cell nucleus stained with 4′,6-diamidino-2-phenylindole (DAPI) could be observed in Figure 5a,d. The cells incubated with D1 alone show no emission fluorescence collected at 820-850 nm.
(Figure 5b and overlay image Figure 5c) upon excitation at 808 nm, indicating a failed endocytosis or aggregation of D1 in cellular environments. Huh-7 cells incubated with \( \text{D1}_{\text{micelle}} \) clearly show high fluorescence intensity at 820-850 nm with a complete distribution in the cytoplasm of the cells. Figures 5e and 5f are the fluorescence images of the cells incubated with \( \text{D1}_{\text{micelle}} \). Together with the low cytotoxicity obtained from the MTT assay and the fluorescence imaging experiments, the promising potential of \( \text{D1}_{\text{micelle}} \) for \textit{in vitro} fluorescence bioimaging was successfully proven.

\textit{In vivo} fluorescence imaging was then carried out using a live seven-week-old severe combined immune deficiency (SCID) female mouse. Belly fur was removed using a depilatory cream prior to imaging. The \textit{in vivo} fluorescence imaging studies on the mouse were performed using IVIS lumina II preclinical imaging system and analyzed using the IVIS Living Imaging 4.4. The animal was anesthetized with isoflurane, and the fluorescence images were acquired from the dorsal side of the mouse (see the Methods section for more details). Fluorescence imaging of the mouse placed in the supine position was performed in epifluorescence mode before and after the administration of \( \text{D1}_{\text{micelle}} \). Same illumination and acquisition settings were used for all fluorescence image acquisitions. Figure 6a shows the fluorescence image of the mouse before the injection of \( \text{D1}_{\text{micelle}} \). No detectable fluorescence signals were observed when imaged with 745/30 nm excitation and 840/20 nm emission settings. Then, \( \text{D1}_{\text{micelle}} \) (200 \( \mu \)L) was injected intravenously, and the \( \text{D1}_{\text{micelle}} \) concentration was calculated as \( 1.47 \times 10^{-5} \) M per intravenous injection after optimizing 4 repeated trials. The thoracic/abdominal area of the mouse was imaged within 35 minutes after the administration. Figure 6b presents the top view fluorescence image of the mouse at 840 nm excitation. After imaging experiments, the mouse was euthanized, its organs were dissected, and the obtained sections were imaged. As evident
from Figure 6c, the accumulation of $D_1_{\text{micelle}}$ was primarily in liver ($2.4 \times 10^8$ p/sec/cm$^2$ upon setting a threshold intensity of $1.5 \times 10^8$ p/sec/cm$^2$). Figure S4 in the Supporting Information shows the photograph of fluorescence reflectance image for the internal organs of mouse imaged at 840 nm. Clear localization of $D_1_{\text{micelle}}$ in the liver (abdominal cavity) was observed, indicating the effective fluorescence tracing of $D_1_{\text{micelle}}$ metabolism and accumulation. In short, in vitro and in vivo fluorescence imaging experiments demonstrate the high stability and potential of $D_1_{\text{micelle}}$ as an NIR imaging probe for fluorescence microscopy and tomographic applications.

**Figure 6.** (a,b) Tracking of $D_1_{\text{micelle}}$ distribution in a live mouse before and after intravenous injection using the IVIS imaging system. Fluorescence reflectance images from dorsal side of female mouse were measured (a) before and (b) after two minutes of intravenous injection with $D_1_{\text{micelle}}$ (200 µL). (c) Quantitative fluorescence reflectance image analysis showing $D_1_{\text{micelle}}$ distribution in major organs (threshold intensity of $1.5 \times 10^8$ p/sec/cm$^2$).

