

Cite this: *Chem. Sci.*, 2018, 9, 1996

A luminescent organic radical with two pyridyl groups: high photostability and dual stimuli-responsive properties, with theoretical analyses of photophysical processes†

Shun Kimura,^a Akira Tanushi,^a Tetsuro Kusamoto,^{*a} Shuntaro Kochi,^b Tohru Sato^{bc} and Hiroshi Nishihara^{*a}

Luminescent monoradicals are expected to show unique properties based on their doublet state, where establishing a method to improve their photostability is an important issue for expanding their photofunctionality. We synthesized a highly photostable luminescent organic radical, the bis(3,5-dichloro-4-pyridyl)(2,4,6-trichlorophenyl)methyl radical (bisPyTM), containing two pyridyl groups on a tris(2,4,6-trichlorophenyl)methyl radical (TTM) skeleton. bisPyTM in dichloromethane exhibited fluorescence with an emission peak wavelength, λ_{em} , of 650 nm. We visually detected an emission ($\lambda_{\text{em}} = 712$ nm) from crystalline bisPyTM at 77 K, which is the first example of definite solid-state emission in a radical. Introducing the two nitrogen atoms into the TTM skeleton was shown to lower the energies of the frontier orbitals. The oscillator strength, f , of the electronic transition between the lowest excited state and the ground state, and the off-diagonal vibronic coupling constants (VCCs) were calculated theoretically for bisPyTM and the (3,5-dichloro-4-pyridyl)bis(2,4,6-trichlorophenyl)methyl radical (PyBTM). The calculated PyBTM to bisPyTM ratios for f or VCC agreed well with experimental radiative and non-radiative rate constants (k_r and k_{nr}) ratios, respectively. This study shows that scaled k_r and k_{nr} can be estimated and compared in this class of radicals using theoretical calculations, greatly advancing the prediction and design of their photofunctionality. The half-life of bisPyTM upon continuous UV light irradiation in dichloromethane was 47 or 3000 times longer than those of PyBTM (which contains one pyridyl group) and TTM (which has no pyridyl rings), respectively. The electrochemical and luminescent properties of bisPyTM were modulated in two stages using protons or $\text{B}(\text{C}_6\text{F}_5)_3$.

Received 14th September 2017
Accepted 9th January 2018

DOI: 10.1039/c7sc04034b

rsc.li/chemical-science

Introduction

Luminescent molecules are attracting increasing attention because of their suitability for a wide variety of scientific and engineering applications, such as organic light-emitting diodes and chemosensors. Several emission mechanisms, such as fluorescence *via* the singlet excited state,¹ phosphorescence from the triplet excited state,² and thermally activated delayed fluorescence,^{3,4} have been used to obtain the desired device

properties. In contrast with conventional luminescent molecules, which are closed-shell with singlet ground states, radicals with an unpaired electron have been the focus of research into next-generation luminophores.^{5–12} Radicals, in which fluorescence occurs from the lowest doublet excited state (D_1) to the doublet ground state (D_0), can exhibit unique characteristics, such as having long emission wavelengths without extended π -conjugation.¹³ Of particular importance in electroluminescent devices is the high efficiency of the internal electron (hole) $\rightarrow$ photon conversion ratio, up to 100%,^{14,15} which has been experimentally suggested recently.¹⁶ Although closed-shell fluorescent emitters suffer from quenching *via* the lowest triplet excited state (T_1), which is statistically formed together with the lowest singlet excited state (S_1) in a ratio of 3 : 1, particular monoradicals do not suffer from this quenching problem because of the absence of excited states between D_1 and D_0 and the spin-allowed $D_1 \rightarrow D_0$ transition.¹⁴

The doublet luminescence of radicals has been investigated much less than that of closed-shell luminescent molecules, because of its rarity and low chemical stability under ambient

^aDepartment of Chemistry, Graduate School of Science, The University of Tokyo, 7-3-1 Hongo, Bunkyo-ku, Tokyo 113-0033, Japan. E-mail: kusamoto@chem.s.u-tokyo.ac.jp; nishihara@chem.s.u-tokyo.ac.jp

^bDepartment of Molecular Engineering, Graduate School of Engineering, Kyoto University, Nishikyo-ku, Kyoto, 615-8510, Japan

^cUnit of Elements Strategy Initiative for Catalysts & Batteries, Kyoto University, Nishikyo-ku, Kyoto, 615-8510, Japan

† Electronic supplementary information (ESI) available: Crystal structure data, ESR spectral data, DFT and TD-DFT calculations data, spectroscopic data, quantum yield and photostability in various solvents, electrochemical data (PDF). CCDC 1559962. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c7sc04034b



conditions or in photoexcited states.^{17,18} Following studies on the emission properties of *in situ*-generated unstable diarylmethyl or triarylmethyl radicals,^{19–22} polychlorinated triarylmethyl (PTM) and tris(2,4,6-trichlorophenyl)methyl (TTM) radicals (Fig. 1) have been developed as luminescent radicals.^{23,24} They show excellent chemical stability under ambient conditions, although they decompose upon photoirradiation.^{17,18} Improving the photostability of radicals is an important step toward developing unique doublet-based photofunctionality or novel phenomena arising from the interplay between luminescent and magnetic (spin) properties.

A drastic increase in the photostability of luminescent radicals has been achieved in the (3,5-dichloro-4-pyridyl)bis(2,4,6-trichlorophenyl)methyl radical (PyBTM; Fig. 1), prepared by our group.¹⁴ PyBTM has a pyridyl ring in the TTM skeleton and showed fluorescence with an absolute quantum yield (ϕ_{em}) of 0.01–0.03 in solution and a ϕ_{em} value of 0.26 in a rigid matrix at room temperature. Importantly, its stability upon UV irradiation was up to 115 times higher than that of TTM. The excellent photostability has expanded the scope of doublet-based photofunctionality in radicals. We have reported on stimuli responsiveness,²⁵ the halogen atom substitution effect,²⁶ and increased ϕ_{em} and photostability upon coordination to metal ions in PyBTM.^{27–29} Studies on PyBTM have suggested that introducing a pyridyl group may lead to radicals with improved photostability, owing to the lowering of energy levels around the frontier orbitals. However, the utility of this method is not well established.

In this study, we developed a new bis(3,5-dichloro-4-pyridyl)(2,4,6-trichlorophenyl)methyl radical (bisPyTM; Fig. 1), which has two pyridyl groups, and is stable and luminescent. We aimed to resolve two issues in understanding the fundamental properties of luminescent triarylmethyl radicals. The first is the effects of introducing an additional pyridyl ring into the PyBTM skeleton on the energies of the frontier orbitals, luminescent properties, and photostability. The increase in the number of pyridyl rings is expected to increase the photostability and to expand the variation of the coordination structure in relation to multistage chemical responses or high-dimensional assemblies.^{30,31}

The second challenge is interpreting the luminescence characteristics and photophysical parameters, including the radiative and non-radiative rate constants (k_r and k_{nr}), through theoretical calculations using density functional theory (DFT) and time-dependent (TD)-DFT. The technical challenges and limited number of compounds mean that joint theoretical and experimental studies on the relaxation processes of luminescent radicals in their excited state are still rare. However, conical

intersection³² and back-electron transfer⁸ have been investigated theoretically as non-radiative relaxation processes of radicals in the excited state. We calculated the excited states and molecular and electronic structures of PyBTM and bisPyTM, and investigated non-radiative relaxation processes *via* vibronic coupling density (VCD) analysis³³ to evaluate the rate of internal conversion. The differences in k_r and k_{nr} between the two radicals can be explained through the difference in oscillator strengths and off-diagonal vibronic coupling constants (VCCs).

Results and discussion

Theoretical methods for estimating photophysical parameters

The relaxation process from the D_1 state to the D_0 state consists of radiative and non-radiative transitions (Scheme 1). Luminescence corresponds to the radiative $D_1 \rightarrow D_0$ transition; thus, k_r is proportional to the oscillator strength, f , of the transition for the D_1 nuclear coordinates. The f value is calculated using TD-DFT in this study. In contrast, the non-radiative transition is expected to consist of internal conversion followed by vibrational relaxation in the radicals studied here (Scheme 1). Intersystem crossing from S_1 to T_n states can also provide a non-radiative relaxation pathway in conventional closed-shell luminescent molecules. However, intersystem crossing from D_1 to other excited states with different multiplicities, such as a quartet state (Q_1), is negligible, because the Q_1 state is energetically much higher than the D_1 state.¹⁴ Thus, k_{nr} is determined mainly by the rate of internal conversion. Internal conversion has been suggested as a major pathway in the non-radiative decay of some radicals,^{21,28,34} however, no theoretical calculations have been performed to clarify the deactivation mechanism. In this study, VCD analysis of the luminescent radicals elucidated the mechanism of their non-radiative decay. This was suitable for our radicals, because it allowed us to evaluate the rate of internal conversion and to determine the vibrational modes that contribute to internal conversion. Internal conversion is caused by off-diagonal vibronic coupling. The rate of internal conversion is proportional to the square of

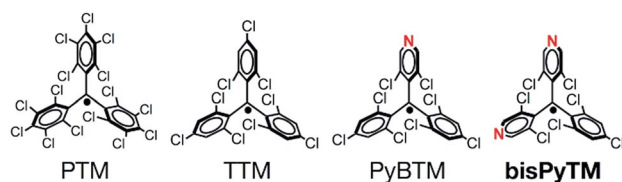
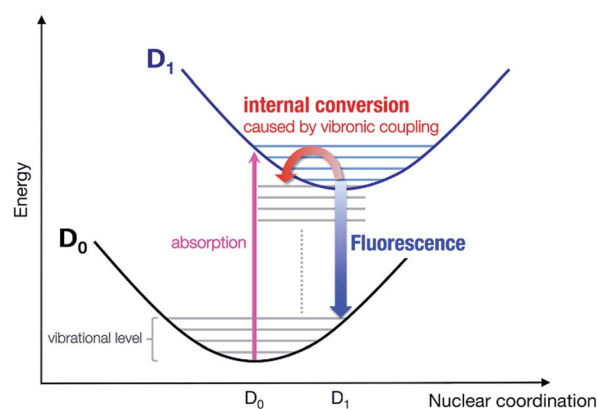


Fig. 1 The molecular structures of PTM, TTM, PyBTM and bisPyTM.



