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Organization and intramolecular charge-transfer enhancement in tripodal tris[(pyridine-4-yl)-phenyl]amine push–pull molecules by intercalation into layered materials bearing acidic functionalities†

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Two new intercalates of tris[4-(pyridin-4-yl)phenyl]amine (**TPPA**) with zirconium hydrogen phosphate and zirconium 4-sulfophenylphosphonate having formulae $Zr(HPO_4)_2{\cdot}0.21(C_{33}H_{24}N_4){\cdot}2.5H_2O$ and $Zr(HO_3SC_6H_4PO_3)_{1.3}(C_6H_5PO_3)_{0.7}{\cdot}0.35(C_{33}H_{24}N_4){\cdot}2.5H_2O$ were prepared and characterized by thermogravimetry, IR spectroscopy, and powder X-ray diffraction. The **TPPA** molecule has been selected as a model tripodal push–pull system with three peripheral basic centers that may undergo protonation. Their protonation/quaternization afforded **HTPPA**/**MeTPPA** molecules with enhanced intramolecular charge-transfer (ICT), which has been documented by electrochemical measurements, UV-Vis spectra and calculated properties such as the HOMO/LUMO levels and the first and second hyperpolarizabilities. Intercalation of **TPPA** into layered zirconium hydrogen phosphate and zirconium 4-sulfophenylphosphonate led to its significant organization and protonation as shown by the IR spectra. From the powder X-ray data we can deduce that the **TPPA** molecules are placed in the interlayer space of both hosts by anchoring two peripheral nitrogen atoms to one host layer and the opposite pyridine-4-yl terminus to the other neighboring host layer. In zirconium 4-sulfophenylphosphonate, the **TPPA** molecules are oriented perpendicularly, while in zirconium phosphate these molecules are slanted with respect to the layers of the host. On dehydration by heating, the interlayer distance of the intercalate decreases, which indicates a further slanting of the **TPPA** molecules. It follows from the UV-Vis spectra that **TPPA** is present in both intercalates in an equilibrium of protonated and non-protonated forms. The described materials represent the first case when a tripodal push–pull system was incorporated into a system with restricted geometry with the aim to influence its optical properties.

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Introduction

Organic π-conjugated compounds are of interest for a wide scientific community due to their unique properties and miscellaneous applications. In an organic molecule with a D-π-A arrangement, intramolecular charge-transfer (ICT) from the donor (D) to the acceptor (A) occurs and the molecule constitutes a dipole. Such push–pull systems are currently tremendously investigated as active molecules for nonlinear optics (NLO) and (opto)electronics.[1,2] Tripodal derived push–pull systems attract attention due to their two-photon absorption properties and thus have promising applications in microscopy, data storage, microfabrication or photodynamic therapy.[3–5] Triphenylamine based compounds represent a $(A\text{-}\pi)_3$-D octupolar type of Y-shaped push–pull molecule featuring a central amino donor and three acceptor-substituted π-branches. For instance, the structure and electronic properties of the metal–organic framework of triphenylamine molecules bearing π-deficient pyridine peripheral acceptors have been recently investigated.[6,7] It is well known that the electron withdrawing ability of pyridine and the related six-membered nitrogen-containing heterocyclic compounds can

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†Electronic supplementary information (ESI) available: Preparation and characterization of **TPPA** and **MeTPPA**, UV-Vis absorption spectra of **TPPA** and **MeTPPA**, ^{1}H NMR and ^{13}C NMR APT spectrum of **MeTPPA**, thermal powder X-ray patterns of **ZrP·TPPA** and **ZrSPP·TPPA**, IR spectrum of partially protonated TPPA, deconvoluted UV-Vis spectra of **ZrP·TPPA** and **ZrSPP·TPPA**, UV-Vis spectrum of **ZrP·TPPA** treated with HCl. See DOI: 10.1039/c4dt00149d

be improved by N-alkylation/protonation.[8] Hence, in such molecules it would be very tempting to perform quaternization (ICT enhancement) in parallel with structure ordering by incorporation into a system with confined geometry.[9–11] Materials prepared in such a way have the advantage, relative to organic hosts, of improved rigidity and thermal stability. An important class of these hybrid materials is intercalation compounds. These materials consist of layered solids (hosts), which contain other species (guests) between the layers, in the interlayer space. Restricted geometry in the interlayer space of layered inorganic solids helps to organize the incorporated species in a way favorable for the improvement of their optical properties.[12]

Up to now, only a few cases of intercalates with NLO properties have been reported.[3,10,13,14] As the host materials, mostly clays[15,16] and layered MPS_3 (M = Mn, Cd, Zn)[9,14] were used. In the case of cationic chromophores derived from stilbazolium intercalated into MPS_3, it is claimed that the confined interlayer space of the host leads to a chromophore aggregation.[9] The formation of J aggregates has been reported to induce SHG properties because of the resulting noncentrosymmetric packing of the interacting chromophores.[17,18]