Photoacoustic imaging capability of $D_1_{\text{micelle}}$ was initially studied using a tissue-mimicking phantom prior to in vivo imaging investigations. Photoacoustic imaging studies were
performed using the multi-spectral optoacoustic tomography (MSOT) system. Detailed descriptions of the imaging system and experimental parameters are given in the Materials and Methods section. A cylindrical tissue-mimicking phantom having 4 mm diameter with two inclusions each was used to evaluate the photoacoustic signal generation from $D_1_{\text{micelle}}$. $D_1_{\text{micelle}}$ in PBS buffer at pH 7.4 was loaded inside the first inclusion of the phantom, and the second inclusion containing the PBS buffer at pH 7.4 served as the control. Photoacoustic signals were then measured for excitation wavelengths ranging from 800 to 900 nm with 15 nm step intervals. Figure 7a displays the absorption spectrum (red dots) of $D_1_{\text{micelle}}$ in PBS buffer at pH 7.4 along with corresponding photoacoustic signal intensities for $D_1_{\text{micelle}}$ (black square) and control (blue triangle) across 800-900 nm. Multispectral scanning experiments show the maximum signal intensity in the range of 815-840 nm, which indicate efficient photoacoustic signal generation capability of $D_1_{\text{micelle}}$ across these excitation wavelengths. Furthermore, the photoacoustic signal intensities obtained from each excitation wavelength match well with the absorption spectrum of $D_1_{\text{micelle}}$, indicating high photostability of $D_1_{\text{micelle}}$ when used for photoacoustic studies. Figure 7b shows the corresponding photoacoustic image of phantom loaded with $D_1_{\text{micelle}}$ (red arrow) and blank control (PBS alone, green arrow). High signal intensity from $D_1_{\text{micelle}}$ could be observed at 830 nm when compared to the control. Similarly, the photoacoustic images of phantom at distinct excitation wavelengths from 800 nm to 900 nm were also generated (Figure 7c). The multispectral photoacoustic images show that the maximum image contrast is obtained at 830 nm, which in turn corresponds to the wavelength at which $D_1_{\text{micelle}}$ exhibits the maximum absorbance. Thus, the photoacoustic image contrast varies linearly according to the absorption spectrum of $D_1_{\text{micelle}}$. 
**Figure 7.** Multispectral photoacoustic imaging of tissue-mimicking phantom containing D1\textsubscript{micelle}.

a) Black squares and blue triangles represent multispectral photoacoustic mean intensity of D1\textsubscript{micelle} and blank control (PBS at pH 7.4) respectively inside tissue-mimicking phantom. Red spectrum shows corresponding absorption of D1\textsubscript{micelle} in PBS at pH 7.4. b) Single wavelength photoacoustic image of phantom acquired at 830 nm (scale bar: 3 mm). Red arrow and green arrow indicate the channels containing D1\textsubscript{micelle} and blank solution, respectively. c) Spectrally resolved MSOT images of D1\textsubscript{micelle} distribution in the phantom at excitation wavelength of i) 815, ii) 830, iii) 840, and iv) 860 nm.

To further investigate the *in vivo* photoacoustic imaging capability of D1\textsubscript{micelle}, we carried out MSOT on an adult female mouse before and after intravenous injection of D1\textsubscript{micelle}. 
Transverse slices of the mouse were formed by averaging photoacoustic signals acquired from 5 laser pulses for each excitation wavelength ranging from 800 to 900 nm with 15 nm step intervals. A tomographic view of 172° could be achieved with the curvature of detector containing 64 element array transducer having 5 MHz central frequency. During imaging, the mouse was translated across a distance of 150 mm with 1 mm step size to form a transverse slice of the mouse at each position. Photoacoustic signals were acquired using a multi-channel digitizer and then processed using standard machine integrated ViewMSOT software. Photoacoustic images were reconstructed using a model-based approach and the photoacoustic signals from D1micelle were spectrally resolved using principal component analysis (PCA) based spectral unmixing technique. Images were acquired before (pre-scan) and after (post scan) intravenous (tail vein) injection of D1micelle.

Figure 8 depicts spectrally resolved photoacoustic images (selected) at 840 nm of wavelength from D1micelle within 35 min after the intravenous injection of D1micelle. Schematic diagrams showing the scanned area and the anatomy of the section with the highest contrast are given in Figure 8a and 8b. No obvious contrast was observed for D1micelle during pre-scan (Figure 8c). Figure S5a in the Supporting Information shows the corresponding three-dimensional (3D) rendering of images during the pre-scan. A full set of images containing scans at 1 mm distance is provided in Figure S6 of the Supporting Information (also see Movie M1). After 35 min of D1micelle injection, a clear contrast difference could be visualized in the abdominal region (Figure 8d) for D1micelle. Corresponding 3D rendering images are shown in Figure S5b of the Supporting Information. A full set of images containing scans at 1 mm distance is provided in Figure S7 of the Supporting Information (also see Movie M2). The
obvious post injection images provide solid evidence in favor of photoacoustic imaging capability of D1_micelle for \textit{in vivo} imaging applications.

