

Electronic Supporting Information

Effect of *N*-alkyl chain length on photophysical properties of indene-1,3-dionemethylene-1,4-dihydropyridine derivatives

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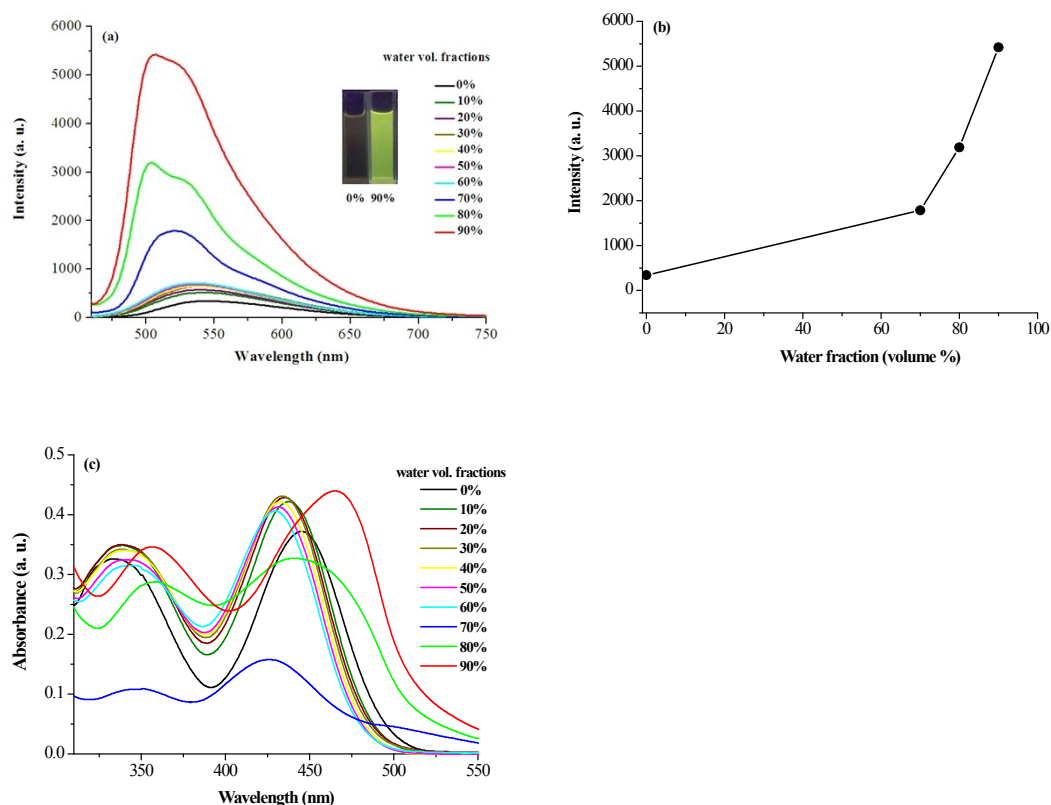


Fig. S1 (a) Fluorescence spectra of **4b** at 10 μM in THF/water mixtures with different f_w values ($\lambda_{ex} = 440$ nm). Insets: fluorescence images with water contents of 0% and 90% under a 365 nm UV lamp. (b) The effect of f_w values ($f_w = 0, 70\%, 80\%$, and 90%) on the maximum emission intensity of **4b** in THF/water mixtures at a concentration of 10 μM. (c) Absorption spectra of **4b** at 10 μM in THF/water mixtures with f_w values.

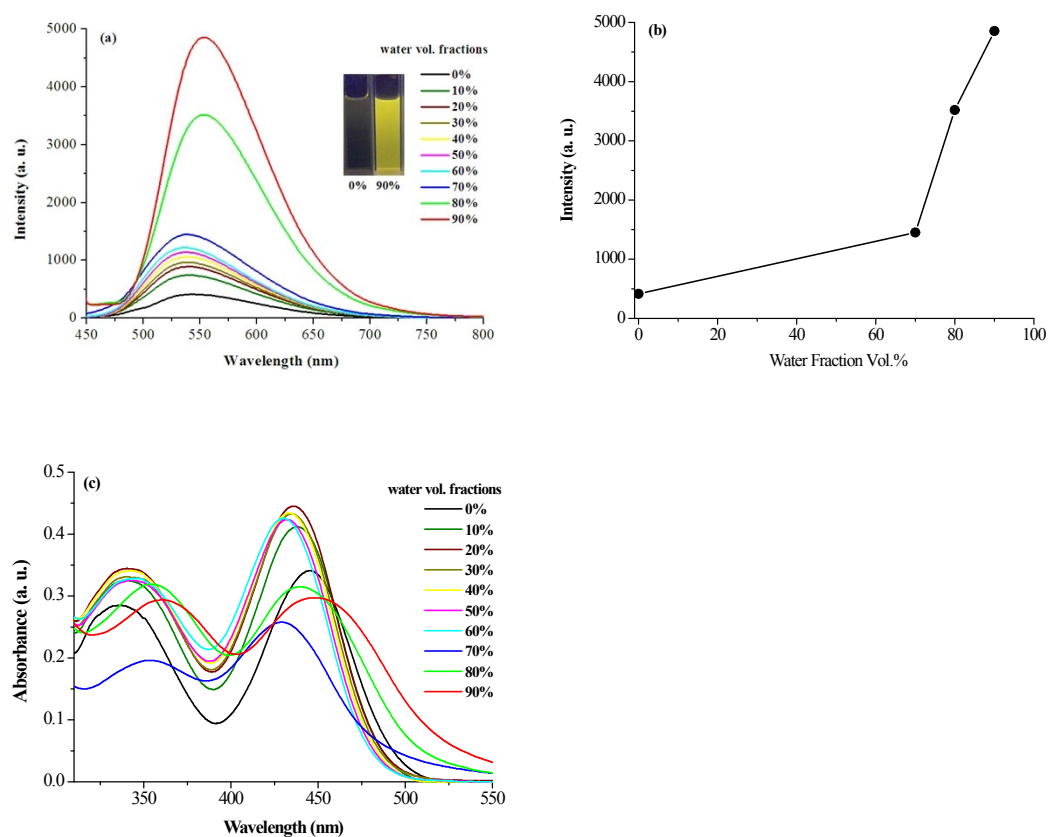


Fig. S2 (a) Fluorescence spectra of **4c** at 10 μM in THF/water mixtures with different f_w values ($\lambda_{\text{ex}} = 430 \text{ nm}$). Insets: fluorescence images with water contents of 0% and 90% under a 365 nm UV lamp. (b) The effect of f_w values ($f_w = 0, 70\%, 80\%, \text{ and } 90\%$) on the maximum emission intensity of **4c** in THF/water mixtures at a concentration of 10 μM . (c) Absorption spectra of **4c** at 10 μM in THF/water mixtures with f_w values.

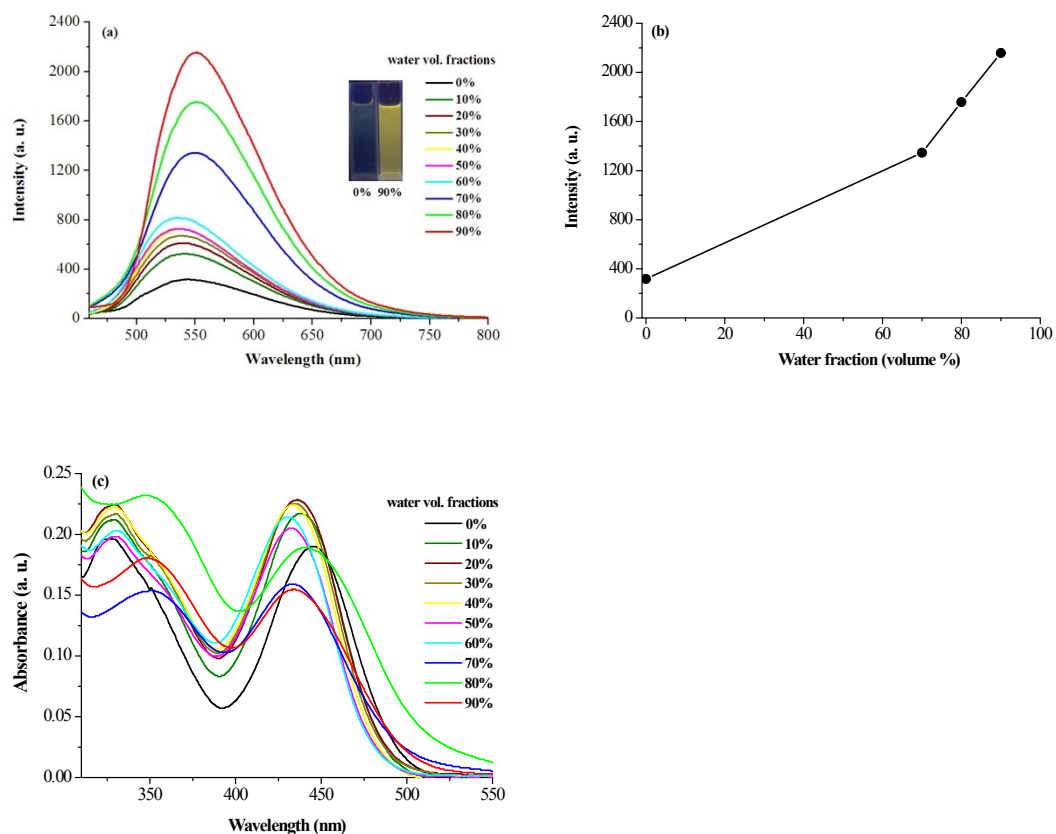


Fig. S3 (a) Fluorescence spectra of **4d** at 10 μM in THF/water mixtures with different f_w values ($\lambda_{\text{ex}} = 440$ nm). Insets: fluorescence images with water contents of 0% and 90% under a 365 nm UV lamp. (b) The effect of f_w values ($f_w = 0, 70\%, 80\%$, and 90%) on the maximum emission intensity of **4d** in THF/water mixtures at a concentration of 10 μM . (c) Absorption spectra of **4d** at 10 μM in THF/water mixtures with f_w values.

