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Conjugated metallo-supramolecular polymers containing a phosphole unit

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Supporting Information Placeholder

ABSTRACT: A new building block, **TPT**, composed of the substituted phosphole ring surrounded by two thiophene rings with 2,2':6',2''-terpyridine-4'-yl (*tpy*) end-groups, is prepared and assembled with metal ions (Co^{2+} , Cu^{2+} , Fe^{2+} , Ni^{2+} and Zn^{2+}) into metallo-supramolecular polymers (MSPs) and properties of both **TPT** and the MSPs are compared with those of their counterparts with terthiophene central blocks. A distinct red shift of the UV/vis band about 60 to 100 nm proves the decrease in the bandgap energy due to replacing the thiophene-2,5-diyl with a phosphole-2,5-diyl central unit, which is due to the lowered aromaticity of phosphole ring compared to the aromaticity of thiophene ring. Assembling of **TPT** with metal ions gives oligomeric chains comprising up to ten unimeric units in dilute solutions. MSPs with Fe^{2+} and Ni^{2+} ion couplers exhibit very slow constitutional dynamics, while those with Cu^{2+} and Zn^{2+} ion couplers quite fast constitutional dynamics. A metal to ligand charge transfer is observed only for Fe^{2+} -MSPs, while luminescence only for Zn^{2+} -MSPs, mainly at an excess of Zn^{2+} ions, which indicates a positive effect of the end-capping of MSP chains with these ions.

INTRODUCTION

The development of semiconducting organic and organic/inorganic materials for applications in photovoltaics, light-emitting devices, field effect transistors and molecular electronics is an area of permanently high interest.^{1–3} Supramolecular chemistry based on the self-assembly of polymers from properly designed building blocks has become an efficient tool for the construction of soft functional materials having specific optical and/or redox properties.⁴ Metallo-supramolecular polymers (MSPs) based on building blocks with chelating end-groups linked to chains through reversible metal-ligand coordination represents an important class of soft organic/inorganic supramolecular materials.^{5,6} These macromolecular metal-containing systems may exhibit the properties of standard organic polymers (viscosity, processability, etc.), but the incorporation of metallic complexes in the polymer chain opens new perspectives giving access to magnetic, redox, optical, electrochromic, or specific mechanical properties; In addition, they can reversibly disassemble and reassemble at increased temperature in solution and exchange metal ions (ion couplers) and/or building blocks with other MSPs under these conditions leading to the development of environment-adaptable materials.^{7,8} Owing to their unique properties, MSPs are promising systems for the development of functional self-organized architectures with tunable properties.

The physical properties of these constitutionally dynamic coordination polymers or, in short, dynamers⁹ can be tuned by (i) the structure of central parts of their unimers,¹⁰ (ii) choice of their chelate end-groups (ion selectors), and (iii) choice of ions (ion couplers) connecting unimers species into chains.^{11,12} The most popular chelating end-group is the tridentate

2,2':6',2''-terpyridine because it is able to form well-defined octahedral complexes with a wide range of transition metal cations. Furthermore, the structure of the central part of the unimer plays an important role since it can introduce a great variety of structural arrangements and/or physicochemical properties (e.g., viscosity, optical, mechanical properties, etc.) to the targeted metallo-polymer. For example, various conjugated MSPs have been prepared from linear unimers with oligophenylene^{13–15}, oligofluorene^{16–18} and oligothiophene^{19–24} central blocks capped with tridentate 2,2':6',2''-terpyridine-4'-yl end-groups assembled with metal ions such as Ru^{2+} , Fe^{2+} , Zn^{2+} and Co^{2+} .^{11,12,25} In general, central blocks of these unimers are composed of σ -bonded highly aromatic rings, which doesn't lead to an efficient delocalization of π -electrons along the whole molecule owing to the rather high autonomy of strongly aromatic systems. Hence the introduction of less aromatic conjugated rings into the unimer structure can be expected to give the desired unimers and MSPs with significantly decreased energy band-gaps. For this purpose, the phosphole ring is a good candidate since it possesses a weak aromaticity and a low-lying LUMO level. Effectively, the phosphorus atom in the ring namely adapts a pyramidal geometry²⁶ where its lone electron pair almost does not interact with the endocyclic diene π -electrons. Therefore, delocalization of electrons in a phosphole ring stems from the so called σ - π hyperconjugation involving the exocyclic P-R σ -bond and the endocyclic diene π -system.²⁷ As a result, phosphole is less aromatic compared to pyrrole and thiophene.²⁸ Furthermore, mixed phosphole-thiophene derivatives are the most widely investigated phosphole based π -conjugated systems^{27,29–40} since (i) this combination of heteroles allows to obtain low

HOMO-LUMO gap derivatives due to maximized orbital interactions, (ii) a built in phosphole ring represents a new center for potential structural variations of such chains mainly by choice of the P-atom substituent R, and oxidation of the ring P-atom (e.g., to P=S or P=O), and (iii) it is possible to exploit the rich chemistry of thiophene to readily obtain novel derivatives *via* Pd-catalyzed C-C coupling reactions or electropolymerization.^{27,29,36,41}

Considering these appealing functions of mixed phosphole-thiophene derivatives, there is a great interest for the development of unimers and MSPs based on mixed phosphole-thiophene structures in order to gain more insight into structure-property relationship and to further develop MSPs for specific applications.

MSPs composed of a unimer that comprises the phosphole-2,5-diyl unit has not yet been reported. Only a simple complex of $[\text{ZnL}_2]^{2+}$, where the ligand L is mono-*tpy* derivative of dithieno[3,2-b:2',3'-d]phosphole, has been reported by Bai et al.⁴² Therefore, in order to examine the above-outlined idea, a new unimer composed of a substituted phosphole-2,5-diyl unit surrounded with two thiophene-2,5-diyl units capped with *tpy* end-groups has been prepared, characterized, and assembled with various metal ions (Co^{2+} , Cu^{2+} , Fe^{2+} , Ni^{2+} and Zn^{2+}) into corresponding MSPs (**Scheme 1**). Optical spectral properties of this new phosphole unimer, hereafter denoted **TPT**, and from it derived MSPs are discussed and compared with the properties of recently reported related unimers (**Chart 1**) and MSPs with terthiophene central blocks.²²

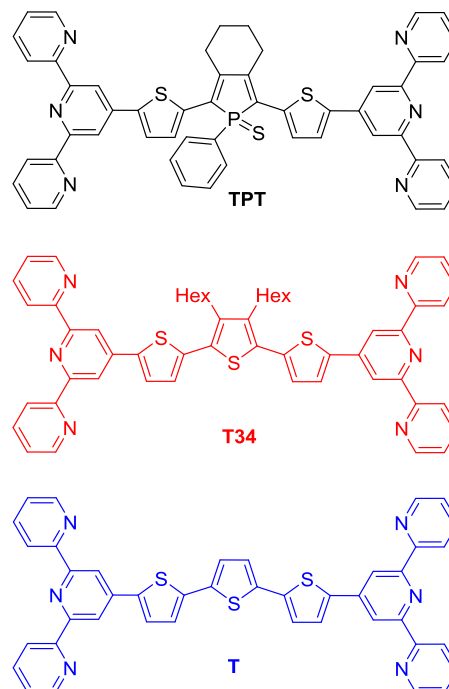
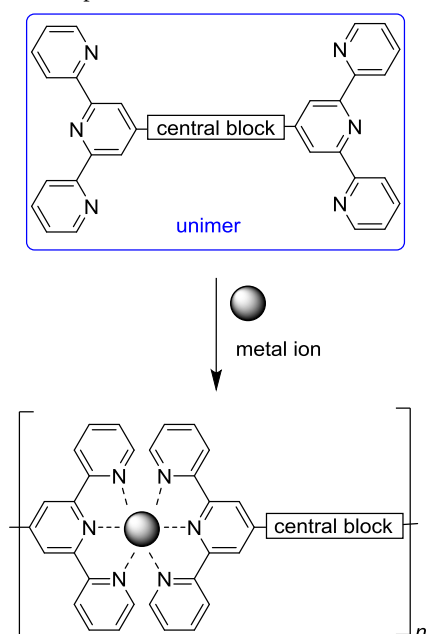


