

## Compared to Purpurinimides, the Pyropheophorbide Containing an Iodobenzyl Group Showed Enhanced PDT Efficacy and Tumor Imaging ( $^{124}\text{I}$ -PET) Ability

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Two positional isomers of purpurinimide, 3-[1'-(3-iodobenzoyloxyethyl)] purpurin-18-*N*-hexylimide methyl ester **4**, in which the iodobenzyl group is present at the top half of the molecule (position-3), and a 3-(1'-hexyloxyethyl)purpurin-18-*N*-(3-iodo-benzylimide)] methyl ester **5**, where the iodobenzyl group is introduced at the bottom half (*N*-substituted cyclicimide) of the molecule, were derived from chlorophyll-*a*. The tumor uptake and phototherapeutic abilities of these isomers were compared with the pyropheophorbide analogue **1** (lead compound). These compounds were then converted into the corresponding  $^{124}\text{I}$ -labeled PET imaging agents with specific activity  $>1\text{ Ci}/\mu\text{mol}$ . Among the positional isomers **4** and **5**, purpurinimide **5** showed enhanced imaging and therapeutic potential. However, the lead compound **1** derived from pyropheophorbide-*a* exhibited the best PET imaging and PDT efficacy. For investigating the overall lipophilicity of the molecule, the 3-*O*-hexyl ether group present at position-3 of purpurinimide **5** was replaced with a methyl ether substituent, and the resulting product **10** showed improved tumor uptake, but due to its significantly higher uptake in the liver, spleen, and other organs, a poor tumor contrast in whole-body tumor imaging was observed.

### INTRODUCTION

Positron emission tomography (PET) has wide appeal for research at the drug development stage as it allows studying the drug distribution noninvasively (1). Dedicated animal PET systems whose resolution could reach near 1 mm have intensified this field by enabling drug studies with murine disease models. In recent years,  $^{18}\text{F}$ -fluorodeoxyglucose ( $^{18}\text{F}$ -FDG) has been the primary PET tracer. It is being used in the evaluation of several neoplasms, both before and after therapy, as well as the planning of the radiotherapy in various cancers. Its use in the assessment of cancer after therapy, including restaging tumors and monitoring tumor response has been of particular interest for oncologists. However,  $^{18}\text{F}$ -FDG suffers from pitfalls in cases such as where tumors are not metabolically active enough. Additionally, a short half-life of  $^{18}\text{F}$ -isotope (110 min) limits its use in studies involving antibodies and photosensitizers (PS) related to porphyrins for use in photodynamic therapy (PDT), which take considerably longer time to accumulate in a tumor in high concentrations (2). In this respect,  $^{124}\text{I}$  is a better choice due to its half-life of 4.2 days (3, 4). The labeling technique for  $^{124}\text{I}$ -nuclide is now well established, and this approach is continuously being followed to label a variety of biologically active molecules (5–12). In the past few years, various porphyrin-based photosensitizers have been labeled/

chelated with  $^{111}\text{In}$ ,  $^{113}\text{Sn}$ , and  $^{99\text{m}}\text{Tc}$  radionuclide for the purpose of scintigraphy (SPECT) (13–21). The metalation of the porphyrin core, however, alters the physicochemical characteristics of the molecule. The clinically effective PS have also been labeled with  $^{131}\text{I}$ ,  $^{14}\text{C}$ , and  $^3\text{H}$  mainly for the purpose of understanding their pharmacokinetic and pharmacodynamic characteristics (22–26). There are a few reports in the field of PDT where various F-18 based radiotracers such as  $^{18}\text{F}$ -FDG (27–29),  $^{18}\text{F}$ -FHBG{9-(4- $^{18}\text{F}$ -fluoro-3-hydroxymethyl-butyl)guanine} (30),  $^{18}\text{F}$ -FLT (3'-deoxy-3'- $^{18}\text{F}$ -fluorothymidine) (31), and  $^{99\text{m}}\text{Tc}$ -Annexin V (32) have been used to monitor cellular events during and post-PDT in vivo.

One of the main reasons for using the porphyrin-based compounds in phototherapy is their ability to retain in tumor (33–35). The tumor localizing ability of certain porphyrins has also been explored in developing multifunctional agents, where these compounds were used as vehicles to deliver the desired imaging moieties (fluorescence, MRI, and PET) to tumors (36–41).

For quite some time, one of the objectives of our laboratory has been to develop photosensitizers with long absorption wavelengths  $>700$ . Irradiation of tumors with light in this range should allow deeper tissue penetration, which may help in treating the large and deeply seated tumors. To achieve our goal, we modified the in vivo unstable purpurin-18 and bacteriopurpurin systems, and synthesized a series of tumor-avid *N*- and *O*-alkyl- or the corresponding trifluoromethylbenzyl substituted purpurinimides (700 nm) and bacteriopurpurinimides (800 nm) with variable lipophilicity (42, 43). Some of the synthetic analogues showed promising in vivo activity. Interestingly, among the compounds with similar lipophilicity, the position of the substituents at various peripheral positions of the tetrapyrrolic system showed a significant difference in long-term tumor response.

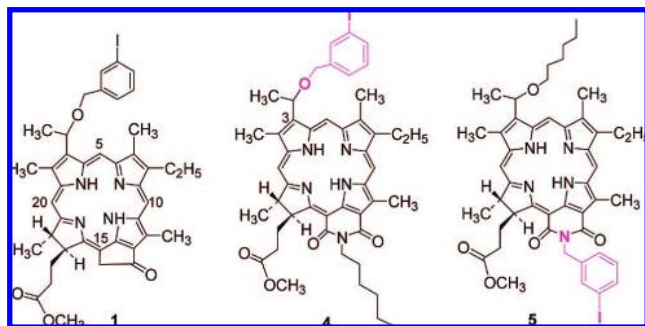
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**Figure 1.** Structures of 3-(1'-*m*-iodobenzyloxy) pyropheophorbide **1** and the iodobenzyl purpurinimides **4** and **5** (isomers).

Herein, we report the synthesis of the iodobenzyl substituted purpurinimides **4** and **5** (positional isomers; Figure 1) and their significant difference in PDT efficacy. The comparative bio-distribution properties of the corresponding  $^{125}\text{I}$ -analogues as well as their tumor imaging (PET) abilities are also discussed.