A host material with thoroughly studied intercalation chemistry is layered zirconium phosphate. This phosphate exists in two modifications: α-$Zr(HPO_4)_2{\cdot}H_2O$ and γ-$Zr(PO_4)(H_2PO_4){\cdot}2H_2O$. Both phosphates contain acidic OH groups, whose interaction with basic guest molecules serves as a driving force for the intercalations.[19] Both zirconium phosphates are transparent and their intercalation compounds with basic molecules are stable. An NLO active material made by intercalation of a $(4'$-(dimethylamino)-1-methylstilbazolium$)^+$ cation into layered γ-$ZrPO_4(H_2PO_4){\cdot}2H_2O$ phases has been reported by Coradin *et al.*[20] Another group of host materials is represented by zirconium phosphonates, derived from the phosphates by replacing the OH group with an organic rest.[21] When such an organophosphonate is functionalized with an appropriate acidic group on the organic backbone, it can be used as a suitable host material. Recently, we have prepared zirconium 4-sulfophenylphosphonate with a general formula $Zr(HO_3SC_6H_4PO_3)_2{\cdot}yH_2O$,[22] and tested its intercalation ability.[23] Zirconium 4-sulfophenylphosphonate proved to be a very good host material for the intercalation of nitrogen-containing basic organic molecules.

In this paper we report on intercalation of a tris[4-(pyridin-4-yl)phenyl]amine (**TPPA**) guest molecule into α-modification of zirconium phosphate and into zirconium 4-sulfophenylphosphonate. The prepared materials were characterized by thermogravimetry, IR spectroscopy and powder X-ray diffraction. The presumed arrangement and electronic properties of the intercalated species are discussed.

Experimental

Synthesis of zirconium phosphate

Well-crystallized α-modification of $Zr(HPO_4)_2{\cdot}H_2O$ (further denoted as **ZrP**) was obtained according to the method proposed by Alberti and Torracca.[24] A clear solution was prepared by dissolving $ZrOCl_2{\cdot}8H_2O$ (10.1 g), hydrofluoric acid (40% w/w, 8 mL) and H_3PO_4 (85% w/w, 92 mL) in water (160 mL). The solution was heated at 80 °C for 4 days, maintaining a constant volume by continuously adding water. The **ZrP** precipitate was washed with de-ionized water and dried in air.

Synthesis of a ZrP intercalate with ε-aminocaproic acid

ε-Aminocaproic acid (further denoted as **ACA**) was intercalated by refluxing **ZrP** (10 g) in an aqueous solution of ε-aminocaproic acid (0.25 M, 400 mL) for 7 days. The solid product (further denoted as **ZrP·ACA**) was separated by filtration, washed with water and ethanol and dried in air. Its formula was determined to be $Zr(HPO_4)_2{\cdot}(H_2N(CH_2)_5COOH)_{0.80}{\cdot}1.5H_2O$.[25]

Preparation of a ZrP intercalate with tris[4-(pyridin-4-yl)-phenyl]amine

The **ZrP·ACA** intercalate (0.05 g) was mixed with tris(4-(pyridin-4-yl)phenyl)amine (0.08 g) in water (9 mL). The mixture was placed in a Teflon-lined 23 mL Parr acid digestion bomb and heated under autogenous pressure at 130 °C for about 20 hours. The product (further denoted as **ZrP·TPPA**) was separated by filtration, washed with water and dried in air. For optical measurements the product was extracted in a Soxhlet extractor until the extract was colorless. Elemental analysis calcd/found for $Zr(HPO_4)_2{\cdot}0.21(C_{33}H_{24}N_4){\cdot}2.5H_2O$, M_r = 428.30; C, 19.43/19.85 ± 0.04; H, 2.83/2.78 ± 0.01; N, 2.75/2.69 ± 0.02.

Synthesis of zirconium 4-sulfophenylphosphonate

Zirconium 4-sulfophenylphosphonate (further denoted as **ZrSPP**) was prepared according to the previously described procedure.[22] 4-Sulfophenylphosphonic acid (2.38 g) and $ZrOCl_2{\cdot}8H_2O$ (2.58 g) were added to a mixture of 1 M HF (50 mL) and 1 M HCl (50 mL) in a 300 mL PP beaker. The reaction mixture was heated to 80 °C in an oil bath overnight during which it evaporated to a half of the volume. After that the mixture was evaporated to dryness at 80 °C. The solid was suspended in 1 M HCl and then centrifuged. This process was repeated three times and the obtained slurry was dried in a rotary evaporator at 70 °C to remove hydrochloric acid. The product was dried in a desiccator over NaOH. The sulfophenyl group was partially desulfonated during the synthesis, therefore the formula of the obtained product was $Zr(HO_3SC_6H_4PO_3)_{1.3}(C_6H_5PO_3)_{0.7}{\cdot}2H_2O$.