Chart 1. Structures of unimers under discussion

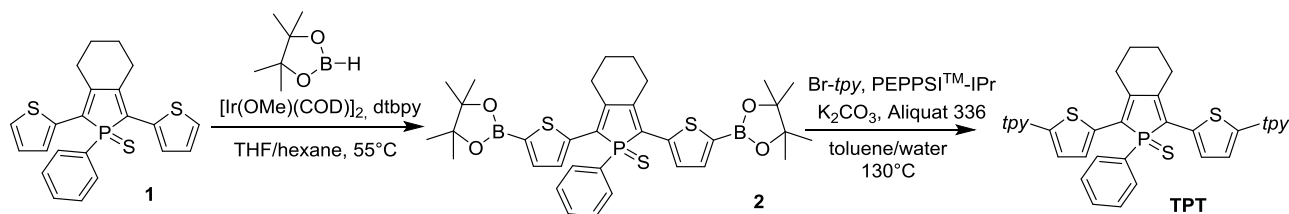


Scheme 1. Metallo-supramolecular polymerization of bis-terpyridines

RESULTS AND DISCUSSION

Synthesis of phosphole unimer TPT and its polymers

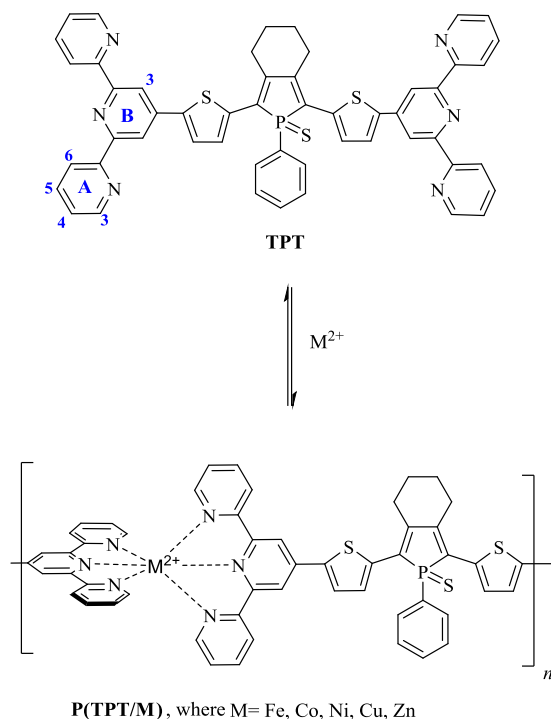
The **TPT** unimer was synthesized from 1-phenyl-2,5-bis(thiophen-2-yl)thioxophosphole (**1**) by attaching the *tpy* end-groups using a Suzuki coupling strategy (**Scheme 2**). The central block **1**, which has been previously described,³⁶ was borylated using a direct C-H activation approach in the presence of an iridium catalyst⁴³ (**Scheme 2**). The reaction conditions used for this C-H reaction are those currently applied for the borylation of the thiophene ring. In addition, the reaction could be monitored by phosphorus NMR since the signals of the borylated species and starting molecule occur at slightly different chemical shifts (^{31}P NMR: $\delta = +52.1$ ppm for compound **1** and $+51.8$ ppm for compound **2**) (**Scheme 2**). When the characteristic chemical shift of compound **1** disappears, the mixture is diluted with dichloromethane (20 mL) and water (20 mL). The compound **2** is then isolated from the organic phase by evaporation of the organic solvent followed by re-precipitation.



Scheme 2. Synthesis of unimer TPT

The α,ω -bis(borylated) central block **2** was then reacted with Br-*tpy* using modified Suzuki coupling conditions. The best yields were obtained using the PEPPSITM-IPr catalyst in a toluene/water (1/1, v/v) biphasic system in the presence of Aliquat 336 phase transfer catalyst and potassium carbonate (**Scheme 2**). Though the reaction mixture was kept at high temperature and in alkaline environment, the formation of a phospholene type side product, which can sometimes takes place at such temperatures,⁴⁴ was not observed under the applied conditions. The unimer **TPT** was purified by column chromatography and isolated as an air-stable powder in modest yield (15%). (For overall scheme of the synthesis see **Scheme S1** in the Supporting Information).

Metallo-supramolecular polymers were prepared by spontaneous assembly of **TPT** with metal ions in a chloroform/acetonitrile solvent mixture (1/1 by vol.) at room temperature. A solution containing equimolar amounts of **TPT** and particular metal(II) (Fe, Co, Cu, Ni, Zn) perchlorate was allowed to equilibrate for one day and the formed MSP isolated by evaporation of solvents. Prepared MSPs are further denoted by abbreviations: **P(TPT/Fe)**, **P(TPT/Co)**, **P(TPT/Cu)**, **P(TPT/Ni)** and **P(TPT/Zn)** or, in general, **P(TPT/M)** (**Scheme 3**) and their spectral characteristic are discussed below.



Scheme 3. Assembly of unimer TPT and metal ions into MSPs

Optical and redox properties of the unimer TPT

In order to have an understanding of the characteristics of this novel π -conjugated unimer **TPT**, we studied its optical and electrochemical properties. The UV/vis absorption spectrum in solution of **TPT** (**Figure 1a**) shows a band centered at 281 nm that is mainly contributed by transitions in *tpy* end-groups⁴⁵ and a band centered at $\lambda_{\text{abs}} = 481$ nm that is contributed by HOMO to LUMO transitions that are spread over the unimer central block and adjacent central rings of *tpy* end-groups^{20-22,24,42,45} (see diagrams obtained by the DFT calculations, **Figure S1**). The fluorescence spectrum of **TPT** (**Figure 1b**) shows a structured emission band centered at $\lambda_{\text{F}} = 603$ nm. As expected, the presence of an extended π -conjugation in **TPT** induces a significant red-shift in the absorption and emission maxima compared to the corresponding bands of the central block without *tpy* end-groups ($\lambda_{\text{abs}} = 432$ nm; $\lambda_{\text{F}} = 548$ nm).³⁴

Furthermore, the UV/vis as well as the luminescence band of **TPT** is substantially red shifted compared to the corresponding band of **T** as well as **T34** (see also **Table 1**, **Figure 1**), two related terthiophene unimers possessing identical number of conjugated double bonds and similar structure, only the phosphole central ring is formally replaced by the unsubstituted (in unimer **T**) or dihexylated (**T34**) thiophene ring. These observations show a substantial decrease in the band-gap energy caused by replacing the middle high aromatic thiophene ring in the unimer central block with a low aromatic phosphole ring. The decreased energy bandgap of **TPT** was confirmed by cyclic voltammetry, 2.74 eV for **TPT** and 3.08 eV for **T**.²¹ The DFT calculations provided slightly lower band-gap energy values for both **TPT** (2.62 eV) and **T** (2.97 eV), but the same band-gap energy difference between **TPT** and **T** equal to ca -0.25 eV. Comparing the thin film spectra of the unimers, **TPT** shows the main UV/vis band ($\lambda_{\text{abs}} = 520$ nm) considerably red shifted compared to the bands of **T34** ($\lambda_{\text{abs}} = 455$ nm) and **T** ($\lambda_{\text{abs}} = 426$ nm), thus confirming a decrease in the band-gap energy even in the thin film of **TPT**. (The fact that **T** shows λ_{abs} lower than **T34** has been explained by the damping of thermal twisting of thiophene rings by the attached hexyl groups which are partly anchored in solid films²²). The observed red-shift in UV/vis might but need not directly relate to increased delocalization of electrons. The results of DFT calculations (**Figure S1a**) namely show that the HOMO of **TPT** is delocalized rather equally over all three central rings, while the LUMO is localized primarily on the phosphole ring with more minor contributions from thiophene rings. This indicates that the observed spectral red shift is very likely contributed by charge-transfer interactions within **TPT** molecules.

