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Michael J. Therien *et al.*
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Driving high quantum yield NIR emission through proquinoidal linkage motifs in conjugated supermolecular arrays†

Erin J. Peterson, Wei Qi, Ian N. Stanton, Peng Zhang and Michael J. Therien*

High quantum yield NIR fluorophores are rare. Factors that drive low emission quantum yields at long wavelength include the facts that radiative rate constants increase proportional to the cube of the emission energy, while nonradiative rate constants increase in an approximately exponentially with decreasing S_0-S_1 energy gaps (in accordance with the energy gap law). This work demonstrates how the proquinoidal BTD building blocks can be utilized to minimize the extent of excited-state structural relaxation relative to the ground-state conformation in highly conjugated porphyrin oligomers, and shows that 4-ethynylbenzo[*c*][1,2,5]thiadiazole (E-BTD) units that terminate *meso*-to-*meso* ethyne-bridged (porphyrinato)zinc (PZn_n) arrays, and 4,7-diethynylbenzo[*c*][1,2,5]thiadiazole (E-BTD-E) spacers that are integrated into the backbone of these compositions, elucidate new classes of impressive NIR fluorophores. We report the syntheses, electronic structural properties, and emissive characteristics of neoteric PZn -(BTD- PZn) $_n$, PZn_2 -(BTD- PZn_2) $_n$, and BTD- PZn_n -BTD fluorophores. Absolute fluorescence quantum yield (ϕ_f) measurements, acquired using a calibrated integrating-sphere-based measurement system, demonstrate that these supermolecules display extraordinary ϕ_f values that range from 10–25% in THF solvent, and between 28–36% in toluene solvent over the 700–900 nm window of the NIR. These studies underscore how the regulation of proquinoidal conjugation motifs can be exploited to drive excited-state dynamical properties important for high quantum yield long-wavelength fluorescence emission.

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Introduction

Low band gap π -conjugated molecules and oligomers are central to function in electro-optic applications that span excitonic solar cells,^{1–7} field-effect transistors,^{8,9} optical power limiting (OPL),¹⁰ dye-sensitized solar cells (DSSCs),^{11,12} photon-upconversion (UC) technologies,^{13–17} long-wavelength light-emitting diodes,^{18–20} nonlinear optics (NLO),^{21–24} and biological imaging.^{25–27} Desirable low band gap materials for these applications typically feature singlet manifold transitions ($S_0 \rightarrow S_1$, $S_0 \rightarrow S_n$, and $S_1 \rightarrow S_n$) that possess large absorptive oscillator strengths covering broad spectral domains that include the near-infrared (NIR).

Driving high fluorescence quantum yields is perhaps the most demanding design challenge for low band gap chromophores. Engineering augmented $S_0 \rightarrow S_1$ transition oscillator strength is not sufficient to realize correspondingly high $S_1 \rightarrow S_0$ fluorescence quantum yields (ϕ_f values), particularly as the

optical band gap is diminished: while it is often the case that a large $S_0 \rightarrow S_1$ absorptive extinction coefficient is correlated with a correspondingly large $S_1 \rightarrow S_0$ radiative rate constant congruent with the Strickler–Berg relationship,²⁸ most strongly



Chart 1 Chemical structures for (A) benchmark PZn_n , (B) BTD- PZn_n -BTD and (C) PZn_2 -(BTD- PZn_2) $_n$ and PZn -(BTD- PZn) $_n$ fluorophores. R = [2,6-bis(3,3-dimethyl-1-butoxy)phenyl].

Department of Chemistry, French Family Science Center, Duke University, 124 Science Drive, Durham, North Carolina 27708-0346, USA. E-mail: michael.therien@duke.edu

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absorbing NIR chromophores are not impressive emitters, due to the substantial S_1 state non-radiative rate constants (k_{nr} values) that ensue from the energy gap law.²⁹

With respect to high oscillator strength NIR absorbers, *meso-to-meso* ethyne-bridged (porphinato)zinc (**PZn_n**) arrays (Chart 1A)^{17,30–46} constitute a notable exception to these generalizations. These supermolecules – conjugated structures realized from strong coupling of simple chromophoric building blocks – manifest such strong electronic and excitonic interactions between the constituent (porphinato)zinc oscillators that entirely unique photophysical entities are defined. **PZn_n** structures, for example, possess fluorescence quantum yields comparable to the most impressive NIR laser dyes in the 750–900 nm regime,⁴¹ and rival the highest reported for organic dyes in this spectral window;^{47–49} importantly, **PZn_n** emitters do not suffer from the commonly cited drawbacks of tricarbo-cyanine dyes that include poor photostability and fluorescence quantum yields that drop precipitously with decreasing solvent polarity.^{25,50,51} **PZn_n** chromophores feature low energy Q-state derived π – π^* excited-states that are polarized exclusively along the long molecular axis, and lowest energy transitions that gain in intensity and progressively red-shift with increasing numbers of PZn units. Extended conjugation in these highly polarizable chromophores drives red-shifting of **PZn_n** fluorescence spectra. Because triplet excitons are more spatially confined than singlet excitons in these systems,^{31,34,35,39,52,53} an unusual dependence of ϕ_f magnitude upon increasing emission wavelength [$\lambda_{max}(S_1 \rightarrow S_0)$] is manifest that confounds that anticipated simply from the energy gap law.⁴¹ Because of this disparity in lowest energy singlet (S_1) and triplet (T_1) state delocalization, diminished S_1 – T_1 intersystem crossing rate constants (k_{isc} values) track with increasing conjugation in