Scheme 1 Monoradical deactivation processes from the lowest excited (D_1) state.



the off-diagonal VCCs.³³ The off-diagonal VCC of mode α between electronic states m and n is defined by

$$V_{mn,\alpha} = \left\langle \Psi_m(r, R_0) \left| \left(\frac{\partial H(r, R)}{\partial Q_\alpha} \right)_{R_0} \right| \Psi_n(r, R_0) \right\rangle, \quad (1)$$

where $H(r, R)$ is the molecular Hamiltonian for a set of electronic coordinates $r = \{r_1, \dots, r_i, \dots, r_N\}$ and a set of nuclear coordinates $R = \{R_1, \dots, R_A, \dots, R_M\}$. Ψ_m and Ψ_n are the electronic wavefunctions of states m and n , respectively. Q_α is the normal coordinate of mode α , and R_0 denotes a reference nuclear configuration. We took the geometries after the vibrational relaxations as reference nuclear configurations.

The origin of a vibronic coupling in terms of the electronic and vibrational states can be visualized using VCD analysis,³⁴ which provides insight into the vibronic coupling in a molecule. Furthermore, we can control vibronic couplings to suppress internal conversion. For instance, based on the VCD analysis of a non-emitting molecule, triphenylamine (TPA), Sato *et al.* reduced the off-diagonal VCCs by introducing non-emitting substituents to make the TPA moiety fluorescent.^{35,36} The VCD is defined by

$$\eta_{mn,\alpha}(r_i) = \rho_{mn}(r_i) \times \nu_\alpha(r_i), \quad (2)$$

where ρ_{mn} is the overlap density

$$\rho_{mn}(r_i) = N \int \Psi_m^*(r, R) \Psi_n(r, R) dr_1 ds_1 \dots dr_N ds_N \quad (3)$$

and ν_α is the potential derivative with respect to Q_α

$$\nu_\alpha(r_i) = \left[\frac{\partial}{\partial Q_\alpha} \left(\sum_{A=1}^M \frac{-Z_A e^2}{4\pi\epsilon_0 |r_i - R_A|} \right) \right]_{R_0}. \quad (4)$$

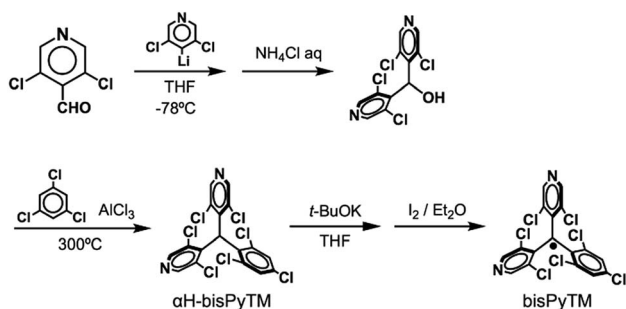
The spatial integral of the VCD is equal to the VCC:

$$V_{mn,\alpha} = \int \eta_{mn,\alpha}(r_i) dr_i. \quad (5)$$

Using eqn (2) and (5), the origin of a VCC can be described using functions of η , ρ , and ν , which depend on position.

Synthesis and characterization

bisPyTM was prepared according to Scheme 2, and characterized *via* electron spin resonance (ESR) spectroscopy, high-



Scheme 2 The synthetic route for bisPyTM.

resolution electrospray ionization mass spectrometry, and elemental analysis. The $S = 1/2$ spin per bisPyTM molecule was confirmed by comparing the value of the second integral of the ESR signal intensity with that of the reference compound, 4-hydroxy-TEMPO. The molecular structure of bisPyTM in the crystalline state was analyzed *via* single-crystal X-ray diffraction (Fig. 2). The central carbon atom, C1, is sp^2 hybridized and sterically shielded by six halogen atoms. C1 and its three neighboring carbon atoms (C4, C9, and C15) lie in the same plane. The three aromatic rings are held in a propeller-like conformation due to steric hindrance from the chlorine atoms. The bond angles and dihedral angles (Table S1†) are similar to those of PyBTM.

ESR spectroscopy at 177 K

An ESR spectrum of bisPyTM in dichloromethane was measured at 177 K to estimate the spin density distribution over the bisPyTM skeleton (Fig. 3a). The spectrum showed a characteristic hyperfine structure with a g value of 2.004, similar to that of PyBTM. The spectrum was reproduced *via* computational simulation, considering hyperfine coupling with six ^1H and two ^{14}N atoms. The obtained hyperfine coupling constants (hccs) were qualitatively similar to those calculated *via* DFT (Table S2†). The results indicate that the spin density was delocalized over the molecule, including the central C1 atom and two pyridyl nitrogen atoms (Fig. 3b). The simulated hcc of ^{14}N (0.108 mT) was comparable to that of PyBTM (0.115 mT). This result indicates that the spin density on the N atom was similar in bisPyTM and PyBTM;¹⁴ the densities estimated from the simulated hcc and DFT were 0.06 and 0.069, respectively.

Electronic structure in the ground state (D_0 state)

The electronic properties of bisPyTM were calculated *via* DFT, to reveal the effects of introducing two pyridyl rings to the TTM framework on the distribution and energies around the frontier orbitals. The unrestricted B3LYP level of theory with 6-31G(d,p) basis sets was used, and the solvent effect (dichloromethane) was treated using the polarizable continuum model. The calculated molecular orbitals (MOs) of bisPyTM are shown in Fig. 4. The next highest occupied molecular orbital (NHOMO; 120α and β) and NHOMO-1 (119α and β) were assumed to be the bonding and antibonding combination of wavefunctions at

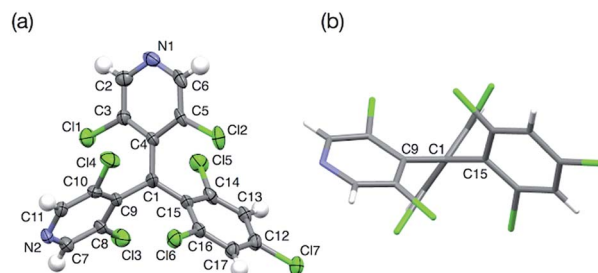


Fig. 2 (a) The molecular structure of bisPyTM in the crystalline state with thermal ellipsoids set at 50% probability, and (b) the view along the C1–C4 bond.



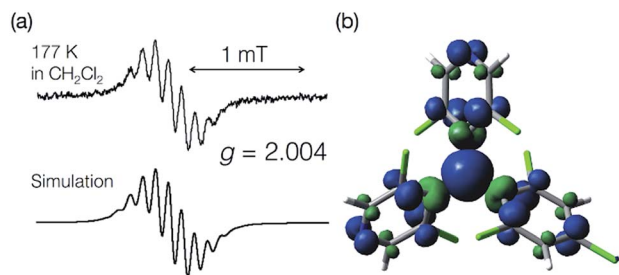


Fig. 3 (a) An ESR spectrum of bisPyTM in CH_2Cl_2 at 177 K (top), and a computational simulation (bottom). (b) The spin density distribution on bisPyTM calculated *via* DFT (UB3LYP/6-31G(d,p)).

the two pyridyl rings, which were distributed mainly on the two pyridyl rings. These two MOs were almost degenerate. The next lowest unoccupied molecular orbital (NLUMO; 122α and β) of bisPyTM was delocalized on the two pyridyl rings, whereas the singly occupied molecular orbital (SOMO; 121α and β) was located mainly on the central carbon atom with non-negligible delocalization on the three aromatic rings. The distribution of the MOs was similar to that in PyBTM, where the NHOMO and NLUMO were distributed mainly on the pyridyl ring, and the SOMO was located on the central carbon atom and extended to the aromatic rings.

The energy diagram shows that the energies of the frontier orbitals (119α – 122α and 119β – 122β) of the radicals all decreased as the number of pyridyl rings increased; bisPyTM had energies 0.10–0.20 eV lower than PyBTM. This decrease is expected to change the optical and redox properties and stabilities of the compounds, because the frontier orbital energies directly or indirectly determine the optical and redox properties.²⁵ The lowering of the energy of the β -SOMO (121β for bisPyTM and 121β for PyBTM) was confirmed *via* cyclic voltammetry.

