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Double Doping of a Low-Ionization-Energy Polythiophene with a Molybdenum Dithiolene Complex

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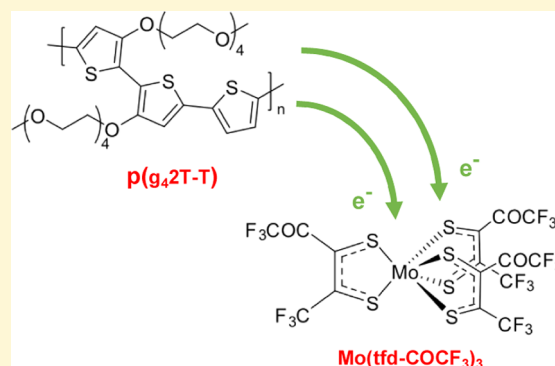


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ABSTRACT: Doping of organic semiconductors is crucial for tuning the charge-carrier density of conjugated polymers. The exchange of more than one electron between a monomeric dopant and an organic semiconductor allows the polaron density to be increased relative to the number of counterions that are introduced into the host matrix. Here, a molybdenum dithiolene complex with a high electron affinity of 5.5 eV is shown to accept two electrons from a polythiophene that has a low ionization energy of 4.7 eV. Double p-doping is consistent with the ability of the monoanion salt of the molybdenum dithiolene complex to dope the polymer. The transfer of two electrons to the neutral dopant was also confirmed by electron paramagnetic resonance spectroscopy since the monoanion, but not the dianion, of the molybdenum dithiolene complex features an unpaired electron. Double doping allowed an ionization efficiency of 200% to be reached, which facilitates the design of strongly doped semiconductors while lessening any counterion-induced disruption of the nanostructure.



INTRODUCTION

Organic semiconductors attract a great deal of attention since they enable the development of lightweight, flexible, and biocompatible electronic devices for applications from energy harvesting and storage to bioelectronics, which cannot be achieved with inorganic semiconductors alone.^{1,2} Doping is a widely used tool to tune the charge-carrier density, which allows the electrical properties of organic semiconductors to be optimized.^{3–5} The process typically involves the addition of a dopant, such as an organic molecule, a metal–organic complex, or a metallic salt, that undergoes an electron transfer with the organic semiconductor.⁵ Electron transfer readily occurs in the case of a favorable energetic offset between the two materials. In the case of p-doping, for example, it is beneficial if the dopant has an electron affinity (EA) that is larger than the ionization energy (IE) of the semiconductor, i.e., $EA_{\text{dopant}} \geq IE_{\text{semiconductor}}$. Here, it is important to note that the EA and IE, which are often estimated from electrochemical methods, can take on very different values once the dopant and semiconductor are mixed.⁶ Furthermore, the structural and energetic disorder inherent to many organic semiconductors leads to a broad density of states, which may facilitate some degree of electron transfer despite an unfavorable energy offset.^{7,8} Provided that each p-dopant accepts one electron from the semiconductor, i.e., the pair undergoes integer charge transfer, one polaron is created per dopant molecule, resulting in an ionization efficiency of 100%. The ionization efficiency is

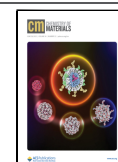
reduced to less than 100% in the case that only partial charge transfer occurs, if there is an unfavorable energetic offset between EA_{dopant} and $IE_{\text{semiconductor}}$ or if the dopant molecules aggregate. The presence of excess dopants and dopant aggregates unduly disrupts the nanostructure of the semiconductor, which tends to negatively affect the electrical properties.^{9–11}

We have recently reported that quinodimethane-type dopants with a sufficiently high EA_{dopant} can accept two electrons from a low-IE conjugated polymer, resulting in an ionization efficiency of 200%.¹² This type of double doping was observed for a thienothiophene-based copolymer with tetraethylene glycol side chains, for which $IE_{\text{polymer}} = 4.5$ eV, doped with either 2,3,5,6-tetrafluoro-7,7,8-tetracyanoquinodimethane ($F_4\text{TCNQ}$), for which $EA_{F_4\text{TCNQ}} = 5.2$ eV, or 1,3,4,5,7,8-hexafluorotetracyanonaphthoquinodimethane ($F_6\text{TCNNQ}$), for which $EA_{F_6\text{TCNNQ}} = 5.3$ eV (electron affinity estimated from half-wave potential $E_{1/2}$ vs ferrocene/ferrocenium, Fc/Fc^+ ; $EA_{\text{dopant}} = 5.1$ eV + $E_{1/2}$).¹² The electron

