**E/Z isomerization effects on aggregation-enhanced emission of tetraphenylethene derivatives assisted by host–guest recognition†**

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Pure stereoisomers of tetraphenylethene derivatives functionalized with dibenzylamine groups, E and Z isomers, were successfully synthesized. Their aggregation-enhanced emissions, which occurred through host–guest recognition with a dibenzo-24-crown-8 based tetraphenylethene under an acid condition, showed recognizable isomerization effects.

In nature, there are many fascinating E and Z isomers, both of which possess unique and elegant properties. *Cis–trans* isomers have the same chemical formula but with different geometrical positions in space, leading to very different functionalities. Therefore, their synthesis and separation are of great importance in medicinal industry,† mechanical devices,‡ and information storage.§ Tetraphenylethene (TPE) is intensively emissive in solid or aggregated states, but not in dilute solution.¶ This aggregation-induced emission (AIE) active molecule can be easily functionalized at the para position of the phenyl groups and has been utilized as a building block to develop AIE or aggregation-enhanced emission (AEE) materials. However, TPE derivatives are commonly collected as a stereoisomeric mixture containing both E and Z conformers. This situation does not permit scientists to distinguish the steric effects on the AIE or AEE properties. In only a few case studies,‖∞ the E isomer of TPE derivatives has shown a different AIE behavior from the Z isomer. The remaining challenge is how to efficiently separate the pure E and Z stereoisomers of TPE derivatives.

Recently, Sun’s group†‡ and we⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥⊥epermanently changed the way we think about light, and how we can use it to create new materials with fascinating properties.

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corresponded to the aromatic proton $H_f$. This signal was clearly characterized by $^1H$, $^{13}C$ NMR (Fig. S1). The intermediate compounds and final products were fully characterized by $^1H$, $^{13}C$ NMR (Fig. S1–S21). All of the geometries of the $E$ and $Z$ isomers were further confirmed by NOESY and COSY NMR spectroscopy (Fig. S13–S27).

The $^1H$ NMR spectra of $E$-2CHO–TPE and $Z$-2CHO–TPE and their stereoisomer mixture were displayed in Fig. 1a–c. From $E$ to $Z$ stereoisomer, the resonances for the aromatic protons $H_e$ and $H_d$ showed recognizable downfield shifts of 0.03 and 0.02 ppm, respectively. For the protons $H_c$, the resonance signal was shifted upfield with 0.05 ppm. Similar resonance shifts were also found in the $^1H$ NMR spectra of $E$-2DBA–TPE and $Z$-2DBA–TPE (Fig. S28). Both $E$-2DBA–TPE and $Z$-2DBA–TPE were further analyzed by using the NOESY and COSY NMR spectra (Fig. S21–S27 and 1d). For $E$-2DBA–TPE, the resonance at 7.57 ppm was correlated with the aromatic proton $H_e$. This signal was clearly correlated with the resonances at $\delta = 7.44$ and 7.41 (Fig. S22 and S23). This observation allowed us to assign them to the $H_e$ and $H_d$ protons, respectively. Subsequently, the signal correlation of $H_d$ with the resonance at $\delta = 7.17$ led to the clear attribution of this peak to $H_{a,d,d',d''}$ in A and B phenyl rings. Following these assignments, the proton resonance at 7.12 was due to $H_{b,c',c''}$ in the rings of C and D. Accordingly, the resonances at $\delta = 7.16$–7.12 ppm were assigned to the $H_e$ and $H_d$ protons. As marked by the black arrows in Fig. 1d, the red cross peaks revealed a clear correlation between $H_{c,c}$ and $H_{d,d',d''}$ in $Z$-2DBA–TPE (Fig. S24–S27). This was reasonable when one considered that the rings of A and C were far from B and D in the $Z$ isomer.

$E$-2DBA–TPE and $Z$-2DBA–TPE showed typical AIE properties in the solid states (Fig. S32 and Table S1†) and THF/water mixtures (Fig. S33–S37†). When the water fraction was 99%, the fluorescence intensities reached their maximum values with enhancement factors of 134 and 163, respectively. Therefore, their AIE can be efficiently distinguished by the degree of fluorescence enhancements in the THF/water mixture (Fig. S34†). This was consistent with previously reported AIE properties of TPE-based stereoisomers. A similar AIE feature was also found in the case of $4DB24C8$–TPE (Fig. S34†) but with a maximum fluorescence enhancement of only 35 at the water fraction of 95%.

When 2.4 equivalents of HPF$_6$ were added into a mixture of $4DB24C8$–TPE and $E$-2DBA–TPE in CD$_2$Cl$_2$, in which the DBA groups were protonated into dibenzylammoniums, significant changes occurred in the $^1H$ NMR spectra (Fig. 2a and b). All of the proton signals were greatly broadened and complicated. Four new broad bands appeared at $\delta = 4.48$, 4.62, 4.91, and 5.07 ppm (Fig. 2b). According to previously reported $^1H$ NMR result on host–guest recognition between DB24C8 and dibenzylammonium groups, these resonances were clearly assigned to the benzylic methylene protons ($H_{lc} + H_{mc}$) adjacent to the NH$_2$ centers hosted by the DB24C8 rings. These shifted, broadened, and complicated resonances suggested that a supramolecular hyperbranched polymer (SHP)$^9$ formed in solution by the $A_4$
(4DB24C8–TPE) and B<sub>2</sub> (E-2DBA–TPE) monomers through host–guest recognition of the DB24C8 rings with the dialkylammonium ions. With reference to the previous 1H NMR studies on this class of SHPs, the isolated and broad signal at 6.30 ppm was due to the complexed aromatic proton H<sub>Cl</sub>. This resonance was independent of those signals for the complexed benzylic methylene protons (H<sub>B</sub> + H<sub>Cl</sub>). This situation allowed us to determine the percentage recognition (p) and polymerization degree (n) according to the following equation:

\[ p = n/(n+1) = A(H_q + H_{lm})/(nA(H_q)) \]

where A(H<sub>q</sub>) and A(H<sub>lm</sub> + H<sub>Cl</sub>) are the average integrals of H<sub>q</sub> and (H<sub>lm</sub> + H<sub>Cl</sub>) respectively. Accordingly, the p and n values were calculated to be 99% and 99 at a DB24C8 concentration of 2.0 \times 10^{-3} \text{ mol L}^{-1}. This permitted the calculation of the molecular weight (M<sub>n</sub>) to be 4.5 \times 10^{5} \text{ g mol}^{-1}. The subsequent addition of 2.8 equivalents of N,N,N′,N″,N‴,N‴″-hexamethylphosphorimidic triamide (P<sub>T</sub>-tBu), a strong base in the same solution caused a fully recovered 1H NMR spectrum (Fig. 2c), indicating that the SHP formed reversibly in solution. A similar acid–base controlled SHP was also fabricated by mixing 4DB24C8–TPE and Z-2DBA–TPE in CD<sub>2</sub>Cl<sub>2</sub> (Fig. S38†). The calculated values of p, n, and M<sub>n</sub> were 95%, 19 and 7.9 \times 10^{5} \text{ g mol}^{-1}, much smaller than those in the case of E-2DBA–TPE, respectively. This result suggested that the protonated E-2DBA–TPE was more easily and fully connected with 4DB24C8–TPE than the protonated Z-2DBA–TPE through the host–guest recognition, consistent with the different geometries of the stereoisomers. This reversible supramolecular polymerization was further supported by the DOSY NMR experiments (Fig. S40 and S41†).

With these results in mind, we continued to investigate acid–base controlled AEE features and aggregate morphologies in solution. Alternating treatments of the THF mixture solution of 4DB24C8–TPE and E-2DBA–TPE or Z-2DBA–TPE with HCl and NaOH resulted in consecutive fluorescence enhancement and attenuation (Fig. 3). In the case of E-2DBA–TPE, the fluorescence intensities increased with a factor of 86, 125 and 157 at the first, second, and third acidification, respectively. In contrast, the fluorescence of Z-2DBA–TPE was enhanced only to 21, 78, and 97 folds. Both of them showed a stepwise increase in the fluorescence intensities with the acid treatments. With the corresponding base treatments, however, the enhanced fluorescence almost returned back to the original levels. Concomitant with these fluorescence enhancements, the emissive bands were blue-shifted from 530 to 477 (53), 468 (62) and 463 nm (67 nm) for E-2DBA–TPE, while the blue shifts of 26, 28 and 38 nm were clearly observed in the case of Z-2DBA–TPE. This observation was in excellent agreement with much easier supramolecular polymerization of 4DB24C8–TPE than the protonated Z-2DBA–TPE as stated above.

To understand the observed photophysical phenomena, the above solutions were subjected to dynamic light scattering (DLS) and transmission electron microscopy (TEM) measurements. The DLS plot of the initial mixture solution of E-2DBA–TPE or Z-2DBA–TPE with 4DB24C8–TPE offered a hydrodynamic diameter (D<sub>h</sub>) of 1.8 nm (Fig. 4a). This value was consistent with the molecular sizes of those TPE derivatives. Upon three-cycle acid treatments with HCl, the value of D<sub>h</sub> increased to 563, 2485, and 6014 nm for E-2DBA–TPE, respectively, while in the case of Z-2DBA–TPE, the D<sub>h</sub> value was 157, 1838, and 2431 nm. The stepwise increasing D<sub>h</sub> values coincided with the increasing trends of the fluorescence enhancements and blue shifts with the acidification cycles as addressed above. Therefore, these aggregates suppressed the intramolecular rotation of phenyl groups in both E/Z-2DBA–TPE and 4DB24C8–TPE in solution, leading to stepwise fluorescence enhancements and blue shifts (Fig. 3). In total, the sizes of the aggregates formed by the protonated E-2DBA–TPE with 4DB24C8–TPE were much bigger than those by the protonated Z-2DBA–TPE with 4DB24C8–TPE. These DLS data were in line with the aforementioned 1H NMR, AEE, and blue shift results. TEM images showed that the SHP
fabricated by the protonated 6,2DBA–TPE and 4DB24C8–TPE self-assembled into sheet like aggregates, while in the case of Z-2DBA–TPE, ruptured spheres formed. The more ordered forms of TPE derivatives in the nanosheets explained well the much larger blue shifts and fluorescence enhancements in the case of E-2DBA–TPE. All of these distinct behaviors stemmed from the E/Z isomerization effect of the stereoisomers, 6,2DBA–TPE and Z-2DBA–TPE during the host–guest recognition process under acid conditions.

In summary, we have isolated and characterized TPE-based stereoisomers, 6,2DBA–TPE and Z-2DBA–TPE. Upon consecutive treatments with acid and base, they can be reversibly co polymerized with 4DB24C8–TPE into SHPs through host– guest recognition, leading to reversible fluorescence enhancements and blue shifts. However, both the supramolecular polymerization and corresponding photophysical changes were more significant in the case of 6,2DBA–TPE than in Z-2DBA–TPE. This difference is originated from their different geometries leading to distinguishable percentage recognitions and polymerization degrees. The present supramolecular system on the basis of TPE stereoisomers displays tunable photophysical properties and morphological features and are of great importance to potential applications in the fields of chemical and biological sensors and optoelectronics.

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Notes and references