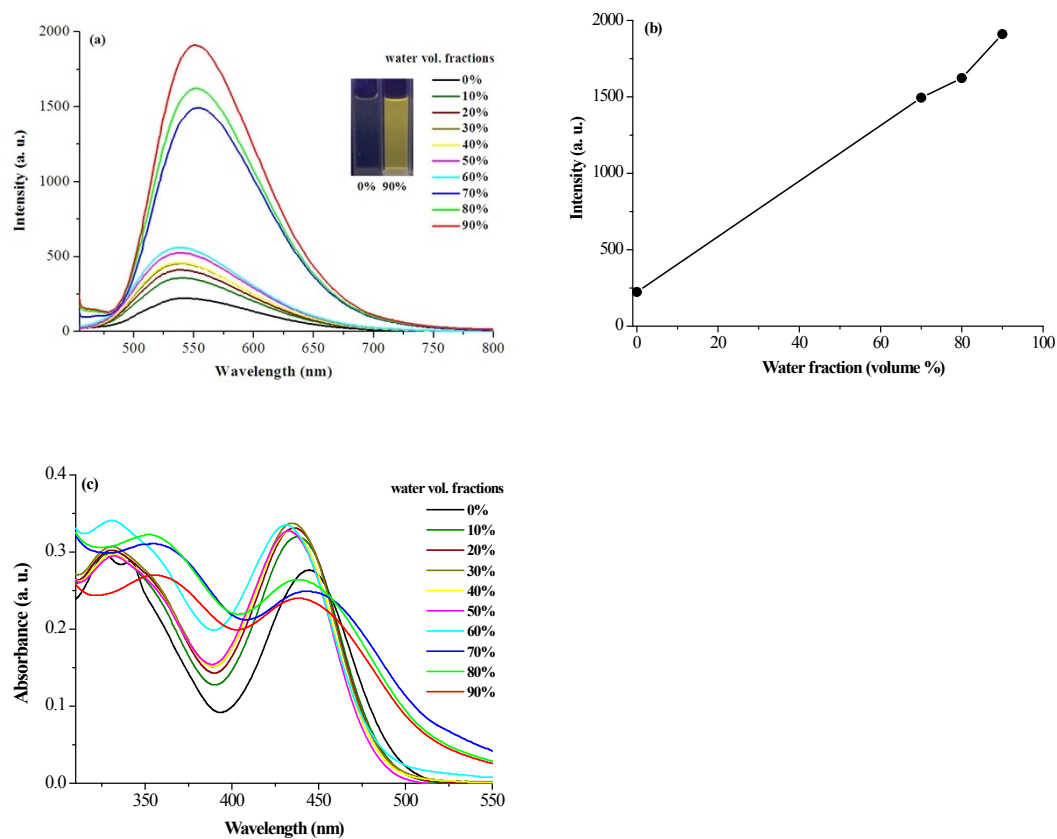


Fig. S4 (a) Fluorescence spectra of **4e** at 10 μM in THF/water mixtures with different f_w values ($\lambda_{\text{ex}} = 435 \text{ nm}$). Insets: fluorescence images with water contents of 0% and 90% under a 365 nm UV lamp. (b) The effect of f_w values ($f_w = 0, 70\%, 80\%, \text{ and } 90\%$) on the maximum emission intensity of **4e** in THF/water mixtures at a concentration of 10 μM . (c) Absorption spectra of **4e** at 10 μM in THF/water mixtures with f_w values.

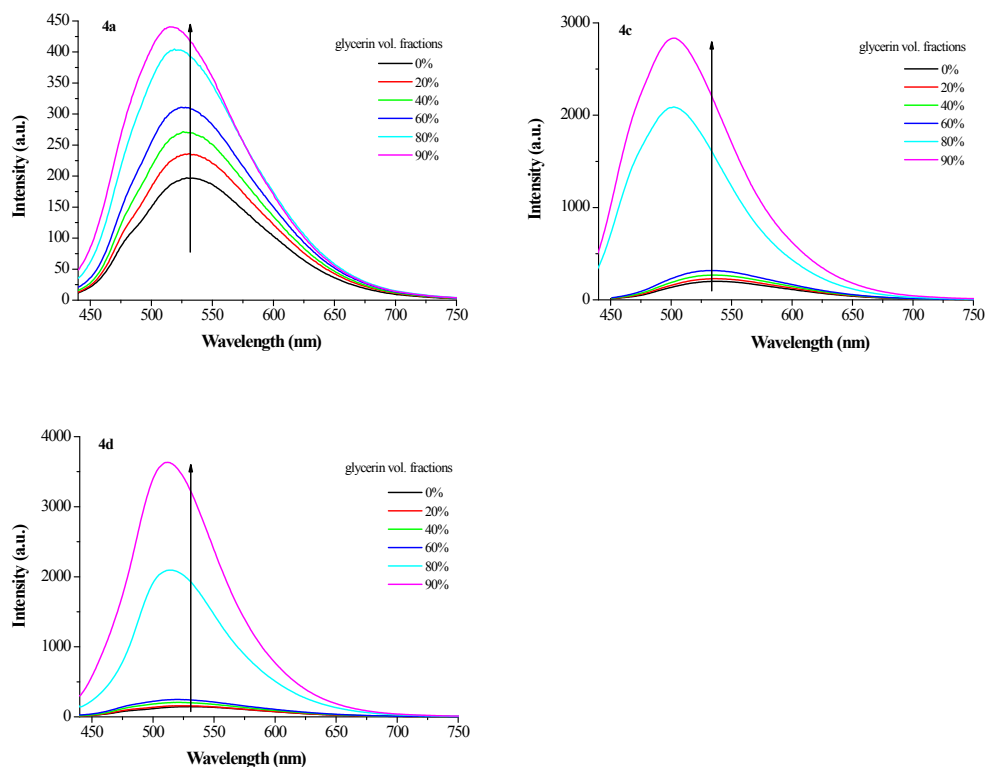


Fig. S5 Fluorescence spectra of **4a** ($\lambda_{\text{ex}} = 420$ nm), **4c** ($\lambda_{\text{ex}} = 440$ nm), and **4d** ($\lambda_{\text{ex}} = 420$ nm) in methanol/glycerol mixtures (10 μM , containing 0.5 vol % THF) with different glycerol volume fractions.

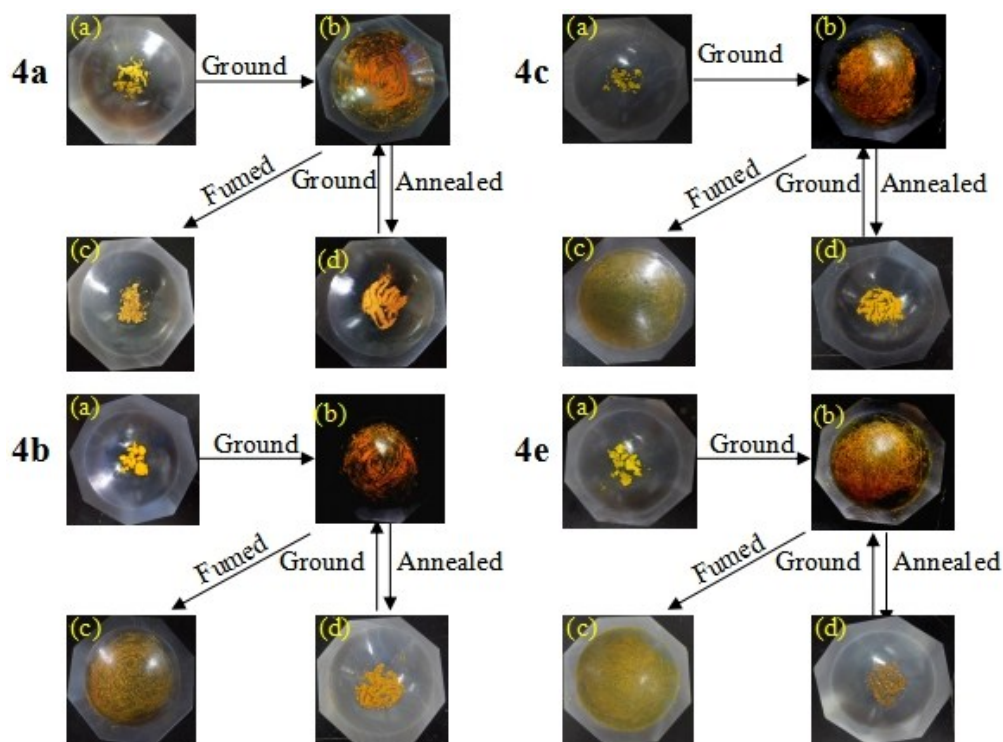


Fig. S6 Images of **4a–4c** and **4e** solid samples taken under natural light. (a) as-synthesized samples; (b) ground samples; (c) fumed samples using EA vapor; (d) annealed samples.