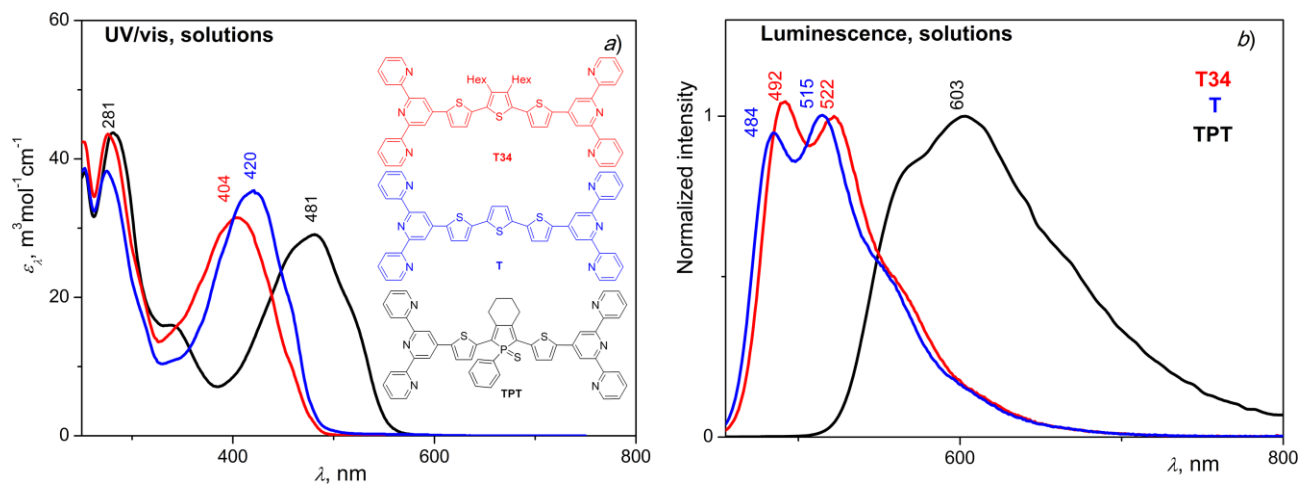


Figure 1. Solution UV/vis (a) and fluorescence (b) spectra of the phosphole unimer TPT and terthiophene unimers T, T34.

Table 1. Spectroscopic data for unimers TPT, T and T34 and related MSPs.

	λ_{abs} , nm		λ_{F} , nm		Φ ,%		$\Delta\nu_{\text{s}}$, cm ⁻¹	
	solution	film	solution	film	solution	film	solution	film
TPT	482	520	603	685	19	0.7	4 160	4630
P(TPT/Zn)	507	509	634	641	-	0.5	3 950	4 100
P(TPT/Ni)	509	526	-	-	-	-	-	-
P(TPT/Cu)	514	531	-	-	-	-	-	-
P(TPT/Co)	529	554	-	-	-	-	-	-
P(TPT/Fe)	491	499 (λ_{MLCT} =625 nm)			-	-	-	-
T	420*	426	484/515*	730			3 150/4 390*	9780
T34	404*	455	492/522*	600			4 430/5 600*	5310
P(T/Zn)	479*	469*	635*	686*			5 130*	6 750*
P(T34/Zn)	480*	495*	620*	640*			4 700*	4 600*

Solvent chloroform/acetonitrile (1/1 by vol.); films casted on a quartz glass; Φ luminescence quantum efficiency; $\Delta\nu_{\text{s}}$ Stokes shift calculated by formula: $1/\lambda_{\text{abs}} - 1/\lambda_{\text{F}}$; *data taken from ref.²² (solutions in THF); λ_{MLCT} stands for the wavelength of the absorption maximum of the band of the metal-to-ligand charge transfer (MLCT) transitions (an electron transfer from a metal atom orbital to the ligand π^* orbital). For a comparison, we provide here λ_{abs} values for chloroform/acetonitrile solutions of $[\text{M}(\text{T-tpy})_2]^{2+}$ complexes (where **T-tpy** stands for 4'-(2-thienyl)-2,2':6',2''-terpyridine) of corresponding M^{2+} ions: 351 nm (for M = Ni), 352 nm (Cu), 333 nm (Co) (all our new data); 344 nm (Zn) and 320 nm (Fe; $\lambda_{\text{MLCT}} = 551$ nm), both from ref.⁴⁵

The maximum of the thin film photoluminescence band of unimer **TPT** ($\lambda_F = 685$ nm, see **Figure S2**) is red shifted compared to that of unimer **T34** ($\lambda_F = 597$ nm) but blue shifted compared to that of unimer **T** ($\lambda_F = 731$ nm), for which the Stokes shift approaches 10^4 cm⁻¹ (i.e., 1.24 eV; see **Table 1**). Such large Stokes shift proves extensive conformational relaxation of excited **T** molecules, which could be due to their regular packing in aggregates allowing so high relaxation, as it has been reported for oligothiophenes.⁴⁶ Large Stokes shifts accompanied with low fluorescence quantum yields are typical of H-aggregates.⁴⁷ To examine this hypothesis more in detail, the photoluminescence spectra of **T**, **T34** and **TPT** were recorded on samples of unimers dispersed at various concentrations in KBr pellets as matrix.