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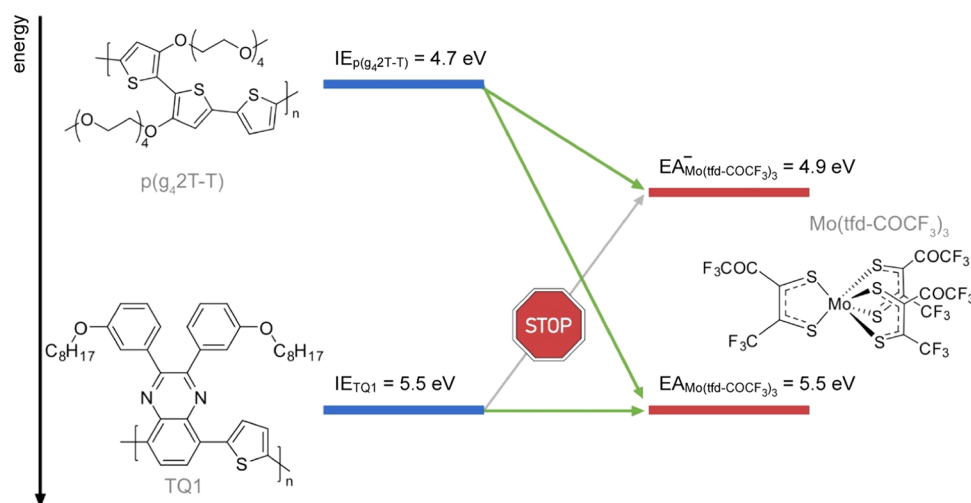


Figure 1. Energy diagram showing the energy levels that are relevant for charge transfer from polymers $p(g_42T-T)$ and TQ1 to the p-type dopant $Mo(tfd-COCF_3)_3$.

affinities of the anions of these two quinodimethane-type dopants, EA_{dopant}^- are estimated to remain sufficiently low that $EA_{dopant}^- > IE_{polymer}$ allowing each dopant to accept a second electron from the polymer. Other dopants that can give rise to more than one polaron include the dimers formed by certain 19-electron organometallic sandwich compounds or by benzoimidazoline radicals, the overall reaction of which with a host results in the formation of two monomeric dopants and the release of two electrons.⁷ These dopant dimers, however, have approximately twice the molecular volume of related monomeric dopants, which reduces the benefit of double doping because it doubles the overall volume that is occupied by counterions. Likewise, multivalent radical cation salts that comprise two or four triphenylamine units can accept two or even four electrons from a conjugated polymer^{13,14} but have a close to 2–4 times larger volume compared to the corresponding monovalent radical anion salt tris(4-bromophenyl)ammoniumyl hexachloroantimonate ($[TPA-Br_3]^+[SbCl_6]^-$; Magic Blue). We argue that double doping is preferably achieved without increasing the size of the dopant so that fewer dopant molecules must be added to obtain a certain polaron density, which reduces the risk of dopant aggregation and lessens the impact of doping on the nanostructure of the polymer.

Here we ask whether double doping is unique to quinodimethane-type monomeric dopants or whether it is a generic concept that can also be observed for other species. In particular, we investigate doping using the dithiolene complex $Mo(tfd-COCF_3)_3$,¹⁵ for which $EA_{Mo(tfd-COCF_3)_3}$ can be estimated as ca. 5.5 eV (reduction potential $E_{red} \approx +0.39$ V vs decamethylferrocene/decamethylferrocenium, $DmFc/DmFc^+$; $EA_{Mo(tfd-COCF_3)_3} = 5.1$ eV + E_{red}).¹⁶ We note that electrochemically based estimates of the electron affinity are approximate because solvation effects and solid-state polarization effects likely vary between molecules of different sizes and shapes. However, the value of $EA_{Mo(tfd-COCF_3)_3} = 5.5$ eV is in reasonable agreement with a value of 5.6 eV measured in the solid state by inverse photoelectron spectroscopy for a related molecule, $Mo(tfd)_3$, for which $E_{red} \approx +0.28$ V vs $DmFc/DmFc^+$.¹⁷ $Mo(tfd-COCF_3)_3$ has been used to dope a wide range of conjugated polymers including poly(3-hexylthiophene)