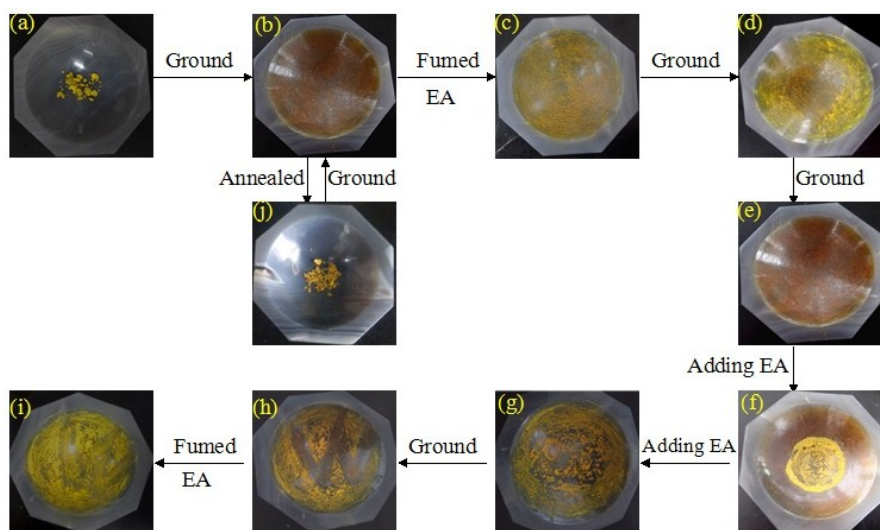


Fig. S7 Images of solid samples of **4d** taken under natural light: (a) as-synthesized samples; (b) ground samples; (c) fumed samples; (d) the central part of fumed samples was ground; (e) all fumed samples were ground; (f) several drops of EA were dropped onto the ground samples; (g) all ground samples were soaked with EA; (h) and (i) **4d** is used to write “W” with a metal spatula and then fumed using EA; (j) annealed samples.

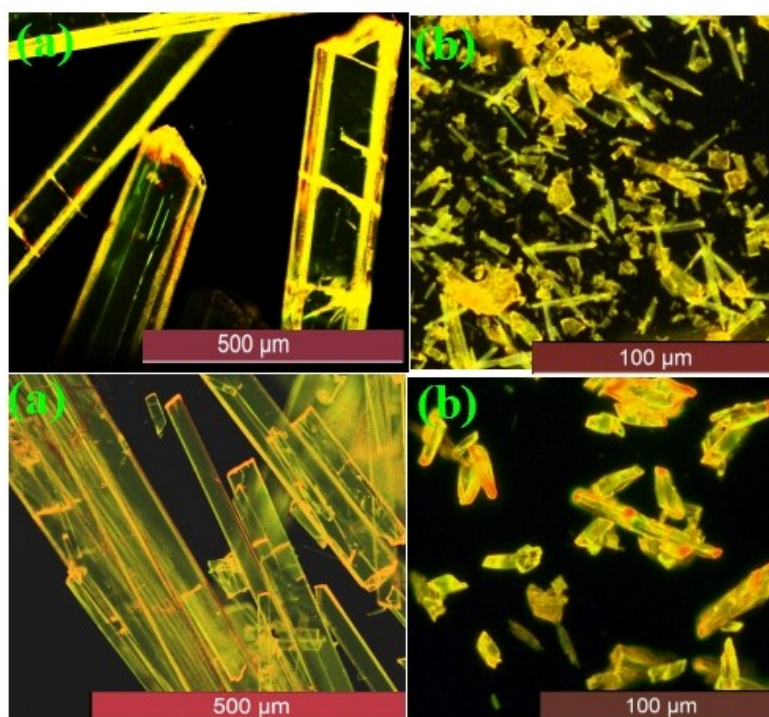


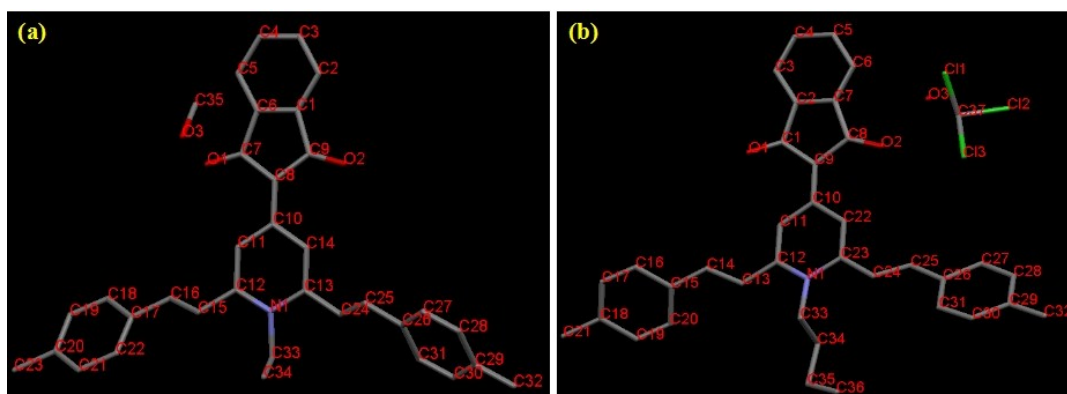
Fig. S8 The fluorescence microscope images of **4a** (top) and **4b** (bottom): (a) single crystals; (b) as-synthesized samples.

Table S1 Selected torsion angles [deg] for 3b and 3c

3b		3c	
C(12)-N(1)-C(17)-C(18)	87.75(12)	C(12)-N(1)-C(17)-C(18)	88.7(2)
C(13)-N(1)-C(17)-C(18)	-89.57(12)	C(13)-N(1)-C(17)-C(18)	-90.1(2)

Table S2 Selected torsion angles [deg] for 4a and 4b

4a		4b	
C(12)-N(1)-C(33)-C(34)	-101.2(4)	C(12)-N(1)-C(33)-C(34)	101.3(4)
C(13)-N(1)-C(33)-C(34)	82.6(4)	C(23)-N(1)-C(33)-C(34)	-92.3(4)
N(1)-C(13)-C(24)-C(25)	-143.8(3)	C(22)-C(23)-C(24)-C(25)	-26.2(5)
C(14)-C(13)-C(24)-C(25)	38.4(4)	N(1)-C(23)-C(24)-C(25)	155.8(3)
C(24)-C(25)-C(26)-C(31)	6.0(5)	C(24)-C(25)-C(26)-C(27)	-179.9(4)
C(24)-C(25)-C(26)-C(27)	-174.9(3)	C(24)-C(25)-C(26)-C(31)	-5.2(6)
C(11)-C(12)-C(15)-C(16)	28.8(4)	C(11)-C(12)-C(13)-C(14)	-28.6(5)
N(1)-C(12)-C(15)-C(16)	-152.0(3)	N(1)-C(12)-C(13)-C(14)	154.8(3)
C(15)-C(16)-C(17)-C(22)	4.0(5)	C(13)-C(14)-C(15)-C(16)	176.3(3)
C(15)-C(16)-C(17)-C(18)	-177.2(3)	C(13)-C(14)-C(15)-C(20)	-5.7(6)

**Fig. S9** Single-crystal structure of **4a** (a) with a CH₃OH molecule and **4b** (b) containing a CHCl₃ molecule and a water molecule. Hydrogen atoms are omitted for clarity.