PZn_n chromophores, and serve to partly counter-balance the impact of augmented Franck–Condon mediated internal conversion rate constants (k_{ic} values) that typically accompany diminished S_0 – S_1 energy gaps in the NIR. These impressive **PZn_n** photophysical properties have been exploited in a wide range of fluorescence imaging applications.^{54–61}

The photophysical properties of **PZn_n** chromophores may be extensively modulated *via* incorporation of proquinoidal units into these emblematic conjugated arrays. For example, proquinoidal spacer (Sp) moieties such as 4,7-diethynylbenzo [c][1,2,5]thiadiazole (**E-BTD-E**), 6,13-diethynylpentacene (**E-PC-E**), 4,9-diethynyl-6,7-dimethyl[1,2,5]thiadiazolo[3,4-*g*]quinoxaline (**E-TDQ-E**), and 4,8-diethynylbenzo[1,2-*c*:4,5-*c'*]bis[1,2,5]thiadiazole (**E-BBTD-E**) that link bis[(porphinato)zinc] units can be utilized to modulate the relative degrees of quinoidal character that characterize the ground and electronically excited singlet states in these highly conjugated supermolecules.⁶² Recent work highlights how electronic modulation of proquinoidal conjugation motifs can also be exploited as a powerful means to modulate independently the dynamics of excited-state relaxation pathways in these systems, enabling, for example, exceptional NIR absorbers that possess long-lived electronically excited triplet states ($\tau_{T_1} > \mu s$) that are generated at unit quantum yield.⁶³ In this contribution, we (i) show how proquinoidal BTD building blocks can be utilized to minimize the extent of excited-state structural relaxation relative to the ground-state conformation in highly conjugated porphyrin oligomers, and (ii) demonstrate that 4-ethynylbenzo[c][1,2,5]thiadiazole (**E-BTD**) units that terminate **PZn_n** chromophores (Chart 1B), and **E-BTD-E** spacers that are integrated into the backbone of these compositions (Chart 1C), enable elucidation of new classes of impressive NIR fluorophores.



Scheme 1 Synthetic routes for **PZn-(BTD-PZn)_n** compounds.



Scheme 2 Synthetic routes for BTD-PZn_n-BTD compounds.

Results and discussion

Synthesis

Schemes 1–3 outline the synthetic strategies for the fabrication of BTD-PZn_n-BTD, PZn-(BTD-PZn)_n, and PZn₂-(BTD-PZn₂)_n chromophores. These structures were synthesized by palladium (Pd)-mediated cross-coupling reactions involving appropriately substituted (porphinato)zinc(II) (PZn) compounds, BTD spacers,

and terminal BTD units. These structures exploit [2,6-bis(3,3-dimethyl-1-butoxy)phenyl] groups as 10- and 20-*meso*-porphyrin substituents, which facilitate solubility and straightforward assignment of ¹H-NMR spectra.^{37,64} The nature of the functionalized PZn, PZn_n, and BTD moieties used in the syntheses of these BTD-PZn_n-BTD, PZn-(BTD-PZn)_n, and PZn₂-(BTD-PZn₂)_n arrays varied with the degree of conjugation in the precursor molecules. Synthetic details may be found in the ESI.†

Scheme 3 Synthetic routes for PZn₂-(BTD-PZn₂)_n compounds.

Steady-state absorption and emission spectroscopy of $\text{BTD-PZn}_n\text{-BTD}$, PZn-(BTD-PZn)_n , and $\text{PZn}_2\text{-(BTD-PZn}_2)_n$ chromophores

Steady-state electronic absorption and fluorescence spectra recorded for reference PZn_n chromophores (PZn_2 , PZn_3 , and PZn_5 ; Fig. 1A),^{17,30–46,54–56,58,59,61,63,65} as well as PZn-(BTD-PZn)_n [PZn-BTD-PZn ,⁶² PZn-(BTD-PZn)_2 , and PZn-(BTD-PZn)_4 ; Fig. 1B], $\text{PZn}_2\text{-(BTD-PZn}_2)_n$ [$\text{PZn}_2\text{-BTD-PZn}_2$ and $\text{PZn}_2\text{-(BTD-PZn}_2)_2$; Fig. 1C], and $\text{BTD-PZn}_n\text{-BTD}$ [BTD-PZn-BTD ,⁶⁶ $\text{BTD-PZn}_2\text{-BTD}$, $\text{BTD-PZn}_3\text{-BTD}$, and $\text{BTD-PZn}_5\text{-BTD}$; Fig. 1D] arrays, are shown in Fig. 1. The overall characteristics of the electronic absorption spectra for these PZn-(BTD-PZn)_n , $\text{PZn}_2\text{-(BTD-PZn}_2)_n$, and $\text{BTD-PZn}_n\text{-BTD}$ chromophores resemble those described previously for dimeric and multimeric PZn compounds that feature a *meso-to-meso* ethyne-linkage topology (PZn_n arrays); these spectra exhibit two distinct intense absorption manifolds that are derived from the porphyrin B- (Soret, $S_0 \rightarrow S_2$) and Q-band ($S_0 \rightarrow S_1$) transitions.^{30,32–34,37,38,40}

