Sc$_3$N@$_7$C$_{80}$ based donor–acceptor conjugate: role of thiophene spacer in promoting ultrafast excited state charge separation†

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Light induced charge separation in a newly synthesized triphenylamine–thiophene–Sc$_3$N@$_7$C$_{80}$ donor–acceptor conjugate and its C$_{60}$ analog, triphenylamine–thiophene–C$_{60}$ conjugate is reported, and the significance of the thiophene spacer in promoting electron transfer events is unraveled.

Endohedral metallofullerenes (EMF) are compounds of great interest due to their fascinating structure and unique electronic properties, while the early efforts were dedicated to their electronic structure, their chemical functionalization has aroused significant interest in recent years, not only to utilize their interesting properties but also to prepare new compounds with potential applications in photovoltaics, biomedical and materials science applications. Due to their higher lowest unoccupied molecular orbital (LUMO) compared to those of empty fullerene cages, EMFs are excellent acceptor materials in organic photovoltaics, capable of increasing the open circuit voltage of the devices. To this end, a few covalent donor–acceptor systems comprising EMFs have been prepared, although the higher stability of the radical-ion pair in comparison with analogous empty fullerenes has been described in an elegant study showing that the isoxazoline ring is fused to a [5,6]-bond junction in the endohedral cage.

Here, we describe the synthesis and photophysical study employing femtosecond transient absorption (fs-TA) spectroscopy, of a donor–acceptor conjugate formed by a triphenylamine moiety as a donor group and Sc$_3$N@$_7$C$_{80}$-isoxazoline-C$_{60}$ as the acceptor, linked by a thiophene spacer (Fig. 1). The analogous C$_{60}$ derivative was also prepared for comparison. Here we show that the central thiophene ring promotes $\pi$-conjugation between the donor and acceptor entities thus facilitating charge separation processes in these conjugates, useful for optoelectronic applications.

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Up until now, several methods have been developed to functionalize EMFs such as Diels–Alder, Bingel–Hirsch reaction, cycloaddition of carbenes or benzynes etc. Among these, 1,3-dipolar cycloaddition of azomethine ylides is perhaps the most used to functionalize EMFs, although other dipoles as nitrile imines have been used as well. On the other hand, nitrile oxides react with alkenes to afford isoxazolines and can be prepared from aldoximes by treatment with a chlorinated agent and a weak base, generally prepared in situ to circumvent dimerization. Cycloadditions of nitrile oxides on fullerenes, carbon nanotubes or graphene are known but only one example of the cycloaddition of a stable nitrile oxide to Sc$_3$N@$_7$C$_{80}$ has been described in an elegant study showing that the isoxazoline ring is fused to a [5,6]-bond junction in the endohedral cage.

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Fig. 1. Structure of the newly synthesized triphenylamine–thiophene–Sc$_3$N@$_7$C$_{80}$ endohedral fullerene conjugate, (3a) and its C$_{60}$ analog derivative (3b).
Scheme 1 shows the synthetic approach for both donor-acceptor conjugates. Reaction of aldehyde 1 with hydroxylamine hydrochloride in ethanol/pyridine afforded oxime 2, as a mixture of Z-E isomers, in 77% yield. Then, 2 was reacted with N-chlorosuccinimide (NCS) and after 30 min a solution of Sc₃N@C₆₀ in 1,2-dichlorobenzene (o-DCB) and triethylamine was added and stirred for 72 h at room temperature. The progress of the reaction was followed by HPLC, which showed that a new single peak appears as that for Sc₃N@I₉-C₆₀ disappears. After purification by column chromatography, followed by preparative HPLC, conjugate 3a was obtained in 25% yield. A similar procedure was followed to prepare the C₆₀-based compound 3b (see the Experimental section in the ESI for details†). Compound 3a was characterized by means of MALDI-TOF mass analysis showing the molecular ion peak at 1677.28 amu. The ¹H NMR spectrum contains all the expected signals of the addend. The thiophene H-atoms appear at 7.96 and 7.24 ppm, deshielded compared to those in the oxime due to the presence of the fullerene cage. Note that this signal is broad as a consequence of the hindered rotation. In the ¹³C NMR spectrum, the most relevant signals are those of the sp³ C-atoms of the functionalized fullerene cage at 97.7 and 70.3 ppm. Similar features are observed in the spectrum of the C₆₀-based compound 3b. Finally, the appearance of a broad UV-Vis absorption around 700 nm and the lack of absorption in the 800 nm region, expected for the [6,6]-isomer, confirmed the formation of the [5,6]-isomer; a result that agrees well with the previously described examples of 1,3-dipolar cycloadditions on formation of the [5,6]-isomer; a result that agrees well with the features are observed in the spectrum of the C₆₀-based compound 3b. Additionally, a stronger absorption is observed around 690 nm, which is red-shifted to 412 nm in cycloadducts 3a and 3b, respectively, suggesting the overlap of two reduction processes.