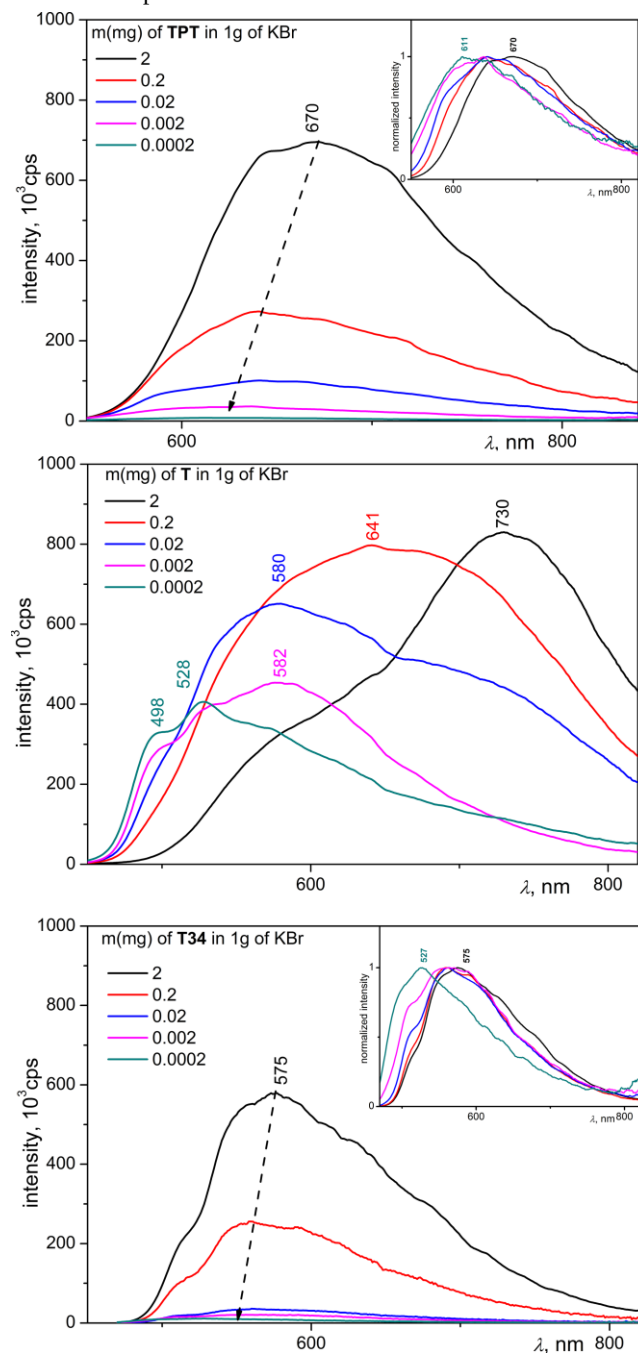


Figure 2: The solid-state luminescence spectra of unimers **T**, **T34** and **TPT** dispersed at various concentrations in KBr pellets; insets show normalized spectra.

The unimer **T** showed a gradual blue-shift of the emission band (up to -232 nm) and a relatively low decrease in the band intensity with decreasing concentration of **T** in the matrix (**Figure 2**). On the contrary, emission maxima of dispersed unimers **T34** and **TPT** showed a small blue-shift (up to 60 nm) and rather high intensity decrease upon dilution. These observations are consistent with the hypothesis of regular packing of the **T** molecules and poor packing of the **TPT** and **T34** molecules in solid films. For the unimer **TPT**, the pyramidal geometry of the phosphorus atom precludes a regular cofacial molecular packing in the solid state.³⁴ While, for unimer **T34**, the hexyl side groups can be regarded as the reason for irregular molecular packing.

Optical properties of metallo-supramolecular polymers

The UV-Vis spectra recorded for the different synthesized MSPs unambiguously show the formation of polymer chain including bis(*tpy*)M²⁺ units. According to the optical spectral features (see **Figure 3** and **Table 1**), the MSPs derived from **TPT** can be divided in three categories. The first category concerns the **P(TPT/Co)**, **P(TPT/Ni)** and **P(TPT/Cu)** MSPs, which show neither luminescence nor a band of the metal-to-ligand charge transfer (MLCT) transitions. Only the absorption band of the π - π^* transitions is red-shifted about 27 to 47 nm (change in band-gap energy, E_g , from -0.14 to -0.23 eV) in solutions and 6 to 34 nm (-0.03 to -0.15 eV) for films compared to those of unimer **TPT**. The highest band shift as well as the band broadening is observed for **P(TPT/Co)**, which might be due to the increased contribution of the metal centered d-d transitions, as Kurth et al. have suggested for the cobalt(II) MSP of 1,4-bis(*tpy*)benzene.⁴⁸ The polymer **P(TPT/Zn)**, representative of the second category, shows luminescence but not the MLCT band. The position as well as intensity of the optical band of **P(TPT/Zn)** is similar to the bands of the first category of MSPs for the spectra taken from solution. However, the solid state spectra show the main-chain band blue shifted (-11 nm, i.e., 0.05 eV) and attenuated compared to the band of free **TPT**. Finally, the UV/vis absorption spectrum of **P(TPT/Fe)**, representative of the third category, shows a very intense band of the MLCT transitions which are typical of [Fe(*tpy*)₂]²⁺ species.^{25,49} This band occurs at the wavelength $\lambda_{MLCT} \cong 625$ nm that is about 20 to 30 nm red-shifted compared to λ_{MLCT} of Fe-MSPs derived from α,ω -bis(*tpy*)ter- and quater-thiophenes.²³ The difference between positions of the main-chain bands of **P(TPT/Fe)** and free **TPT** is positive though very small (+9 nm) for the solution but negative (-21 nm) for the solid-state spectra. In addition, this band intensity reaches only less than half of the intensity of corresponding bands for the type (i) **P(TPT/M)**s. These differences in the main-chain band position and intensity can be explained by the shift of some main-chain electronic transitions in **P(TPT/Fe)** to the spectral region of the MLCT band (*vide infra* Raman spectra).

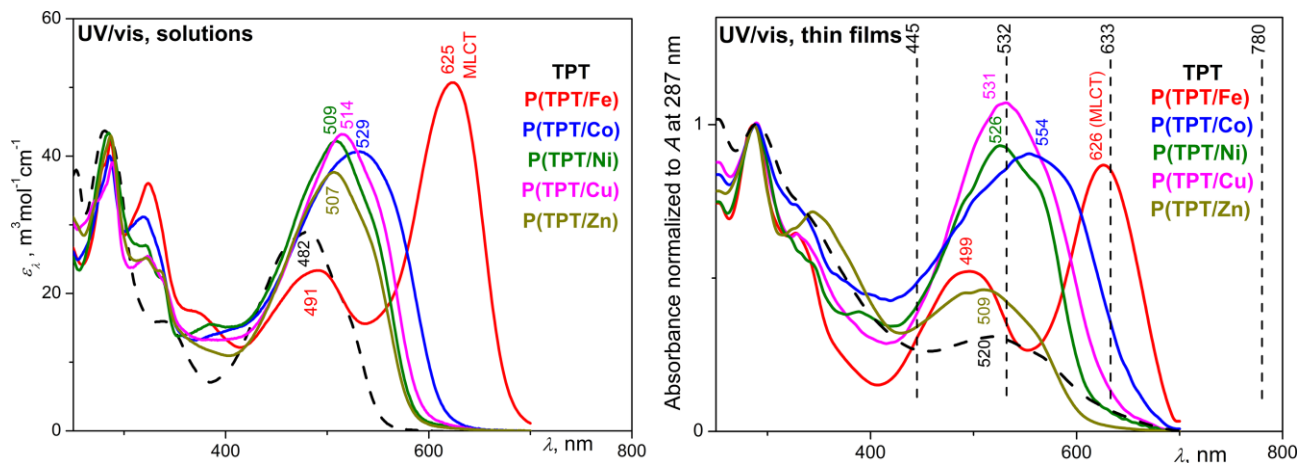


Figure 3: Solution and solid-state UV/vis spectra of unimer TPT and its MSPs. Doted verticals mark excitation wavelengths used in the Raman spectra measurement.