The observed perturbations from these benchmark PZn_n spectra trace their origin to the proquinoidal BTD units that are connected to the porphyrin macrocycle *meso* carbons *via* ethynyl moieties. Previous investigations demonstrate that the nature of proquinoidal unit conjugation to the PZn macrocycle exerts a pronounced impact on the magnitude of B- and Q-state mixing,^{62,63,66} for example, for $\text{PZn-(proquinoidal Sp)-PZn}$ chromophores, the long axis-polarized Q state (Q_x) absorption maxima can be modulated from 689 to 1006 nm, depending upon the extent of the quinoidal resonance contribution to the electronically excited singlet state.⁶² A combination of semi-empirical electronic structure calculations and electrochemical data underscore the cardinal role that PZn and proquinoidal fragment orbital energy differences play in fixing the radical cation and anion state energy levels in these structures.^{62,63}

The principles that informed design of these PZn-(BTD-PZn)_n , $\text{PZn}_2\text{-(BTD-PZn}_2)_n$, and $\text{BTD-PZn}_n\text{-BTD}$ chromophores stem from prior work that examined the photophysics of (porphyrinato)metal-(proquinoidal Sp)-(porphyrinato)metal supermolecules.⁶³ This work highlighted the central importance of the magnitude of the energy separation between the (porphyrinato)metal and proquinoidal Sp fragment frontier orbitals in determining whether radiative, internal conversion, or inter-system crossing decay channels dominated the relaxation dynamics of the initially prepared electronically excited states of these complexes. When the (porphyrinato)metal and proquinoidal Sp fragment LUMO levels featured energy separations on the order of a few tenths of an eV, as they are for the 5-ethynyl- PZn , 4-ethynyl- BTD , and 4,7-diethynyl- BTD fragments of these PZn-(BTD-PZn)_n , $\text{PZn}_2\text{-(BTD-PZn}_2)_n$, and $\text{BTD-PZn}_n\text{-BTD}$ structures, chromophores having multiconfigurational S_1 states characterized by a modest degree of charge transfer (CT) character are anticipated; such supermolecules were shown to display large $S_1 \rightarrow S_0$ radiative rate constants and substantial fluorescence quantum yields. Other factors that informed the blueprint of the Fig. 1B–D supermolecules included insights derived from monomeric PZn complexes in which ethynyl- BTD

units were used to expand conjugation at the macrocycle *meso*-carbon position;⁶⁶ these designs led to PZn chromophores characterized by enhanced transfer of B-to-x-polarized Q-state oscillator strength, intensified Q_x absorption bands, chromophore structural rigidification, spectrally narrow fluorescence emission bands, small magnitude Stokes shifts, and enhanced radiative rate constant magnitudes relative to simple benchmark PZn complexes.

Fluorescence emission metrics of $\text{BTD-PZn}_n\text{-BTD}$, PZn-(BTD-PZn)_n , and $\text{PZn}_2\text{-(BTD-PZn}_2)_n$ chromophores

Long wavelength absorption maxima (λ_{max} , $S_0 \rightarrow S_1$), corresponding extinction coefficient measurements, fluorescence ($S_1 \rightarrow S_0$) emission maxima, full-width at fluorescence half-