## EXPERIMENTAL PROCEDURES

**Chemistry.** All chemicals were of reagent grade and used as such. Solvents were dried using standard methods. Reactions were carried out under nitrogen atmosphere and were monitored by pre-coated (0.20 mm) silica TLC plastic sheet (20 × 20 cm) strips (POLYGRAM SIL N-HR) and/or UV-visible spectroscopy. Silica gel 60 (70–230 mesh, Merck) was used for column chromatography. Melting points were determined on a Fisher–Johns melting point apparatus. UV-visible spectra were recorded on a Varian (Cary-50 Bio) spectrophotometer.  $^1\text{H}$  NMR spectra were recorded on a Bruker AMX 400 MHz NMR spectrometer at 303 K. Proton chemical shifts ( $\delta$ ) are reported in parts per million (ppm) relative to  $\text{CDCl}_3$  (7.26 ppm), pyridine- $d_5$  (7.22 ppm, most downfield), or TMS (0.00 ppm). Coupling constants ( $J$ ) are reported in Hertz (Hz), and s, d, t, q, p, m, and br refer to singlet, doublet, triplet, quartet, pentet, multiplet, and broad, respectively. HRMS data were obtained from the Mass Spectroscopy facility of Michigan State University. Analytical HPLC was used to assess the purity of compounds. A Waters (Milford, MA) system including a Waters 600 Controller, a Delta 600 pump, and a 996 Photodiode Array Detector was used. A reverse phase, Symmetry C18, 5  $\mu\text{m}$ , 4.6 × 150 mm column (Waters, Made in Ireland) was used under an isocratic setting of MeOH/ $\text{H}_2\text{O}$  for final compounds (**1**, **4**, **5**, and the corresponding trimethyl stannyl analogues). The solvent flow rate was kept constant at 1.00 mL/min, and the detector was set at 254, 410, 535, and 660 nm (for pyropheophorbides-*a* **1**, and its trimethyl stannyl derivative) and 254, 415, 545, and 700 nm (for purpurinimides **4**, **5**, and their trimethyl stannyl derivatives). All final products were found to be >95% pure, and their retention time is reported in the characterization section. Reactions were first carried out for nonradioactive iodine and analyzed in the above HPLC system. However, in the case of final I-124 radiolabeling HPLC data obtained from the above system were transferred to another system comprising a Chrom Tech Iso-2000 pump, Hitachi L-4000 UV detector, and a radiation detector. These detectors are connected to a computer with HP Chemstation software via an HP 35900E interface. A Bioscan system 200 imaging scanner was used for thin layer chromatography of the radiolabeled compounds.

**Synthesis of 3-{1'-(3-Iodobenzyloxy)ethyl}pyropheophorbide-*a* Methyl Ester (**1**).** It was synthesized by following our previously reported procedure (40). Yield: 77%. MP = 112–114 °C; analytical RP HPLC (95/5: MeOH/ $\text{H}_2\text{O}$ ):  $t_R$  = 20.97 min, >96%. UV-vis (MeOH): 662 ( $4.75 \times 10^4$ ), 536 ( $1.08 \times 10^4$ ),

505 ( $1.18 \times 10^4$ ), 410 ( $1.45 \times 10^5$ ).  $^1\text{H}$  NMR ( $\text{CDCl}_3$ ; 400 MHz):  $\delta$  9.76, 9.55 and 8.56 (all s, 1H, meso-H); 7.76 (s, 1H, ArH); 7.64 (d,  $J$  = 6.8, 1H, ArH); 7.30 (d,  $J$  = 8.0, 1H, ArH); 7.05 (t,  $J$  = 8.2, 1H, ArH); 6.00 (q,  $J$  = 6.9, 1H, 3<sup>1</sup>-H); 5.28 (d,  $J$  = 19.8, 1H, 13<sup>2</sup>-CH<sub>2</sub>); 5.13 (d,  $J$  = 19.8, 1H, 13<sup>2</sup>-CH<sub>2</sub>); 4.70 (d,  $J$  = 12.0, 1H, OCH<sub>2</sub>Ar); 4.56 (dd,  $J$  = 3.2, 11.6, 1H, OCH<sub>2</sub>Ar); 4.48–4.53 (m, 1H, 18-H); 4.30–4.33 (m, 1H, 17-H); 3.72 (q,  $J$  = 8.0, 2H, 8-CH<sub>2</sub>CH<sub>3</sub>); 3.69, 3.61, 3.38 and 3.21 (all s, all 3H, for 17<sup>3</sup>-CO<sub>2</sub>CH<sub>3</sub> and 3 × ring CH<sub>3</sub>); 2.66–2.74, 2.52–2.61 and 2.23–2.37 (m, 4H, 17<sup>1</sup>-H and 17<sup>2</sup>-H); 2.18 (dd,  $J$  = 2.8, 6.4, 3H, 3<sup>1</sup>-CH<sub>3</sub>); 1.83 (d,  $J$  = 8.0, 3H, 18-CH<sub>3</sub>); 1.72 (t,  $J$  = 7.6, 3H, 8-CH<sub>2</sub>CH<sub>3</sub>); 0.41 (brs, 1H, NH); -1.71 (brs, 1H, NH). HRMS for C<sub>41</sub>H<sub>43</sub>N<sub>4</sub>O<sub>4</sub>I: 783.2329 (calculated, M + 1); found, 783.2407. Anal. Calcd. for C<sub>41</sub>H<sub>43</sub>N<sub>4</sub>O<sub>4</sub>I: C, 62.91; H, 5.54; N, 7.16; I, 16.21. Found: C, 62.60; H, 5.59; N, 7.13; I, 16.45.

**Synthesis of 3-{1'-(3-Iodobenzyloxy)ethyl}purpurin-18-*N*-hexylimide Methyl Ester (**4**).** 30% Hydrobromic acid (HBr) in acetic acid (2 mL) was added to purpurin-18-*N*-hexylimide methyl ester (100 mg, 0.15 mmol) (**23**), and the reaction was stirred at room temperature for 2 h. After evaporating the acids under high vacuum (0.1 mmHg), excess of 3-iodobenzyl alcohol (0.45 mL, 20-fold excess), dry dichloromethane (5 mL), and anhydrous potassium carbonate (40 mg) were added. The reaction mixture was stirred under nitrogen atmosphere for 45 min. It was then diluted with dichloromethane (200 mL), washed with aqueous sodium bicarbonate solution (100 mL), and then with water (2 × 200 mL). The dichloromethane layer was dried over anhydrous sodium sulfate, concentrated, and treated with diazomethane. Evaporation of the solvent gave a syrupy residue, which was chromatographed over a silica column using (1:4) ethyl acetate/hexane as eluant to remove excess 3-iodobenzyl alcohol, followed by (1:1) ethyl acetate/hexane to yield 110 mg (81%) of the desired compound **4**, which is sticky in nature. Analytical HPLC (Symmetry C18; 99/1: MeOH/ $\text{H}_2\text{O}$ ):  $t_R$  = 38.71 min, >96%. UV-vis (MeOH): 701 ( $4.31 \times 10^4$ ), 545 ( $2.15 \times 10^4$ ), 508 ( $7.32 \times 10^3$ ), 414 ( $1.31 \times 10^5$ ).  $^1\text{H}$  NMR ( $\text{CDCl}_3$ ; 400 MHz):  $\delta$  9.72 (s, 1H, meso-H); 9.66 (s, 1H, meso-H); 8.58 (s, 1H, meso-H); 7.75 (s, 1H, ArH); 7.64 (d,  $J$  = 9.2, 1H, ArH); 7.29 (d,  $J$  = 7.2, 1H, ArH); 7.06 (dt,  $J$  = 2.4, 7.4, 1H, ArH); 5.88 (q,  $J$  = 6.2, 1H, 3<sup>1</sup>-H); 5.41 (d,  $J$  = 8.8, 1H, 17-H); 4.68 (dd,  $J$  = 2.6, 12.6, 1H, OCH<sub>2</sub>Ar); 4.55 (dd,  $J$  = 3.2, 12.0, 1H, OCH<sub>2</sub>Ar); 4.45 (t,  $J$  = 6.8, 2H, NCH<sub>2</sub>(CH<sub>2</sub>)<sub>4</sub>CH<sub>3</sub>); 4.37 (q,  $J$  = 7.2, 1H, 18-H); 3.84 (s, 3H, 12-CH<sub>3</sub>); 3.68 (q,  $J$  = 7.6, 2H, 8-CH<sub>2</sub>CH<sub>3</sub>); 3.56 (s, 3H, 17<sup>2</sup>-CO<sub>2</sub>CH<sub>3</sub>); 3.31 (split s, 3H, 2-CH<sub>3</sub>); 3.14 (s, 3H, 7-CH<sub>3</sub>); 2.63–2.72 (m, 1H, 17<sup>1</sup>-H); 2.38–2.48 (m, 1H, 17<sup>2</sup>-H); 2.26–2.36 (m, 1H, 17<sup>1</sup>-H); 2.12 (dd,  $J$  = 2.2, 6.6, 3H, 3<sup>1</sup>-CH<sub>3</sub>); 1.95–2.05 (m, 3H, 17<sup>2</sup>-H, and NCH<sub>2</sub>(CH<sub>2</sub>)<sub>3</sub>CH<sub>3</sub>); 1.77 (d,  $J$  = 7.2, 3H, 18-CH<sub>3</sub>); 1.68 (t,  $J$  = 7.6, 3H, 8-CH<sub>2</sub>CH<sub>3</sub>); 1.56–1.61 (m, 2H, N(CH<sub>2</sub>)<sub>2</sub>CH<sub>2</sub>(CH<sub>2</sub>)<sub>2</sub>-CH<sub>3</sub>); 1.38–1.50 (m, 4H, N(CH<sub>2</sub>)<sub>3</sub>(CH<sub>2</sub>)<sub>2</sub>CH<sub>3</sub>); 0.95 (t,  $J$  = 7.0, 3H, N(CH<sub>2</sub>)<sub>5</sub>CH<sub>3</sub>); -0.14 (brs, 1H, NH); -0.19 (brs, 1H, NH). HRMS for C<sub>47</sub>H<sub>54</sub>N<sub>5</sub>O<sub>5</sub>I: 896.3171 (calculated, M + 1); found, 896.3241.