Vibrational spectra of TPT and P(TPT/M)

Raman spectra can provide valuable information related to the electronic structure of conjugated compounds due to the Raman resonance effect which consists in the selective enhancement of vibrational bands pertaining to groups that absorb light at wavelengths around the excitation wavelength λ_{exc} .^{50–52}

For the unimer **TPT**, only off resonance Raman spectrum (taken with $\lambda_{\text{exc}} = 780$ nm which does not match any absorption band, **Figures S3 and S4**) could be obtained owing to strong luminescence of **TPT** when using excitations at lower wavelengths ($\lambda_{\text{exc}} = 435, 532$ and 633 nm). The DFT calculated spectrum is in a good agreement with the experimental one (**Figure S3**). As can be expected for a large molecule composed of different constitutional units, the calculations revealed that all strong spectral bands consist of various stretching and deformation vibrational modes of the units. The spectrum of **TPT** is dominated by the bands of the thiophene and phosphole units ($1400 - 1500 \text{ cm}^{-1}$) while the bands of *tpy* end-groups ($1520 - 1600 \text{ cm}^{-1}$) are very weak. Such spectral pattern is typical of the off-resonance spectra of all α,ω -bis(*tpy*)oligothiophenes.^{17–21,39} Compared to the spectra of **T** and **T34**, the spectrum of **TPT** shows enhanced bands in the region from 1400 to 1460 cm^{-1} .

Incorporation of the **TPT** unimer into a metallo-supramolecular polymer results in the occurrence of stretching (1610 cm^{-1}) and breathing ($\sim 1030 \text{ cm}^{-1}$) Raman bands of coordinated *tpy* end-groups (that underwent an *anti*-to-*syn* conformational transition during coordination to the metal ions, see Fig. 3), and restructuring and slight band shifts assigned to the central block (**Figure S3**). The off-resonance spectra of **P(TPT/M)** are similar to each other, except for the spectrum of **P(TPT/Fe)** which shows stronger bands for the stretching (1610 cm^{-1}) and in-plane deformation (1364 cm^{-1}) modes of pyridine rings and the ring-breathing band shifted to 1039 cm^{-1} .

The resonance Raman spectra of **P(TPT/Fe)** taken with different excitation wavelengths, λ_{exc} , each matching different region of the polymer optical spectrum (**Figure 2**) are shown in **Figure S5**. The spectrum taken with $\lambda_{\text{exc}} = 445$ nm is dominated by the bands of the thiophene and phosphole rings and the *tpy* end-group bands being almost absent. This proves that the optical band of **P(TPT/Fe)** at around 445 nm is largely contributed by electronic transitions within the central blocks.

In contrast, the spectrum taken with $\lambda_{\text{exc}} = 532$ nm displays attenuated bands of the central-block rings, strong bands of *tpy* end-groups and a series of combined bands and overtones in the region above 2000 cm^{-1} . Such spectral pattern is typical of $\text{Fe}^{2+}(\text{terpyridine})_2$ species²⁴ showing $\lambda_{\text{MLCT}} \cong 550 \text{ nm}$.^{45,53} This proves that the optical absorption of **P(TPT/Fe)** at 532 nm is dominantly contributed by the electronic transitions in coordinated *tpy* groups.⁵³ Perhaps the most interesting is the Raman spectrum of **P(TPT/Fe)** taken with $\lambda_{\text{exc}} = 633$ nm, matching the maximum of the MLCT band. This spectrum shows strong bands of the *tpy* groups as well as the central block units. This proves that the electronic transitions from the central blocks of enchaind **TPT** units substantially contribute to the MLCT absorption band. This simultaneously explains why the main UV/vis band of the **TPT** units enchaind in **P(TPT/Fe)** is very weak and blue-shifted to 500 nm .

The IR spectra of **TPT** and related polymers are shown in **Figure S6**, and the experimental and DFT-calculated spectra of **TPT** are compared in **Figure S3**. In the IR spectra, the incorporation of **TPT** into a metallo-polymer is mainly reflected by the occurrence of a strong stretching band from the *tpy* units at 1610 cm^{-1} and the development of a strong broad and unresolved band in the region of deformation modes ($1200 - 1000 \text{ cm}^{-1}$); a similar band has not been observed for MSPs derived from α,ω -bis(*tpy*)oligothiophenes.^{17–20} The IR spectra of MSPs are similar to each other except for the spectrum of **P(TPT/Fe)**, which should be related to the presence of the MLCT transitions in **P(TPT/Fe)** molecules.

Assembly of polymers in solutions

Assembly of MSPs from **TPT** and metal ions (M^{2+}) in a chloroform/acetonitrile (1/1 by vol.) solvent mixture was monitored by the UV/vis and luminescence spectroscopies. A set of solutions of the constant **TPT** concentration of $2 \cdot 10^{-5} \text{ M}$ and the ion-to-unimer mole ratio $r = [\text{M}^{2+}]/[\text{TPT}]$ increasing from 0 to 3, was prepared for each M^{2+}/TPT couple and the solutions were allowed to equilibrate for one day before monitoring their spectra. The UV/vis spectral changes (see **Figures S7** and dependences in **Figure 4**) indicate three stages of assembling, which are usually observed for similar systems.^{23,24}

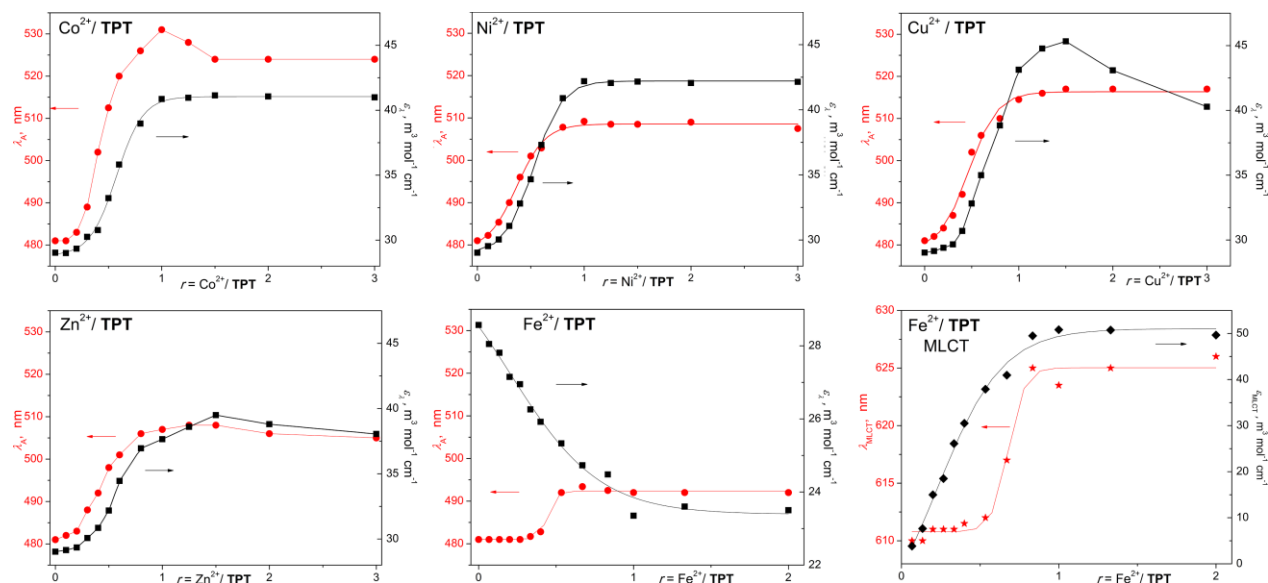


Figure 4: Wavelength (λ_A) and molar absorption coefficient (ϵ) of the main UV/vis band as a function of the ion-to-unimer mole ratio ($r = [M^{2+}]/[TPT]$) of the studied systems and *dto* for the MLCT band of the Fe^{2+}/TPT system.