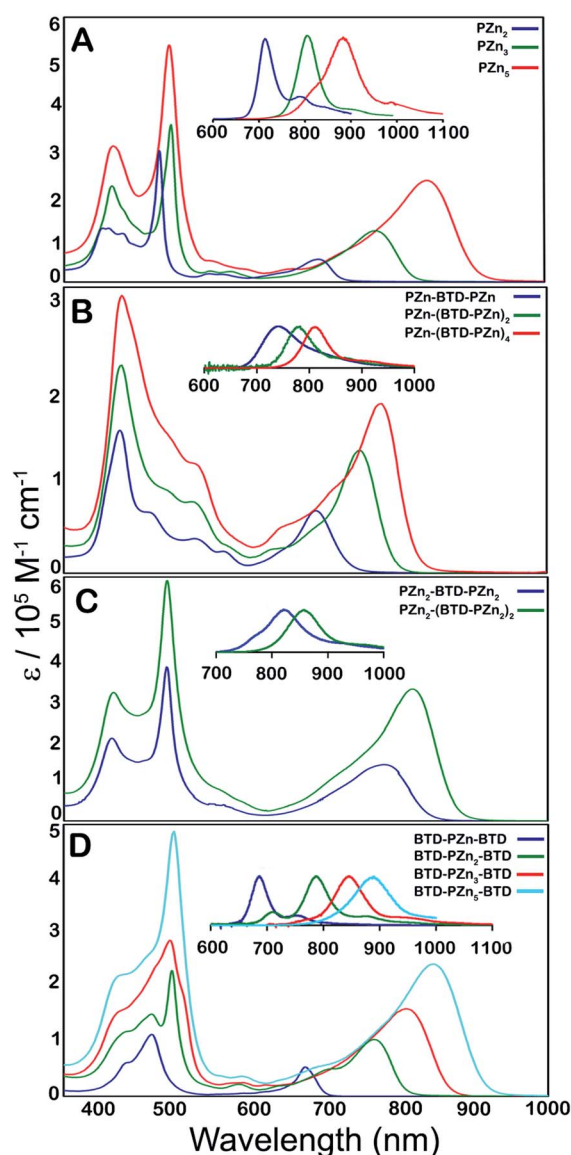


Fig. 1 Electronic absorption and emission spectra (figure panel insets) recorded for: (A) benchmark PZn_n chromophores (see ref. 30 and 41), and (B) PZn-(BTD-PZn)_n , (C) $\text{PZn}_2\text{-(BTD-PZn}_2)_n$, and (D) $\text{BTD-PZn}_n\text{-BTD}$ compositions. Experimental conditions: $T = 298$ K, THF solvent.



Table 1 Electronic absorption and emission data in THF solvent for **BTD-PZn_n-BTD**, **PZn-(BTD-PZn)_n**, and **PZn₂-(BTD-PZn₂)_n** chromophores relative to corresponding **PZn_n** benchmarks

	λ_{\max} (S ₀ → S ₁) ^a [nm]	$\epsilon_{\text{g}}@ \lambda_{\max}$ (S ₀ → S ₁) [M ⁻¹ cm ⁻¹]	λ_{\max} (S ₁ → S ₀) [nm]	FWHM (S ₁ → S ₀) [cm ⁻¹]	Stokes shift ^b (cm ⁻¹)	$\phi_{\text{f}}^{\text{c,d}}$	$\tau_{\text{F}}^{\text{e}}$ [ns]	k_{f}^{f} [×10 ⁸ s ⁻¹]	k_{nr}^{f} [×10 ⁸ s ⁻¹]
PZn-BTD-PZn	689 (1194)	64 600	741	1605	1018	0.25 (0.31)	1.6	1.6	4.7
PZn-(BTD-PZn)₂	745 (1178)	132 700	784	977	543	0.22 (0.36)	1.1	2.0	7.1
PZn-(BTD-PZn)₄	777 (1558)	184 000	811	846	490	0.16 (0.29)	0.8	2.0	11
BTD-PZn-BTD	674 (672)	84 000	687	727	281	0.13 (0.13)	1.3	1.0	6.7
BTD-PZn₂-BTD	765 (1243)	101 600	787	802	365	0.23 (0.30)	1.4	1.6	5.5
BTD-PZn₃-BTD	809 (1663)	157 500	846	872	510	0.16 (0.29)	0.8	2.0	11
BTD-PZn₅-BTD	847 (1675)	238 300	888	1008	587	0.10 (0.28)	0.6	1.7	15
PZn₂-BTD-PZn₂	780 (1822)	145 400	822	926	688	0.14 (0.34)	0.6	2.3	14
PZn₂-(BTD-PZn₂)₂	816 (1582)	340 000	857	1154	586	0.12 (0.36)	0.5	2.4	18
PZn₂	695 (1082)	51 400	711	810	324	0.14 (0.16)	1.09	1.3	7.9
PZn₃	770 (1386)	116 000	806	875	580	0.19 (0.27)	1.13	1.7	7.1
PZn₅	842 (1562)	230 000	883	955	551	0.09 (0.11)	0.45	2.0	20

^a Numbers in parentheses are spectral breadths (full-widths at half-maximum, FWHM) of the respective transitions in units of cm⁻¹. ^b Stokes shifts values correspond to the difference in energy between the low energy (Q_x) absorption (S₀ → S₁) and fluorescence (S₁ → S₀) band maxima.

^c Fluorescence quantum yields (ϕ_{f} values) were determined using an integrating sphere-based absolute emission quantum yield measurement system (see ESI†). ^d Values in parentheses represent those determined in toluene solvent. ^e These values were determined using S₀ → S₂ excitation (483 nm). Fluorescence lifetimes were measured *via* time-correlated single-photon-counting using a picosecond fluorescence lifetime measurement system. ^f Excited-state relaxation constants were calculated based on the following equations: $\tau_{\text{S1}} = 1/(k_{\text{r}} + k_{\text{nr}})$, $\phi_{\text{f}} = k_{\text{r}} \times \tau_{\text{S1}}$.

maximum (FWHM), Stokes shifts, fluorescence lifetimes (τ_{F}), radiative rate constant (k_{r}), non-radiative rate constant (k_{nr}), and fluorescence quantum yield (ϕ_{f}) data, are tabulated in Table 1 for the **PZn_n**, **PZn-(BTD-PZn)_n**, **PZn₂-(BTD-PZn₂)_n**, and **BTD-PZn_n-BTD** supermolecules.

These fluorescence quantum yield data correspond to absolute measurements, acquired using a calibrated integrating-sphere-based measurement system, which collects photons emitted from the sample following excitation through NIR

wavelengths that range up to 1100 nm (see ESI† for a complete system description). Note in this regard that minor differences between the Table 1 ϕ_{f} data acquired for **PZn_n** chromophores and those reported originally⁴¹ stem from the fact that these earlier ϕ_{f} values were determined *via* the reference method. Table 2 displays comparative integrated oscillator strengths of the B- and Q-band spectral regions of **BTD-PZn_n-BTD**, **PZn-(BTD-PZn)_n**, and **PZn₂-(BTD-PZn₂)_n** chromophores relative to corresponding **PZn_n** benchmarks.