**Synthesis of 3-{1'-(Hexyloxy)ethyl}purpurin-18-*N*-(3-iodo)benzylimide Methyl Ester (**5**).** Hydrobromic acid (HBr) (30%) in acetic acid (2 mL) was added to purpurin-18-*N*-(3-iodo)benzylimide methyl ester (100 mg, 0.125 mmol) (**23**), and the reaction was stirred at room temperature for 2 h. After evaporating the acids under high vacuum (0.1 mmHg), an excess of *n*-hexanol (0.5 mL, 30-fold excess), dry dichloromethane (5 mL), and anhydrous potassium carbonate (25 mg) were added to the residue. The reaction mixture was stirred under nitrogen atmosphere for 45 min. It was then diluted with dichloromethane (200 mL), washed with aqueous sodium bicarbonate solution (100 mL), and then with water (2 × 200 mL). The dichloromethane layer was dried over anhydrous sodium sulfate,

concentrated, and treated with diazomethane. Evaporation of the solvent gave a syrupy residue, which was chromatographed over a silica column using 1% acetone in dichloromethane as eluant to yield 95 mg (85%) of the desired compound **5**, which is sticky in nature. Analytical HPLC (Symmetry C18; 99/1: MeOH/H<sub>2</sub>O):  $t_R = 48.97$  min, >96%. UV-vis (MeOH): 700 ( $4.31 \times 10^4$ ), 545 ( $2.05 \times 10^4$ ), 508 ( $7.76 \times 10^3$ ), 414 ( $1.29 \times 10^5$ ). <sup>1</sup>H NMR (CDCl<sub>3</sub>; 400 MHz):  $\delta$  9.75 (splits, 1H, meso-H); 9.62 (s, 1H, meso-H); 8.53 (s, 1H, meso-H); 8.09 (s, 1H, ArH); 7.70 (d,  $J = 7.6$ , 1H, ArH); 7.66 (d,  $J = 8.0$ , 1H, ArH); 7.60 (d,  $J = 7.6$ , 1H, ArH); 7.32 (d,  $J = 8.4$ , 1H, ArH); 7.10 (t,  $J = 7.8$ , 1H, ArH); 5.77 (q,  $J = 6.8$ , 1H, 3<sup>1</sup>-H); 5.61 (s, 2H, NCH<sub>2</sub>Ar); 5.37 (d,  $J = 8.4$ , 1H, 17-H); 4.35 (q,  $J = 7.2$ , 1H, 18-H); 3.81 (s, 3H, 12-CH<sub>3</sub>); 3.65 (m, 4H, 8-CH<sub>2</sub>CH<sub>3</sub>, OCH<sub>2</sub>(CH<sub>2</sub>)<sub>4</sub>CH<sub>3</sub>); 3.56 (s, 3H, 17<sup>2</sup>-CO<sub>2</sub>CH<sub>3</sub>); 3.30 (s, 3H, 2-CH<sub>3</sub>); 3.18 (s, 3H, 7-CH<sub>3</sub>); 2.62–2.72 (m, 3H, OCH<sub>2</sub>-CH<sub>2</sub>(CH<sub>2</sub>)<sub>3</sub>CH<sub>3</sub> and 17<sup>1</sup>-H); 2.30–2.45 (m, 2H, 17<sup>2</sup>-H and 17<sup>1</sup>-H); 2.05 (dd,  $J = 2.4$ , 6.8, 3H, 3<sup>1</sup>-CH<sub>3</sub>); 1.95–2.02 (m, 1H, 17<sup>2</sup>-H); 1.70–1.80 (m, 5H, 18-CH<sub>3</sub> and O(CH<sub>2</sub>)<sub>2</sub>CH<sub>2</sub>(CH<sub>2</sub>)<sub>2</sub>CH<sub>3</sub>); 1.67 (t,  $J = 8.0$ , 3H, 8-CH<sub>2</sub>CH<sub>3</sub>); 1.23 (m, 4H, O(CH<sub>2</sub>)<sub>3</sub>(CH<sub>2</sub>)<sub>2</sub>CH<sub>3</sub>); 0.78 (m, 3H, O(CH<sub>2</sub>)<sub>5</sub>CH<sub>3</sub>); 0.04 (brs, 1H, NH); –0.07 (brs, 1H, NH). HRMS for C<sub>47</sub>H<sub>54</sub>N<sub>5</sub>O<sub>5</sub>I: 896.3171 (calculated, M + 1); found, 896.3241.