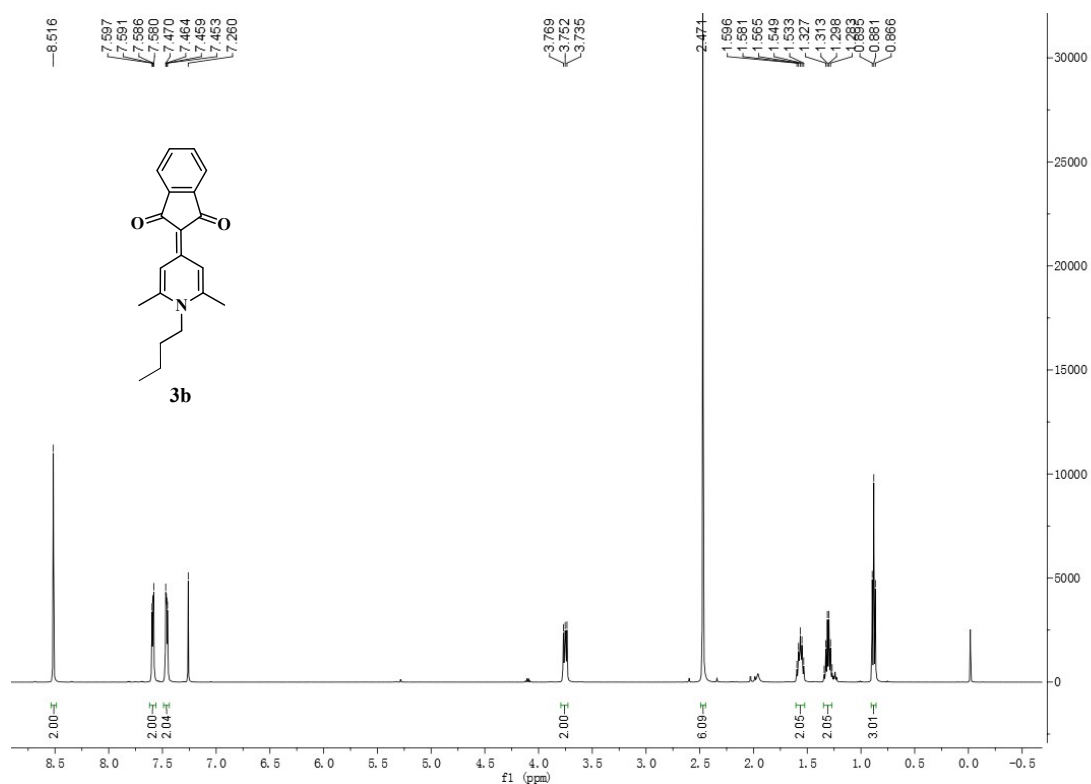


Fig. S10 ¹H NMR of **3b** (CDCl₃, 500 MHz).

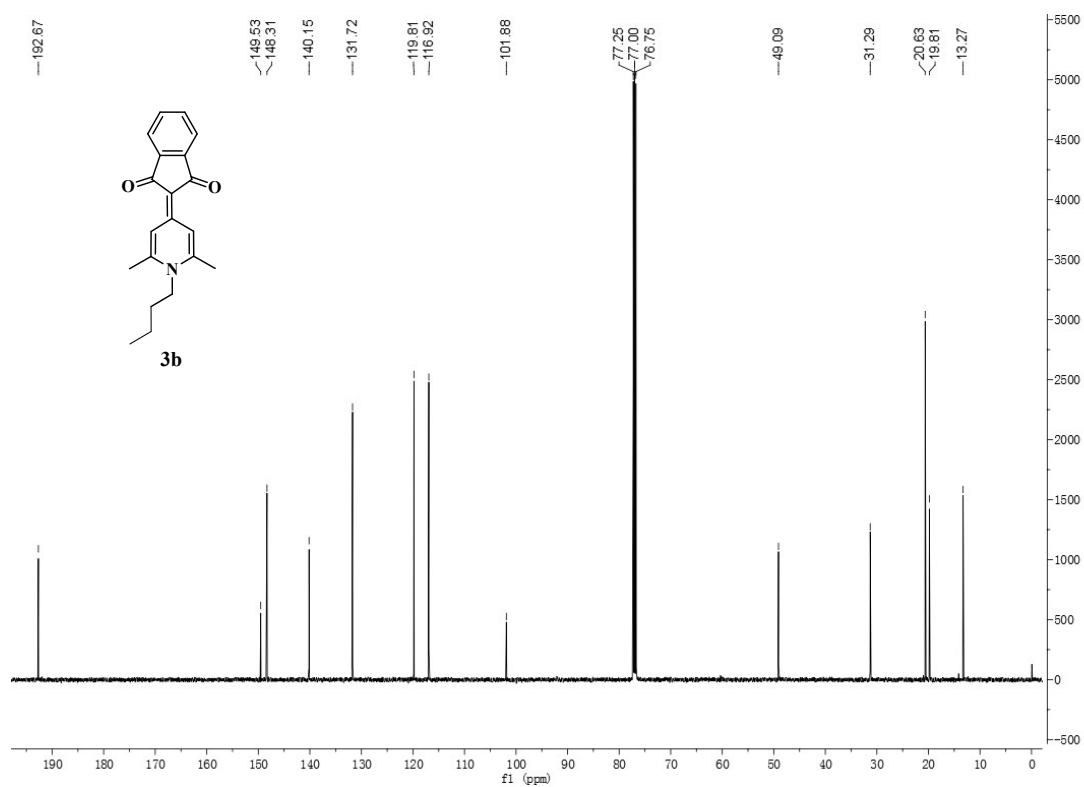


Fig. S11 ¹³C NMR of **3b** (CDCl₃, 125 MHz).

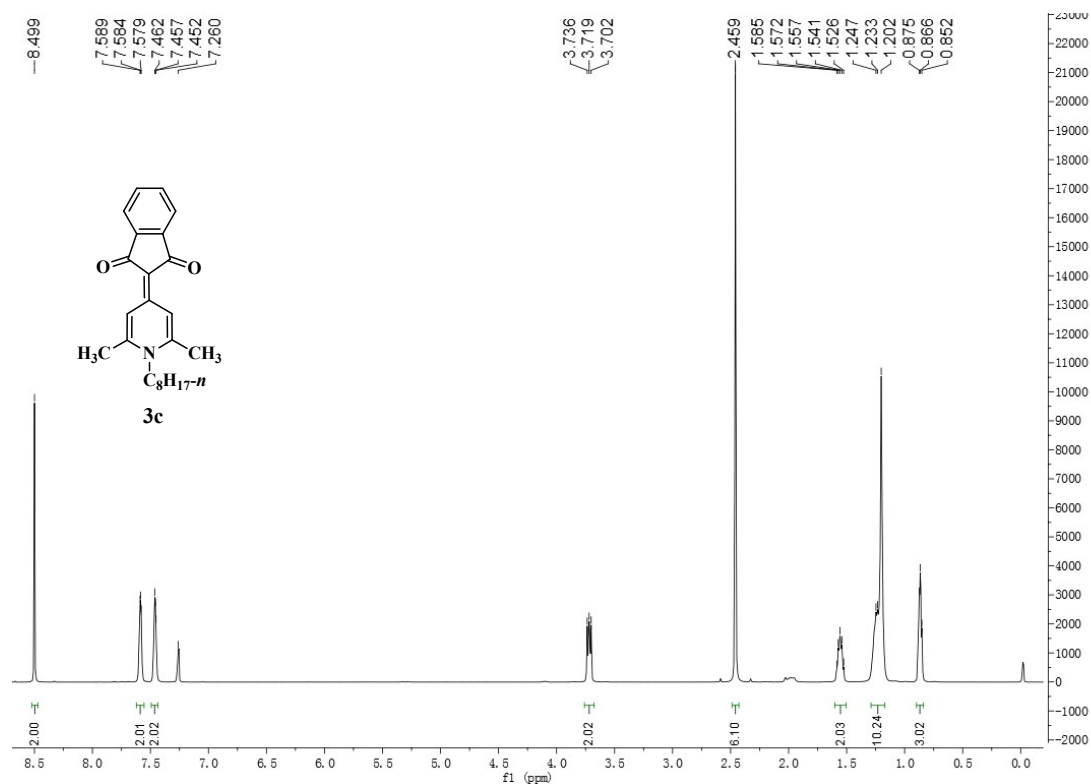


Fig. S12 ¹H NMR of **3c** (CDCl₃, 500 MHz).

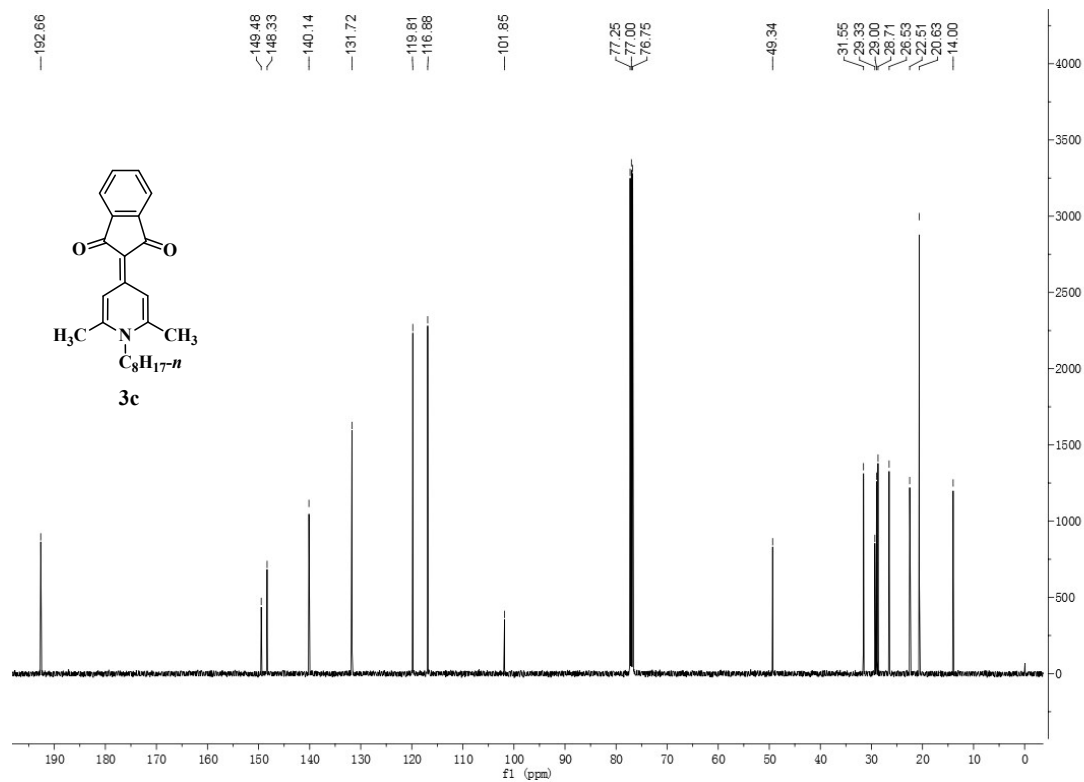


Fig. S13 ¹³C NMR of **3c** (CDCl₃, 125 MHz).