Cyclic voltammetry

The redox properties of the radicals were investigated *via* cyclic voltammetry. The cyclic voltammogram of bisPyTM in 0.1 M $n\text{-Bu}_4\text{NClO}_4/\text{CH}_2\text{Cl}_2$ (Fig. 5a and S1†) showed a reversible reduction wave at $E^0 = -0.57$ V vs. the ferrocenium/ferrocene (Fc^+/Fc) redox couple, corresponding to the reduction of bisPyTM (Scheme 3a). This process was explained through the injection of an electron into the β -SOMO. The E^0 value of bisPyTM was more positive than those of PyBTM (-0.74 V) and TTM (-0.99 V) under identical conditions, suggesting that increasing the number of pyridyl rings increases the electron-accepting ability of the radical and decreases the energy level of the β -SOMO. This observation can be explained by the higher electronegativity of nitrogen compared to carbon.¹⁴

Optical properties

The UV/vis absorption spectrum of bisPyTM in dichloromethane displayed a strong near-UV band ($\lambda_{\text{abs}} = 355$ nm, transition from 121α to 122α and 123α) and a low-lying band (Fig. 6a). The lowest-energy transition band (transition from

120β to 121β) that formed the lowest excited state (D_1 state) was observed at a peak wavelength of 536 nm. These transitions were assigned based on TD-DFT calculations (ESI†). The relative wavelengths and oscillator strengths for the calculated transition bands agreed with the absorption spectrum obtained experimentally, supporting the orbital assignments and the validity of the calculations. A β -electron-centered $120\beta \rightarrow 121\beta$ electronic transition induced the D_1 state, as in TTM, PyBTM, and PyBTM derivatives.^{14,25–29}

bisPyTM in dichloromethane displayed fluorescence, with an emission maximum wavelength, λ_{em} , of 650 nm upon excitation at $\lambda_{\text{ex}} = 355$ nm (Fig. 6a and c). ϕ_{em} and the lifetime (τ) were measured as being 0.009 and 3.6 ns, respectively. This emission wavelength was strongly red-shifted compared with PyBTM and TTM, indicating the large structural change in bisPyTM in the excited state.

We estimated k_r and k_{nr} of bisPyTM and PyBTM using eqn (6) and (7) (Table 1). k_r of bisPyTM was smaller than that of PyBTM, whereas k_{nr} was considerably larger. The observed PyBTM to bisPyTM ratio of the k_r values was consistent with results from the theoretical calculation described in the next section. The greater k_{nr} of bisPyTM was explained by accelerated internal conversion, which is also described in the next section.

$$\phi_{\text{em}} = k_r / (k_r + k_{\text{nr}}) \quad (6)$$

$$\tau = 1 / (k_r + k_{\text{nr}}) \quad (7)$$

Surprisingly, bisPyTM exhibited luminescence both in solution and in the solid state. Deep red luminescence with $\lambda_{\text{em}} = 712$ nm was observed in the crystalline state (Fig. 6a and d) at 77 K. To the best of our knowledge, this is the first reported organic radical showing distinct luminescence in its crystalline state.³⁷ We propose that quenching *via* intermolecular interactions in the crystalline state is weaker for bisPyTM than for the other radicals, explaining the luminescence in the crystalline state. Solid-state luminescence was also observed when we doped bisPyTM into the αH -bisPyTM crystal, the precursor for bisPyTM (Scheme 2). The emission wavelength of the 3 wt%-doped sample was shorter than that of bisPyTM in dichloromethane, which would be caused by reduced reorientation energy due to rigid packing. The emission wavelength was red-shifted, and the intensity decreased with increasing bisPyTM concentration (Fig. 6b and e), implying that an excimer was formed under highly doped conditions.³⁸

Molecular and electronic structures in the lowest excited state (D_1 state)

The molecular and electronic structures of bisPyTM in the D_1 state were calculated *via* TD-DFT (UB3LYP/6-31G(d,p)) to clarify the characteristics of the radiative and non-radiative processes that occurred from the D_1 state.

In the optimized molecular structure of bisPyTM in its D_1 state, the bond lengths between the central carbon atom and the aryl moieties (C1–C4, C1–C9, C1–C15) and between carbon atoms at the *ortho* and *meta* positions (C5–C6, C2–C3, C7–C8,



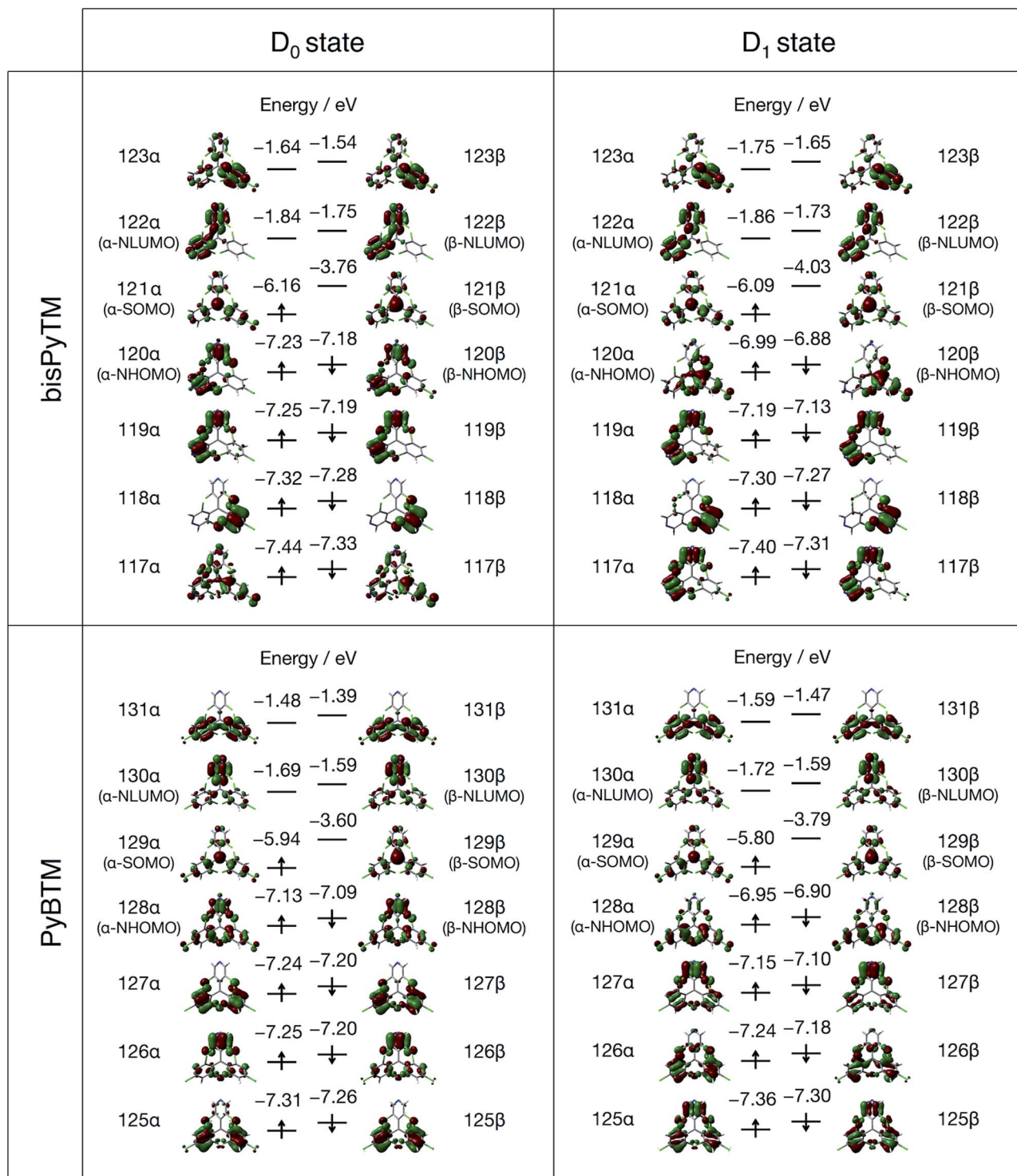


Fig. 4 The molecular orbitals of bisPyTM and PyBTM in the D₀ and D₁ states, calculated *via* DFT (UB3LYP/6-31G(d,p)). The solvent (dichloromethane) effect is treated using the polarizable continuum model.

C10–C11, C16–C17, C13–C14) were shorter than those in the D₀ state, whereas the other C–C and C–N bonds were longer (Fig. 7a). The N1–C4–C1 and N2–C9–C1 bond angles in the two pyridyl rings decreased (Fig. 7b). These results indicate the increased quinoid character of the aryl groups as the aromaticity of the pyridyl rings decreased in the D₁ state.

The MOs of bisPyTM in the D₁ state in dichloromethane are shown in Fig. 4. The distribution of each MO is similar to that in the D₀ state, except the α(β)-NHOMO (120α(β)). The distribution of the α(β)-NHOMO was similar to that of the α(β)-NHOMO–3 in the D₀ state. The structural change between the D₁ and the D₀ states induced the energy level reordering of the energetically closed MOs.



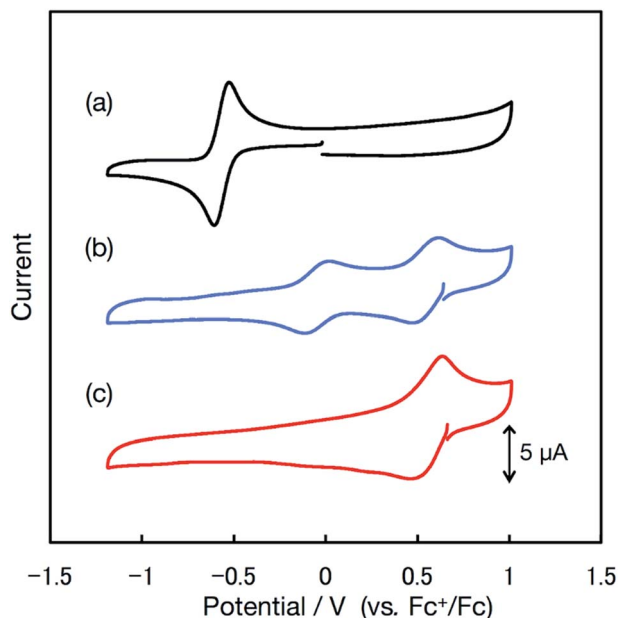
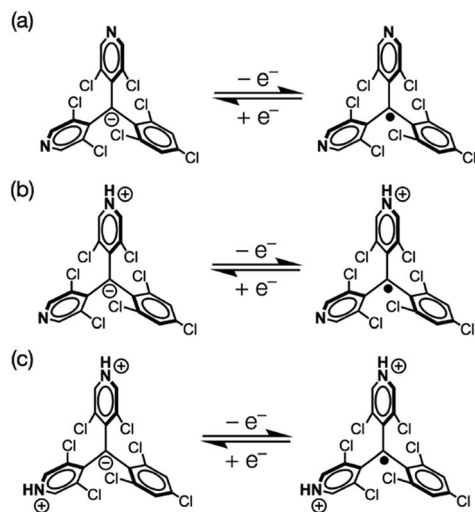


Fig. 5 Cyclic voltammograms of bisPyTM (0.5 mM) in 0.1 M $n\text{Bu}_4\text{NClO}_4/\text{CH}_2\text{Cl}_2$ at a scan rate of 0.1 V s^{-1} (a) with no additives and (b) with 2.0 eq. and (c) 5.0 eq. of TfOH.