**Synthesis of 3-{1'-(3-Trimethylstannylbenzyloxy)ethyl}purpurin-18-N-hexylimide Methyl Ester (6).** To a solution of 3-{1'-(3-iodobenzyloxy)ethyl}purpurin-18-N-hexylimide methyl ester (**4**) (15 mg, 0.017 mmol) in dry THF (10 mL) were added hexamethylditin (15  $\mu$ L, 0.072 mmol) and bis-(triphenylphosphine)palladium (II) dichloride (5 mg), and the reaction mixture was stirred at 60 °C for 2 h. The reaction mixture was rotavapoed to dryness, and the crude product was purified over a silica gel column using CH<sub>2</sub>Cl<sub>2</sub> as eluant to yield 12 mg (77%) of title compound **6**. Analytical HPLC (Symmetry C18; 99/1: MeOH/H<sub>2</sub>O):  $t_R = 48.19$  min, >95%. <sup>1</sup>H NMR (CDCl<sub>3</sub>; 400 MHz):  $\delta$  9.73 (s, 1H, meso-H); 9.66 (s, 1H, meso-H); 8.56 (s, 1H, meso-H); 7.42 (m, 2H, ArH); 7.34 (m, 2H, ArH); 5.90 (m, 1H, 3<sup>1</sup>-H); 5.41 (dd,  $J = 2.6$ , 9.0, 1H, 17-H); 4.76 (dd,  $J = 4.4$ , 11.6, 1H, OCH<sub>2</sub>Ar); 4.55 (dd,  $J = 1.2$ , 12.0, 1H, OCH<sub>2</sub>Ar); 4.45 (m, 2H, NCH<sub>2</sub>(CH<sub>2</sub>)<sub>4</sub>CH<sub>3</sub>); 4.36 (q,  $J = 7.3$ , 1H, 18-H); 3.85 (s, 3H, 12-CH<sub>3</sub>); 3.68 (q,  $J = 7.6$ , 2H, 8-CH<sub>2</sub>CH<sub>3</sub>); 3.55 (s, 3H, 17<sup>2</sup>-CO<sub>2</sub>CH<sub>3</sub>); 3.31 (split s, 3H, 2-CH<sub>3</sub>); 3.11 (s, 3H, 7-CH<sub>3</sub>); 2.63–2.72 (m, 1H, 17<sup>1</sup>-H); 2.38–2.48 (m, 1H, 17<sup>2</sup>-H); 2.26–2.36 (m, 1H, 17<sup>1</sup>-H); 2.12 (dd,  $J = 2.4$ , 6.8, 3H, 3<sup>1</sup>-CH<sub>3</sub>); 1.95–2.05 (m, 3H, 17<sup>2</sup>-H and NCH<sub>2</sub>CH<sub>2</sub>(CH<sub>2</sub>)<sub>3</sub>CH<sub>3</sub>); 1.77 (d,  $J = 7.2$ , 3H, 18-CH<sub>3</sub>); 1.68 (t,  $J = 7.6$ , 3H, 8-CH<sub>2</sub>CH<sub>3</sub>); 1.61 (m, 2H, N(CH<sub>2</sub>)<sub>2</sub>CH<sub>2</sub>(CH<sub>2</sub>)<sub>2</sub>CH<sub>3</sub>); 1.38–1.50 (m, 4H, N(CH<sub>2</sub>)<sub>3</sub>(CH<sub>2</sub>)<sub>2</sub>CH<sub>3</sub>); 0.95 (t,  $J = 7.2$ , 3H, N(CH<sub>2</sub>)<sub>5</sub>CH<sub>3</sub>); 0.18 (s, 9H, Sn(CH<sub>3</sub>)<sub>3</sub>); –0.09 (brs, 1H, NH); –0.16 (brs, 1H, NH).

**Synthesis of 3-{1'-(Hexyloxy)ethyl}purpurin-18-N-(3-trimethylstannyl)benzylimide Methyl Ester (7).** It was synthesized following the procedure described above for **6** from the respective compound 3-{1'-(hexyloxy)ethyl}purpurin-18-N-(3-iodo)benzylimide methyl ester (**5**). Yield: 75%. Analytical HPLC (Symmetry C18; 99/1: MeOH/H<sub>2</sub>O):  $t_R = 53.13$  min, >95%. <sup>1</sup>H NMR (CDCl<sub>3</sub>; 400 MHz):  $\delta$  9.75 (splits, 1H, meso-H); 9.64 (s, 1H, meso-H); 8.53 (s, 1H, meso-H); 7.89 (s, 1H, ArH); 7.66 (d,  $J = 8.0$ , 1H, ArH); 7.39 (d,  $J = 7.2$ , 1H, ArH); 7.32 (m, H, ArH); 5.78 (q,  $J = 6.6$ , 1H, 3<sup>1</sup>-H); 5.68 (q,  $J = 13.8$ , 2H, NCH<sub>2</sub>Ar); 5.38 (d,  $J = 8.0$ , 1H, 17-H); 4.35 (m, 1H, 18-H); 3.83 (s, 3H, 12-CH<sub>3</sub>); 3.57–3.68 (m, 4H, 8-CH<sub>2</sub>CH<sub>3</sub>, OCH<sub>2</sub>(CH<sub>2</sub>)<sub>4</sub>CH<sub>3</sub>); 3.54 (s, 3H, 17<sup>2</sup>-CO<sub>2</sub>CH<sub>3</sub>); 3.30 (s, 3H, 2-CH<sub>3</sub>); 3.18 (s, 3H, 7-CH<sub>3</sub>); 2.62–2.72 (m, 3H, OCH<sub>2</sub>CH<sub>2</sub>(CH<sub>2</sub>)<sub>3</sub>CH<sub>3</sub> and 17<sup>1</sup>-H); 2.30–2.45 (m, 2H, 17<sup>2</sup>-H and 17<sup>1</sup>-H); 2.05 (dd,  $J = 2.6$ , 6.6, 3H, 3<sup>1</sup>-CH<sub>3</sub>); 1.95–2.02 (m, 1H, 17<sup>2</sup>-H); 1.73 (d,  $J = 7.6$ , 3H, 18-CH<sub>3</sub>); 1.67 (t,  $J = 7.6$ , 3H, 8-CH<sub>2</sub>CH<sub>3</sub>);

1.25 (m, 6H, O(CH<sub>2</sub>)<sub>2</sub>(CH<sub>2</sub>)<sub>3</sub>CH<sub>3</sub>); 0.78 (m, 3H, O(CH<sub>2</sub>)<sub>5</sub>CH<sub>3</sub>); 0.27 (s, 9H, Sn(CH<sub>3</sub>)<sub>3</sub>); –0.01 (brs, 1H, NH); –0.13 (brs, 1H, NH).

**Synthesis of 3-{1'-(Methoxy)ethyl}purpurin-18-N-(3-iodo)benzylimide Methyl Ester (10).** It was prepared by following the method described for compound **5** except the intermediate bromo-derivative was reacted with methanol, instead of *n*-hexanol. Pure product was obtained by column chromatography over a silica column using 1% acetone in dichloromethane as eluant. Analytical HPLC (Symmetry C18; 99/1: MeOH/H<sub>2</sub>O):  $t_R = 31.42$  min, >98%. UV-vis (CH<sub>2</sub>Cl<sub>2</sub>): 700 ( $4.31 \times 10^4$ ), 545 ( $2.12 \times 10^4$ ), 414 ( $1.29 \times 10^5$ ). <sup>1</sup>H NMR (CDCl<sub>3</sub>; 400 MHz):  $\delta$  9.66 (s, 1H, meso-H); 9.64 (s, 1H, meso-H); 8.54 (s, 1H, meso-H); 8.08 (s, 1H, ArH); 7.70 (d,  $J = 8.0$ , 1H, ArH); 7.60 (d,  $J = 8.0$ , 1H, ArH); 7.10 (t,  $J = 8.0$ , 1H, ArH); 5.73 (q,  $J = 6.9$ , 1H, 3<sup>1</sup>-H); 5.63 (s, 2H, NCH<sub>2</sub>Ar); 5.38 (dd,  $J = 1.6$ , 8.4, 1H, 17-H); 4.35 (q,  $J = 7.2$ , 1H, 18-H); 3.84 (s, 3H, 12-CH<sub>3</sub>); 3.67 (q,  $J = 7.4$ , 2H, 8-CH<sub>2</sub>CH<sub>3</sub>); 3.56 (s, 3H, 17<sup>2</sup>-CO<sub>2</sub>CH<sub>3</sub>); 3.54 (d,  $J = 2.8$ , 3H, OCH<sub>3</sub>); 3.32 (s, 3H, 2-CH<sub>3</sub>); 3.19 (s, 3H, 7-CH<sub>3</sub>); 2.62–2.72 (m, 1H, 17<sup>1</sup>-H); 2.30–2.45 (m, 2H, 17<sup>2</sup>-H and 17<sup>1</sup>-H); 2.06 (dd,  $J = 2.2$ , 6.6, 3H, 3<sup>2</sup>-CH<sub>3</sub>); 1.92–2.02 (m, 1H, 17<sup>2</sup>-H); 1.76 (d,  $J = 6.0$ , 3H, 18-CH<sub>3</sub>); 1.68 (t,  $J = 7.6$ , 3H, 8-CH<sub>2</sub>CH<sub>3</sub>); 0.03 (brs, 1H, NH); –0.07 (brs, 1H, NH). HRMS for C<sub>42</sub>H<sub>44</sub>N<sub>5</sub>O<sub>5</sub>I: 826.2387 (calculated, M + 1); found, 826.2470.