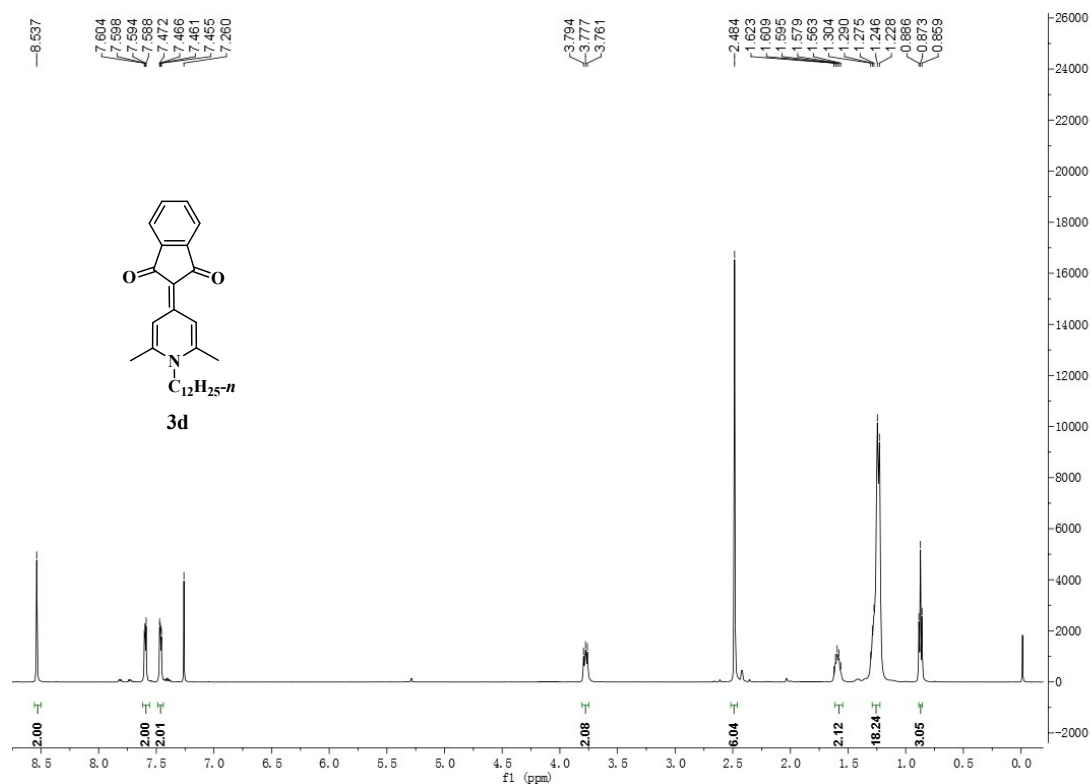


Fig. S14 ¹H NMR of **3d** (CDCl₃, 500 MHz).

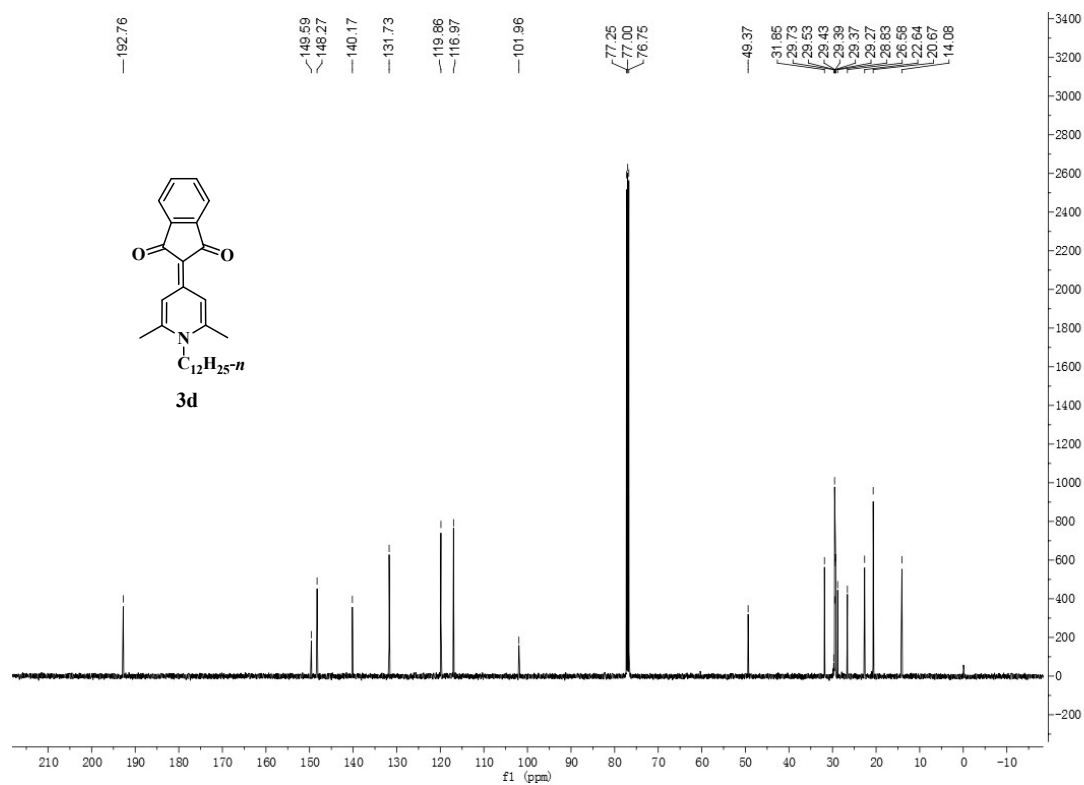


Fig. S15 ¹³C NMR of **3d** (CDCl₃, 125 MHz).

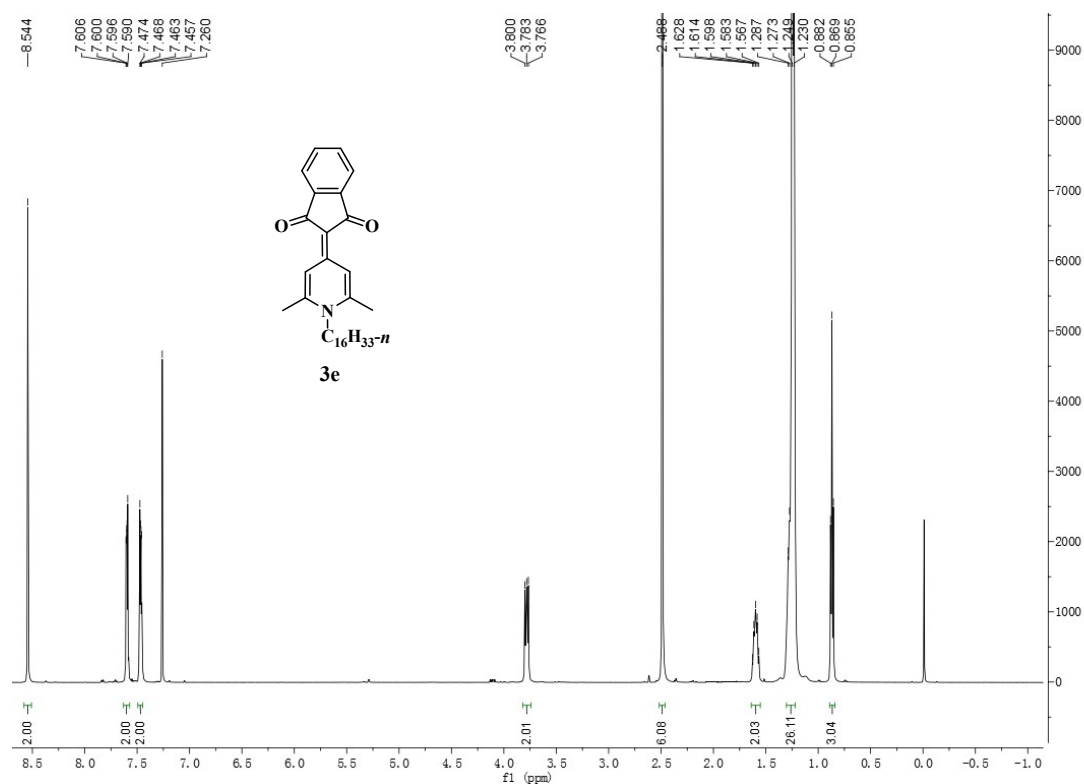


Fig. S16 1H NMR of **3e** ($CDCl_3$, 500 MHz).