Scheme 3 Redox behaviors of (a) bisPyTM, (b) $[\text{HN-bisPyTM}]^+$ and (c) $[(\text{HN})_2\text{-bisPyTM}]^{2+}$.

The MOs of the reference compound, PyBTM, in the D_1 state in dichloromethane were calculated (Fig. 4). The MO distributions were similar to those in the D_0 state in the unoccupied orbitals, whereas modulation of the distribution was detected in the $\alpha(\beta)\text{-NHOMO}$ level. The $\alpha(\beta)\text{-NHOMO}$ was delocalized on the pyridyl group and two trichlorophenyl groups in the D_0 state, whereas it was delocalized on the two trichlorophenyl groups in the D_1 state. Introducing the solvent effect using the polarizable continuum model caused the reordering of energy levels and altered the distribution of MOs, including the $\alpha(\beta)\text{-NHOMO}$. The $\alpha(\beta)\text{-NHOMO}$ was located at the pyridyl ring in the D_1 state without the solvent effect, whereas it was

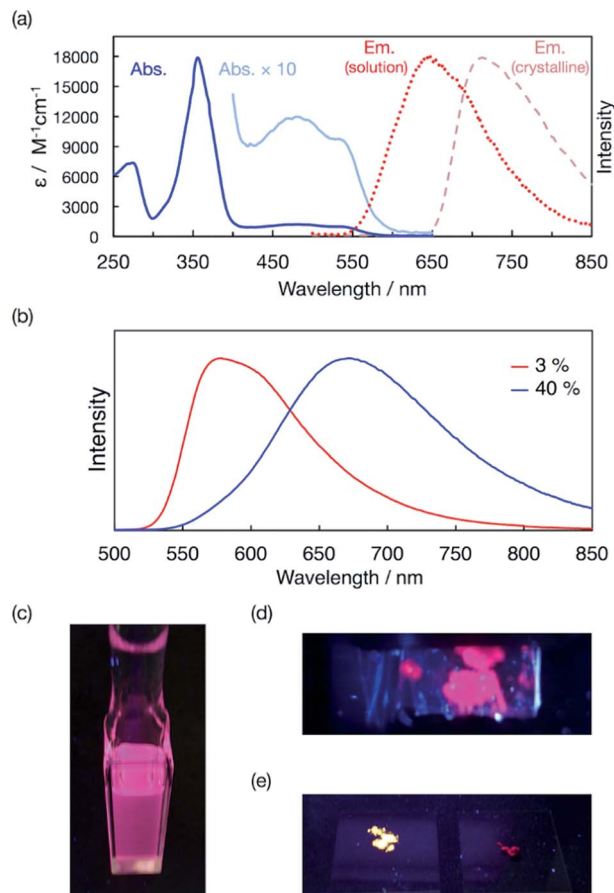


Fig. 6 (a) Absorption (blue line) and emission (red dotted line, $\lambda_{\text{ex}} = 355 \text{ nm}$) spectra of bisPyTM in dichloromethane (4.5×10^{-5} and $1.2 \times 10^{-5} \text{ M}$, respectively) at room temperature, and an emission spectrum in the crystalline state at 77 K (pale red dashed line, $\lambda_{\text{ex}} = 355 \text{ nm}$). An enlarged portion of the absorption spectrum (10 fold) is shown for $\lambda = 400$ to 650 nm (pale blue line). (b) Emission spectra of $\alpha\text{H-bisPyTM}$ crystals doped with 3 (red) and 40 wt% (blue) bisPyTM. (c) Photograph of bisPyTM solution in dichloromethane under UV light at $\lambda = 365 \text{ nm}$. (d) Photograph of bisPyTM crystals at 77 K under UV light at $\lambda = 365 \text{ nm}$. (e) Photographs of $\alpha\text{H-bisPyTM}$ crystals doped with 3 (left) and 40 wt% (right) bisPyTM under UV light at $\lambda = 365 \text{ nm}$.

distributed on the trichlorophenyl moieties with negligible extension to the pyridyl ring with the solvent effect (Fig. 4 and S2†). Because the $\beta\text{-NHOMO}$ was directly involved in forming the D_1 state, the change in the distribution on the $\beta\text{-NHOMO}$ should affect the photophysical properties. These results demonstrate the importance of considering the solvent effect in these calculations.²⁹ Therefore, we performed theoretical analysis considering the effects of dichloromethane.

Theoretical analyses of photophysical parameters

We estimated the rates of photophysical processes, k_r and k_{nr} , for PyBTM and bisPyTM to explain the experimentally observed difference in ϕ_{em} . k_r and k_{nr} can be evaluated by calculating the f values and off-diagonal VCCs between the D_1 and the D_0 states, as mentioned in the section 'Theoretical methods for estimating photophysical parameters'. Off-diagonal VCDs



Table 1 The photophysical parameters of the radicals

	$\lambda_{\text{abs1}}^{\text{exp}}/\text{nm}$	$\lambda_{\text{abs2}}^{\text{exp}}/\text{nm}$	$\lambda_{\text{em}}^{\text{exp}}/\text{nm}$	$\lambda_{\text{abs2}}^{\text{calc}}/\text{nm}$ (<i>f</i>)	$\lambda_{\text{em}}^{\text{calc}}/\text{nm}$ (<i>f</i>)	ϕ_{em}	τ/ns	$k_{\text{r}}/\times 10^6 \text{ s}^{-1}$	$k_{\text{nr}}/\times 10^8 \text{ s}^{-1}$
bisPyTM	355	536	650	496 (0.0153)	643 (0.0290)	0.009(0)	3.6	2.5	2.8
PyBTM	370	541	585	487 (0.0349)	572 (0.0457)	0.025(0)	6.4	3.9	1.5
TTM	373	542	570	—	—	0.02	7.0	3	1.4

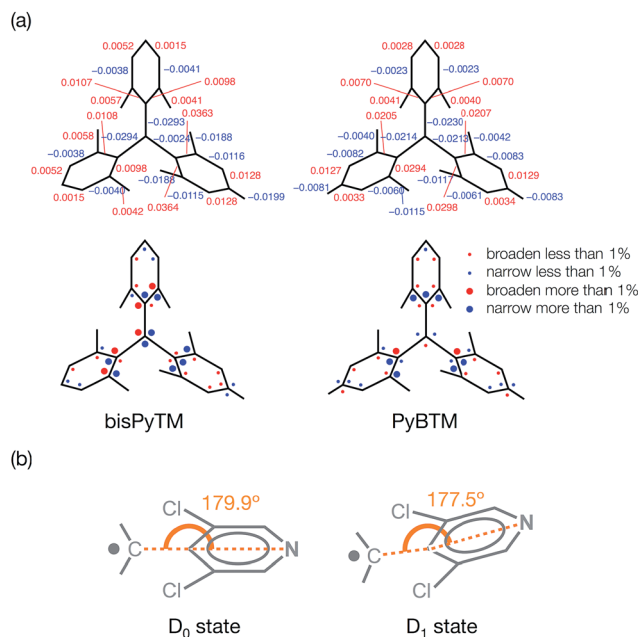


Fig. 7 A schematic diagram of the structural changes from the D₀ to D₁ states of bisPyTM and PyBTM: (a) changes in the bond lengths, in units of Å, and angles; and (b) changes in the N1–C4–C1 and N2–C9–C1 angles of bisPyTM.

indicate the vibrational modes that contribute to internal conversion.

The *f* values for the D₁ nuclear coordinates calculated using TD-DFT, considering the effects of the solvent (dichloromethane), were 0.0290 and 0.0457 for bisPyTM and PyBTM, respectively. The PyBTM to bisPyTM ratio of *f* calculated was 1.58. Neglecting the effects of Franck–Condon factors on the radiative rate constants,³³ we compared this ratio with the observed ratio of the radiative rate constants. The observed ratio of PyBTM to bisPyTM was 1.6. The calculated *f* values reproduced the ratio of the observed *k_r* values for the two radicals well.