\textbf{Figure 8.} Single wavelength photoacoustic images of live mouse anatomy at 840 nm. a) Schematic diagram of completely scanned area. b) Schematic section corresponding to a single area under analysis. c) Individual anatomy sections of live mouse before intravenous injection of
D1$_{\text{micelle}}$ (scale bar: 3 mm). d) Individual anatomy sections recorded after 35 min post-intravenous injection of D1$_{\text{micelle}}$ (scale bar: 3 mm).

CONCLUSIONS

In summary, we have designed, synthesized and characterized an NIR active 4-methylquinolinium based dicyanovinyl substituted squaraine dye with oxyethylene side chains (D1). D1 was further encapsulated within micelles formed from a biocompatible non-ionic surfactant (Pluronic F-127) to achieve D1$_{\text{micelle}}$. Easy aggregation of D1 in aqueous conditions was prevented via the micelle encapsulation strategy. Thus, D1$_{\text{micelle}}$ exhibited intense absorption and fluorescence properties from 800 to 900 nm in aqueous conditions with a maximum absorption at 840 nm. The low cytotoxicity, high stability in aqueous conditions, and reserved NIR photophysical properties make D1$_{\text{micelle}}$ an exceptional probe for fluorescence and photoacoustic bimodal imaging at 840 nm when compared to recently reported NIR probes (Table S1 in the Supporting Information). The in vitro and in vivo fluorescence imaging and in vivo photoacoustic imaging studies show that the developed D1$_{\text{micelle}}$ possesses tremendous capabilities for fluorescence and photoacoustic bimodal imaging.

In this work, we have successfully validated the strategy of micelle encapsulation in order to maintain the photophysical properties of monomeric squaraine dye in aqueous conditions for in vivo fluorescence and photoacoustic bimodal imaging. This strategy could be generalized to other dye molecules having similar issues by selecting suitable encapsulation platforms. In the follow-up studies, we would like to incorporate targeting ligands within the dye-encapsulated micelles for targeted fluorescence and photoacoustic bimodal imaging. In this case, the dye-encapsulated micelles could be precisely localized in certain types of cells or organs for imaging.
Such targeted imaging could guide the disease treatment, which is especially important for tumor therapy. Thus, the advanced development of the bimodal imaging system will be translated from mice to human for clinical uses in the near future.

METHODS

All chemical reagents, unless otherwise specified, were purchased from Sigma-Aldrich Co. All solvents were of reagent grade and were purchased from local companies. All solvents were dried and distilled prior to use by following standard procedures. \(^1\)H NMR spectra were recorded on a Bruker 300 MHz FT-NMR (model: Advance-DPX 300) spectrometer at 25°C. The chemical shift (\(\delta\)) data and coupling constant (J) values were given in parts per million (ppm) and Hertz, respectively, unless otherwise mentioned. Electrospray ionization mass spectrometry (ESI MS) measurements were carried out on a Waters QTof-Micro instrument. The high-resolution time-of-flight mass spectrometry (TOF MS) was performed on a Waters Q-tof Premier MS spectrometer. Elemental analysis was performed on a EuroVector Euro EA elemental analyzer. UV/Vis spectra were obtained by using a Shimadzu UV-3600 UV-vis-NIR spectrometer. Steady-state emission spectra at room temperature were obtained using a Shimadzu RF-5301PC spectrofluorimeter. Transmission electron microscopy (TEM) images were recorded on a JEM 1400 electron microscope (120 kV) equipped with slow scan CCD using cold cathode field emission as the gun. The samples were prepared by dropping a droplet of the sample solution onto a TEM grid (copper grid, 300 meshes, coated with carbon film) and allowing air-dry. Dynamic light scattering (DLS) measurements were carried out with a Zetasizer Nano ZS instrument from Malvern Instruments Ltd. at 298 K using a 633 nm “red” laser. The mean hydrodynamic size was calculated with Zetasizer software. Fluorescence
tomography studies were performed using IVIS lumina II preclinical imaging system and analyzed using the IVIS Living Imaging 4.4 software (PerkinElmer Inc., Alameda, CA, USA). Photoacoustic imaging studies of the tissue mimicking phantom and live mouse were performed using the multispectral optoacoustic tomography (MSOT) system (MSOT insight 64, iThera Medical GmbH).