**Synthesis of 3-{1'-(Methoxy)ethyl}purpurin-18-N-(3-trimethylstannyl)benzylimide Methyl Ester (11).** The title compound was synthesized following the procedure described above for compound **7** from the respective compound 3-{1'-(methoxy)ethyl}purpurin-18-N-(3-iodo)benzylimide methyl ester (**10**). Yield: 80%. Analytical HPLC (Symmetry C18; 99/1: MeOH/H<sub>2</sub>O):  $t_R = 35.79$  min, >98%. <sup>1</sup>H NMR (CDCl<sub>3</sub>; 400 MHz):  $\delta$  9.68 (s, 1H, meso-H); 9.65 (s, 1H, meso-H); 8.56 (s, 1H, meso-H); 7.90 (s, 1H, ArH); 7.68 (d,  $J = 7.6$ , 1H, ArH); 7.40 (d,  $J = 6.8$ , 1H, ArH); 7.34 (t,  $J = 7.2$ , 1H, ArH); 5.60–5.80 (m, 3H, 3<sup>1</sup>-H and NCH<sub>2</sub>Ar); 5.38 (dd,  $J = 2.0$ , 6.8, 1H, 17-H); 4.36 (q,  $J = 7.2$ , 1H, 18-H); 3.84 (s, 3H, 12-CH<sub>3</sub>); 3.67 (q,  $J = 7.4$ , 2H, 8-CH<sub>2</sub>CH<sub>3</sub>); 3.56 (s, 3H, 17<sup>2</sup>-CO<sub>2</sub>CH<sub>3</sub>); 3.55 (d,  $J = 2.8$ , 3H, OCH<sub>3</sub>); 3.33 (s, 3H, 2-CH<sub>3</sub>); 3.20 (s, 3H, 7-CH<sub>3</sub>); 2.62–2.72 (m, 1H, 17<sup>1</sup>-H); 2.30–2.50 (m, 2H, 17<sup>2</sup>-H and 17<sup>1</sup>-H); 2.08 (dd,  $J = 2.2$ , 6.6, 3H, 3<sup>2</sup>-CH<sub>3</sub>); 1.92–2.02 (m, 1H, 17<sup>2</sup>-H); 1.76 (d,  $J = 7.2$ , 3H, 18-CH<sub>3</sub>); 1.68 (t,  $J = 7.8$ , 3H, 8-CH<sub>2</sub>CH<sub>3</sub>); 0.28 (s, 9H, Sn(CH<sub>3</sub>)<sub>3</sub>); –0.03 (brs, 1H, NH); –0.12 (brs, 1H, NH).

**Radioactive Labeling.** <sup>124</sup>I-analogues of **4**, **5**, and **10** were prepared from the corresponding trimethylstannyl analogues **6**, **7**, and **11**, respectively, by following the procedure as described below for the <sup>124</sup>I-analogue of compound **4**.

**Synthesis of <sup>124</sup>I-Labeled Analogue of 3-{1'-(3-iodobenzyl)oxyethyl}purpurin-18-N-hexylimide Methyl Ester (4).** The trimethyltin analogue **6** (50  $\mu$ g) was dissolved in 50  $\mu$ L of 5% acetic acid in methanol. Then, 100  $\mu$ L of 5% acetic acid in methanol was added to Na<sup>124</sup>I in 10  $\mu$ L of 0.1 N NaOH. The two solutions were mixed, and an IODOGEN bead (Pierce Biotechnology, Inc., Rockford, IL 61106) was added. The reaction mixture was incubated at room temperature for 15 min, the iodobead was removed, and the reaction mixture was injected on an HPLC column (Symmetry C18 5  $\mu$ m, 150  $\times$  4.6 mm), which was eluted with an isocratic 99/1 MeOH/H<sub>2</sub>O at a flow rate of 1 mL/min. The UV detector was set at 254 nm wavelength. The labeled product (**4**) eluted at 46.7 min was collected, and the solvent was evaporated to dryness under a stream of N<sub>2</sub> at 60 °C. The product was formulated in saline containing 10% ethanol for in vivo experiments. RadioTLC confirmed the radiochemical purity (>95%) of the product. A standard curve was generated between peak area versus mass by injecting known mass of carrier **4** onto the column. The mass

associated with the labeled product was calculated by relating the peak area of the UV absorbance peak of **4** in the labeled product to the standard curve. The specific activity was obtained by dividing the activity of the labeled product collected by the calculated mass in micromoles. Specific activity of the radio labeled product was  $>1$  Ci/ $\mu$ mol. The radiochemical yield was found to be 40%.

**PET Imaging.** Mice were imaged in the microPET FOCUS 120, a dedicated 3D small-animal PET scanner (Concorde Microsystems Incorporated) at State University of New York at Buffalo (south campus) under the Institutional Animal Care and Use Committee (IACUC) guidelines. The C3H mice were subcutaneously injected with  $3 \times 10^5$  RIF cells in 30  $\mu$ L complete  $\alpha$ -MEM (into the axilla), and tumors were grown until they reached 4–5 mm in diameter (approximately 5 days). All tumored C<sub>3</sub>H mice were injected via the tail vein 50–200  $\mu$ Ci of **1**, **8**, **9**, and **12**.

After 24, 48, 72, and 96 h postinjection, the mice were anesthetized by inhalation of isoflurane/oxygen, placed head first prone for imaging, and the acquisition time was set for 30 min. Radioiodine uptake by the thyroid or stomach was not blocked.

**Biodistribution Studies.** All studies were performed as per IACUC guidelines. The mice were injected with 50–200  $\mu$ Ci of **1**, **8**, **9**, and **12** via tail vein, and 3 or 4 mice each at 24, 48, 72, and 96 h time interval were sacrificed and body organs (tumor, heart, liver, spleen, kidney, lung, muscle, etc.) removed immediately. After weighing, the amount of radioactivity in the tumor (50–150 mg), body organs, and blood was measured by a gamma well counter. Radioactivity uptake was calculated as the percentage of the injected dose per gram of the tissue (%ID/g). Statistical analyses and data (%ID/g vs time point) were plotted using Microsoft Excel.