The first stage of the MSP assembly appears for r up to ca 0.5 and is characterized by: (i) narrowing and a small red shift (4 to 7 nm) of the band of *tpy* groups (281 nm for free **TPT**), (ii) red shift (ca 20 nm) and enhancement of the absorption band involving transitions centered on the central-block of the **TPT** units (except for the Fe^{2+}/TPT system showing a decrease in the intensity), and (iii) appearance of three isosbestic points (IP) at around 300, 415 and 480 nm for each M^{2+}/TPT system. The presence of IPs indicates a direct conversion of the **TPT** molecules into another defined species which, according to the stoichiometry, should be “butterfly dimer” species **TPT**- M^{2+} -**TPT**. The second stage of assembly (r from ca 0.5 to 1) is characterized by the continued red shift and intensity increase of the main absorption band, but without occurrence of a clear IP. These features are typical of the prolongation of MSP chains.^{22–24,54,55} The third stage of the assembly ($r > 1$) is characterized by a small change in the spectral pattern that can be attributed to end-capping of MSP chains with metal ions and/or equilibrium dissociation of longer chains to shorter ones induced by superfluous metal ions.^{42,56,57}

As indicated above, the Fe^{2+}/TPT system behaves differently from the other ones: only a small red shift (+ 11 nm) and a continuous attenuation of the main optical band is observed if the value of r increases, but together with progressive development of the MLCT band. The main band attenuation can be ascribed to the shift of corresponding electronic transitions to the MLCT band as suggested previously based on the resonance Raman spectra of **P(TPT/Fe)**. Moreover, the MLCT band appears at 612 nm for dimers while it appears at 625 nm for mature **P(TPT/Fe)**, which supports this hypothesis.

The luminescence spectra show continuous attenuation of the luminescence with increased ratio r to its total quenching at $r \approx 0.8$. This behavior can be ascribed to the fact that metal ions with an open shell electronic structure usually exhibit strong metal-to-ligand orbital interaction resulting in non-emissive quenching of electronically excited states.⁵⁸ The only exception is the Zn^{2+}/TPT system for which the luminescence starts to intensify for r values above 0.6, finally becoming stronger than that of the free unimer (**Figure S8**) at $r > 1$,

where end-capping of the Zn-MSP chains take part. This suggests that the luminescence of **P(TPT/Zn)** is mainly contributed by terminal *tpy* groups capped with Zn^{2+} ions, the contribution of internal -*tpy*- Zn^{2+} -*tpy*- linkages being less important.

Dynamics and degree of polymerization of MSPs in solution

The size exclusion chromatography (SEC) method can provide qualitative information on the constitutional dynamics of MSPs in the solvent used as a mobile phase.^{24,59} If an MSP with fast establishing assembly equilibrium (fast constitutional dynamics) is analyzed by SEC, only a peak for the free unimer is observed due to the fast disassembly of MSP upon multiple dilution of the injected solution inside columns (injected 20 μ L, elution volume of ca 40 mL). However, if an MSP with slow constitutional dynamics is injected, its chains do not completely or substantially dissociate during the SEC analysis and an SEC record resembling the record of an oligomer or polymer is obtained, from which the average degree of polymerization can be calculated. If the SEC device is fitted with a diode-array UV/vis detector, it also provides the spectra of SEC fractions from which the type of chains present in the fractions (dimers, longer or end-capped chains) can be identified or at least ascertained.

For the SEC analysis, one day old solution of **TPT** (0.5 mM) and M^{2+} ions (r from 0 to 1.3) was injected and a chloroform/acetonitrile (1/1 by volume) solvent mixture containing $Bu_4N^+PF_6^-$ (0.5 % by weight; to suppress aggregation) was used as the mobile phase. Samples with Co^{2+} ions were not analyzed since they provided opalescent solutions indicating the presence of aggregates. The Zn^{2+}/TPT and Cu^{2+}/TPT systems provided SEC records showing just a peak of free **TPT**, which proves fast constitutional dynamics for **P(TPT/Zn)** and **P(TPT/Cu)**. This agrees with earlier observations made for other MSPs and complexes derived from *tpy* and Zn^{2+} or Cu^{2+} ions.⁶⁰ The systems Fe^{2+}/TPT and Ni^{2+}/TPT provided SEC records similar to covalent polymers proving slow constitutional dynamics of these MSPs. Good resolved SEC records were given by the systems with $r \leq 1$ (**Figure 5** and **S9**). The systems with $r \geq 1$ gave poorly resolved rec-

ords showing a decrease in the elution peak area with increasing value of r , which should be ascribed to the retention of chains end-capped with M^{2+} ions in the SEC columns.

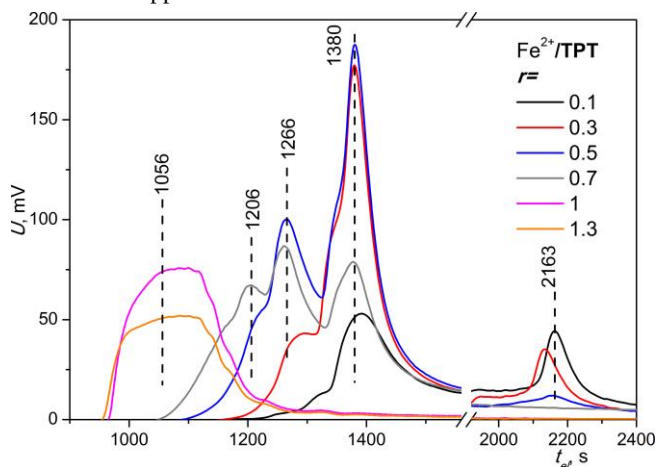


Figure 5. SEC records for different ratios Fe^{2+}/TPT

The UV/vis spectra of SEC fractions within Fe^{2+}/TPT system (**Figure S10**) show a continuous development of the spectral pattern which is typical of long **P(TPT/Fe)** chains for the first eluted SEC fractions to the pattern for butterfly dimers in the last fraction. The presence of longer chains in the systems with a stoichiometric lack of Fe^{2+} ions ($r = 0.2$ and 0.5) can be attributed to the transiently locally increased concentration of the ions during mixing their solution with the solution of **TPT**. Another spectral pattern difference is seen for the SEC fractions with identical elution times but pertaining to systems of different composition ratio r (differences in the fractions of dimers are also shown in **Figure S12**). In this case the differences can be ascribed to the end-capping of molecules with Fe^{2+} ions.²⁴

The number-average, $\langle X \rangle_n$, and weight-average, $\langle X \rangle_w$, degrees of polymerization of the **P(TPT/Fe)** chains formed in the solution with $r = 1$ have been estimated using the PMMA calibration: $\langle X \rangle_n = 4$ and $\langle X \rangle_w = 5.7$. These values should be underestimated owing to the retention of polymer higher fractions in the SEC columns. However, a recent time-resolved transient optical absorption study on solutions of **P(Zn/T)** and its derivative **P(Zn/T16)** (with hexyl groups on carbons next to the *tpy* groups), in which equilibrium fractions of the free unimers **T** and **T16** have been determined,⁶¹ gave values of $\langle X \rangle_n = 11$ and $\langle X \rangle_w = 21$ for **P(Zn/T)** and $\langle X \rangle_n = 5$ and $\langle X \rangle_w = 9$ for **P(Zn/T16)** calculated using the Carothers equations. The latter values are close to those found for **P(TPT/Fe)**.

CONCLUSIONS

A novel linear conjugated unimer, **TPT**, of the bis(*tpy*) family has been prepared with the aim to verify the theory that a replacement of a thiophene unit with the phosphole unit in a terthiophene central block gives a unimer with significantly decreased band-gap energy. A feasible synthetic path to **TPT** has been developed and the photophysical properties of **TPT** have been compared to related bis(*tpy*)terthiophene unimers. The obtained results confirm that the substitution of thiophene ring with a phosphole ring significantly enhances the delocalization of electrons along the unimer molecule. One small drawback is that P-substituent needed for the good chemical

stability of the phosphole ring prevents the regular packing of **TPT** molecules in films.