Fig. 8 shows the off-diagonal VCCs and VCDs of bisPyTM and PyBTM. The VCCs of bisPyTM were smaller than those of common organic compounds. For instance, the maximum VCC in a triphenylamine derivative that is known as a non-fluorescent molecule is *ca.* 8×10^{-4} a.u.,³⁵ while that in bisPyTM is less than 7×10^{-4} a.u. The square sum of the VCCs of bisPyTM was 1.9×10^{-6} a.u., which was larger than that of PyBTM (1.4×10^{-6} a.u.), reproducing the greater *k_{nr}* value of bisPyTM. The observed ratio of *k_{nr}*, *k_{nr}* (bisPyTM)/*k_{nr}* (PyBTM), was 1.9 (Table 1). Neglecting the effects of Franck–Condon factors on *k_{nr}*,³³ we calculated the ratio of *k_{nr}* values from the square sums of the off-diagonal VCCs. The ratio of the square sum for bisPyTM to that for PyBTM is 1.36, which is reasonably consistent with the observed ratio. This difference in off-

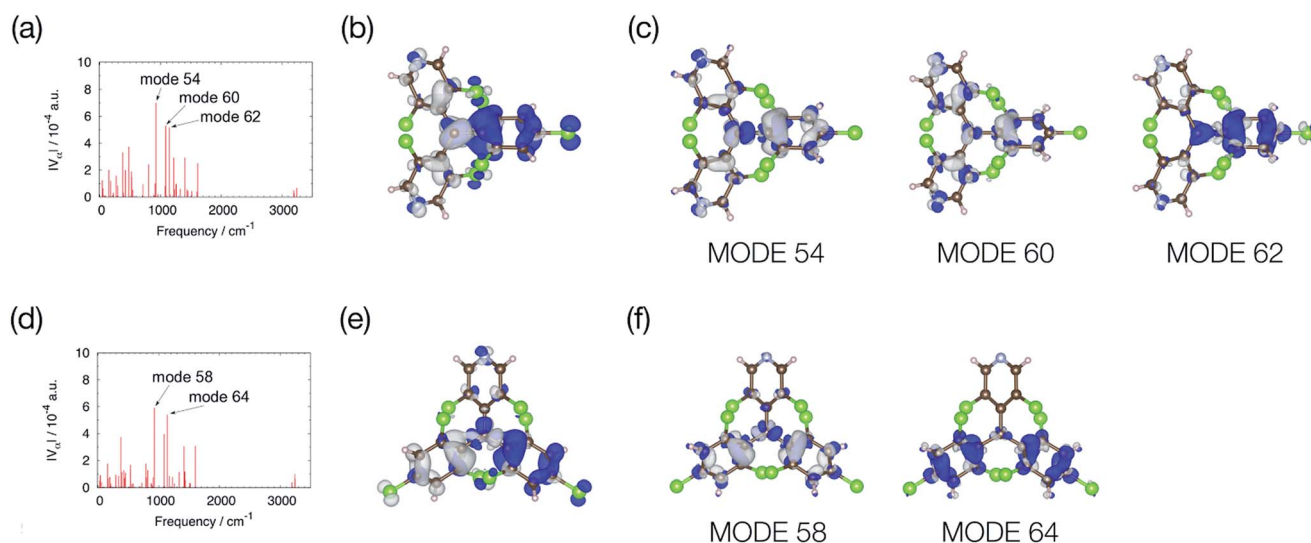


Fig. 8 Off-diagonal VCCs between the adiabatic D₁ and D₀ states for the normal mode of (a) bisPyTM and (d) PyBTM in dichloromethane. The overlap density (isovalue of $\rho = 0.01$) of (b) bisPyTM and (e) PyBTM, and the off-diagonal VCD between the adiabatic D₁ and D₀ states for selected vibrational modes of (c) bisPyTM and (f) PyBTM (isovalue of 4×10^{-6} a.u.).



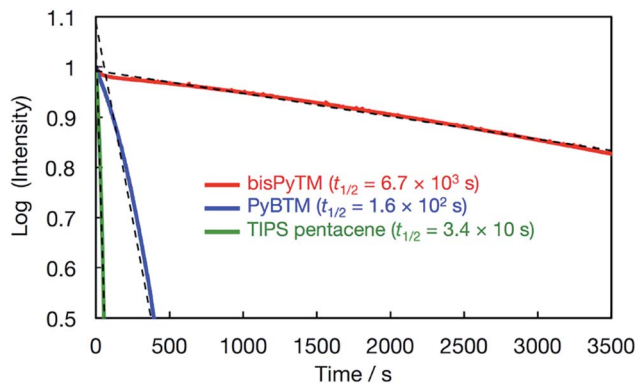


Fig. 9 Plots showing the emission decay of bisPyTM, PyBTM, and TIPS pentacene in dichloromethane under continuous excitation, with light at $\lambda_{\text{ex}} = 355$ nm (bisPyTM), 370 nm (PyBTM), and 310 nm (TIPS pentacene). The emission intensities are normalized.

diagonal VCCs in bisPyTM and PyBTM came from the difference in overlap density, which arose from the distribution of the NHOMOs. The overlap density was distributed on the two trichlorophenyl moieties in PyBTM, whereas it was on the only trichlorophenyl moiety in bisPyTM; the greater localization in bisPyTM resulted in its larger VCD.

These results suggest that internal conversion was mainly attributed to a non-radiative decay process, and that the theoretical methods shown here enable us to estimate k_r and k_{nr} for this class of radicals, greatly advancing the prediction and design of their photofunctionality.

Photostability

The low photostability of luminescent radicals is an important problem that must be overcome. The decay of fluorescence intensity was monitored and compared for bisPyTM and PyBTM upon continuous irradiation with light at each maximum absorption wavelength. Fig. 9 shows the decay in luminescence intensity for each compound, and the degree of decay represents the decomposition rate upon exposure to light. The estimated half-life ($t_{1/2}$) in dichloromethane was 6.7×10^3 s; this value was 43 times larger than that of PyBTM (1.6×10^2 s).

Considering our previous results,¹⁴ bisPyTM is expected to show a half-life 3000 times longer than TTM. When the photostability is defined as a product of the absorption coefficients (ϵ) at the irradiation wavelengths and $t_{1/2}$, the photostability of bisPyTM was 30 times higher than PyBTM. In other aprotic solvents, bisPyTM was 4 to 44 times more stable (6 to 71 times longer half-lives) than PyBTM (Table 2). These results suggest that the photostability is greatly improved as the number of pyridyl groups in the TTM framework increases and the energy levels of the frontier orbitals are lowered.

$t_{1/2}$ values in the solvents were not simply scaled by solvent parameters such as the dielectric constant or dipole moment (Fig. S3†). This trend suggests that there is a specific interaction between the radical and solvent molecules that modulates the photostability. For example, exceptionally low $t_{1/2}$ values in alcohols were detected for both PyBTM and bisPyTM. The low values

Table 2 The half-lives ($t_{1/2}$) of compounds upon light irradiation

	bisPyTM ^a		PyBTM ^b		TIPS pentacene ^c		bisPyTM/ PyBTM ^d		bisPyTM/TIPS pentacene ^d	
	$t_{1/2}/\text{s}$	$\epsilon/\text{M}^{-1}\text{cm}^{-1}$	$t_{1/2} \times \epsilon$	$t_{1/2}/\text{s}$	$\epsilon/\text{M}^{-1}\text{cm}^{-1}$	$t_{1/2} \times \epsilon$	$t_{1/2}/\text{s}$	$t_{1/2} \times \epsilon$	$t_{1/2}/\text{s}$	$t_{1/2} \times \epsilon$
Dichloromethane	$6.7(3) \times 10^3$	1.78×10^4	1.2×10^8	$1.56(2) \times 10^2$	2.54×10^4	4.0×10^6	43	3.0×10^6	2.0×10	23
Acetonitrile	$1.4(1) \times 10^4$	2.19×10^4	3.2×10^8	$3.07(7) \times 10^3$	2.47×10^4	7.6×10^6	47	41	38	—
Acetone	$4.0(4) \times 10^3$	1.36×10^4	5.5×10^7	$6.4(1) \times 10^3$	2.26×10^4	1.4×10^7	6.3	3.8	35	21
Hexane	$7.8(5) \times 10^3$	1.85×10^4	1.4×10^8	$1.10(1) \times 10^2$	3.90×10^4	4.3×10^6	71	34	4.1	0.6
Chloroform	$7.3(5) \times 10^3$	1.65×10^4	1.2×10^8	$1.20(3) \times 10^3$	2.25×10^4	2.7×10^6	60	44	6.6×10	61
Ethanol	$2.0(2) \times 10$	1.48×10^4	3.0×10^5	$1.46(4) \times 10$	2.09×10^4	3.1×10^5	1.4	1.0	—	—
Methanol	$1.8(2) \times 10$	1.23×10^4	2.2×10^5	$2.0(3) \times 10$	2.25×10^4	4.4×10^5	0.90	0.49	—	—

^a $\lambda_{\text{ex}} = 355$ nm. ^b $\lambda_{\text{ex}} = 370$ nm. ^c $\lambda_{\text{ex}} = 310$ nm in dichloromethane, hexane, and chloroform; $\lambda_{\text{ex}} = 330$ nm in acetonitrile; and $\lambda_{\text{ex}} = 330$ nm in acetone. ^d A comparison of $t_{1/2}$ and $t_{1/2} \times \epsilon$ for the compounds. ^e Molar absorption coefficients at the irradiation wavelengths.

may result from hydrogen bonds formed between nitrogen atoms on the radicals and hydrogen atoms in the EtOH or MeOH molecules, which accelerate decomposition in the photoexcited state.

For industrial applications, the photostabilities of bisPyTM and 6,13-bis(triisopropylsilyl)ethynyl)pentacene (TIPS pentacene) were compared. TIPS pentacene is a closed-shell molecule commonly used in organic semiconductors as a benchmark material due to its high photostability and solubility.^{39–41} Its fluorescence peak maximum ($\lambda_{\text{em}} = 650$ nm upon excitation at $\lambda_{\text{ex}} = 360$ nm) was similar to those of PyBTM and bisPyTM. Fig. 9 and Table 2 provide clear evidence for the superior photostability of bisPyTM; the photostability of the radical was 0.6 to 61 times higher (4 to 660 times larger $t_{1/2}$ values) than TIPS pentacene. This result demonstrates the potential of the compound for practical applications.