Table 2 Comparative integrated oscillator strengths of the B- and Q-band spectral regions of **BTD-PZn_n-BTD**, **PZn-(BTD-PZn)_n**, and **PZn₂-(BTD-PZn₂)_n** chromophores relative to corresponding **PZn_n** benchmarks^a

	Oscillator strength B-band region ^b	Oscillator strength Q-band region ^c	Total oscillator strength
PZn-BTD-PZn	2.11	0.72	3.52
PZn-(BTD-PZn)₂	4.65	1.03	5.67
PZn-(BTD-PZn)₄	5.98	1.58	7.57
BTD-PZn-BTD	2.24	0.37	2.63
BTD-PZn₂-BTD	3.54	0.89	4.42
BTD-PZn₃-BTD	5.23	1.53	6.76
BTD-PZn₅-BTD	7.64	2.10	9.74
PZn₂-BTD-PZn₂	6.04	1.30	7.35
PZn₂-(BTD-PZn₂)₂	10.49	2.84	13.32
PZn₂^d	2.134	0.303	2.438
PZn₃^d	3.240	0.716	3.956
PZn₅^d	5.986	1.622	7.608

^a Integrated oscillator strengths (f) were calculated based on the following expression: $f = 4.3 \times 10^{-9} \int \epsilon d\nu$, where ϵ is the experimental extinction coefficient, and ν is the energy (in wave numbers) of the absorption. Values noted derive from electronic absorption spectra recorded in THF solvent.

^b Oscillator strengths calculated over the following wavelength domains: **PZn-BTD-PZn** (~360 to 600 nm); **PZn-(BTD-PZn)₂** (~360 to 600 nm); **PZn-(BTD-PZn)₄** (~360 to 605 nm); **BTD-PZn-BTD** (~360 to 560 nm); **BTD-PZn₂-BTD** (~360 to 560 nm); **BTD-PZn₃-BTD** (~360 to 560 nm); **BTD-PZn₅-BTD** (~360 to 560 nm); **PZn₂-BTD-PZn₂** (~360 to 610 nm); **PZn₂-(BTD-PZn₂)₂** (~360 to 610 nm). ^c Oscillator strengths calculated over the following wavelength domains: **PZn-BTD-PZn** (~600 to 760 nm); **PZn-(BTD-PZn)₂** (~600 to 820 nm); **PZn-(BTD-PZn)₄** (~600 to 900 nm); **BTD-PZn-BTD** (~560 to 720 nm); **BTD-PZn₂-BTD** (~560 to 850 nm); **BTD-PZn₃-BTD** (~560 to 910 nm); **BTD-PZn₅-BTD** (~560 to 1050 nm); **PZn₂-BTD-PZn₂** (~610 to 860 nm); **PZn₂-(BTD-PZn₂)₂** (~610 to 920 nm). ^d Ref. 41.



Table 1 highlights that **E-BTD** units that terminate **PZn_n** chromophores (Fig. 1A) give rise to **BTD-PZn_n-BTD** supermolecules (Fig. 1D) in which $\lambda_{\max}(S_1 \rightarrow S_0)$ for **BTD-PZn₂-BTD**, **BTD-PZn₃-BTD**, and **BTD-PZn₅-BTD** redshift by 1358, 587, and 64 cm^{-1} , respectively, relative to their **PZn₂**, **PZn₃**, and **PZn₅** benchmarks. Note that as a function of the number of PZn units in these arrays, **BTD-PZn_n-BTD** and **PZn_n** supermolecules display similarly narrow fluorescence FWHM values as well as modest Stokes shifts; these energy differences between the low energy (Q_x) absorption ($S_0 \rightarrow S_1$) and fluorescence ($S_1 \rightarrow S_0$) band maxima of these chromophores range from 325–585 cm^{-1} . A key spectroscopic ramification of terminal ethynyl-BTD conjugation to these **PZn_n** frameworks is highlighted in the data chronicled in Table 2, which contrasts the integrated oscillator strengths of the B- and Q-band regions of the electronic absorption spectra of these chromophores. Note that the integrated Q-state oscillator strengths of these **BTD-PZn_n-BTD** chromophores are augmented by more than 50% relative their corresponding **PZn_n** benchmark. These factors, coupled with a diminished nonradiative rate constant, play important roles in driving the substantial, long-wavelength emission quantum yield manifest by **BTD-PZn₅-BTD** ($\phi_f = 0.28$) relative to that for the parent **PZn₅** chromophore ($\phi_f = 0.11$) in low dielectric toluene solvent (Table 1).