Reagents and conditions: (a) hydroxylamine hydrochloride, pyrididine; (b) NCS, pyridine (c) Et₃N, Sc₃N@I₉-C₆₀ or C₆₀, r.t.

Table 1 Redox potentials (in V vs. Fc/Fc⁺) determined by OSWV of the processes observed for 2, 3a–b and reference fullerences

| Compound | E¹²⁻⁻⁻²⁻⁻⁻²⁻⁻¹⁻⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁻¹⁺⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻⁻
existence of some electronic interactions with the respective fullerene cage. In the cathodic region, the first reduction potential of 3a is positively shifted by 270 mV when compared to that of pristine Sc$_3$N@I$_2$-C$_{80}$. This behavior is similar to what was previously observed for an isoxazoline Sc$_3$N@I$_2$-C$_{80}$ derivative, confirming the strong influence that functionalization has on the electrochemical properties of the EMF cage. However, the first reduction of 3b and C$_{80}$ showed similar values (−1.05 V and −1.03 V, respectively) as described in other donor-acceptor isoxazolinofullerene derivatives. This behavior is different from that for C$_{80}$ derived pyrrolidinofullerenes, where the first reduction is about 120 mV more negative than for pristine C$_{80}$ as a result of the saturation of one of the double bonds.

From the optical absorption and emission, electrochemical redox potential and structural parameters, Gibbs free-energy changes associated with charge separation (CS) and charge recombination (CR) were estimated according to eqn (i) and (ii)

$$-\Delta G_{CR} = E_{ox} - E_{red} + \Delta G_S \quad (i)$$

$$-\Delta G_{CS} = \Delta E_{0,0} - (-\Delta G_{CR}) \quad (ii)$$

where $\Delta E_{0,0}$ corresponds to the energy of the $^1$TPA-Tf state calculated from luminescence peak maximum at 2.72 eV. The $E_{ox}$ and $E_{red}$ are the first oxidation potential of TPA-Tf and the first reduction potential of the fullerene in the conjugate. The term $\Delta G_S$ refers to the electrostatic energy calculated according to the dielectric continuum model. The calculated $\Delta G_S$ in o-DCB for 3a and 3b were found to be −0.09 and −0.11 eV, respectively. Incorporation of these values into eqn (i) and (ii) resulted in $\Delta G_{CS}$ and $\Delta G_{CR}$ of −1.59 and −1.13 eV for 3a and −1.46 and −1.26 for 3b, respectively.

The energy level diagram shown in Fig. 3b was constructed to visualize the possible photochemical events. Excitation of TPA in the dyads leads to a population of its lowest excited singlet state, $^1$TPA-Tf$. The $^1$TPA-Tf state can undergo a thermodynamically allowed CS to the appended fullerene. The fluorescence spectra in Fig. 2b reveals that attaching Sc$_3$N@I$_2$-C$_{80}$ or C$_{60}$ to TPA via the thiophene spacer leads to quantitative quenching and no new peaks due to fullerene emission suggesting no energy transfer. The data suggest that CS is the most likely quenching mechanism. The high $\Delta G_{CS}$ and $\Delta G_{CR}$ values and the spatial closeness between the donor and acceptor entities along with the strong electronic coupling by through the thiophene spacer are anticipated to accelerate both the CS and CR processes. Further, fs-TA studies were performed in solvents of different polarity to clarify these predictions. Prior to this, in an effort to spectrally characterize electron transfer products, TPA-Tf (compound 2) was chemically oxidized using NOBF$_4$ in benzonitrile, as shown in Fig. S11.† Increased additions of the oxidizing agent decreased the intensity of the main peak at 390 nm and resulted in the appearance of new peaks at 375, 486, 731 and 967 nm, corresponding to (TPA-Tf)$^+$. Spectral characteristics of Sc$_3$N@I$_2$-C$_{80}$ and C$_{60}$ with peaks around 1000 nm have been reported earlier.\textsuperscript{54,55}