(P3HT),^{18,19} poly[2,5-bis(3-tetradecylthiophen-2-yl)thieno-[3,2-*b*]thiophene] (PBTTT),^{20,21} and a benzodithiophene-based copolymer.^{9,22,23} We chose $Mo(tfd-COCF_3)_3$ because it can exist both as an anion and dianion, meaning that it can sustain two electron transfer events, and both of these species are known to have reasonable chemical stability (indeed, the neutral dopant is obtained by chemical oxidation of the dianion).¹⁶ The $Mo(tfd-COCF_3)_3$ anion electron affinity, $EA_{Mo(tfd-COCF_3)_3}^-$, can be estimated as ca. 4.9 eV ($E_{red} \approx -0.16$ V vs Fc/Fc^+),¹⁶ although the potential separation between subsequent redox processes in electrochemical experiments can be highly medium-dependent,²⁴ meaning there will be a larger uncertainty in this estimate than in those of the electron affinity of neutral molecules. Nevertheless, if $EA_{Mo(tfd-COCF_3)_3}^-$ is indeed ca. 4.9 eV, then polymers with an $IE_{polymer} < 4.9$ eV should be able to experience double doping when brought in contact with $Mo(tfd-COCF_3)_3$. Doping of two polymers was investigated, the polythiophene $p(g_42T-T)$ with tetraethylene glycol side chains and a low $IE_{p(g_42T-T)} = 4.7$ eV (oxidation potential $E_{ox} \approx -0.44$ V vs Fc/Fc^+ ; $IE_{p(g_42T-T)} = 5.1$ eV + E_{ox})²⁵ and the thiophene-quinoxaline copolymer TQ1,^{26,27} which features a large $IE_{TQ1} = 5.5$ eV ($E_{ox} \approx 0.37$ V vs Fc/Fc^+ ; see Figure 1 for chemical structures and energy levels). We here show that $p(g_42T-T)$ readily undergoes double doping by $Mo(tfd-COCF_3)_3$.

RESULTS AND DISCUSSION

The dopant investigated here can exist as a neutral complex, $Mo(tfd-COCF_3)_3$, as well as a salt comprising either its anion or dianion. The dianion salt $Mo(tfd-COCF_3)_3(Et_4N)_2$ is a precursor for the synthesis of $Mo(tfd-COCF_3)_3$, as reported previously.^{15,16} Neutral $Mo(tfd-COCF_3)_3$ was prepared by oxidizing the dianion salt using NOPF₆. The anion salt, instead, was prepared by the comproportionation of equimolar amounts of $Mo(tfd-COCF_3)_3$ and the dianion salt $Mo(tfd-COCF_3)_3(Et_4N)_2$. All three species give rise to distinct ultraviolet–visible–near-infrared (UV–vis–NIR) absorbance spectra (Figure 2a). These absorptions, however, occur at wavelengths where either neat $p(g_42T-T)$ and TQ1 absorb or where polaronic absorbance peaks tend to arise upon doping. Hence, it is challenging to extract information about the

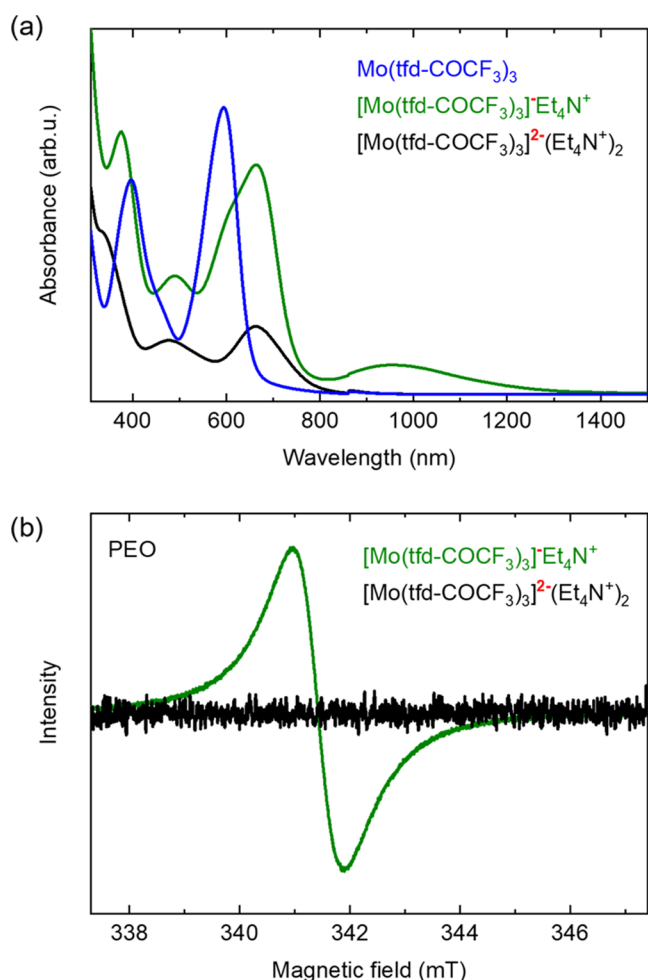


Figure 2. (a) UV-vis spectra of Mo(tfd-COCF₃)₃, [Mo(tfd-COCF₃)₃]⁻Et₄N⁺, and [Mo(tfd-COCF₃)₃]²⁻(Et₄N⁺)₂ dissolved in dichloromethane (DCM). (b) EPR spectra of [Mo(tfd-COCF₃)₃]⁻Et₄N⁺ and [Mo(tfd-COCF₃)₃]²⁻(Et₄N⁺)₂ dispersed in a matrix of poly(ethylene oxide) (PEO; see Figure S4 for nonsmoothed EPR spectra).

presence of different species from optical spectroscopy. Instead, we chose to use electron paramagnetic resonance (EPR) spectroscopy to detect the presence of different species.