**In Vitro Photosensitizing Efficacy.** The photosensitizing activity of **4**, **5**, and **10** was determined in the RIF tumor cell line. The RIF tumor cells were grown in  $\alpha$ -MEM with 10% fetal calf serum, L-glutamine, penicillin, streptomycin, and neomycin. Cells were maintained in 5% CO<sub>2</sub>, 95% air, and 100% humidity. For determining the PDT efficacy, these cells were plated in 96-well plates at a density of  $5 \times 10^3$  cells/well in complete media. After an overnight incubation at 37 °C, the photosensitizers were added at variable concentrations and incubated at 37 °C for 24 h in the dark. Prior to light treatment, the cells were replaced with drug-free complete media. Cells were then illuminated with an argon-pumped dye laser set at 700 nm at a dose rate of 3.2 mW/cm<sup>2</sup> for 0–6 J/cm<sup>2</sup>. After PDT, the cells were incubated for 48 h at 37 °C in the dark. Following the 48 h incubation, 10  $\mu$ L of 5.0 mg/mL solution of 3-[4,5-dimethylthiazol-2-yl]-2,5-diphenyltetra-zoliumbromide (MTT) dissolved in PBS (Sigma, St. Louis, MO) was added to each well. After a 4 h incubation at 37 °C, the MTT and media were removed, and 100  $\mu$ L DMSO was added to solubilize the formazan crystals. The 96-well plate was read on a microtiter plate reader (Miles Inc. Titertek Multiscan Plus MK II) at an absorbance of 560 nm. The results were plotted as percent survival of the corresponding dark (drug no light) control for each compound tested, and each experiment was done with 4 replicate wells.

**In Vivo Photosensitizing Efficacy.** The in vivo PDT experiments were performed in C3H mice when tumors grew to 4–5 mm in diameter (approximately day 5 post inoculation). The day before laser light treatment, all hair was removed from the inoculation site, and the mice were injected intravenously with varying photosensitizer concentrations. At 24 h postinjection, the mice were restrained without anesthesia in Plexiglas holders and then treated with laser light from an argon-pumped dye laser tuned to emit drug-activating wavelengths as set by the monochromator (665 nm for **1**, 705 nm for **4**, **5**, and **10**). The

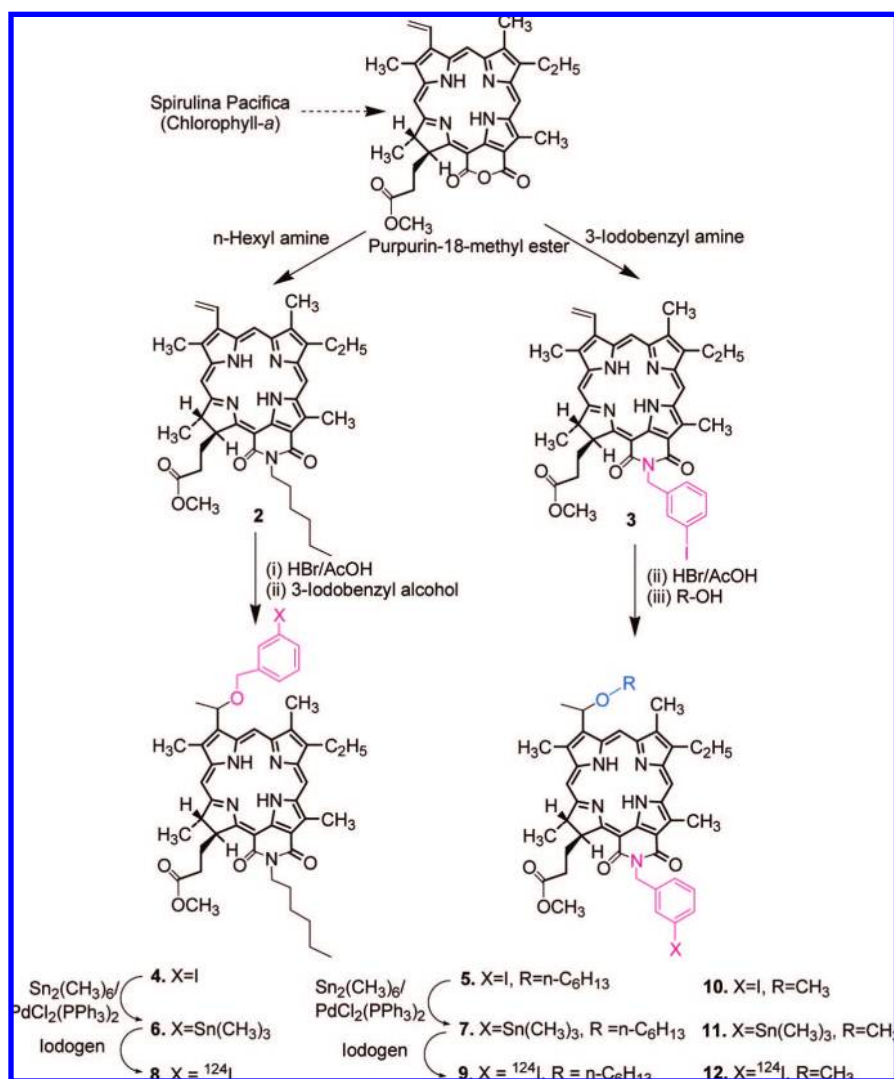
compounds were treated with light under similar treatment parameters under the fluence rate of 75 mW/cm<sup>2</sup> with a light dose of 135 J/cm<sup>2</sup>. The mice were observed daily for signs of weight loss, necrotic scabbing, or tumor regrowth. If tumor growth appeared, the tumors were measured using two orthogonal measurements *L* and *W* (perpendicular to *L*), and the volumes were calculated using the formula  $V = (L \times W^2)/2$  and recorded. Mice were considered cured if there was no sign of tumor regrowth by day 60 post-PDT treatment.

## RESULTS AND DISCUSSION

**Chemistry.** For the synthesis of desired compounds, methylpheophorbide-*a*, isolated from *Spirulina pacifica* was converted into purpurin-18 methyl ester by following the known methodology (44). For the preparation of 3-(1'-*m*-iodobenzoyloxyethyl) analogue **4**, it was first reacted with *n*-hexylamine, the intermediate amide derivative (isomeric mixture) so obtained on intramolecular cyclization under basic reaction conditions, and gave **2** in excellent yield. Further reaction of purpurinimide **2** with HBr/AcOH at room temperature produced the intermediate bromo-analogue, which was dried under vacuum and immediately reacted with *m*-iodobenzyl alcohol to afford **4** as a mixture of methyl ester and the corresponding carboxylic acid, which on treating with diazomethane produced the methyl ester derivative **4** in  $>70\%$  yield. For the synthesis of the related isomer **5**, the purpurin-18 methyl ester was first refluxed with *m*-iodobenzyl amine, and the intermediate **3** thus obtained on reacting with 1-hexanol by following the approach depicted in Scheme 1 gave the desired photosensitizer in modest yield. Compound **10** was synthesized from **3** following the methodology outlined for **5** and by replacing *n*-hexanol with methanol. For the preparation of the corresponding <sup>124</sup>I-analogues **8**, **9**, and **12**, the trimethylstannyl substituted analogues **6**, **7**, and **11** on electrophilic aromatic iodination with Na<sup>124</sup>I in the presence of iodogen beads afforded the <sup>124</sup>I-labeled purpurinimides **8**, **9**, and **12** with  $>95\%$  radioactive specificity. The purity of the final compounds (**4**, **5**, and **10**) was confirmed by NMR (experimental section) and HPLC analysis (see Supporting Information).