PZn-(BTD-PZn)_n chromophores (Fig. 1B) contrast the electronic spectral properties of **BTD-PZn_n-BTD** supermolecules; **PZn-BTD-PZn**, **PZn-(BTD-PZn)₂**, and **PZn-(BTD-PZn)₄** display Q_x state absorption maxima that are blue-shifted 126, 436, and 994 cm^{-1} with respect to the $\lambda_{\max}(S_0 \rightarrow S_1)$ transitions of their respective **PZn₂**, **PZn₃**, and **PZn₅** benchmarks (Table 1, Fig. 1); this effect derives from the fact that 4,7-diethynylbenzo[*c*][1,2,5]thiadiazole provides diminished PZn–PZn electronic coupling relative to the ethynyl linker. While **PZn-BTD-PZn** displays an augmented Stokes shift (1018 cm^{-1}) relative to **PZn₂** (324 cm^{-1}) due to the greater cumulenlic character in its relaxed electronically excited S_1 state,⁶² the Stokes shifts manifest for **PZn-(BTD-PZn)₂**, **PZn-(BTD-PZn)₄**, **PZn₃**, and **PZn₅** are similar in magnitude ($\sim 560 \text{ cm}^{-1}$), congruent with more modest structural differences between the relaxed S_0 and S_1 states for these supermolecules (*vide infra*).

PZn-BTD-PZn, **PZn-(BTD-PZn)₂**, and **PZn-(BTD-PZn)₄** display fluorescence emission maxima centered at 741, 784, and 811 nm, respectively, and substantial fluorescence quantum yields in THF solvent [$\phi_f(\text{PZn-BTD-PZn}) = 0.25$; $\phi_f(\text{PZn-(BTD-PZn)}_2) = 0.22$; $\phi_f(\text{PZn-(BTD-PZn)}_4) = 0.16$]. Similar to that observed for **BTD-PZn_n-BTD** chromophores, fluorescence quantum yields are significantly enhanced in nonpolar solvent; note in this regard that $\phi_f(\text{PZn-(BTD-PZn)}_4)$ is amplified to 29% in toluene (Table 1).

Fig. 1C highlights the absorptive and emissive spectral properties that ensue when **PZn₂** chromophores are linked by 4,7-diethynylbenzo[*c*][1,2,5]thiadiazole units. These **PZn₂-(BTD-PZn)_n** chromophores display dramatic Q_x absorption band intensification with increasing conjugation (Tables 1 and 2); note in this regard that **PZn₂-(BTD-PZn)_n** is an exceptional long-wavelength absorber ($\epsilon(816 \text{ nm}) = 340\,000 \text{ M}^{-1} \text{ cm}^{-1}$; Table 1). **PZn₂-BTD-PZn₂** and **PZn₂-(BTD-PZn)₂** emit

respectively at 822 and 857 nm in THF solvent, with corresponding quantum yields of 14 and 12%. As demonstrated for both **PZn-(BTD-PZn)_n** and **BTD-PZn_n-BTD** chromophores, these fluorescence quantum yields are dramatically enhanced in toluene solvent [$\phi_f(\text{PZn₂-BTD-PZn₂)} = 0.34$; $\phi_f(\text{PZn₂-(BTD-PZn₂)}_2) = 0.36$]. These data acquired for **PZn₂-(BTD-PZn)_n** designs suggest additional approaches to realize related frameworks that make possible high quantum yield NIR emission that include electronic modulation of the proquinoidal units that both terminate supermolecules that utilize **PZn_n** building blocks, as well as those that are integrated into the conjugated backbones of these compositions.

Computed electronic structures of BTD-PZn_n-BTD, PZn-(BTD-PZn)_n, and PZn₂-(BTD-PZn)_n chromophores

The natures of the low energy S_1 states of **BTD-PZn_n-BTD**, **PZn-(BTD-PZn)_n**, and **PZn₂-(BTD-PZn)_n** chromophores were probed through frontier orbital (FO) population and transition matrix

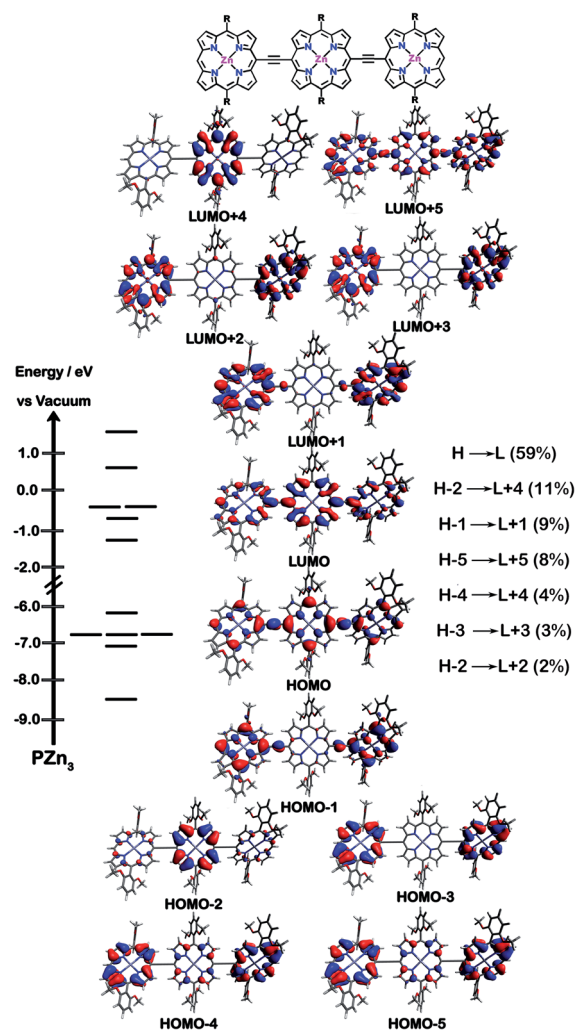


Fig. 2 Calculated frontier molecular orbitals, energy levels, and relative one-electron configurations that contribute to the lowest energy (Q_x) transition for the **PZn₃** benchmark. Calculations were performed at the M11/6-311g(d) theory level.