Preparation of a ZrSPP intercalate with tris[4-(pyridin-4-yl)-phenyl]amine

Zirconium 4-sulfophenylphosphonate dihydrate (0.05 g) was mixed with tris[4-(pyridin-4-yl)phenyl]amine (0.08 g) in a mixture of water (6 mL) and ethanol (3 mL). The mixture was placed in a Teflon-lined 23 mL Parr acid digestion bomb and heated under autogenous pressure at 130 °C for about 20 hours. The product was separated by filtration, washed with an ethanol–water mixture (1/2, v/v), and then with ethanol and dried in air. For optical measurements the product was

extracted in a Soxhlet extractor until the extract was colorless. Elemental analysis calcd/found for $Zr(HO_3SC_6H_4PO_3)_{1.3}(C_6H_5PO_3)_{0.7}\cdot 0.35(C_{33}H_{24}N_4)\cdot 2.5H_2O$, M_r = 719.30; C, 39.32/37.66 ± 0.05; H, 3.28/2.86 ± 0.01; N, 5.80/6.25 ± 0.02; S, 5.80/6.25 ± 0.03. The compound in the further text is denoted as **ZrSPP·TPPA**.

Powder X-ray diffraction data were obtained with a D8 Advance diffractometer (Bruker AXS, Germany) with a Bragg–Brentano θ–θ geometry (40 kV, 30 mA) using Cu Kα radiation with a secondary graphite monochromator. The diffraction angles were measured at room temperature from 2 to 70° (2θ) in 0.02° steps with a counting time of 15 s per step. Powder X-ray diffraction measurements at 210 ± 1 °C were carried out on a heated brass block equipped with a thermocouple in the range from 2 to 35° (2θ) in 0.025° steps with a counting time of 15 s per step. The size of the crystallites of the intercalates was calculated according to the Scherrer formula[26] using EVA software.[27]

Thermogravimetric measurements (TGA) were done using home-made apparatus constructed of a computer-controlled oven and a Sartorius BP210 S balance. The measurements were carried out in air between 30 and 960 °C at a heating rate of 5 °C min^{-1}.

Infrared spectra in the range of 600–4000 cm^{-1} were recorded at 64 scans per spectrum at 2 cm^{-1} resolution using a HATR adapter on a Perkin-Elmer FTIR Spectrum BX spectrometer on neat samples. All spectra were corrected for the presence of moisture and carbon dioxide in the optical path.

UV-Vis diffuse reflectance spectra of diluted (2% w/w by Al_2O_3) powder materials were recorded in the range from 210 to 800 nm using a UV-Vis Lambda 20 spectrometer (Perkin-Elmer, USA) equipped with a diffuse reflectance attachment with a 3 inch integrating sphere with Al_2O_3 as a reference. The reflectance values were re-calculated using the Schuster–Kubelka–Munk equation, $F(R_\infty) = (1 - R_\infty)^2/2R_\infty$, where R_∞ is the diffuse reflectance from a semi-infinite layer. For details see ref. 28 and 29.

Results and discussion

Preparation and properties of TPPA/HTPPA/MeTPPA

The tris[4-(pyridin-4-yl)phenyl]amine (**TPPA**) guest molecule was prepared in a threefold Suzuki–Miyaura cross-coupling reaction of tris(4-iodophenyl)amine with pyridine-4-ylboronic acid (see also the ESI†).[6,7] This molecule possesses three basic pyridine-4-yl (Py) groups that can easily be quaternized (Fig. 1a). Whereas in the non-protonated **TPPA** molecule only diminished ICT from the central amino donor to peripheral Py groups takes place, protonation/methylation of the Py groups afforded **HTPPA**/**MeTPPA** with a significantly enhanced electron withdrawing ability and ICT efficiency. The *N*-methyl derivative **MeTPPA** is chemically well-accessible by the reaction of **TPPA** with an excess of iodomethane. For its synthesis and full spectral characterization see the ESI.† The enhanced charge transfer in **HTPPA**/**MeTPPA** can be demonstrated by

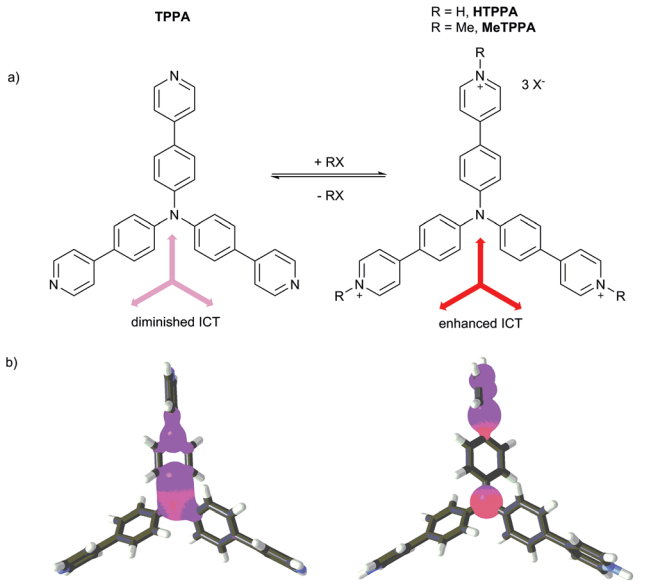

Fig. 1 Quaternization of **TPPA** and its impact on the ICT (a) and HOMO (red) and LUMO (blue) localizations in **TPPA** and **HTPPA**, respectively (b). Counter ions were omitted for clarity.