**Biological Studies. Comparative Imaging and Biodistribution of <sup>124</sup>I-Labeled Purpurinimide Isomers **8**, **9**, and **12**.** The PET imaging and biodistribution study of the radioactive purpurinimides **8** were performed in C3H mice bearing RIF tumors. In a typical experiment, 12 tumored mice were injected with each compound (50–200  $\mu$ Ci), and 3 mice/group were imaged at 24, 48, 72, and 96 h for 30 min with microPET (Siemens Preclinical Solutions, Knoxville, TN), and finally sacrificed after the 96 h time point. Mice used in longitudinal imaging received higher activity (150–200  $\mu$ Ci) compared to the mice used in biodistribution alone (50–100  $\mu$ Ci).

For the biodistribution studies, selected organs [tumor, muscle, kidney, lungs, intestine (gut), stomach, spleen, heart, and lung] were removed, weighed, and measured in a gamma well counter. The tail was also taken into consideration in biodistribution studies to determine the accuracy of the injection. Interestingly, the two isomers showed a remarkable difference in imaging and biodistribution characteristics. The imaging and biodistribution data also correlated well with each other. Between the two isomers, isomer **8** (*O*-iodobenzoyloxyethyl purpurinimide) had a higher background (liver and spleen), and the tumor was not visualized. Although the tumor was also not significantly visualized with isomer **9** (*N*-(3-iodobenzyl purpurinimide), compared to compound **8**, the background images were not as high (Figure 2). If compared with lead compound **1** (<sup>124</sup>I-labeled), purpurinimide **9** also exhibited higher tumor uptake at 72 and 96 h PI, and unfortunately, higher background uptake negated tumor visualization (Figure 3). However, in

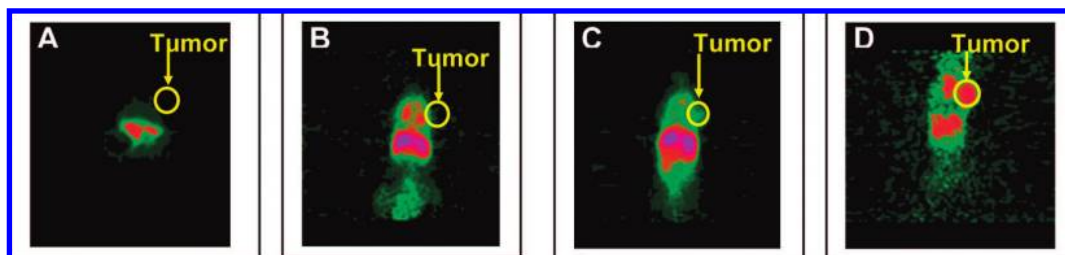
Scheme 1<sup>a</sup>

<sup>a</sup> Purpurinimides **4**, **5** (isomers), **10** and the corresponding <sup>124</sup>I-analogues **8**, **9**, and **12**, respectively.

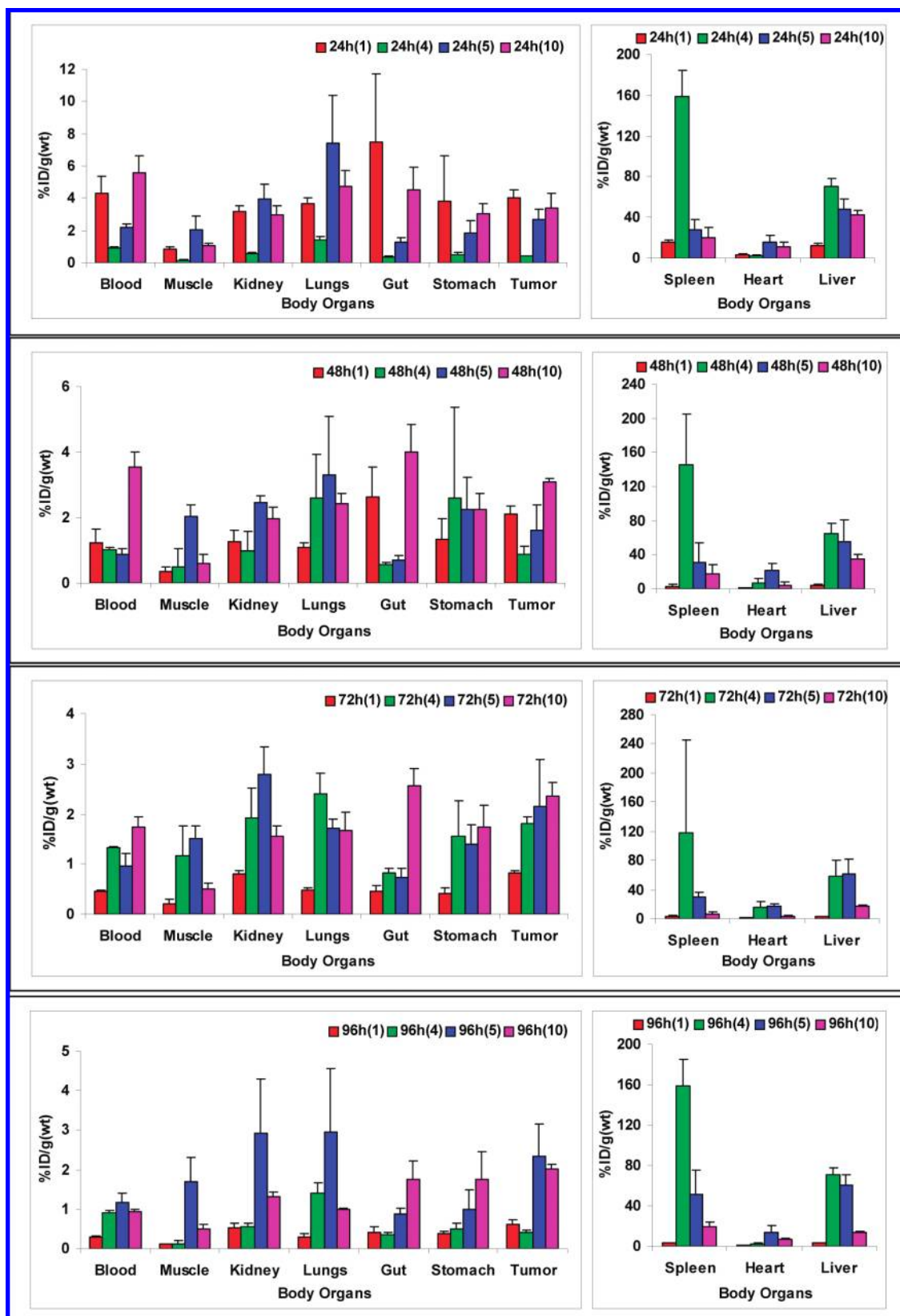
comparison with the <sup>124</sup>I-analogue of pyropheophorbide **1**, both purpurinimide isomers **8** and **9** showed exceptionally high liver (32-fold and 8-fold, respectively) and spleen uptake (85-fold and 3-fold, respectively) at 24 h PI. The significantly high uptake of purpurinimides **8** and **9** in the liver and spleen resulted in nonvisualization of the tumor and thus produced a poor contrast in whole body PET-imaging. Among isomers **8** and **9**, isomer **9** was selected for further modifications. The 1'-hexyloxyethyl group present at the top half of the molecule (position-3) was replaced with a methyl substituent. The resulting product **10**, with a reduced overall lipophilicity, was labeled with <sup>124</sup>I (compound **12**) and the PET imaging biodistribution data were performed in C3H mice bearing RIF tumors. The results

obtained from the biodistribution studies suggest that the reduction in the overall lipophilicity of the molecule substantially reduces the uptake of purpurinimide **12** in the spleen and liver and therefore enhances its tumor imaging capability at 96 h postinjection. The biodistribution and whole-body PET imaging results obtained from compounds **1**, **8**, **9**, and **12** suggest that for an efficient tumor imaging agent it is of utmost important to have a high uptake of the contrast agent in the tumor with a faster clearance profile from other organs. This characteristic possibly explains the improved imaging capability of lead compound **1** over that of the other agents studied so far.