**Preparation of D1micelle**: In a typical experiment, D1 (1 mg mL\(^{-1}\)) and concentrated PF-127 surfactant (1.5 \times 10^{-4}\) M) were dissolved in dichloromethane (10 mL) in an open round bottom flask. An equal volume of PBS buffer at pH 7.4 was then added to the mixture, which was allowed to stir at room temperature for 24 h. Complete evaporation of dichloromethane led to the formation of dye encapsulated micelles (D1micelle). As-prepared D1micelle was then filtered and dialyzed against water for 24 h in order to remove impurities and big particles.

**In vitro cytotoxicity assay**: MTT assay test was carried out by following standard procedures. Huh-7 cells were seeded into a 96-well plate (1×10^4 cells per well) in DMEM (Dulbecco’s Modified Eagle’s Medium) cell culture medium containing 10% fetal bovine serum and grown under a humidified atmosphere with 5% CO\(_2\) at 37 °C. After 12 h incubation, the media in the wells were replaced with fresh DMEM (100 µL per well) containing D1micelle with different concentrations, and the cells were further incubated for 6h. Then, the medium was changed by DMEM (100 µL per well) containing MTT (0.5 mg mL\(^{-1}\)), followed by incubation for another 4h. The culture medium was removed and frozen crystals were dissolved with freshly prepared DMSO (100 µL). Before the cytotoxicity measurement, the plate was agitated gently for 15 min, and then the absorbance intensity at 560 nm was recorded by a micro plate reader. The relative cell viability (%) for each sample related to the control well was finally calculated.
**In vivo fluorescence tomography:** In vivo fluorescence tomography studies of the live mouse was performed using IVIS lumina II preclinical imaging system and analyzed using the IVIS Living Imaging 4.4 software. A back-thinned back illuminated CCD camera having 2048 x 2048 pixels cooled to -90°C was used as the detector. Similar illumination and acquisition settings were used for acquiring all the fluorescence images. Images were acquired with binning factor 8, 4 sec exposure time, and 12.9 cm field of view under 745/30 nm excitation and 840/20 nm emission filters. Mouse was then placed on a temperature-controlled stage inside the closed chamber of the imaging system equipped with an integrated isoflurane based anaesthesia system.

**Multispectral optoacoustic studies:** This study conforms to the Guide for the Care and Use of Laboratory Animals published by the National Institutes of Health, USA and protocol approved by the Institutional Animal Care and Use Committee (IACUC), National University of Singapore. Photoacoustic imaging studies of the tissue-mimicking phantom and live mouse were performed using the MSOT system. The imaging system typically consists of an illumination source, a spherical array ultrasound probe, parallel data acquisition system and a computer. An optical parametric oscillator based laser, generating 5 ns duration pulses having energy of 20 mJ and 10 Hz repetition rate, serves as the illumination source. The output of the laser source could be tuned from 680 nm to 980 nm on a per pulse basis. The output beam of the laser was coupled into a fiber bundle and was guided onto the surface of the imaging sample. Photoacoustically generated volumetric data were detected using the 64 element piezoelectric transducer array having a spherical geometry, which can provide a tomographic view of 172°. Belly fur of the mouse was removed using the depilatory cream prior to any imaging procedure. By following previously explained anesthesia procedure, the animal was anesthetized and placed in supine position over a thin polyethylene membrane, which was then positioned inside the imaging
chamber filled with deionized water. The polyethylene membrane prevents the direct contact of mouse with water and permits excellent acoustic coupling between the mouse and detector array. The water bath was temperature controlled, and the mouse holder could be translated across the imaging plane to obtain multiple transverse image slices of the mouse. Photoacoustic images were acquired with an in-plane resolution of approximate 150 μm.

ASSOCIATED CONTENT

Supporting Information. Additional experimental details. This material is available free of charge via the Internet at http://pubs.acs.org.

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ACKNOWLEDGMENT

This research is supported by the National Research Foundation (NRF), Prime Minister’s Office, Singapore under its NRF Fellowship (NRF2009NRF-RF001-015) and Campus for Research Excellence and Technological Enterprise (CREATE) Programme–Singapore Peking University Research Centre for a Sustainable Low-Carbon Future, as well as the NTU-A*Star Silicon Technologies Centre of Excellence under program grant no. 112 351 0003. We thank Dr. A. K. Mandal and Dr. S. K. Maji for their assistance during the synthesis and characterization. We are grateful to Dr. Tingchao He for his assistance during the fluorescence measurements. We thank www.designs4research.com for the graphical representation.
REFERENCES AND NOTES


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NIR Squaraine Encapsulated Micelles

Mode 1: Fluorescence Imaging

Mode 2: Photoacoustic Imaging