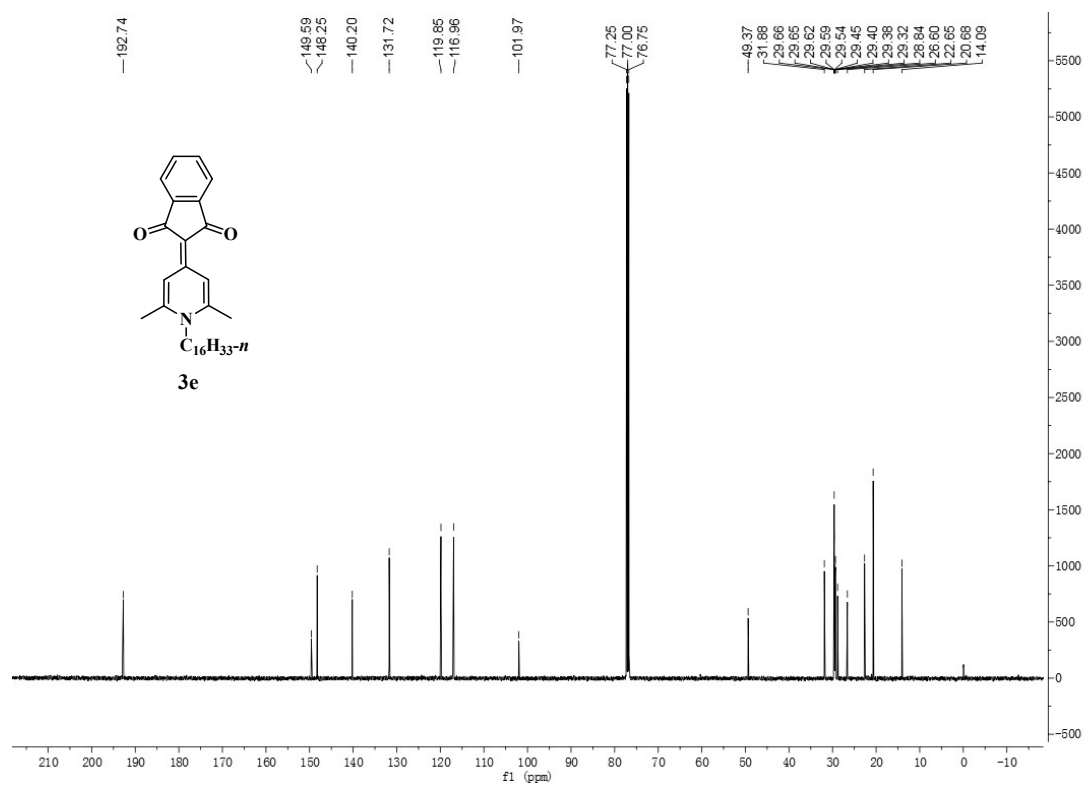


Fig. S17 ^{13}C NMR of **3e** ($CDCl_3$, 125 MHz).

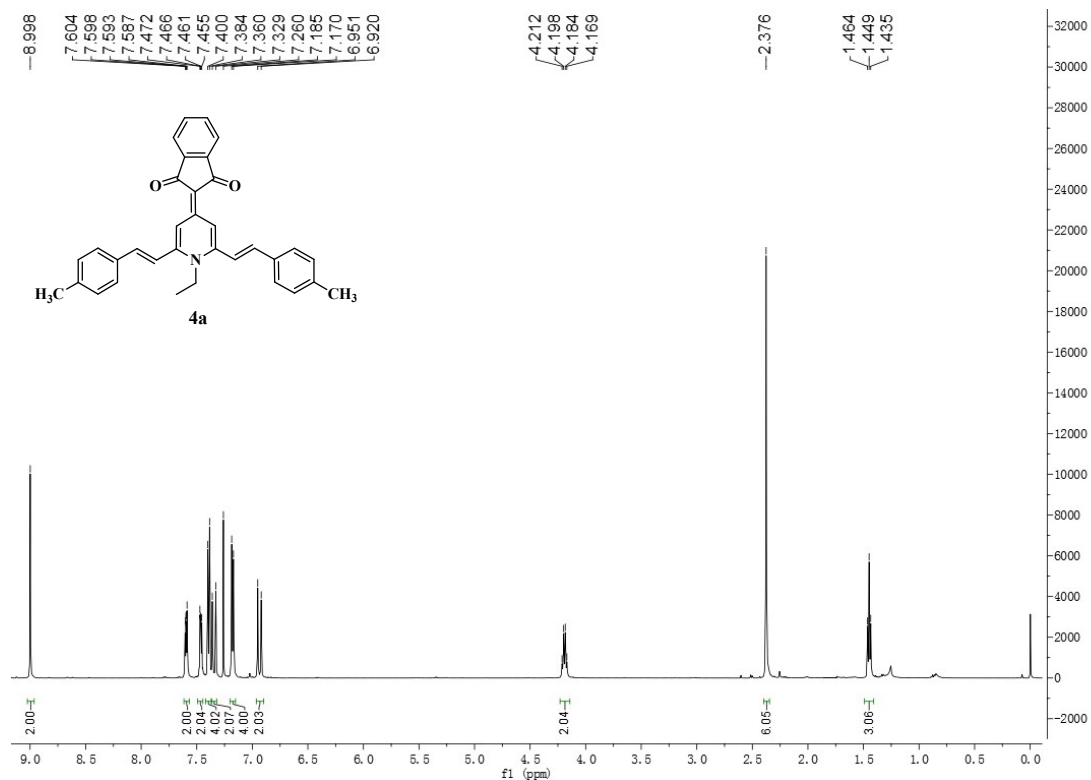


Fig. S18 ^1H NMR of **4a** (CDCl₃, 500 MHz).

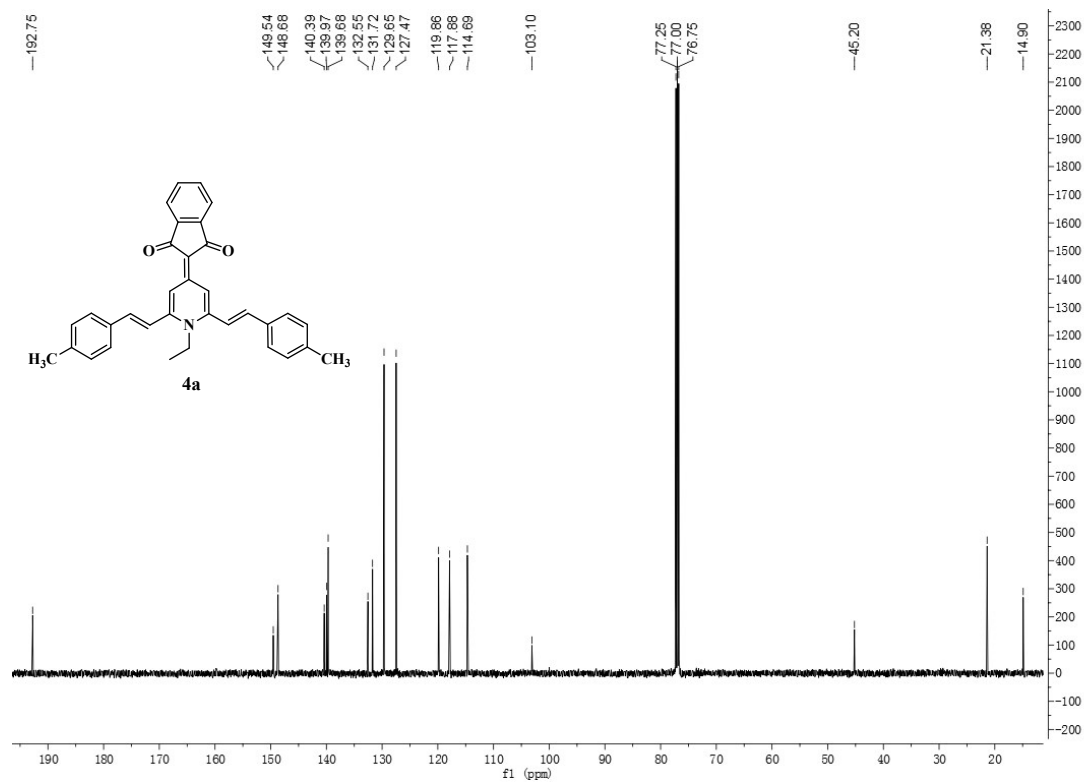


Fig. S19 ^{13}C NMR of **4a** (CDCl₃, 125 MHz).