eigenvector analyses derived using TD-DFT methods. Fig. 2–4 show FO diagrams for **BTD-PZn₃-BTD** and **PZn-(BTD-PZn)₂**, along with those for the **PZn₃** benchmark, and highlight the prominent one-electron configurations that contribute to their respective lowest energy Q_x transitions; related data for the **PZn₂-(BTD-PZn)₂** chromophore is presented in the ESI.†

The natures and energy separations between the **PZn₃**, **BTD-PZn₃-BTD**, and **PZn-(BTD-PZn)₂** FOs underscore the x-polarized nature of the lowest energy excited state for these supermolecules, and the globally delocalized character of their respective S_1 excited states. The low-lying excited states of these supermolecules are described by extensive configuration interaction (CI). The $S_0 \rightarrow S_1$ transitions of **PZn₃**, **BTD-PZn₃-BTD**, and **PZn-(BTD-PZn)₂** have large contributions ($\sim 55\%$) from the one electron HOMO \rightarrow LUMO (H \rightarrow L) configuration, and highlight the importance of quinoidal resonance contributions to this low-lying excited state; this resonance contribution plays

a key role in the solvent-dielectric dependent fluorescence quantum yields evidenced in THF and toluene solvent.

For **PZn₃** (Fig. 2), seven single excitation configurations describe the transition matrix eigenvector; in contrast, nine and six single excitation configurations describe respectively the transition matrix eigenvectors for **BTD-PZn₃-BTD** and **PZn-(BTD-PZn)₂**. For **PZn₃**, the transition eigenvector is dominated by configurations (representing an $\sim 82\%$ weighting in the CI expansion) that do not redistribute to a significant degree electron density. For **BTD-PZn₃-BTD** and **PZn-(BTD-PZn)₂**, the relative importance of such single excitation configurations that do not redistribute significantly electron density drops respectively to 74 and 22%, highlighting the increased importance of single-excitation configurations that redistribute electron density in the CI expansions that describe the $S_0 \rightarrow S_1$ transition eigenvectors for these supermolecules. While the bandgap represented by $\Delta E_{\text{HOMO-LUMO}}$ varies to a minor degree and spans ~ 4.5 to 4.8 eV, note that the FO bandwidth in these Fig. 2–4 fluorophores varies significantly: $\Delta E_{\text{H-5-L+5}}$ for **PZn₃** is 9.77 eV but only 7.76 eV for **PZn-(BTD-PZn)₂**, a decrease of 2.01 eV.



Fig. 3 Calculated frontier molecular orbitals, energy levels, and relative one-electron configurations that contribute to the lowest energy (Q_x) transition for **BTD-PZn₃-BTD**. Calculations were performed at the M11/6-311g(d) theory level.



Fig. 4 Calculated frontier molecular orbitals, energy levels, and relative one-electron configurations that contribute to the lowest energy (Q_x) transition for **PZn-(BTD-PZn)₂**. Calculations were performed at the M11/6-311g(d) theory level.



Note that as supermolecular conjugation evolves from PZn_3 to $\text{BTD-PZn}_3\text{-BTD}$ and PZn-(BTD-PZn)_2 , an increased degeneracy of energy eigenvalues is manifest in the Fig. 2–4 FOs. This increased density of states (DOS) evident in $\text{BTD-PZn}_3\text{-BTD}$ and PZn-(BTD-PZn)_2 derives from the mixing of BTD and PZn frontier orbitals. Congruent with Fermi's Golden Rule, transition probability correlates with increased DOS. This larger DOS near the HOMO and LUMO levels serve to increase the multi-reference nature of the S_1 state wave function; S_1 configurations that involve these delocalized and mixed frontier orbitals will thus exhibit enhanced S_1 state wave function spatial delocalization. This computational trend is in line with the experimentally determined transition moments, wherein Q_x -transition oscillator strengths for $\text{BTD-PZn}_3\text{-BTD}$ and PZn-(BTD-PZn)_2 exceed that of PZn_3 . Note also that this dispersion of FO energies, which decreases progressively from PZn_3 to $\text{BTD-PZn}_3\text{-BTD}$ to PZn-(BTD-PZn)_2 (Fig. 2–4), is correlated with increased π conjugation which is reflected in the computed electronic delocalization range function for these fluorophores (ESI†); due to the nature of these orbitals and their diminished energy gaps within the filled and empty regimes of the FO manifold, a greater weighted fraction of single excitation configurations having charge resonance character is manifest, congruent with the augmented Q_x absorption oscillator strengths observed for $\text{BTD-PZn}_n\text{-BTD}$ and PZn-(BTD-PZn)_n relative to PZn_n (Fig. 1, Tables 1 and 2).