In a first set of experiments, we recorded EPR spectra of the monoanion salt and the dianion salt dispersed in poly(ethylene oxide) (PEO) to mimic the solid-state environment that the two species will experience when dispersed in a p(g₄2T-T) matrix. The UV-vis-NIR spectra of the monoanion and dianion salt dissolved in dichloromethane (DCM) or in PEO are comparable (Figure S1), which suggests that the PEO matrix does not strongly alter the electronic state of the two species. The monoanion salt gives rise to a distinct EPR signal centered at $B_0 = 341.4$ mT, in agreement with reports for other monoanionic molybdenum tris(dithiolene) complexes,^{28–30} while the dianion salt appears spin-silent (Figures 2b and S2–S4), consistent with the sharp NMR spectra reported for [Mo(tfd-COCF₃)₃]²⁻,¹⁵ as well as with magnetic susceptibility³⁰ and computational studies³¹ of other molybdenum tris(dithiolene) dianions.

We next studied doped TQ1, which has a high $IE_{TQ1} = 5.5$ eV that is comparable to $EA_{Mo(tfd-COCF_3)_3}$. Judging by the ionization energy of TQ1, the polymer can be expected to

undergo some electron transfer with neutral Mo(tfd-COCF₃)₃, but the energy levels of TQ1 and [Mo(tfd-COCF₃)₃]⁻ should not allow for a second charge transfer event since $IE_{TQ1} \gg EA_{Mo(tfd-COCF_3)_3}$. UV-vis-NIR spectra of TQ1 sequentially doped with Mo(tfd-COCF₃)₃ reveal clear polaronic absorption bands confirming that electron transfer has indeed occurred (Figure 3a). The EPR spectra for TQ1 doped with Mo(tfd-COCF₃)₃ consist of two overlapping signals, one at a magnetic field of 342.2 mT and one at a lower magnetic field. We assign the signal at a lower magnetic field to the monoanion of Mo(tfd-COCF₃)₃ since it appears at the same position as the EPR signal observed for [Mo(tfd-COCF₃)₃]⁻Et₄N⁺ (Figures 3b, S5, and S6). This observation suggests that doping of TQ1 with Mo(tfd-COCF₃)₃ primarily occurs through a single electron transfer event per dopant, resulting in Mo(tfd-COCF₃)₃ anions that can be detected with EPR spectroscopy.

To confirm the plausibility of assigning the EPR signal at 342.2 mT to the polymer polaron, we also recorded EPR spectra of TQ1 doped with Magic Blue, which is a stronger oxidant than Mo(tfd-COCF₃)₃ and in which the radical cation portion can accept an electron from TQ1 because neutral tris(4-bromophenyl)amine (TPA-Br₃) has a very high $IE_{TPA-Br_3} \approx 5.7$ eV ($E_{1/2} = +0.7$ V vs DmFc/DmFc⁺),³² i.e., its radical cation [TPA-Br₃]⁺ can be reduced by TQ1 since 5.7 eV $> IE_{TQ1}$. Doping of TQ1 with Magic Blue is confirmed by the appearance of a strong polaron absorption band at 900 nm in recorded UV-vis spectra (Figure 3a).³³ The tris(4-bromophenyl)ammoniumyl cation [TPA-Br₃]⁺ becomes neutral as it receives an electron from the polymer during the doping process, and hence ends up spin-silent. Consequently, the EPR spectrum of doped TQ1 exclusively features a polaron signal. The similarity of the signal at ca. 342.2 mT seen for Mo(tfd-COCF₃)₃-doped TQ1 to the EPR spectrum of TQ1 doped with Magic Blue confirms that this signal arises from polarons on TQ1 (Figures 3c, S7, and S8).

In a further set of experiments, we studied sequential doping of p(g₄2T-T) with neutral Mo(tfd-COCF₃)₃ as well as its anion salt, i.e., the two species were allowed to ingress from a dopant solution into a solid film of the polymer (see the Experimental Section for details). UV-vis-NIR spectroscopy reveals stronger polaron absorption bands at 900 nm and in the near-infrared region for p(g₄2T-T) doped with the neutral complex, which indicates a higher degree of doping with Mo(tfd-COCF₃)₃ compared to doping with the monoanion salt (Figure 4a). Moreover, the fact that we can dope p(g₄2T-T) with the monoanion suggests that double doping is possible for this system. Doping of p(g₄2T-T) with Mo(tfd-COCF₃)₃ gives rise to an electrical conductivity of $\sigma = (19.6 \pm 0.6)$ S cm⁻¹, whereas doping with the monoanion salt gives a value of $\sigma = (11.9 \pm 0.3)$ S cm⁻¹. The higher electrical conductivity for p(g₄2T-T) doped with the neutral complex is also an indication that we have a higher degree of doping, i.e., a larger number of charge carriers in this sample compared to p(g₄2T-T) doped with the monoanion salt. EPR spectra of neat p(g₄2T-T) as well as the polymer doped with either neutral Mo(tfd-COCF₃)₃ or the anion salt all reveal a polaron signal at $g = 342.2$ mT (Figures 4b and S9). The signal in the case of the neat p(g₄2T-T) sample likely arises from adventitious oxygen doping of the polymer. In the case of doping with the neutral complex, the recorded EPR spectrum features a broad shoulder at a lower magnetic field around 341 mT, which we explain with the presence of monoanions, i.e.,

