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Subphthalocyanine-based porous organic polymers†

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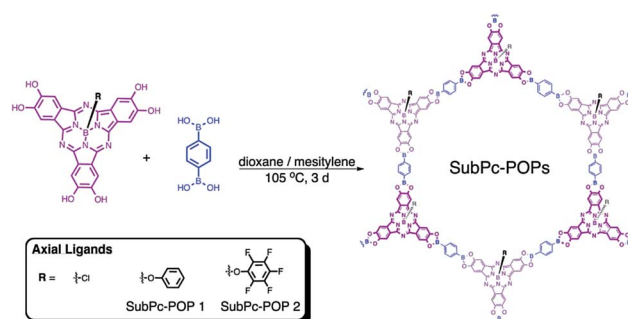
Subphthalocyanines (SubPcs) are bowl-shaped chromophores that possess optical properties desirable for organic electronics. Herein, we present the synthesis and characterization of two SubPc-based porous organic polymers (POPs). The SubPc-POPs exhibit respectable surface areas, and Q-bands that are significantly red-shifted in the solid-state.

Porous organic polymers (POPs) are a diverse group of materials that have garnered interest for their lightweight, highly porous architectures covering a wide range of applications including catalysis,^{1–3} sensing,^{4–7} and separations.^{8–11} Within the broader umbrella of POPs¹² exist many subgroups, including but not limited to porous aromatic frameworks (PAFs),¹³ porous polymer networks (PPNs),^{14–16} and conjugated microporous polymers (CMPs).¹⁷ The incorporation of π -conjugated monomers allows for tuning of the bulk properties of the polymer and increases their potential utility in devices. Inclusion of chromophores with extended π -systems, such as boron-dipyrromethenes (BODIPYs), porphyrins and phthalocyanines (Pcs), into porous polymers has allowed for the creation of materials amenable to optoelectronic applications, such as photocatalysis,^{18,19} organic light emitting diodes (OLEDs),^{20,21} and photovoltaic materials.^{22–24} Despite the incorporation of Pcs into linear polymers^{25–27} and two-dimensional (2D) porous polymers,^{28–36} no porous material to date has incorporated subphthalocyanine (SubPc) monomers.

SubPcs, which are smaller homologues of Pcs, contain three N-fused 1,3-diiminoisoindole units that are chelated to a boron atom with a protruding axial ligand. Our inspiration to pursue the C₃-symmetric bowl-shaped SubPc monomer evolved from its non-planar 14 π -electron conjugated structure which could provide access to unique porous polymers with small Stokes shifts and low reorganizational energies. These features have enabled SubPcs to achieve respectable power conversion efficiencies as both donor and acceptor materials in small molecule-based organic photovoltaic devices (OPVs).^{37–40} However, 2D polymeric materials based on SubPcs are essentially unknown, with the exception of their inclusion into self-assembled SubPc films.^{41–43} While these thin layer 2D systems could be useful for information storage or liquid crystalline technologies, only slow progress has been made in this

direction. Polymers successfully incorporating SubPc monomers are rare and only a few examples of styrene copolymers with pendant SubPc groups have been reported.^{44,45} However, this post polymerization strategy was problematic, and did not afford a high incorporation of SubPc into the polymeric material. It also lacks the added functionality of porosity as a way to interact with guest molecules. Our strategy opted for a mild co-condensation reaction to form a boronate ester linkage, which can be constructed without an external catalyst at moderate temperatures, while simultaneously incorporating the SubPc monomeric units into a covalently linked polymeric network. Such nonplanar polymeric systems incorporating SubPcs have yet to be reported.

Herein, we present the synthesis and characterization of SubPc-POP 1 and 2 (Scheme 1). The POPs were constructed by reacting SubPcs **2b** and **3b** (Fig. 1) with 1,4-benzene diboronic acid (BDBA) in a 1 : 1 (v/v) mixture of dioxane and mesitylene to yield dark purple solids. These optimal reaction conditions were obtained by thoroughly screening solvent ratios and reaction temperatures (Table 1, ESI†). It should be noted that our initial polymer design aimed to utilize **1b** to form a 2D POP with axial chloride ligands (Fig. 1). However, compound **1b** and the TIPS-protected **1a** were highly sensitive to acidic or basic environments and ambient light, which made them difficult to isolate in large quantities. We believe that this rapid



Scheme 1 Synthesis of the SubPc-POPs.