Dual responsive properties

We aimed to control the optical and electrochemical properties of bisPyTM by using the two pyridyl nitrogen atoms in the molecular structure as chemical stimulus-responsive sites. First, we investigated the effects of adding a Brønsted–Lowry acid. The reversible response of the radical to protonation and deprotonation was confirmed *via* UV/vis absorption spectroscopy during an acid–base titration (Fig. 10a and c). A new transition band at $\lambda = 409$ nm appeared as the intensity of the absorption at $\lambda = 355$ nm decreased upon the addition of trifluoromethanesulfonic acid (TfOH). The initial spectral shape was almost recovered by adding triethylamine (NEt_3). Protonation affected the luminescence properties; the fluorescence intensity decreased substantially upon addition of TfOH (Fig. 10b and d). PyBTM showed similar absorption and emission spectra upon protonation and deprotonation.¹⁴

The redox properties, examined *via* cyclic voltammetry, also changed drastically upon protonation. Upon the successive addition of TfOH, two new redox waves at $E^{0'} = -0.05$ and 0.55 V appeared, and only one redox wave at $E^{0'} = 0.55$ V was observed after the addition of excess TfOH (Fig. 5c). This result suggests a two-stage increase in the electron-accepting ability of the molecule resulting from single-to-double protonation of the two pyridyl rings. Accordingly, the redox waves at $E^{0'} = -0.05$ and 0.55 V were ascribed to the $[\text{HN-bisPyTM}]^+ / [\text{HN-bisPyTM}]$ and $[(\text{HN})_2\text{-bisPyTM}]^{2+} / [(\text{HN})_2\text{-bisPyTM}]^+$ redox couples, respectively, where $[\text{HN-bisPyTM}]^+$ and $[(\text{HN})_2\text{-bisPyTM}]^{2+}$ are monoprotonated and diprotonated bisPyTM, respectively (Scheme 3). Monoprotonation or diprotonation shifted the reduction potential of bisPyTM by 0.52 or 1.12 V in the positive direction, respectively. Similar positive shifts were observed in the protonation or methylation of PyBTM,^{14,25} and were explained by a decrease in electron density upon protonation, which increased the electron-accepting ability.

The reduction potential of $[(\text{HN})_2\text{-bisPyTM}]^{2+}$ was higher than that of F_4TCNQ (2,3,5,6-tetrafluoro-7,7,8,8-tetracyanoquinodimethane, $E^{0'} = 0.16$ V),⁴² which is commonly used as an electron acceptor or hole dopant in molecule-based devices and materials,^{43,44} showing that $[(\text{HN})_2\text{-bisPyTM}]^{2+}$ functions as a superior electron acceptor (*i.e.*, oxidizing agent).

Next, we used $\text{B}(\text{C}_6\text{F}_5)_3$, which is a Lewis acid capable of binding to nitrogen atoms, as an external stimulus. Upon the addition of $\text{B}(\text{C}_6\text{F}_5)_3$, two isosbestic points ($\lambda = 371$ and 378 nm for 0–1 and 1–3 eq., respectively) were obtained in the UV/vis spectra. These two points can be assigned as the equilibrium of bisPyTM and one $\text{B}(\text{C}_6\text{F}_5)_3$ adduct, and that of one and two adducts, indicating the stepwise response of the two nitrogen atoms to $\text{B}(\text{C}_6\text{F}_5)_3$ (Fig. 10e). The emission characteristics of bisPyTM also showed a two-stage response to $\text{B}(\text{C}_6\text{F}_5)_3$. The

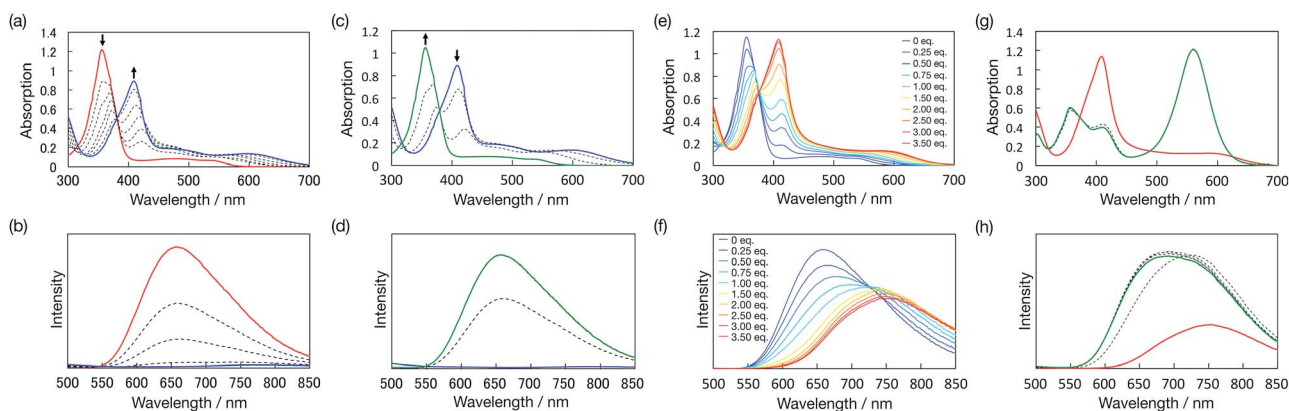


Fig. 10 (a) UV/vis absorption spectra of bisPyTM in CH_2Cl_2 following addition of TfOH: bisPyTM (red) + 0.5, 1.0, 1.5, 2.0, 2.5, and 3.0 (blue) eq. of TfOH. (b) Emission spectra of bisPyTM in CH_2Cl_2 following addition of TfOH ($\lambda_{\text{ex}} = 380$ nm, isosbestic point wavelength): bisPyTM (red) + 0.5, 1.0, 2.0, and 3.0 (blue) eq. of TfOH. (c) UV/vis absorption spectra and (d) emission spectra of bisPyTM in CH_2Cl_2 following addition of NEt_3 ($\lambda_{\text{ex}} = 380$ nm): bisPyTM + 3.0 eq. of TfOH (blue) + 1.0, 2.0, and 3.0 (green) eq. of NEt_3 . (e) UV/vis absorption spectra and (f) emission spectra of bisPyTM in CH_2Cl_2 with addition of $\text{B}(\text{C}_6\text{F}_5)_3$ ($\lambda_{\text{ex}} = 375$ nm, average wavelength of two isosbestic points): bisPyTM (blue) + 0.5, 1.0, 1.5, 2.0, 2.5, 3.0, and 3.5 (red) eq. of $\text{B}(\text{C}_6\text{F}_5)_3$. Fresh solutions were prepared for each measurement. (g) UV/vis absorption spectra and (h) emission spectra of bisPyTM in CH_2Cl_2 with addition of NEt_3 ($\lambda_{\text{ex}} = 375$ nm): bisPyTM + 3.5 eq. of $\text{B}(\text{C}_6\text{F}_5)_3$ (red) + 0, 0.5, 1, 2, 3 and 4 (green) eq. of NEt_3 . Fresh solutions were prepared for each measurement.



emission maximum wavelengths were gradually red-shifted upon the addition of $\text{B}(\text{C}_6\text{F}_5)_3$, and the resulting emission band at $\lambda = 756 \text{ nm}$ extended over 850 nm (Fig. 10f), suggesting the near-IR emission of $[(\text{C}_6\text{F}_5)_3\text{BN}]_2\text{-bisPyTM}$. The mechanism of the red shift is similar to that for $(\text{C}_6\text{F}_5)_3\text{BN-PyBTM}$.²⁵ Upon the addition of NEt_3 , the emission and UV/vis absorption spectra did not return to the initial states, indicating the incomplete recovery and partial decomposition of bisPyTM (Fig. 10g and h). These results show the partly reversible response of the radical to $\text{B}(\text{C}_6\text{F}_5)_3$ and NEt_3 .

Conclusions

We prepared bisPyTM, a luminescent stable organic radical that has two pyridyl moieties. Upon continuous UV light irradiation, it exhibited a half-life in dichloromethane that was 43 and 200 times longer than those of PyBTM and TIPS pentacene, respectively. The photostabilities of the radicals were greatly improved as the number of pyridyl groups in the TTM skeleton increased. We confirmed the luminescence of crystalline bisPyTM at 77 K , which would be the first example of visible solid-state emission from a radical. The two nitrogen atoms of bisPyTM can act as chemical stimulus-responsive sites and react with TFOH and $\text{B}(\text{C}_6\text{F}_5)_3$ to modulate the optical and electrochemical characteristics. The differences in the k_r and k_{nr} values of bisPyTM and PyBTM were reproduced through theoretical calculations based on DFT, TD-DFT, and VCD analyses. The VCDs indicated that the difference in the off-diagonal VCCs in the two radicals comes from the difference in the overlap density, which originates from the distribution of the NHOMOs.

Our results showed that introducing a pyridyl ring increases the photostability by lowering the energy levels of frontier orbitals. We also demonstrated that the scaled k_r and k_{nr} values, important parameters that determine the photophysical properties, can be estimated and compared in this class of radicals *via* theoretical calculations. These findings contribute greatly to predicting the photophysical properties of luminescent radicals, thereby achieving high emission efficiency and controlled photofunctionality.