Fs-TA studies were performed in three solvents of varying polarity. Fig. 4 shows fs-TA spectra of 2, 3a and 3b at the indicated delay times in o-DCB while those recorded in toluene and benzonitrile are shown in Fig. S12 and S13,\textsuperscript{†} respectively. In all these solvents, immediately after excitation, the instantaneously formed $^1$TPA-Tf$^*$ (″TPA-Tf″) revealed a negative peak in the 450–550 nm region due to contributions of ground state bleaching (GSB) and stimulated emission (SE). Positive peaks in the 600 and 1240 nm region due to excited state absorption (ESA) were also observed. With time, the near-IR peak revealed

Fig. 3 (a) Cyclic voltammograms of 3b (black) and 3a (blue) in o-DCB : acetonitrile (4 : 1) (b) energy level diagram showing the different photochemical events occurring for 3a and 3b due to TPA excitation. Energies of different states were evaluated from spectral and electrochemical studies. Solid arrows indicate major photo-processes, dashed arrow indicates minor photo-processes. CS = charge separation, CR = charge recombination.

Fig. 4 Fs-TA spectra at the indicated delay times of (a) 2 ($\lambda_{ex} = 390$ nm), (b) 3a ($\lambda_{ex} = 500$ nm), and (c) 3b ($\lambda_{ex} = 390$ nm) in o-DCB. The DAS are shown in the right hand panels.
a blue-shift, likely due to solvation or vibrational cooling, and at longer delay times, the recovery and decay of negative and positive peaks was associated with emergence of new peaks in the 735 and 1000 nm range likely due to $^{3}\Pi^*$, formed via intersystem crossing (ISC).

For 3a and 3b ultrafast electron transfer events were observed. As shown in Fig. 4b and c, the recovery and decay of peaks corresponding to $^{1}(TPA-Tf)^*$ were rapid with new peaks developing within a few picoseconds characteristic of $(TPA-Tf)^+\text{Sc}_3\text{N}@\text{C}_{80}$ and $(TPA-Tf)^+\text{C}_{60}^-\text{Ih}$ in the case of 3a and $(TPA-Tf)^+\text{C}_{60}^-\text{Ih}$ in the case of 3b. These are signature peaks of $\text{Sc}_3\text{N}@\text{I}_\text{h}\text{C}_{80}$ and $\text{C}_{60}^-\text{Ih}$ in the near-IR region and $(TPA-Tf)^+\text{I}_\text{h}$ in the visible region. Further, decay associated spectra (DAS) were generated from global analysis to evaluate time constants for CS and CR, as shown in the right-hand panels of Fig. 4. Changing the solvent revealed solvent polarity dependent electron transfer kinetics, showing that polar solvents facilitated the charge separation process. The measured time constants and rates are given in Table 2.

The data presented in Table 2 reveals the following: (i) the $k_{\text{CS}}$ values are generally higher for 3a compared to 3b which could be attributed to the free-energy changes (see Fig. 3b) where facile charge separation for 3a over 3b was noted. (ii) Although some charge stabilization is observed in nonpolar toluene, in general $k_{\text{CS}}$ values are also much faster on the order of $10^{10}$ to $10^{14}$ s$^{-1}$. (iii) A comparison of electron transfer rates between the present system and the previously reported TPA-fullerene conjugate$^{34}$ where charge stabilization was observed to some extent suggests that the thiophene entity enhances the $\pi$-conjugation between the donor and acceptor entities in 3a and 3b, thus facilitating electron transfer events in both directions.

### Conclusions

In summary, new donor–acceptor conjugates, TPA-Tf-$\text{Sc}_3\text{N}@\text{I}_\text{h}\text{C}_{80}$, and its $\text{C}_{60}$ analog, TPA-Tf-$\text{C}_{60}$, were synthesized by a 1,3-dipolar cycloaddition reaction of nitrile oxides. The thiophene unit promotes the $\pi$-conjugation between the donor and acceptor entities. An energy level diagram constructed using spectral and electrochemical data revealed an exothermic excited state charge separation, more so for the TPA-Tf-$\text{Sc}_3\text{N}@\text{I}_\text{h}\text{C}_{80}$ conjugate. Photoexcitation of the TPA-Tf entity in the conjugates promoted charge separation within a few picoseconds, attributed to the $\pi$-linker, thiophene, connecting the donor and acceptor units, accessible redox potentials and spatial closeness of the donor and acceptor entities of the conjugates. Further studies to fully explore the potential of the current synthetic methodology to functionalize endohedral fullerenes for photochemical and optoelectronic applications are underway in our laboratories.

### Conflicts of interest

Authors declare no conflict of interest.

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