Photophysical properties of MSPs assembled from **TPT** strongly depend on the metal ion linkages. The MSP with Fe^{2+} ion couplers exhibits a very slow constitutional dynamics, is not luminescent, and shows a strong MLCT band centered at an unusually high wavelength (625 nm) compared to the Fe-MSPs derived from unimers with other central blocks. The Raman excitation profile measurements prove that the red arm of the MLCT band of **P(TPT/Fe)** is significantly contributed with transitions within the central blocks of coordinated **TPT** units. On the other hand, the MSP with Zn^{2+} ion couplers exhibits a very fast constitutional dynamics, is luminescent and does not show an MLCT band. The highest luminescence intensity is observed with an excess of Zn^{2+} ions, which indicates positive effect of end-capping of Zn-MSP chains with Zn^{2+} ions. The MSPs with Co^{2+} , Cu^{2+} and Ni^{2+} ion couplers exhibit variable dynamics and show neither luminescence nor MLCT band. This conclusively indicates that the quenching of luminescence of MSPs is not related to the formation of the MLCT complex that depletes the electronically excited states.

EXPERIMENTAL

Materials

Bis(1,5-cyclooctadiene)di- μ -methoxydiiridium(I) ($[Ir(OMe)(COD)]_2$), 4,4'-di-*tert*-butyl-2,2'-dipyridyl (dtbpy), 4,4,5,5-tetramethyl-1,3,2-dioxaborolane (HBpin), [1,3-bis(2,6-diisopropylphenyl)imidazol-2-ylidene](3-chloropyridyl)palladium(II) dichloride (PEPPSI-IPr), sulfur, zinc(II) perchlorate hexahydrate, nickel(II) perchlorate hexahydrate, copper(II) perchlorate hexahydrate, cobalt(II) perchlorate hexahydrate, iron(II) perchlorate hydrate (all Aldrich), potassium carbonate, magnesium sulfate (VWR) and 4'-bromo-2,2':6',2''-terpyridine (TCI) were used as received, *P,P*-dichlorophenylphosphine (Aldrich) was distilled trap to trap before use. Tetrahydrofuran (THF) (Aldrich) was dried by $LiAlH_4$ or sodium/benzophenone or using MBraun drying solvents system (SPS-800). Toluene (Lachner) was distilled from sodium/benzophenone and water was bubbled with argon before use. Diethylether (Et_2O), dichloromethane, pentane (all VWR or Lachner), acetonitrile (ACN), (all Aldrich), hexane, chloroform (Lachner) were used as obtained. 2,5-di(2-thienyl)-1-phenylthioxophosphole (**1**) was prepared as described in the literature.³⁶

Measurements

1H and ^{13}C NMR spectra were recorded on a Varian UNITY INOVA 400 or Varian SYSTEM 300 instruments in CD_2Cl_2 or $CDCl_3$ and referenced to the solvent signal: 7.25 ppm ($CDCl_3$) or 5.32 ppm (CD_2Cl_2) for 1H and 77.0 ppm ($CDCl_3$) or 53.84 ppm (CD_2Cl_2) for ^{13}C spectra. Coupling constants, J (in Hz), were obtained by the first-order analysis. Infrared spectra were recorded on a Thermo Nicolet 7600 FTIR spectrometer equipped with a Spectra Tech InspecIR Plus microscopic accessory using KBr diluted samples and diffuse reflectance technique (DRIFT) (128 or more scans at resolution 4 cm^{-1}). Raman spectra were recorded on a DXR Raman microscope (Thermo Scientific) using excitations across the whole visible region ($\lambda_{ex} = 445, 532, 633$ and 780 nm) and usual laser power at the sample 0.1 to 0.4 mW. UV/VIS spectra were recorded on a Shimadzu UV-2401PC instrument and photoluminescence spectra on a Fluorolog 3-22 Jobin Yvon Spex instrument, using four-window quartz cuvette (1 cm) for solutions and quartz glass for thin films. Quantum yields, λ_{ex} , of photoluminescence were determined by an integration sphere Quanta- ϕ F-3029. KBr pellets for fluorescence measurement were made by grounding specific amount of sample and KBr using ball mill Vibrator DDR-GM 9458 Narva and then pressed by hydraulic press H62 TRYSTOM. SEC records were obtained using a Spectra Physics Analytical HPLC pump P1000 with two SEC columns: Polymer Labs (Bristol, USA) Mixed-D and Mixed-E. The system was equipped with a Thermo UV6000 DAD detector. A chloro-8 form/acetonitrile (1/1 by volume) mixed solvent containing

Bu₄N⁺PF₆⁻ (0.5 % by weight; to suppress aggregation) was used as the mobile phase (0.7 mL min⁻¹).

Cyclic voltammetry

Cyclic voltammetry measurements were done on a potentiostat UMtE Eco-Trend using drop-casted (from dichloromethane) films of compounds deposited on a carbon disc electrode of diameter of 1 mm. Ag/AgCl reference electrode, scanning rate of 130 mV s⁻¹ and a solution of (Bu₄N)PF₆ (0.1 M) in acetonitrile as the supporting electrolyte were used.

Calculations

Theoretical calculations were done using the density functional theory (DFT), namely the Becke's three parameter functional with the non-local Lee-Yang-Parr correlation functional (B3LYP) with the standard 6-31G(d) basis set as implemented in Gaussian 09 package.⁶² We employed the Becke's three parameter functional with the non-local Lee-Yang-Parr correlation functional (B3LYP) with the standard 6-31G(d) basis set. Electron density and vibrational spectrum of the unimer **TPT** was calculated for the optimized geometry. The wavenumbers values computed contained systematic errors and were thus scaled by the factor 0.946 for Raman and 0.972 for IR spectra.

Complexation experiments

In a typical complexation experiment a measured volume of a solution of Zn²⁺ or Fe²⁺ perchlorate (2×10⁻³M) in acetonitrile/chloroform (1/1 by vol.) was added into a solution of particular unimer in (2×10⁻⁵M) in the same solvent. The metal to unimer (M²⁺/**TPT**) composition ratio was varied from 0 to 3. The UV/vis absorption and the photoluminescence emission spectra were measured for each solution at room temperature one day after preparation.

Synthesis

1-phenyl-2,5-bis[5-(4,4,5,5-tetramethyl-1,3,2-dioxaboralene-2-yl)thiophen-2-yl]thioxophosphole (2).

[Ir(OMe)(COD)]₂ (4 mg, 3 mol%) and dtbpy (2 mg, 3 mol%) were added to a solution of 1-phenyl-2,5-bis(thiophen-2-yl)thioxophosphole **1** (82.5 mg; 0.2 mmol) in dry THF/hexane (7/5 mL) mixed solvent under argon atmosphere, then HBpin (0.18 mL, 1.2 mmol) was added and the reaction mixture kept under stirring at 55°C for 30 hours. The mixture was then diluted with dichloromethane (20 mL) and water (20 mL), stirred for one hour in an open vessel and, finally, the soluble components were extracted with water (3×50 mL). The organic phase was dried over magnesium sulfate, filtered off and evaporated to give the product **2** (130 mg, ca 0.19 mmol) as a honey-like liquid that was used in the next reaction without further purification (since a significant loss of boronic groups was observed at attempts for chromatographic purification). The compound **2** identity has been verified by mass spectroscopy (HRMS) and purity by NMR spectroscopy.