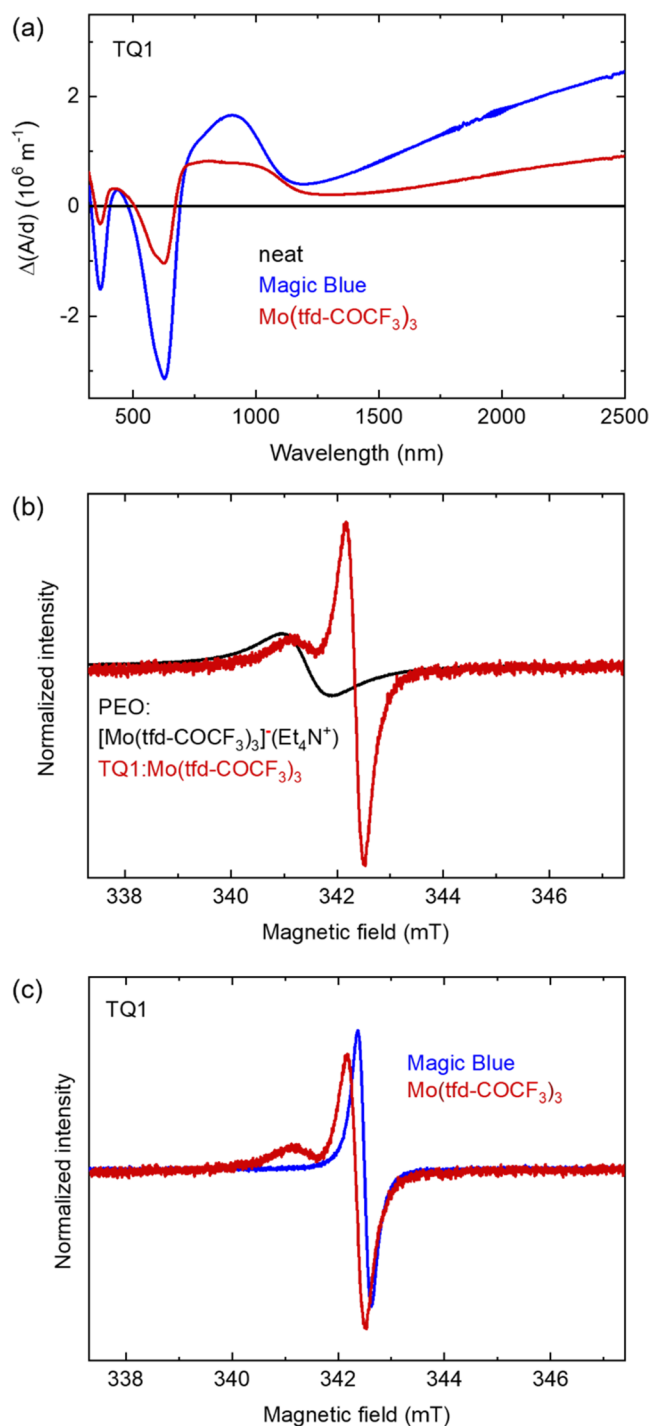


Figure 3. (a) UV-vis-NIR absorbance spectra displaying the difference in thickness-normalized absorbance $\Delta(A/d)$ between the spectra of neat TQ1 and TQ1 doped with Magic Blue or Mo(tfd-COCF₃)₃. (b) EPR spectra of [Mo(tfd-COCF₃)₃]⁺Et₄N⁺ dispersed in a PEO matrix and TQ1 doped with Mo(tfd-COCF₃)₃. The spectra have been normalized to the maximum in the field range 338–341.5 mT. (c) EPR spectra of TQ1 doped with Magic Blue or Mo(tfd-COCF₃)₃. The spectra have been normalized to the same amplitude.

the EPR spectrum is a superposition of the EPR signal of the polaron on the polymer and the EPR signal of the monoanion. The EPR spectrum of p(g₄2T-T) sequentially doped with the monoanion features a signal at a similar field as the neat, oxygen-doped polymer, and the absence of a broad shoulder