Conclusions

We describe a design strategy for (porphinato)zinc-based supermolecules that possess large NIR fluorescence quantum yields. These PZn-(BTD-PZn)_n , $\text{PZn}_2\text{-(BTD-PZn)}_2$, and $\text{BTD-PZn}_n\text{-BTD}$ fluorophores feature either 4-ethynylbenzo[*c*][1,2,5]thiadiazole (**E-BTD**) units that terminate *meso-to-meso* ethyne-bridged (porphinato)zinc (PZn_n) arrays, or 4,7-diethynylbenzo[*c*][1,2,5]thiadiazole (**E-BTD-E**) spacers that are integrated into the backbone of these compositions. PZn-(BTD-PZn)_n , $\text{PZn}_2\text{-(BTD-PZn)}_2$, and $\text{BTD-PZn}_n\text{-BTD}$ chromophores are characterized by enhanced transfer of B-to-x-polarized Q-state oscillator strength relative to their corresponding PZn_n benchmarks, intensified Q_x absorption bands, supermolecule structural rigidification, spectrally narrow fluorescence emission bands, small magnitude Stokes shifts, and large radiative rate constant magnitudes. TD-DFT calculations point to the importance of a greater weighted fraction of single excitation configurations having charge resonance character that describe the $S_0 \rightarrow S_1$ transition matrix eigenvector in PZn-(BTD-PZn)_n , $\text{PZn}_2\text{-(BTD-PZn)}_2$, and $\text{BTD-PZn}_n\text{-BTD}$ emitters relative to PZn_n oscillators in driving these spectroscopic and dynamical properties. Collectively, these systems define an unusual family of intensely absorbing vis-NIR absorbers that display strikingly high fluorescence quantum yields (ϕ_f values) over the 700–900 nm regime of the NIR. These THF ϕ_f values for PZn-(BTD-PZn)_n , $\text{PZn}_2\text{-(BTD-PZn)}_2$, and $\text{BTD-PZn}_n\text{-BTD}$ supermolecules, perhaps without peer in a solvent of this dielectric strength over this spectral window, range from 10–25%; notably these ϕ_f values are dramatically amplified in hydrophobic media and

thus contrast the behavior of classic tricarbocyanine NIR dyes, displaying fluorescence quantum yields ranging from 28–36% in toluene solvent. Because these $\text{BTD-PZn}_n\text{-BTD}$, PZn-(BTD-PZn)_n , and $\text{PZn}_2\text{-(BTD-PZn)}_2$ supermolecules display extraordinarily large NIR fluorescence quantum yields in hydrophobic solvent, these designs underscore new opportunities to evolve NIR-emissive nano- and mesoscale vesicles for fluorescence imaging applications in which the emissive irradiance exceeds the impressive metrics already established for such structures that membrane-disperse PZn_n fluorophores.^{54–61,65}

Experimental section

Synthesis and characterization

The synthetic procedures and corresponding characterization data of all new compounds, complete with the reaction schemes, are given in the ESI.†

Instrumentation

Electronic absorption spectra were recorded on a Shimadzu UV-1700 spectrophotometer. Steady-state emission spectra were recorded on a FLS920 spectrometer that utilized a xenon lamp (Xe900) as the excitation light source and an extended red sensitive PMT (Hamamatsu R2658P side window photomultiplier, spectral range: 200–1010 nm) for detection. Emission spectra were corrected using a calibration curve supplied with the instrument.

Fluorescence lifetime measurements

Time-resolved emission spectra were recorded using a Hamamatsu C4780 picosecond fluorescence lifetime measurement system. This system employs a Hamamatsu Streakscope C4334 as its photon-counting detector; a Hamamatsu C4792-01 synchronous delay generator electronically generated all time delays. The excitation light source chosen was a Hamamatsu 405 nm diode laser. Fluorescence lifetimes were acquired in single-photon-counting mode using Hamamatsu HPD-TA software and analyzed using the Hamamatsu fitting module.

Quantum yield measurements

A Hamamatsu C9920-03 Absolute Quantum Yield Measurement System was employed to make the quantum yield measurements. Excitation initiates from a Xe-lamp, where the wavelength is selected by a monochromator, and passed through a 1 mm optical excitation fiber. The inside of the sphere is coated with Spectralon (Labsphere, Inc.) that has at least 99% reflectance over the 350–1650 nm spectral window. Added detail regarding these measurements is provided in the ESI.†

Time-dependent density functional theory calculations

All electronic structure calculations were performed upon model compounds in which aliphatic chains were truncated to methyl groups (ESI†). Structure optimization and linear response calculations were performed with density functional theory (DFT) using Gaussian 16, revision C.01.⁶⁷ The M11⁶⁸



functional was employed for all calculations. Optimizations were performed with minimal symmetry constraints using tight optimization criteria with the 6-311g(d) basis set implemented. Selected frontier orbital wave functions were plotted as iso-surfaces (iso = 0.02) using Avogadro.⁶⁹ TD-DFT result files were post-processed using the GaussSum package;⁷⁰ this software partitions the wave function amplitudes onto atomic components using Mulliken population analysis,⁷¹ and parses the electronic configurations contributing to each excitation.

Conflicts of interest

No competing financial interests have been declared.

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