HOMO/LUMO localizations shown in Fig. 1b.[30] The HOMO and LUMO are in both structures localized predominantly over one π branch. Degenerated LUMO−1 and LUMO−2 were found on the remaining two branches. Whereas the HOMO/LUMO in **TPPA** are spread over the central amino donor and the adjacent part of the 1,4-phenylene and Py moieties, in **HTPPA** the HOMO and LUMO are obviously separated on the central donor and the peripheral acceptor. This indicates a significant charge-separation. Further electronic parameters of **TPPA** and **MeTPPA** are shown in Table 1.

Electrochemical measurements were carried out in DMF containing 0.1 M Bu_4NPF_6 in a three electrode cell by cyclic voltammetry (CV) and rotating disc voltammetry (RDV) or polarography. The first oxidation and reduction potentials $E_{1/2(ox1)}$ and $E_{1/2(red1)}$ are shown in Table 1. Unfortunately, the first reduction/oxidation potentials of **TPPA** and **MeTPPA**, respectively, are out of the potential window available in DMF and Pt electrodes. However, when going from **TPPA** to **MeTPPA**, the calculated HOMO–LUMO gaps decreased from 7.51 to 5.11 eV. This is mainly caused by a raised HOMO and a decreased LUMO in **MeTPPA** by 0.90 and 1.50 eV. In addition, positions of the longest-wavelength absorption maxima λ_{max} (CT-bands) measured in CH_3OH (see the ESI†) showed a significant bathochromic shift from 363 to 428 nm ($\Delta\lambda_{max}$ = 65 nm) as a result of better D–A interaction in **MeTPPA**. The calculated ground state dipole moments of both molecules are nearly zero due to their centrosymmetry. Hence, the observed enhancement in the calculated first hyperpolarizability β from 0.48 to 4.87×10^{-30} esu must be elucidated as a change in the spatial electron distribution in **MeTPPA**. The second hyperpolarizability γ increased from 179 284 (**TPPA**) to 692 195 × 10^{-39} esu (**MeTPPA**).

Table 1 Experimental and calculated parameters of **TPPA** and **MeTPPA**

Comp.	$E_{1/2(ox1)/HOMO}$ [a] [V/eV]	$E_{1/2(red1)/LUMO}$ [a] [V/eV]	λ_{max} [b] [nm (eV)]	E_{HOMO} [c] [eV]	E_{LUMO} [c] [eV]	ΔE [c] [eV]	β [c] [10^{-30} esu]	γ [c] [10^{-39} esu]
TPPA	1.08/−5.43	—	363 (3.42)	−8.29	−0.78	7.51	0.48	179 284
MeTPPA	—	−1.11/−3.24	428 (2.90)	−7.39	−2.28	5.11	4.87	692 195

[a] $E_{1/2(ox1)}$ and $E_{1/2(red1)}$ are the half-wave potentials of the first oxidation and reduction, respectively, as measured by RDV; $E^{abs}_{HOMO/LUMO} = E_{1/2(ox1/red1)} + 4.35$. [b] Longest-wavelength absorption maxima measured in CH_3OH ($c = 2 \times 10^{-5}$ M). [c] Calculated by MOPAC 2012.

Characterization of the intercalates

The **ZrP·TPPA** intercalate shows a weight decrease of 10% on heating to about 200 °C (Fig. 2). This corresponds to a release of 2.5 molecules of water per formula unit (theoretical weight loss is 10%). A further steep decrease of weight at around 500 °C is due to deintercalation of the amine. The total weight loss observed on heating to 980 °C is 36%, which is close to the theoretical weight loss (37%) calculated for ZrP_2O_7 as the final product of the heating, considering the amount of the intercalate to be 0.21 per formula unit.

The **ZrSPP·TPPA** intercalate is slightly less stable than **ZrP·TPPA**, as can be seen from the comparison of their thermogravimetric curves (Fig. 2). Also for **ZrSPP·TPPA** the first step of the weight loss of 6% corresponds to the release of water from the interlayer space. Further decomposition of the intercalate starts at about 400 °C and is caused by the deintercalation of the amine and decomposition of the organic part of sulfophenylphosphonate. The total weight loss is 66%, which indicates the formation of ZrP_2O_7 as a final product (theoretical weight loss of 63%, considering the amount of the intercalated **TPPA** to be 0.35, as given in the Experimental section).

Infrared spectra

The way of interaction between the host material and the intercalated species was also studied by infrared spectra, as shown in Fig. 3 and 4 for **ZrP** and **ZrSPP** as the hosts, respectively.