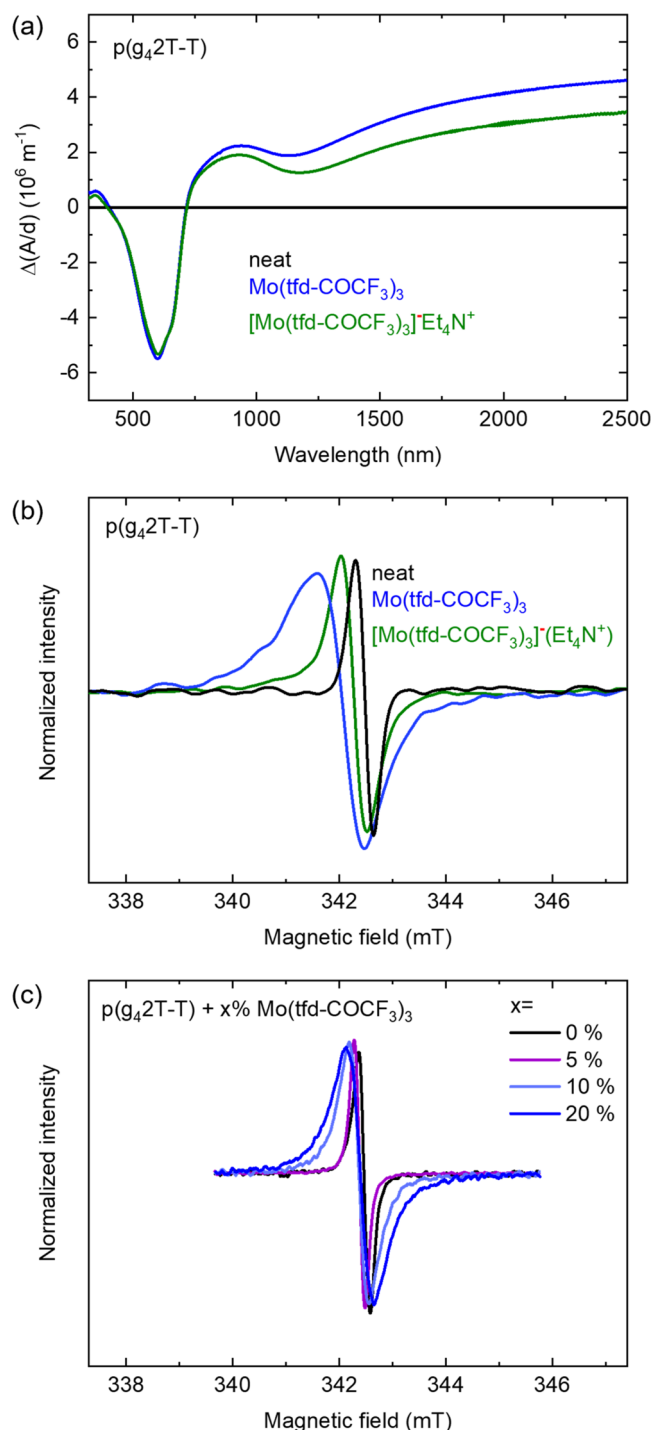


Figure 4. (a) UV-vis-NIR absorbance spectra displaying the difference in thickness-normalized absorbance $\Delta(A/d)$ between the spectra of neat p(g₄2T-T) and p(g₄2T-T) sequentially doped with Mo(tfd-COCF₃)₃ or [Mo(tfd-COCF₃)₃]⁺Et₄N⁺. (b) EPR spectra of neat p(g₄2T-T) and p(g₄2T-T) sequentially doped with Mo(tfd-COCF₃)₃ or [Mo(tfd-COCF₃)₃]⁺Et₄N⁺. (c) EPR spectra of co-processed p(g₄2T-T):Mo(tfd-COCF₃)₃ films (see Figures S9 and S11 for nonsmoothed EPR spectra).

around 341 mT suggests that no major fraction of monoanions is present. Since UV-vis-NIR spectroscopy indicates that the polymer was doped, i.e., polarons were generated (see Figure 4a), the majority of monoanions that have entered the polymer

Table 1. Parameters Quantifying the Extent of Doping and the Resulting Conductivity and Mobility for p(g₄2T-T) Co-Processed with Different Amounts of Mo(tfd-COCF₃)₃; Amount of Dopant in mol %, Calculated Relative to the Molar Mass of the Repeat Unit of p(g₄2T-T), Oxidation Level O_{ox} , i.e., Number of Polarons Per Polymer Thiophene Ring in Percent, Ionization Efficiency η_{ion} , i.e., Number of Polarons Per Added Dopant Molecules in Percent, Number of Polarons N_p , Conductivity σ and Mobility $\mu = \sigma e N_p$, where e is the Elementary Charge

mol % dopant	O_{ox} (%)	η_{ion} (%)	N_p (10^{26} m^{-3})	σ (S cm^{-1})	μ ($\text{cm}^2 \text{ V}^{-1} \text{ s}^{-1}$)
5	3.4 ± 0.2	195 ± 3	0.94 ± 0.02	0.05 ± 0.01	0.003 ± 0.001
10	7.4	200 ^a	2.0	1.1 ± 0.1	0.03 ± 0.01
20	13.1 ± 0.2	158 ± 2.3	3.6 ± 0.1	15 ± 1	0.26 ± 0.04

^aFor 10 mol % dopant, the calculated value of η_{ion} was larger than physically possible and therefore was set to 200%.

film must have accepted an electron from p(g₄2T-T) and thus became spin-silent dianions.