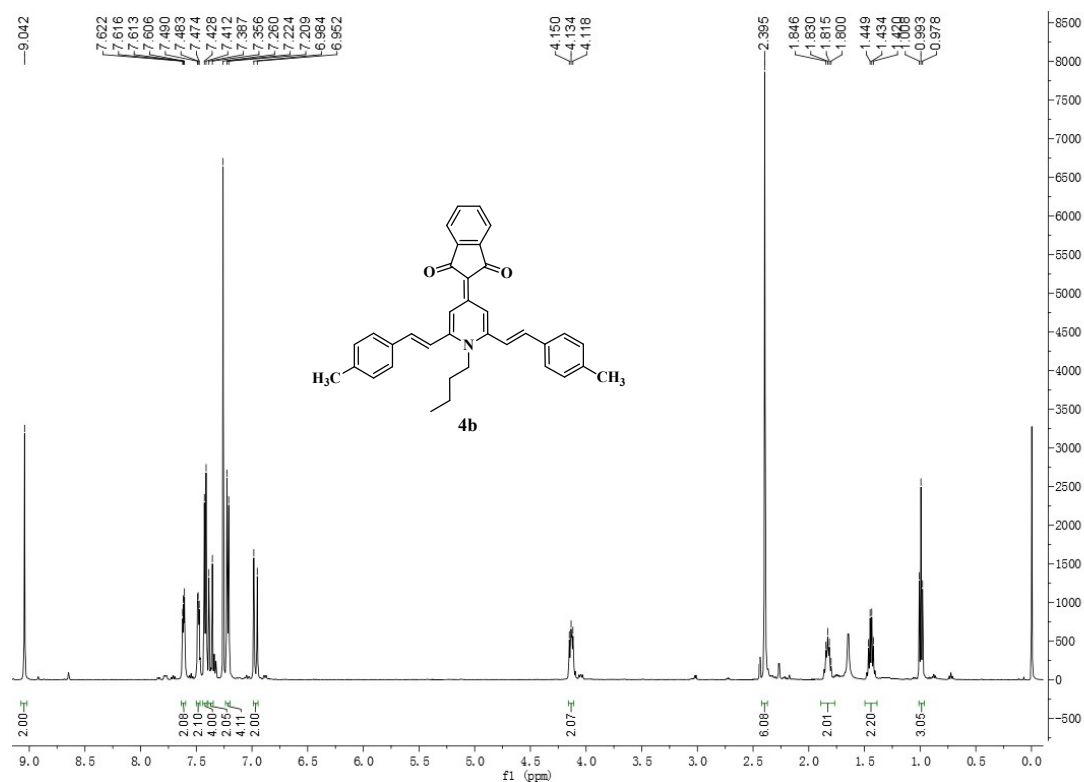


Fig. S20 ^1H NMR of **4b** (CDCl_3 , 500 MHz).

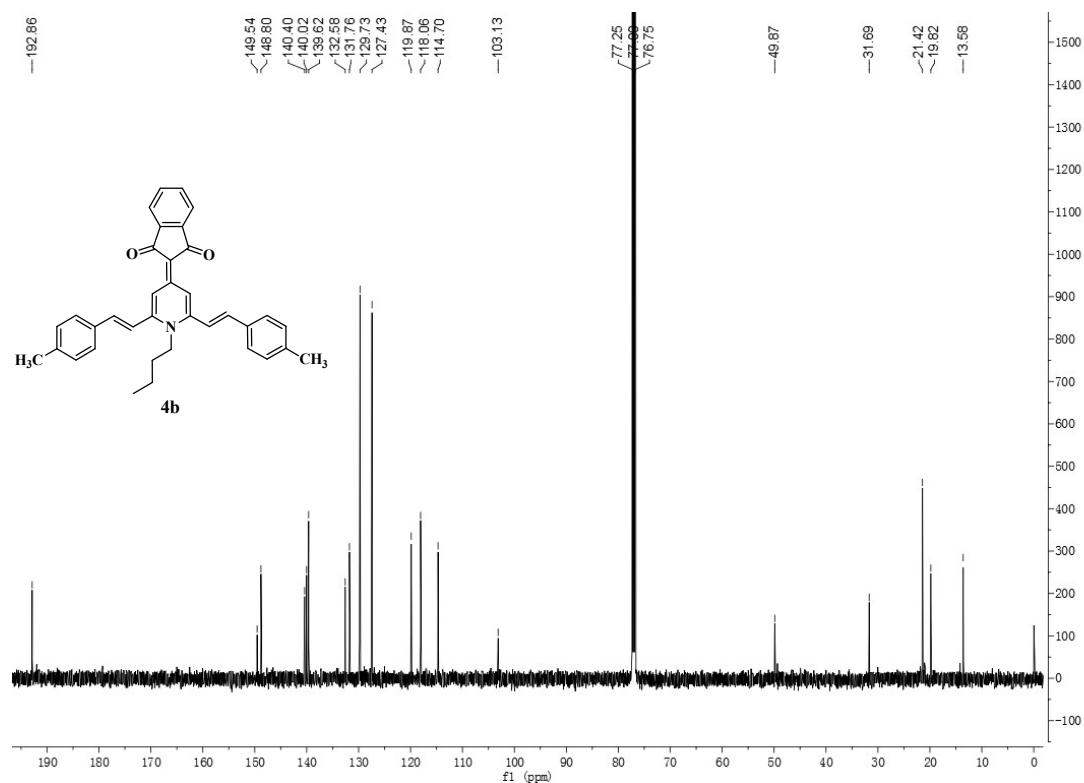


Fig. S21 ^{13}C NMR of **4b** (CDCl_3 , 125 MHz).

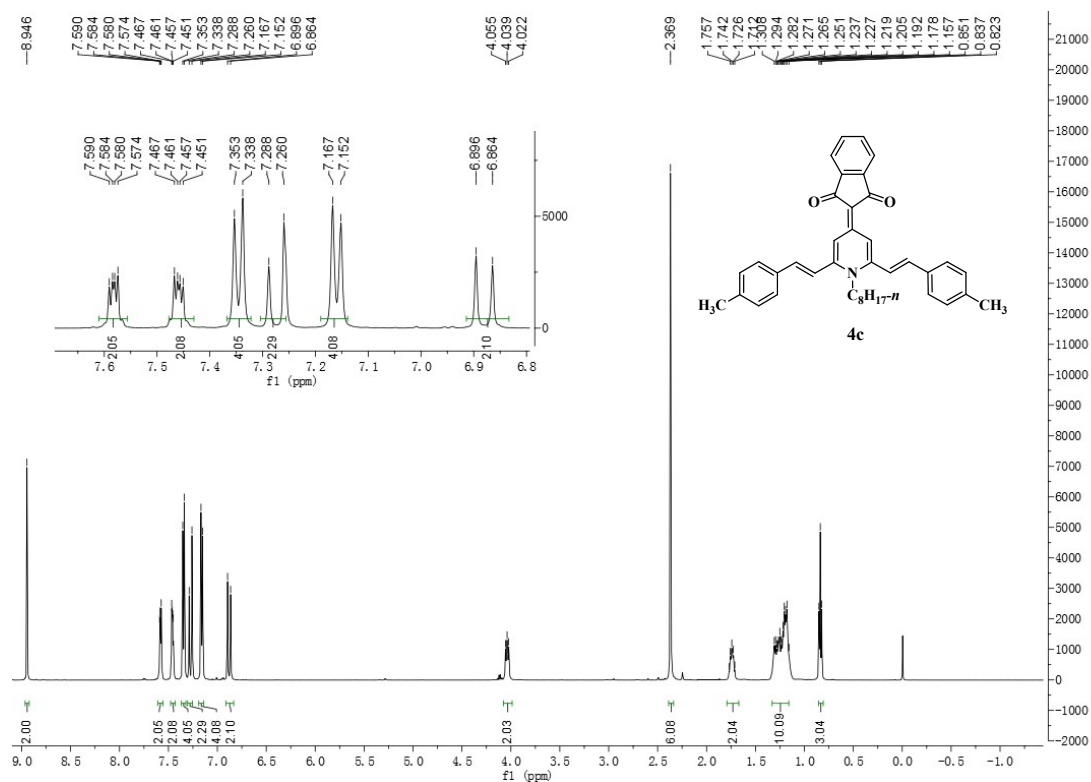


Fig. S22 ¹H NMR of 4c (CDCl₃, 500 MHz).