Experimental

Materials

All chemicals were reagent grade and purchased commercially. Water was purified using an AUTOPURE WD500 system (Yamato Scientific Co., Ltd). The acetonitrile (MeCN), dichloromethane (CH_2Cl_2), diethyl ether (Et_2O), hexane, tetrahydrofuran (THF) and toluene used for syntheses were purified through an organic solvent purifier (Nikko Hansen Co., Ltd).

Equipment

NMR spectra were recorded with a Bruker US500 spectrometer at room temperature. Chemical shifts in ppm were referenced using tetramethylsilane (0.00 ppm) as an internal standard. Microwave assisted synthesis was performed using a Biotage Initiator+. UV-vis-NIR absorption spectra were recorded with

a JASCO V570 spectrophotometer. Steady-state emission spectra were recorded with HITACHI F-4500 and JASCO FP8600 spectrometers. Absolute photoluminescence quantum yields were measured using a Hamamatsu Photonics C9920-02G system. Fluorescence lifetime measurements were measured using a Hamamatsu Photonics QuantaTaurus-Tau C11367-02 system. ESR spectra were recorded with a JEOL JES-FA200 spectrometer. 4-Hydroxy-TEMPO was used as a standard for estimating the spin concentration. Observed g -values were calibrated with an $\text{Mn}^{2+}/\text{MgO}$ marker. Cyclic voltammetry was carried out under an argon atmosphere with a GC working electrode, a platinum wire counter electrode and an Ag^+/Ag reference electrode (10 mM AgClO_4 and $0.1 \text{ mM } ^t\text{Bu}_4\text{NClO}_4$ in MeCN solution), with an ALS-650DT voltammetric analyzer. The platinum working electrode was polished with an alumina abrasive and washed with water and acetone with ultrasonication. The counter electrode was heated with an oxygen flame to remove organic compounds on the surface and washed with acetone. The reference electrode was washed with acetone. A Fc^+/Fc redox couple was observed at 0.188 V under these experimental conditions. The potential of the voltammograms is shown relative to the Fc^+/Fc redox couple.

Single crystal X-ray crystallography

The data for single crystal X-ray diffraction analysis were collected at 113 K on a Rigaku AFC10 diffractometer, with a Rigaku Saturn CCD system equipped with a rotating-anode X-ray generator that emitted graphite-monochromated $\text{Mo K}\alpha$ radiation ($0.7107 \text{ \AA}$). A suitable single crystal was mounted on a looped film (micromount) with liquid paraffin. An empirical absorption correction, using equivalent reflections and Lorentzian polarization correction, was performed with the program Crystal Clear 1.3.6. The structure was solved using SIR-92⁴⁵ and the whole structure was refined against F^2 with SHELXL-97.⁴⁶ All non-hydrogen atoms were refined anisotropically. Hydrogen atoms were located in idealized positions and were refined using a riding model with fixed thermal parameters.

Computational details

DFT calculations were executed using the Gaussian 09 program package.⁴⁷ The geometries of the compounds were optimized with the grid = ultrafine option without symmetry constraints, using the crystal structure coordinates as the starting structure. Calculations were performed using the unrestricted Becke three-parameter hybrid functional with the Lee–Yang–Parr correlation functional (B3LYP)⁴⁸ with the 6-31G(d,p) basis set.⁴⁹ Frequency calculations were carried out to ensure that the optimized geometries were minima on the potential energy surface, in which no imaginary frequencies were observed in any of the compounds. TD-DFT calculations were performed using UB3LYP to calculate the first 15 doublet transitions. The solvent effect was considered using the PCM model.⁵⁰ VCCs and VCDs were calculated using our in-house codes.



Evaluation of the stabilities of bisPyTM, PyBTM and TIPS pentacene under UV light

A solution (*ca.* 5×10^{-6} M, 1.0 mL) in a 1 cm optical path length quartz cell was bubbled with argon, sealed, and set in a JASCO FP8600 spectrometer. The intensity of luminescence at 650 and 585 nm was monitored, exciting with 355 and 370 nm light (the excitation slit was 20 nm, and the shutter control was off), for bisPyTM and PyBTM, respectively. For TIPS pentacene, the intensity of luminescence at 650 nm was monitored, exciting with 310 (in dichloromethane), 300 (in acetonitrile) and 330 (in acetone) nm light. The logarithm of fluorescence intensity *versus* time was plotted and the slope of an approximate line was estimated to be the rate of photolysis.

Synthesis

Synthesis of bis(3,5-dichloro-4-pyridyl)(1,3,5-trichlorophenyl) methane (α H-bisPyTM): under an argon atmosphere, a mixture of bis(3,5-dichloro-4-pyridyl)methanol⁵¹ (7.78 g, 24.0 mmol), 1,3,5-trichlorobenzene (69.4 g, 382 mmol) and aluminum chloride (31.98 g, 240 mmol) was heated to 300 °C for 10 min *via* microwave irradiation. The resulting dark-brown mixture was cooled to room temperature, dissolved in CH₂Cl₂ and added to ice water. The mixture was neutralized using aqueous NaHCO₃ and the organic layer was separated. The water layer was extracted with CH₂Cl₂ (3 times). The organic layers were combined, dried with Na₂SO₄, and evaporated. The crude product was purified *via* silica gel column chromatography (eluent: ethyl acetate/hexane = 1/4, v/v) and recrystallization from CH₂Cl₂/hexane. Preparative GPC yielded the pure product as a pale yellow solid (96 mg, 1%). The steric hindrance of methylene carbon atoms, instability of the reaction intermediate (methyl cation) and lowered electron densities of pyridine rings upon coordination to aluminum chloride can cause this low yield. The major product is bis(3,5-dichloro-4-pyridyl)methane, which would be produced by the decomposition of the starting material due to the severe reaction temperature, whereas the target compound was not formed at lower reaction temperatures. ¹H NMR (CDCl₃, 500 MHz): δ = 6.70 (s, 1H), 7.28 (d, 1H, *J* = 2.5 Hz), 7.41 (d, 1H, *J* = 2.5 Hz), 8.37 (s, 1H), 8.40 (s, 1H), 8.51 (s, 1H), 8.52 (s, 1H). GC-MS: *m/z* = 486 ([C₁₇H₇Cl₇N₂]⁺). Elem. anal. calcd for C₁₇H₆Cl₇N₂: C 41.89, H 1.45, N 5.75; found: C 41.67, H 1.80, N 5.67.

Synthesis of the bis(3,5-dichloro-4-pyridyl)(1,3,5-trichlorophenyl)methyl radical (bisPyTM):¹⁴ under a nitrogen atmosphere, α H-bisPyTM (31.0 mg, 63.8 μ mol) was dissolved in dry THF (3.0 mL). A 1 M solution of ^tBuOK in THF (130 μ L) was added dropwise, and the color of the solution changed to red. The reaction mixture was stirred overnight in the dark. I₂ (121 mg, 476 μ mol) in dry diethyl ether (10.8 mL) was added dropwise and stirred for 2.25 h. The remaining I₂ was reduced by washing with aq. 10% Na₂S₂O₃ 3 times. The water layer was extracted with diethyl ether once, and the combined organic layer was dried with Na₂SO₄. The red solution was filtered, evaporated, purified *via* Al₂O₃ column chromatography (eluent: diethyl ether/hexane = 1/4, v/v) and dried *in vacuo* to afford bisPyTM (28.1 mg, 91%) as a red solid. HRMS (negative ion

mode ESI-TOF) *m/z*: [M][−]. Calcd for C₁₇H₆Cl₇N₂: 484.8322; found 484.8303. Elem. anal. calcd for C₁₇H₆Cl₇N₂: C 41.98, H 1.24, N 5.76; found: C 41.73, H 1.83, N 5.31. ESR: the spin concentration of bisPyTM in toluene (1.2×10^{-5} M) was estimated by comparing the value of the twice-integration of the signal intensity with that of a reference sample (4-hydroxy-TEMPO in toluene; 1.5×10^{-5} M). The existence of *S* = 1/2 spin on one bisPyTM molecule was confirmed.

Conflicts of interest

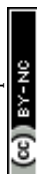
There are no conflicts to declare.

Acknowledgements

The present study was supported by JST CREST grant number JPMJCR15F2. This work was also supported by JSPS KAKENHI grant numbers JP26220801, JP16K13973, JP17H04870, and JP15K05607. T. K. is grateful to the Ogasawara Foundation for the Promotion of Science and Engineering and the Shorai Foundation for Science for financial support. S. K. acknowledges MERIT (the material education program for future leaders in Research, Industry, and Technology) in the MEXT Leading Graduate School Doctoral Program. This research was conducted using the HITACHI SR16000 system (yayoi) in the Information Technology Center, The University of Tokyo.