Finally, we studied a series of p(g₄2T-T) thin films that were prepared by co-processing the polymer with different amounts of the neutral Mo-complex. The conductivity increases with the amount of dopant (Table 1) and the polaron absorbance increases (see UV–vis–NIR spectra in Figure S10), which is consistent with an increasing polaron concentration. For all co-processed samples, the EPR signal is centered at 342.2 mT, i.e., the position assigned to polarons, and we do not observe any broad shoulder at lower magnetic fields (Figures 4c and S11). However, we instead observe a symmetrical broadening with increasing dopant concentration. We interpret the lack of the low-field shoulder as an indication that no significant amount of monoanions is present and hence the EPR signal solely originates from the polarons. The observed symmetrical broadening of the EPR signal with increasing dopant concentration could be attributed to a spin–spin interaction between nearby polarons.³⁴ Another explanation for the observed symmetrical broadening could be a lower degree of interchain charge hopping with increasing dopant concentrations resulting in the transition from Lorentzian to Gaussian absorption features.³⁵

We used the UV–vis–NIR spectra of the co-processed samples to estimate the number of polarons N_p (Table 1). To estimate N_p we compared the thickness-normalized difference in absorbance $\Delta(A_{800}/d)$ between doped samples and the neat polymer at 800 nm, where a pronounced sub-bandgap polaronic absorption peak is situated, with the molar attenuation coefficient of the polaronic absorption obtained from electrochemically doped P3HT at 800 nm, $\epsilon_{800} = (4.1 \pm 0.2) \times 10^3 \text{ m}^2 \text{ mol}^{-1}$.¹⁹ The Beer–Lambert law $\Delta(A_{800}/d) = \epsilon_{800} \times N_p$, where d is the film thickness, was used to estimate N_p . We then calculated the oxidation level according to $O_{ox} = N_p/N_{\text{thiophene}}$ and ionization efficiency according to $\eta_{ion} = N_p/N_{\text{Mo(tfd-COCF}_3)_3}$, where $N_{\text{Mo(tfd-COCF}_3)_3}$ is the number of molybdenum dithiolene complexes (Table 1). We obtain an ionization efficiency of more than 100% for all studied samples, indicating that double doping occurs even for the samples with a higher doping concentration. For 5 and 10 mol % dopants, we estimate that the ionization efficiency is close to 200%, meaning that each Mo(tfd-COCF₃)₃ complex accepts two electrons from p(g₄2T-T). A high degree of double doping explains why we do not observe any clear feature of the monoanion in the EPR spectra of p(g₄2T-T) co-processed with Mo(tfd-COCF₃)₃ (Figure 4c). Most of the monoanions that form undergo a second electron transfer event and thus become dianions, which are spin-silent.

CONCLUSIONS

We conclude that the molybdenum dithiolene complex Mo(tfd-COCF₃)₃ can accept two electrons from the conjugated polymer p(g₄2T-T). The dopant monoanions could be detected with EPR spectroscopy because their signal appears at a lower magnetic field than the signal from the polymer polaron. The absence of a clear monoanion signal in EPR spectra of the polymer co-processed with Mo(tfd-COCF₃)₃ suggests that only dianions are present, i.e., the dopant underwent two electron transfer events with the polymer. The viability of double doping was confirmed by sequential doping of polymer films with the monoanion salt [Mo(tfd-COCF₃)₃]^{2−}Et₄N⁺, which gave rise to a conductivity of $\sigma = (11.9 \pm 0.3) \text{ S cm}^{-1}$ and EPR spectra that did not feature any signal from anions, indicating that each anion had been converted into a spin-silent dianion by accepting one electron from the polymer. Analysis of the UV–vis–NIR spectra of co-processed samples indicated an ionization efficiency of 200% for use of up to 10 mol % Mo(tfd-COCF₃)₃. It can be anticipated that double doping of polymers with suitable monomeric dopants may allow us to achieve high charge-carrier densities while reducing the number of counterions and hence their collective impact on the nanostructure of semiconductor films.