The distinct couple of bands at 3588 and 3507 cm^{-1} found in **ZrP** and corresponding to stretching vibrations of the PO–H bond[31] is replaced by a broad band both in the aminocaproic acid preintercalated host (**ZrP·ACA**) and the final **ZrP·TPPA** intercalate. Also the band at 3112 cm^{-1} corresponding to O–H stretching vibration of water molecules observed in **ZrP** is broadened in both **ZrP·ACA** and **ZrP·TPPA**. The intensive band at 1704 cm^{-1}, corresponding to C=O stretching vibrations of the carboxylic group, which appears after the intercalation of aminocaproic acid, is not present in the **ZrP·TPPA**. This is evidence that all the **ACA** was replaced by **TPPA**.

To determine, whether the **TPPA** guest is present in the intercalate as a protonated or neutral species, the IR spectra of **TPPA** and its methylpyridinium derivative (tris(4-(*N*-methylpyridinium-4-yl)phenyl)amine, **MeTPPA**) were also measured. The most distinct difference between the IR spectra of **TPPA** and **MeTPPA** is a triple of ring stretching vibrations at 1585, 1517,

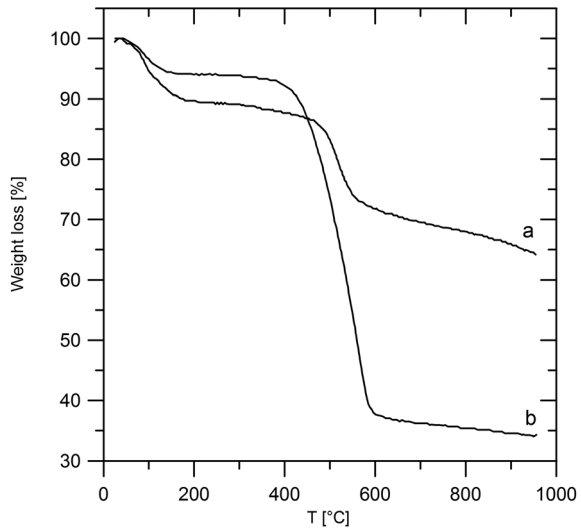

Fig. 2 Thermogravimetric curves of **ZrP·TPPA** (a) and **ZrSPP·TPPA** (b) intercalates.

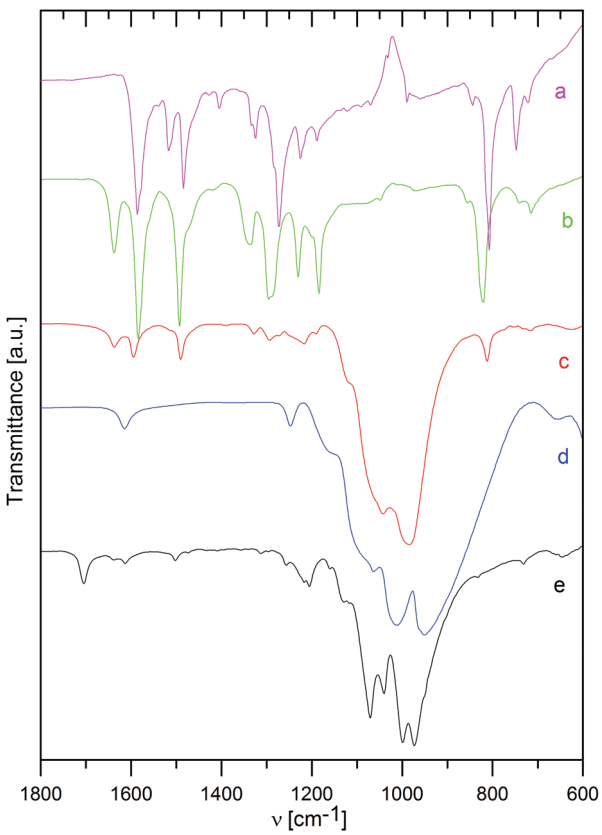

Fig. 3 Infrared spectra of **TPPA** (a), **MeTPPA** (b), **ZrP·TPPA** (c), **ZrP** (d), and **ZrP·ACA** (e).

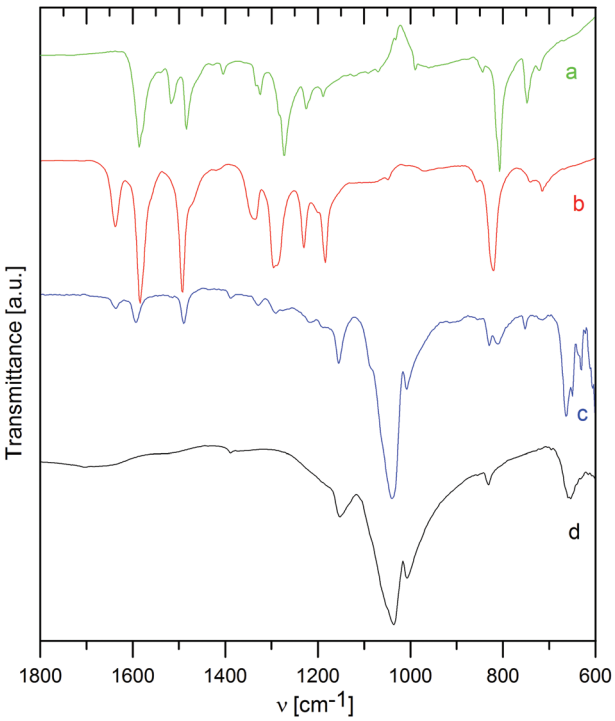

Fig. 4 Infrared spectra of TPPA (a), MeTPPA (b), ZrSPP·TPPA, and ZrSPP (d).