Notes and references

- 1 C. W. Tang and S. A. Vanslyke, *Appl. Phys. Lett.*, 1987, **51**, 913–915.
- 2 M. A. Baldo, D. F. O'Brien, Y. You, A. Shoustikov, S. Sibley, M. E. Thompson and S. R. Forrest, *Nature*, 1998, **395**, 151–154.
- 3 A. Endo, M. Ogasawara, A. Takahashi, D. Yokoyama, Y. Kato and C. Adachi, *Adv. Mater.*, 2009, **21**, 4802–4806.
- 4 H. Uoyama, K. Goushi, K. Shizu, H. Nomura and C. Adachi, *Nature*, 2012, **492**, 234–238.
- 5 V. Gamero, D. Velasco, S. Latorre, F. López-Calahorra, E. Brillas and L. Juliá, *Tetrahedron Lett.*, 2006, **47**, 2305–2309.
- 6 D. Velasco, S. Castellanos, M. López, F. López-Calahorra, E. Brillas and L. Juliá, *J. Org. Chem.*, 2007, **72**, 7523–7532.
- 7 L. Fajari, R. Papoular, M. Reig, E. Brillas, J. L. Jorda, O. Vallcorba, J. Rius, D. Velasco and L. Juliá, *J. Org. Chem.*, 2014, **79**, 1771–1777.
- 8 A. Heckmann, S. Dümmler, J. Pauli, M. Margraf, J. Köhler, D. Stich, C. Lambert, I. Fischer and U. Resch-genger, *J. Phys. Chem. C*, 2009, **113**, 20958–20966.
- 9 E. V. Tretyakov, V. F. Plyusnin, A. O. Suvorova, S. V. Larionov, S. A. Popov, O. V. Antonova, E. M. Zueva, D. V. Stass, A. S. Bogomyakov, G. V. Romanenko and V. I. Ovcharenko, *J. Lumin.*, 2014, **148**, 33–38.
- 10 Y. Gao, A. Obolda, M. Zhang and F. Li, *Dyes Pigm.*, 2017, **139**, 644–650.
- 11 E. Neier, R. Arias, N. Rady, S. Venkatesan, T. W. Hudnall and A. Zakhidov, *Org. Electron.*, 2017, **44**, 126–131.



- 12 Y. Beldjoudi, I. Osorio-Román, M. A. Nascimento and J. M. Rawson, *J. Mater. Chem. C*, 2017, **5**, 2794–2799.
- 13 H. Namai, H. Ikeda, Y. Hoshi, N. Kato, Y. Morishita and K. Mizuno, *J. Am. Chem. Soc.*, 2007, **129**, 9032–9036.
- 14 Y. Hattori, T. Kusamoto and H. Nishihara, *Angew. Chem., Int. Ed.*, 2014, **53**, 11845–11848.
- 15 Q. Peng, A. Obolda, M. Zhang and F. Li, *Angew. Chem., Int. Ed.*, 2015, **54**, 7091–7095.
- 16 A. Obolda, X. Ai, M. Zhang and F. Li, *ACS Appl. Mater. Interfaces*, 2016, **8**, 35472–35478.
- 17 M. A. Fox, E. Gaillard and C. Chen, *J. Am. Chem. Soc.*, 1987, **109**, 7088–7094.
- 18 S. R. Ruberu and M. A. Fox, *J. Phys. Chem.*, 1993, **97**, 143–149.
- 19 M. Sakamoto, X. Cai, M. Hara, S. Tojo, M. Fujitsuka and T. Majima, *J. Am. Chem. Soc.*, 2005, **127**, 3702–3703.
- 20 J. C. Scaiano, L. J. Johnston, W. G. McGimpsey and D. Weir, *Acc. Chem. Res.*, 1988, **21**, 22–29.
- 21 D. Meisel, P. K. Das, G. L. Hug, K. Bhattacharyya and R. W. Fessenden, *J. Am. Chem. Soc.*, 1986, **108**, 4706–4710.
- 22 P. Brodard, A. Sarbach, J. Gumy, T. Bally and E. Vauthey, *J. Phys. Chem. A*, 2001, **105**, 6594–6601.
- 23 M. Ballester, J. Riera, J. Castatier, C. Badfa and J. M. Monsó, *J. Am. Chem. Soc.*, 1971, **93**, 2215–2225.
- 24 O. Armet, J. Veciana, C. Rovira, J. Riera, J. Castaner, E. Molins, J. Rius, C. Miravittles, S. Olivella and J. Brichfeus, *J. Phys. Chem.*, 1987, **91**, 5608–5616.
- 25 T. Kusamoto, S. Kimura, Y. Ogino, C. Ohde and H. Nishihara, *Chem.–Eur. J.*, 2016, **22**, 17725–17733.
- 26 Y. Hattori, T. Kusamoto and H. Nishihara, *RSC Adv.*, 2015, **5**, 64802–64805.
- 27 Y. Hattori, T. Kusamoto and H. Nishihara, *Angew. Chem., Int. Ed.*, 2015, **54**, 3731–3734.
- 28 Y. Hattori, T. Kusamoto, T. Sato and H. Nishihara, *Chem. Commun.*, 2016, **440**, 908–912.
- 29 Y. Ogino, T. Kusamoto, Y. Hattori, M. Shimada, M. Tsuchiya, Y. Yamanoi, E. Nishibori, K. Sugimoto and H. Nishihara, *Inorg. Chem.*, 2017, **56**, 3909–3915.
- 30 H. Iwamura and N. Koga, *Pure Appl. Chem.*, 1999, **71**, 231–238.
- 31 T. K. Maji, R. Matsuda and S. Kitagawa, *Nat. Mater.*, 2007, **6**, 142–148.
- 32 J. Grilj, E. N. Laricheva, M. Olivucci and E. Vauthey, *Angew. Chem., Int. Ed.*, 2011, **50**, 4496–4498.
- 33 M. Uejima, T. Sato, D. Yokoyama, K. Tanaka and J.-W. Park, *Phys. Chem. Chem. Phys.*, 2014, **16**, 14244–14256.
- 34 D. T. Breslin and M. A. Fox, *J. Phys. Chem.*, 1994, **98**, 408–411.
- 35 Y. Kameoka, M. Uebe, A. Ito, T. Sato and K. Tanaka, *Chem. Phys. Lett.*, 2014, **615**, 44–49.
- 36 M. Uebe, A. Ito, Y. Kameoka, T. Sato and K. Tanaka, *Chem. Phys. Lett.*, 2015, **633**, 190–194.
- 37 Emission spectra of radicals in the solid state at 5 K, very low temperature, were reported. (a) R. Beaulac, D. Luneau and C. Reber, *Chem. Phys. Lett.*, 2005, **405**, 153–158; (b) R. Beaulac, G. Bussière, C. Reber, C. Lescop and D. Luneau, *New J. Chem.*, 2003, **27**, 1200–1206.
- 38 D. Blasi, D. M. Nikolaidou, F. Terenziani, I. Ratera and J. Veciana, *Phys. Chem. Chem. Phys.*, 2017, **19**, 9313–9319.
- 39 A. Shimizu, A. Ito and Y. Teki, *Chem. Commun.*, 2016, **52**, 2889–2892.
- 40 A. Maliakal, K. Raghavachari, H. Katz, E. Chandross and T. Siegrist, *Chem. Mater.*, 2004, **16**, 4980–4986.
- 41 I. Kaur, W. Jia, R. P. Kopreski, S. Selvarasah, M. R. Dokmeci, C. Pramanik, N. E. McGruer and G. P. Miller, *J. Am. Chem. Soc.*, 2008, **130**, 16274–16286.
- 42 M. Kivala, C. Boudon, J. P. Gisselbreeht, B. Enko, P. Seiler, B. M. Imke, N. Langer, P. D. Jarowski, G. Geseheidt and F. Diederich, *Chem.–Eur. J.*, 2009, **15**, 4111–4123.
- 43 M. Pfeiffer, K. Leo, X. Zhou, J. S. Huang, M. Hofmann, A. Werner and J. Blochwitz-Nimoth, *Org. Electron.*, 2003, **4**, 89–103.
- 44 K. Walzer, B. Männig, M. Pfeiffer and K. Leo, *Chem. Rev.*, 2007, **107**, 1233–1271.
- 45 A. Altomare, G. Cascarano, C. Giacovazzo, A. Guagliardi, M. C. Burla, G. Polidori and M. Camalli, *J. Appl. Crystallogr.*, 1994, **27**, 435.
- 46 G. M. Sheldrick, *Acta Crystallogr., Sect. A: Found. Crystallogr.*, 2008, **64**, 112–122.
- 47 M. J. Frisch, G. W. Trucks, H. B. Schlegel, G. E. Scuseria, M. A. Robb, J. R. Cheeseman, G. Scalmani, V. Barone, B. Mennucci, G. A. Petersson, H. Nakatsuji, M. Caricato, X. Li, H. P. Hratchian, A. F. Izmaylov, J. Bloino, G. Zheng, J. L. Sonnenberg, M. Hada, M. Ehara, K. Toyota, R. Fukuda, J. Hasegawa, M. Ishida, T. Nakajima, Y. Honda, O. Kitao, H. Nakai, T. Vreven, J. A. Montgomery Jr, J. E. Peralta, F. Ogliaro, M. Bearpark, J. J. Heyd, E. Brothers, K. N. Kudin, V. N. Staroverov, T. Keith, R. Kobayashi, J. Normand, K. Raghavachari, A. Rendell, J. C. Burant, S. S. Iyengar, J. Tomasi, M. Cossi, N. Rega, J. M. Millam, M. Klene, J. E. Knox, J. B. Cross, V. Bakken, C. Adamo, J. Jaramillo, R. Gomperts, R. E. Stratmann, O. Yazyev, A. J. Austin, R. Cammi, C. Pomelli, J. W. Ochterski, R. L. Martin, K. Morokuma, V. G. Zakrzewski, G. A. Voth, P. Salvador, J. J. Dannenberg, S. Dapprich, A. D. Daniels, Ö. Farkas, J. B. Foresman, J. V. Ortiz, J. Cioslowski and D. J. Fox, *Gaussian 09, Revision C.01*, Gaussian, Inc., Wallingford CT, 2010.
- 48 A. D. Becke, *J. Chem. Phys.*, 1993, **98**, 5648–5652.
- 49 P. C. Hariharan and J. A. Pople, *Theor. Chim. Acta*, 1973, **28**, 213–222.
- 50 J. Tomasi, B. Mennucci and R. Cammi, *Chem. Rev.*, 2005, **105**, 2999–3093.
- 51 T. Itoh, A. Takada, K. Hirai and H. Tomioka, *Org. Lett.*, 2005, **7**, 811–814.