EXPERIMENTAL SECTION

Materials. TQ1 (weight- and number-average molecular weights, $M_n = 76 \text{ kg mol}^{-1}$ and PDI = 2.6), p(g₄2T-T) ($M_n = 24 \text{ kg mol}^{-1}$, PDI = 3.3), Mo(tfd-COCF₃)₃, and [Mo(tfd-COCF₃)₃]^{2−}Et₄N⁺ and [Mo(tfd-COCF₃)₃]^{2−}(Et₄N⁺)₂ were prepared according to previously reported procedures.^{15,27,36} Poly(ethylene oxide) (PEO, $M_w = 200 \text{ kg mol}^{-1}$), [TPA-Br₃]⁺[SbCl₆][−] (Magic Blue), dichloromethane (DCM, purity > 99.9%), and anhydrous acetonitrile (AcN, purity > 99.8%) were purchased from Sigma-Aldrich. Chlorobenzene (CB, purity > 99%) and chloroform (CHCl₃, purity > 99.8%) were obtained from Fisher Scientific. All commercial solvents and the Magic Blue dopant were used as received and without further purification.

Sample Preparation. Polymers were dissolved with a concentration of 10 g L^{-1} (TQ1 in CB and p(g₄2T-T) in CHCl₃) and spin-cast onto poly(ethylene terephthalate) (PET) films, yielding a film thickness of 50–100 nm. The polymer films were sequentially doped with Magic Blue dissolved in AcN/CHCl₃ (3:1, v/v; 0.45 g L^{-1}), Mo(tfd-COCF₃)₃ in AcN/CHCl₃ (3:1, v/v; 2.5 g L^{-1}), and [Mo(tfd-COCF₃)₃]^{2−}Et₄N⁺ in AcN/DCM (3:2, v/v; 2.5 g L^{-1}). The dopant solutions were added on top of the films for 30 s, spun off, and finally the films were rinsed with anhydrous AcN to remove any excess dopant on top of the film. To prepare co-processed p(g₄2T-T)/Mo(tfd-COCF₃)₃ samples, p(g₄2T-T) and Mo(tfd-COCF₃)₃ were dissolved in AcN/CHCl₃ (1:1, v/v; 15 and 5 g L^{-1}). Appropriate volumes of AcN/CHCl₃ (1:1, v/v) were then added to the p(g₄2T-T) solutions before the addition of the Mo(tfd-COCF₃)₃ solution to maintain a polymer concentration of $5\text{--}7.5 \text{ g L}^{-1}$ in each polymer/dopant solution while varying the concentration of Mo(tfd-COCF₃)₃.

The co-processed polymer/dopant solutions were spin-cast onto PET films, yielding a film thickness of 80–200 nm. The calculations of molar percentage were based on the molar mass of the polymer repeat unit and the molar mass of the dopant. $\text{Mo}(\text{tfd-COCF}_3)_3$, $[\text{Mo}(\text{tfd-COCF}_3)_3]^- \text{Et}_4\text{N}^+$, and $[\text{Mo}(\text{tfd-COCF}_3)_3]^{2-} (\text{Et}_4\text{N}^+)_2$ were dissolved together with PEO in DCM (4 and 16 g L⁻¹), and the solutions were then spin-cast onto PET films.

UV-vis-NIR Absorption Spectroscopy. Measurements of liquid and solid samples were performed with a PerkinElmer Lambda 1050 spectrometer. $\text{Mo}(\text{tfd-COCF}_3)_3$, $[\text{Mo}(\text{tfd-COCF}_3)_3]^- \text{Et}_4\text{N}^+$, and $[\text{Mo}(\text{tfd-COCF}_3)_3]^{2-} (\text{Et}_4\text{N}^+)_2$ solutions for UV-vis-NIR absorption measurements were prepared at a concentration of 1 g L⁻¹ in DCM.

Electron Paramagnetic Resonance. The polymer films coated on PET films were cut to a size of 25 mm × 3 mm and then sealed under nitrogen in a quartz EPR tube. EPR spectra were recorded at room temperature using a Magnetech MS5000 spectrometer (Freiberg Instruments, now Bruker Biospin). All spectra were corrected for the same microwave frequency (9.6 GHz). Data processing and analysis have been performed using the cwepr Python package.^{37,38}

Electrical Characterization. The electrical resistivity was measured with a four-point probe setup from Jandel Engineering (cylindrical probe head, RM3000) using collinear tungsten carbide electrodes with an equidistant spacing of 1 mm that were held down with a constant weight of 60 g. The electrical conductivity σ was then calculated according to $\sigma = ((V/I)kt)^{-1}$, where V is the voltage, I is the current, $k = 4.53$ is a geometrical correction factor, and t is the thickness.

■ ASSOCIATED CONTENT

SI Supporting Information

The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/acs.chemmater.2c01040>.

UV-vis-NIR spectra as well as angular-dependent and nonsmoothed EPR spectra (PDF)

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Notes

The authors declare no competing financial interest.

EPR raw data and analysis “recipes” used to analyze the EPR data are available online via Zenodo (DOI: [10.5281/zenodo.6592331](https://doi.org/10.5281/zenodo.6592331)).

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