and 1484 cm^{-1} in **TPPA**, which are in **MeTPPA** shifted to 1637, 1581, and 1492 cm^{-1}. These bands are characteristic of aromatic compounds; the observed shift to a higher energy region corresponds to the significantly higher polarization of the **MeTPPA** molecules having a $^{+}$NMe terminus. It was found that this shift occurs also in partially protonated **TPPA**.

In **ZrP·TPPA** these bands are at the same positions (1637, 1594, and 1492 cm^{-1}) as in **MeTPPA**, which confirms that the **TPPA** guest molecules are present in the intercalate in the protonated form. The H–O–H deformation vibration of water molecules observed in **ZrP** and in **ZrP·ACA** is in the **ZrP·TPPA** intercalate most probably masked by the ring stretching vibration bands. The C–H deformation vibration bands present at 1335, 1295, 1230, and 1185 cm^{-1} in **MeTPPA** are very poorly developed in **ZrP·TPPA**. The P–O–H deformation vibration present in **ZrP** at 1247 cm^{-1} is strongly suppressed in **ZrP·TPPA**. A broad and intensive band between 1150 and 900 cm^{-1} belongs to P–O antisymmetric stretching vibrations and P–OH stretching vibrations of the host. In the intercalate as in the guest spectra, there is a distinct band at 820 cm^{-1} of a C–H deformation vibration typical for *para*-substituted benzene derivatives.

In the IR spectrum of **ZrSPP** (see Fig. 4), there is a very weak band at 1386 cm^{-1} corresponding to aromatic C–C ring stretching vibrations. A more distinct band at 1152 cm^{-1} is given by P–O asymmetric stretching vibrations and the most intensive and broad couple of bands appearing at 1039 and 1007 cm^{-1} corresponds to P–O and S–O vibrations. The band at 831 cm^{-1} arises due to 1,4-disubstituted benzene ring vibrations and the band observed at 653 cm^{-1} reflects C–S vibrations of the sulfonic group. All these bands can be observed also in the IR spectrum of **ZrSPP·TPPA**. In addition a distinct triple of bands at 1636, 1594, and 1490 cm^{-1} and less distinct bands at 1328, 1290, and 1216 together with a shoulder at 1180 cm^{-1} are bands characteristic of **MeTPPA**.

In summary, the infrared spectra of both **ZrP·TPPA** and **ZrSPP·TPPA** indicate that the **TPPA** guest is present in its protonated form in the intercalates.

Powder patterns of the intercalates

On intercalation, a distinct enlargement of the interlayer distance was observed for both host materials, see Fig. 5 and 6.

The interlayer distance for the **ZrP·TPPA** intercalate increased to 18.52 Å (Fig. 5) from the original interlayer distance of 7.6 Å in the host material. The peak at 2θ = 33.8° corresponding to the 020 reflection is distinct in the original **ZrP** host but is broadened in **ZrP·TPPA** due to the lower crystallinity of the intercalate. Nevertheless, the fact that this 020 reflection is retained in **ZrP** confirms that the structure of the **ZrP** host is retained in the intercalate. It was reported that the layer thickness of **ZrP** is 6.3 Å.[19] It means that the height of the interlayer space (gallery) in the **ZrP·TPPA** intercalate is 18.5 − 6.3 = 12.2 Å. In the case of the **ZrSPP·TPPA** intercalate, the layer thickness (taken as the distance between the sulfo oxygen atoms of one side of the **ZrSPP** layer and the sulfo oxygen atoms on the other side) was determined to be 16.8 Å,[23] and the interlayer distance in the **ZrSPP·TPPA** intercalate is 33.2 Å (see Fig. 6). Thus, the height of the gallery in this intercalate is 33.2–16.8 = 16.4 Å. The size of the crystallites determined by powder X-ray diffraction is 244 Å for **ZrP·TPPA** and 325 Å for **ZrSPP·TPPA**.