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order (Fig. S9 & S10, ESI†). Initially, we anticipated that a SubPc-POP substituted with chlorine atoms in the axial position would exhibit a concave-to-ligand solid-state packing motif similar to the *trans* binuclear SubPc systems previously reported by Kobayashi and Durfee.⁴⁷ However, these polymeric systems could not be constructed due to the chemical instability of **1b**. As a consequence, we believe the larger axial phenoxy and pentafluorophenoxy substituents of **2b** and **3b** prevent the monomers from forming the concave-to-ligand stacking interactions, and instead lead to the formation of disordered POPs.

The 14- π electron core of SubPcs is capable of absorbing and emitting radiation in the visible spectrum. Normalized UV-Vis and fluorescence spectra of dilute solutions (2×10^{-5} M) of **1b**, **2b**, and **3b** in acetone are shown in Fig. 1b. The absorption maxima range from 566 to 574 nm and are attributed to the typical Q-band absorption peaks for SubPcs. Each absorption spectrum contains distinct shoulder peaks ranging from 547 to 553 nm. The weak high energy absorption bands at ~ 368 nm are possibly attributed to an $n-\pi^*$ transition between the lone pair of the oxygen atoms and the peripheral aromatic ring.⁴⁸ The fluorescence spectra of **1b**, **2b** and **3b** revealed emission peaks at λ_{max} 587, 581 and 580 nm, respectively. In addition, broad shoulders ranging from 616 to 624 nm, which reflected their corresponding absorption spectra, accompanied each emission spectrum.

In contrast to the monomers in solution, SubPc-POPs **1** and **2** did not exhibit any unique fluorescent properties, and were fully quenched in the solid-state. We believe that the disordered packing of the material leads to aggregation-caused quenching (ACQ) due to the rapid thermal decay of the photoexcited state.⁴⁹ However, diffuse reflectance spectra of SubPc-POP **1** and **2** revealed that both materials display peak absorption bands at ~ 583 and 581 nm, respectively, along with weak bands that extend into the near infrared part of the spectrum (Fig. 4). It should be noted that these absorption maxima are considerably red-shifted by ~ 50 – 60 nm compared to the powders of **2b** ($\lambda_{\text{max}} = 530$ nm) and **3b** ($\lambda_{\text{max}} = 520$ nm) (Fig. S24 & S25, ESI†), which suggest the formation of disordered J-aggregates in the solid-state.

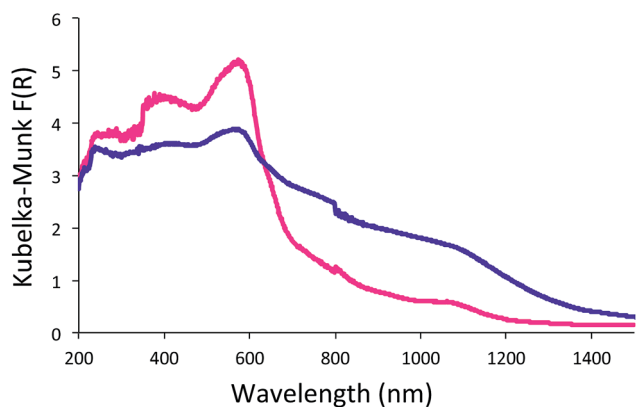


Fig. 4 Diffuse reflectance of SubPc-POP **1** (pink) and SubPc-POP **2** (purple).

Conclusions

In conclusion, we have demonstrated that hexahydroxyl-substituted SubPc monomers can be used to construct nonplanar POPs with respectable pore sizes and surface areas. The diffuse reflectance spectra show that the absorption properties of the SubPc-POPs are significantly red-shifted compared to the monomeric SubPc units. Future studies could involve utilizing axial-substitution to tune the bulk properties of the material in the solid-state.^{50,51} Such investigations could yield functional nonplanar SubPc-based porous polymers with unique optoelectronic properties.

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