¹H NMR (400 MHz, CD₂Cl₂, δ/ppm): 7.87 (ddd, *J*(P,H) = 14.4 Hz, *J*(H,H) = 8.4 Hz, *J*(H,H) = 1.6 Hz 2 H, *o*-H Ph), 7.53 – 7.45 (m, 2H, *m*-H Ph), 7.43 (AB system, 4H, *J*_{AB}(H,H) = 3.7 Hz, *v*_{AB} = 8.7 Hz, H_{thienyl}), 7.38 – 7.33 (m, 1H, *p*-H Ph), 2.98 (m, 4H, C=CCH₂CH₂), 1.90 (m, 4H, C=CCH₂CH₂), 1.24 (s, 24 H, CH₃).

³¹P NMR (162 MHz, CD₂Cl₂, δ/ppm): +51.8 (s).

¹¹B NMR (128 MHz, CD₂Cl₂, δ/ppm): +22.4 (s).

¹³C NMR (101 MHz, CD₂Cl₂, δ/ppm): 146.2 (d, ²*J*(P,C) = 21.0 Hz, PC=C), 135.3 (d, ²*J*(P,C) = 17.4 Hz, C_{thienyl}), 132.6 (d, *J*(P,C) = 3.2 Hz, C_p Ph), 131.2 (d, *J*(P,C) = 11.8 Hz, C_o Ph), 129.5 (d, *J*(P,C) = 75.4 Hz, PC=C), 129.4 (d, *J*(P,C) = 12.6 Hz, C_m Ph), 128.9 (d, *J*(P,C) = 89.0 Hz, C_i Ph), 128.5 (large s, B-C_{thienyl}), 127.7 (s, CH_{thienyl}), 127.4 (s, CH_{thienyl}), 83.6 (large s, CCH₃), 29.7 (d, *J*(P,H) = 12.4 Hz C=CCH₂CH₂), 24.9 (s, CH₃), 23.0 (s, C=CCH₂CH₂).

FT-IR (cm⁻¹): 3409 (s), 2979 (s), 2929(s), 2860(m), 2040(w), 1719(w), 1618 (m), 1571(w), 1514 (s), 1354(s), 1215(s), 1142(s), 983 (m), 957(m), 926(m), 852(s), 746 (m), 666(s), 613 (m), 578 (m), 517 (m), 448 (m), 419(m)

HRMS found *m/z* 685.1996 [M + Na⁺], C₃₄H₄₁B₂O₄PS₃Na requires 685.1988.

1-phenyl-2,5-bis(5-*tpy*-thiophen-2-yl)thioxophosphole (TPT)

1-Phenyl-2,5-bis[5-(4,4,5,5-tetramethyl-1,3,2-dioxaboralene-2-yl)thiophen-2-yl]thioxophosphole (0.13 g, 0.19 mmol), Br-*tpy* (0.12 g, 0.38 mmol), K₂CO₃ (80mg, 0.57 mmol) and Peppsi-IPr (13 mg, 0.10mol%), Aliquat 336 (cat.) were placed into a Schlenk flask and then evacuated and flushed with argon. Then toluene (5 mL) and water (5 mL) were added, and the reaction mixture was kept at 130°C for 18 hours. After cooling to room temperature the reaction mixture was diluted with dichloromethane (20 mL) and washed with water (3×20 mL), then dried with magnesium sulfate, filtered off and evaporated. The crude product was purified by column chromatography (aluminum oxide, hexane/THF =3/2) giving **TPT** as a red powder (isolated yield: 24mg, 0.03 mmol, 15%). Purity and identity of **TPT** has been verified by NMR spectroscopy and mass spectrometry.

¹H NMR (300 MHz, CD₂Cl₂, δ/ppm): 8.73 (dd, *J*(H,H) = 4.9 Hz, *J*(H,H) = 1.0 Hz, 4H, H_{pyridyl}), 8.63 - 8.66 (m, 8 H, H_{pyridyl}), 7.98 (ddd, *J*(P,H) = 14.3 Hz, *J*(H,H) = 7.7 Hz, *J*(H,H) = 1.0 Hz, 2H, *o*-H Ph), 7.89 (*J*(H,H) = 7.5 and *J*(H,H) = 2.00 Hz, 4H, H_{pyridyl}), 7.66 (d, *J*(H,H) = 3.8 Hz, 2H, H_{thienyl}), 7.52 (d, *J*(H,H) = 3.8 Hz, 2H, H_{thienyl}), 7.38 – 7.33 (m, 3H, *p*-H and *m*-H Ph), 7.35 - 7.40 (m, 4H, H_{pyridyl}), 3.11 (m, 4H, C=CCH₂), 2.00 (m, 4H, C=CCH₂CH₂).

³¹P NMR (121 MHz, CD₂Cl₂, δ/ppm): +52.0 (s).

¹³C NMR (101 MHz, CD₂Cl₂, δ/ppm): 156.1 (s, C_{pyridyl}), 155.7 (s, C_{pyridyl}), 149.1 (s, CH_{pyridyl}) 146.8 (d, ²*J*(P,C) = 23.2 Hz, PC=C), 142.8 (s, C_{thienyl}), 142.6 (s, C_{pyridyl}), 136.8 (s, CH_{pyridyl}) 136.5 (d, ²*J*(P,C) = 17.9 Hz, C_{thienyl}), 132.3 (s, C_p Ph), 130.6 (d, *J*(P,C) = 15.6 Hz, C_o Ph), 129.1 (s, CH_{thienyl}), 129.0 (d, *J*(P,C) = 8.1 Hz, C_m Ph), 128.7 (d, *J*(P,C) = 74.6 Hz, PC=C), 126.0 (s, CH_{thienyl}), 124.0 (s, CH_{pyridyl}), 121.1 (s, CH_{pyridyl}), 116.5 (s, CH_{pyridyl}), 29.5 (d, *J*(P,H) = 13.0 Hz C=CCH₂CH₂), 22.3 (s, C=CCH₂CH₂). C, Ph not observed.

FT-IR (cm⁻¹): 3432 (w), 3030 (w), 3051 (w), 3014 (w), 2946 (w), 2867(w), 1598(s), 1582(s), 1567 (s), 1547 (m), 1466 (m), 1449 (s), 1399(s), 1363(w), 1290(w), 1267 (w), 1224(w), 1145(w), 1097(m), 1075 (w), 1042 (w), 1011 (s), 990 (w), 878 (w), 850(w), 788(s), 775 (w), 745 (m), 731 (m), 718 (w) 691 (w), 678 (w), 665 (m), 635 (w), 623 (w), 608 (w), 555 (w), 533 (w), 499 (w), 468(w)

HRMS found *m/z* 873.2059 [M + H⁺], C₅₂H₃₈N₆PS₃ requires 873.20573

ASSOCIATED CONTENT

Supporting Information

Supporting information contains: overall scheme of synthesis, electronic density contours obtained by DFT calculations, Raman and IR spectra, fluorescence spectra in thin films, spectral changes accompanying self-assembly of metallo-supramolecular polymers, SEC records for **P(Ni/TPT)** and UV/vis DAD spectra of SEC fractions of Fe²⁺/**TPT** system, cyclic voltammetry of unimer **TPT**, ¹H and ¹³C NMR spectra of newly synthesized compounds (PDF). The geometry optimized coordinates of DFT calculations are available as supplemental txt file.

The Supporting Information is available free of charge on the ACS Publications website.

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The authors declare no competing financial interests

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