Geometrical considerations on the intercalates

The **TPPA** molecule has a shape of an equilateral triangle with the height of roughly 16.2 Å (see Fig. 7a). We can consider the arrangement of the **TPPA** molecules in the interlayer space of the host in the following ways: (a) the molecules are placed

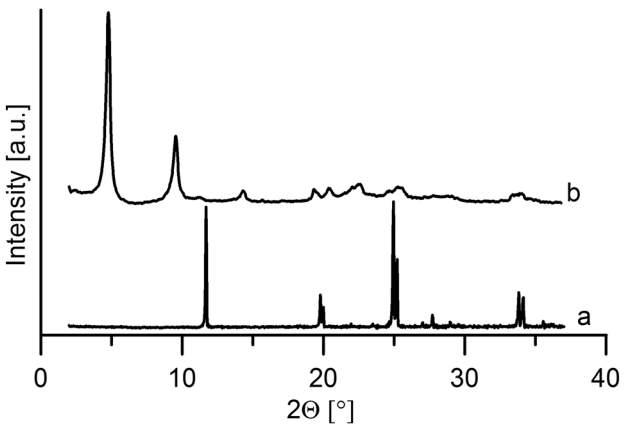

Fig. 5 Diffraction patterns of the host **ZrP** material (a) and its intercalate with **TPPA** (b).

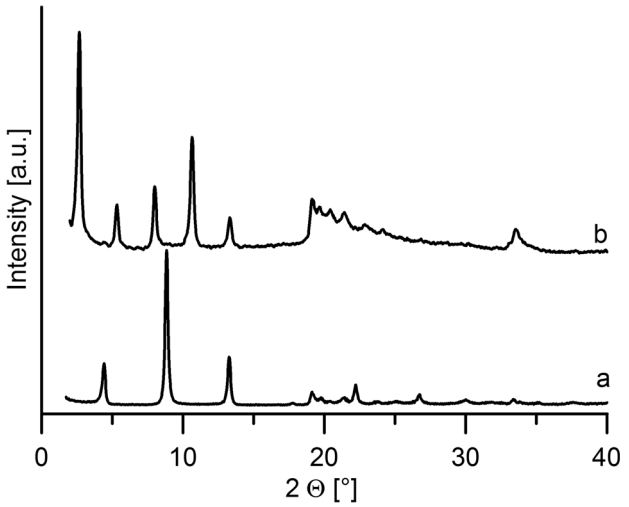

Fig. 6 Diffraction patterns of the host **ZrSPP** (a) and its intercalate with **TPPA** (b).

parallel to the plane of the host layers as monomolecular sheets. In such a case the height of the interlayer space (gallery) should be about 4 Å. (b) The molecules are placed parallel to the host layers in a bimolecular way that is there are two sheets of the **TPPA** molecules placed above each other in the gallery. In such a case the gallery height should be 8 Å. The presence of more sheets of the **TPPA** molecules in the gallery is improbable as in such an arrangement the **TPPA** molecules should be bonded very loosely to the host layers. (c) The **TPPA** molecules are placed perpendicularly to the plane of the host layers. In the most probable arrangement, the **TPPA** molecule is placed such that one of its sides lies on the plane of the host layer while with its remaining corner it is bonded to the neighboring host layer. Then the gallery height should be close to the value given in Fig. 7a, *i.e.* 16.2 Å.

The actual enlargement of the interlayer space on intercalation of **TPPA** into **ZrP** is larger than in the cases (a) and (b) but lower than in the case (c). It means that the **TPPA** molecules are placed in the interlayer space of **ZrP** in the manner described for (c) but with a slightly slanting position. In the case of the **ZrSPP·TPPA** intercalate, the gallery height determined by powder XRD (16.4 Å) is slightly higher than the height of the **TPPA** molecule (16.2 Å). Therefore, in this case we can presume that the **TPPA** molecules are oriented perpendicularly to the host layers of **ZrSPP**.

Let us discuss the way the molecules are placed *on* the host layers; this way influences the amount of the intercalated **TPPA** molecules and consequently also the amount of water present in the intercalate. In the alpha modification of $Zr(HPO_4)_2$ the acidic OH groups of the phosphates are placed uniformly above and below the host layers in an equilateral triangular fashion (see Fig. 7b) with the in-plane distance between them being about 5.3 Å.

The driving force for the intercalation process in **ZrP** is proton transfer from the host HPO_4 group to the intercalated amine. During the intercalation the pyridine nitrogen atoms are protonated, while the OH groups of the phosphate are deprotonated. If we presume the arrangement of the **TPPA** molecule as that described in case (c) above, then the distance of nitrogen atoms in **TPPA** lying on the host plane is about 14.7 Å. The protonated nitrogen atoms should be as close to the acidic oxygen atoms of the phosphate groups as possible. This condition can be fulfilled when the **TPPA** molecules are placed on the host layer in a manner shown in Fig. 7b.

In **ZrSPP** we presumed that its structure is derived from the structure of alpha modification of **ZrP**, with OH phosphate groups being replaced by $C_6H_4SO_3H$ groups, which are oriented perpendicularly to the inorganic sheets of the Zr atoms.[23] It means that the SO_3H groups of the host are placed uniformly at the surface of the host layers in an equilateral triangular fashion as the OH groups in **ZrP**. Similar to **ZrP**, the driving force for the intercalation in **ZrSPP** is a proton transfer from the SO_3H groups to the pyridine nitrogen atoms.