*Comparative in Vitro PDT Efficacy.* The in vitro phototosensitizing ability of the structural isomers **4** and **5** was compared



**Figure 2.** Comparative microPET emission images (coronal view) of C3H mice with RIF tumors at 48 h PI of <sup>124</sup>I-124 labeled purpurinimides **8** (A), **9** (B), **12** (C), and the lead compound **1** (D).

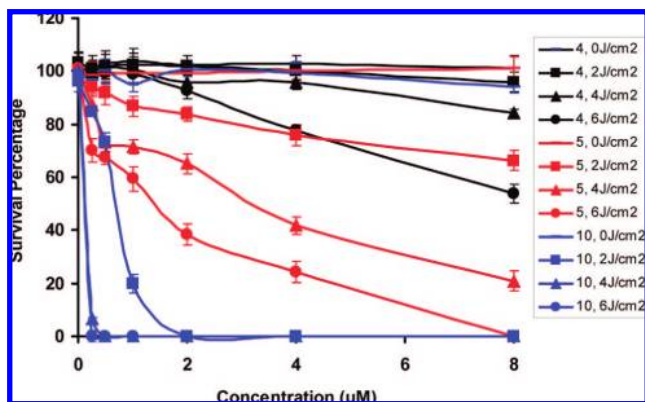


**Figure 3.** Comparative biodistribution of ( $^{124}\text{I}$ -labeled) pyropheophorbide-1 and the purpurinimides-8, 9, and 12 at 24, 48, 72, and 96 h PI in C3H mice bearing RIF tumors (4 mice/time point). Note: compounds 8, 9, and 12 are  $^{124}\text{I}$ -labeled 4, 5, and 10, respectively.

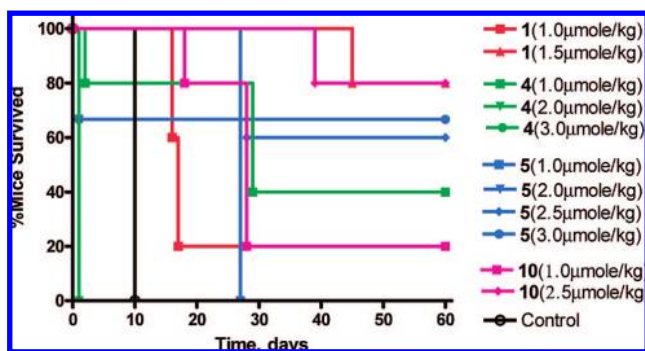
at variable experimental conditions (MTT assay; see Experimental Procedures) in RIF cells. Both isomers were ineffective in vitro at lower light and drug doses. However, as can be seen from Figure 4, at 24 h postincubation and higher light dose ( $6.0 \text{ J/cm}^2$ , drug concentration ( $1.0 \mu\text{M}$  and higher), isomer 5 was more effective than the structural isomer 4. Reducing the overall

lipophilicity of 5, by replacing the O-hexyl group at position-3 with an O-methyl group 10 produced enhanced efficacy over isomers 4 and 5.

*In Vivo PDT Efficiency of Pyropheophorbide-a 1 and Purpurinimide Isomers 4 and 5.* The in vivo PDT efficacy of isomers 4 and 5 was determined at three doses (1.0, 2.0, and



**Figure 4.** Comparative in vitro photosensitizing activity of **4**, **5**, and **10** at variable drug concentrations and light doses in RIF tumor cells at 24 h post incubation.



**Figure 5.** Kaplan–Meier plot for compounds **1**, **4**, **5**, and **10** at various doses. C3H mice bearing RIF tumor (5 mice/group) on the shoulder. Light dose: 135 J/cm<sup>2</sup>, 75 mW/cm<sup>2</sup> mice/group.

3.0  $\mu\text{mol/kg}$ ) and was compared with lead compound **1** in C3H mice bearing RIF tumors (5 mice/group). The tumors were exposed to light ( $\lambda_{\text{max}} = 705 \text{ nm}$  for **4** and **5**; 665 nm for **1**, 135 J/cm<sup>2</sup>, 75 mW/cm<sup>2</sup>) at 24 h postinjection. The tumor regrowth was measured daily. From the results summarized in Figure 5, it can be seen that compound **4**, having the iodobenzyl group at the top half of the molecule, at a dose of 1.0  $\mu\text{mol/kg}$  gave a good response (2/5 mice were tumor free on day 60). At the same dose, the isomer **5** in which the iodobenzyl group is present at the lower half of the molecule did not give any significant long-term tumor response. Under the same treatment parameters, compound **1** produced some tumor response; however, it was more effective at a dose of 1.5  $\mu\text{mol/kg}$ , and 4/5 mice were tumor free on day 60. Among isomers **4** and **5**, at higher doses (e.g., 2  $\mu\text{mol/kg}$ ) compound **4** was toxic, and all mice died within 24 h PDT treatment (Figure 5), whereas compound **5** did not show any significant activity. However, at higher doses (2.5 and 3.0  $\mu\text{mol/kg}$ ) it was found to be quite effective.

To investigate the effect of the overall lipophilicity in PDT efficacy, the hexyl ether group present at position-3 in **5** was replaced with a methyl ether substituent **10**, and the biological efficacy of both the analogues was compared at doses of 1.0 and 2.5 mmol/kg. The tumors were treated with a laser light ( $\lambda_{\text{max}} = 705 \text{ nm}$ , 135 J/cm<sup>2</sup>, 75 mW/cm<sup>2</sup>) at 24 h postinjection. From the results summarized in Figure 5, it can be seen that among the two analogues, methyl ether derivative **10** showed enhanced activity than the corresponding hexyl ether analogue **5**; however, it was at least 1.5-fold less effective over lead compound **1** derived from methyl pyropheophorbide-*a*. For an accurate reflection of the actual therapeutic response, further

studies with a larger group of mice under variable treatment parameters are currently in progress.

## CONCLUSIONS

Our results suggest that the nature and the position of the substituents in purpurinimides make a significant difference in tumor uptake, which also reflects their imaging and PDT potential. Between the two structural isomers **4** and **5**, compound **5** containing an N-iodobenzyl group introduced at the bottom half of the purpurinimide showed improved imaging and phototherapeutic abilities than **4** where the iodobenzyl group was present at position-3 of the molecule. Decreasing the overall lipophilicity of compound **5** by substituting the hexyl ether with a methyl ether group (compound **10**) further improved its PET imaging ability and PDT efficacy. However, for establishing a correlation between the overall lipophilicity and tumor imaging potential, it is necessary to investigate a series of compounds within a particular system, and these studies are currently in progress.

## ACKNOWLEDGMENT

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**Supporting Information Available:** The <sup>1</sup>H NMR spectra of compounds **4**–**7**, **10**, **11**, and the HPLC chromatograms of compounds **1**, **4**, **5**, and **10**. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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