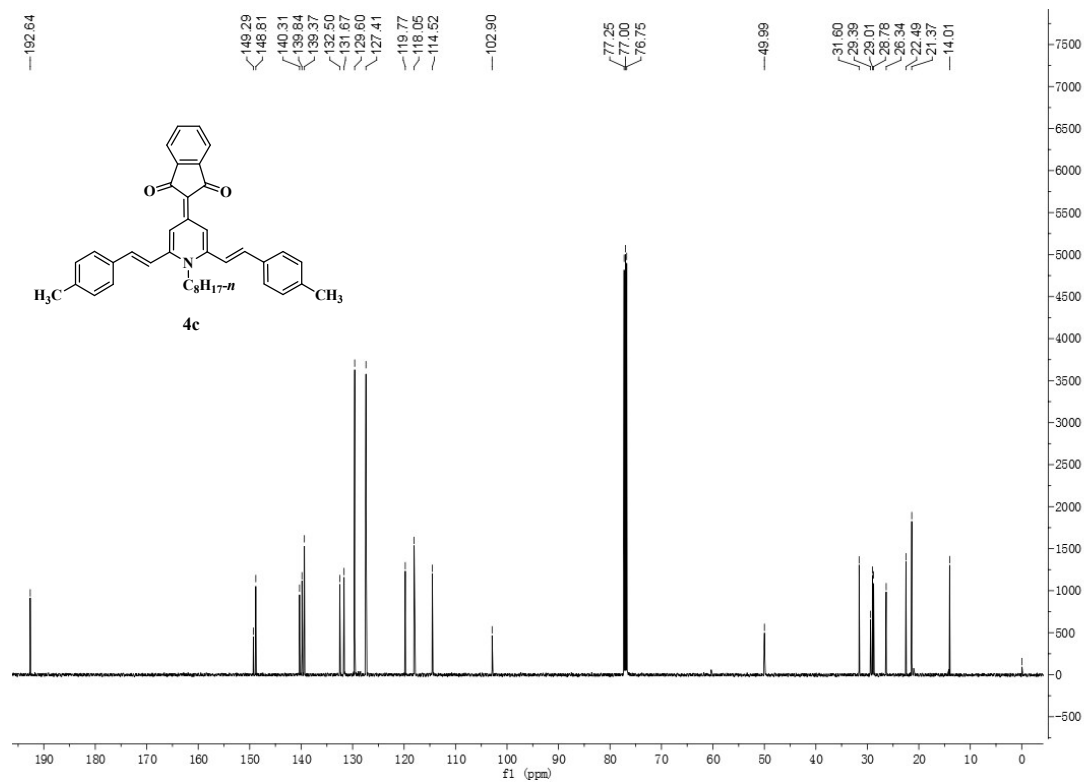
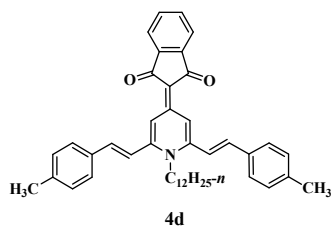


Fig. S23 ¹³C NMR of 4c (CDCl₃, 125 MHz).



Chemical structure of **4d** is shown. The structure consists of a central pyridine ring substituted with a phthalimide group, two 4-methylstyryl groups, and a dodecyl chain ($C_{12}H_{25}$).

^{13}C NMR spectrum (CDCl₃) of **4d** is displayed. The x-axis represents the chemical shift in ppm (f1), ranging from 0 to 190. The y-axis represents intensity. The spectrum shows several peaks corresponding to the structure, including aromatic and carbonyl carbons (120-150 ppm), alkene carbons (137, 135 ppm), and aliphatic carbons of the dodecyl chain (14-31 ppm). The solvent peak (CDCl₃) is visible at 77.00 ppm.

Peak list (ppm): 192.89, 149.54, 148.80, 140.37, 140.02, 139.63, 132.58, 131.77, 129.72, 127.43, 119.88, 118.05, 114.70, 103.12, 77.25, 77.00, 76.75, 60.10, 31.89, 29.61, 29.56, 29.47, 29.39, 29.31, 28.94, 26.45, 22.67, 21.42, 14.10.

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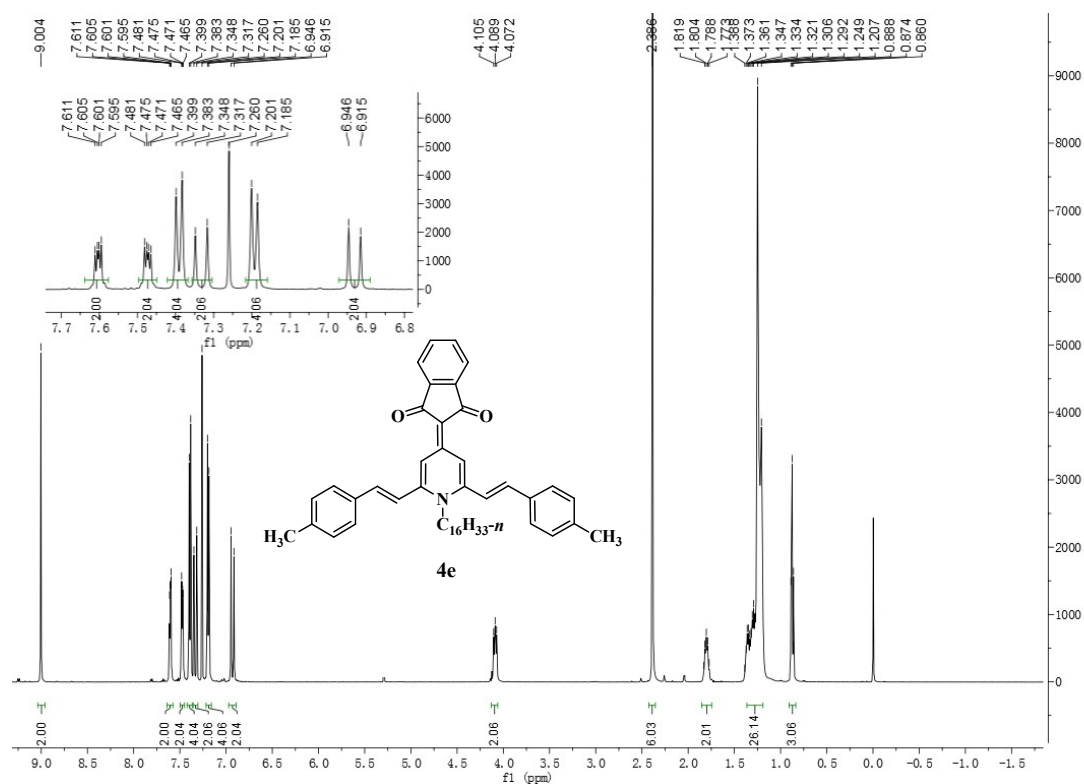


Fig. S26 ¹H NMR of 4e (CDCl₃, 500 MHz).

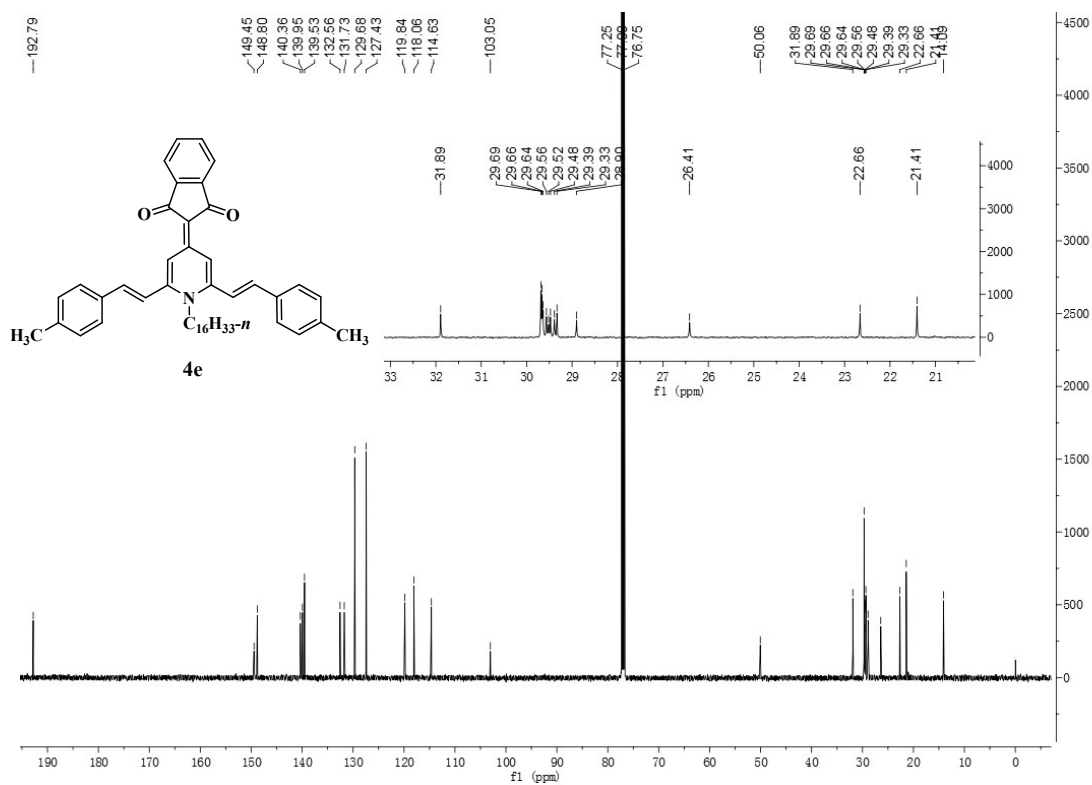


Fig. S27 ¹³C NMR of 4e (CDCl₃, 125 MHz).