From the structure of **ZrP** it follows that on the surface of the host layer there is a "free area" of 24 $Å^2$ associated with each phosphate group.[19] Thus, for each $Zr(HPO_4)_2$ formula unit we have $2 \times 24 = 48$ $Å^2$ free area which can be covered with the guest. Let us consider that the triangular **TPPA** molecule is anchored to one layer of **ZrP** by its base (side) and to the other

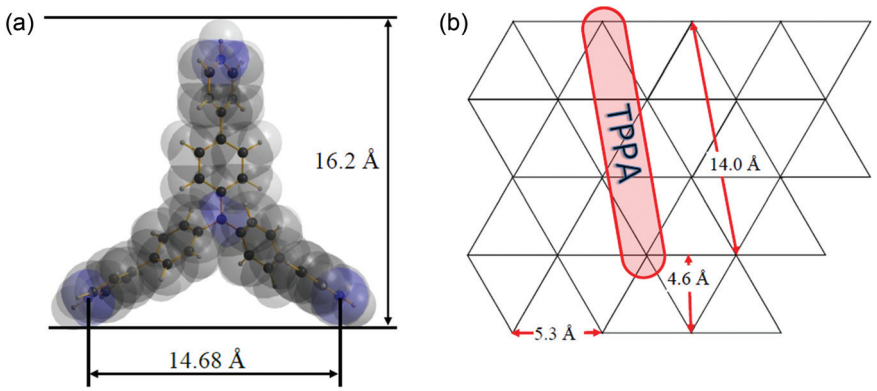

Fig. 7 The **TPPA** molecule and its dimensions (a). The placement of the **TPPA** molecules on the host layer (b).

neighboring layer by its apex (corner). The cross section of the base (the area which is covered by this part of the **TPPA** molecule) is about 116 Å^2. The cross section at the apex of **TPPA** is about 21 Å^2. In summary, the area covered by the **TPPA** molecule in **ZrP·TPPA** is therefore 0.21 × (116 + 21) = 29 Å^2, which is an area much smaller than that provided by the **ZrP** host. In the case of **ZrSPP·TPPA**, the area covered by the **TPPA** molecule is 0.35 × (116 + 21) = 48 Å^2, which indicates that the whole surface of the **ZrSPP** layer is covered by the guest molecules. Therefore the amount of **TPPA** intercalated into **ZrSPP** (0.35 per formula unit) represents the maximum amount which can be intercalated into this type of host due to sterical reasons.

The van der Waals volume of the **TPPA** molecule calculated by Hyperchem software[32] is 462 Å^3. The molecular geometry was optimized by PM3 and PM7 semi-empirical methods implemented in programs ArgusLab[33] and MOPAC2012.[34] The volume accessible for the intercalated entities might be calculated as the "free area" (24 Å^2) multiplied by the gallery height. Thus for the **ZrP·TPPA** the accessible volume is 24 × 12.2 = 293 Å^3 and for **ZrSPP·TPPA** it is 24 × 16.4 = 394 Å^3. In **ZrP·TPPA**, there are 0.21 molecules of **TPPA** per formula unit, and the space occupied by **TPPA** is therefore 0.21 × 462 = 97 Å^3 and in **ZrSPP·TPPA** it is 0.35 × 462 = 162 Å^3. In both cases there is enough space for water molecules to be placed among the **TPPA** molecules.

Thermal behavior of the intercalates

On heating the interlayer distance decreases in both intercalates (see Fig. S4 and S5 in the ESI†). In the case of **ZrP·TPPA** the interlayer distance decreases from 18.5 Å at room temperature to 16.6 Å at 210 °C. In the case of **ZrSPP·TPPA**, this decrease is from 33.2 to 31.5 Å. The change of the interlayer distance can be explained by the release of water. The empty space formed after the dehydration allows further slanting of the **TPPA** molecules in the intercalate resulting in a decrease of the interlayer distance.

The intercalates might be rehydrated. In the case of the **ZrP·TPPA** intercalate, a rehydration by standing the sample at 25–30% relative humidity (RH) and at room temperature for 24 h causes an increase of the interlayer distance to 17.2 Å. The full rehydration close to the original state (18.44 Å) was achieved by standing the sample at room temperature at 100% RH for another 24 h. The rehydration of the **ZrSPP·TPPA** sample was, on the other hand, easier because standing the sample at room temperature at 25–30% RH for 24 h was sufficient to achieve the original value of the interlayer distance.

UV-Vis spectra of the intercalates

Fig. 8 shows UV-Vis spectra of **TPPA**, **MeTPPA**, **ZrP·TPPA** and **ZrSPP·TPPA**. From the comparison of the spectra of **TPPA**, **MeTPPA** and those of the intercalates it follows that both intercalates contain **TPPA** both in protonated and deprotonated forms in an equilibrium. This finding is in discrepancy with the results of the IR spectra measurements, where the bands of the deprotonated **TPPA** were not found in the IR spectra of **ZrP·TPPA** and **ZrSPP·TPPA**. To solve this problem, we measured IR spectra of the partially protonated **TPPA**. We prepared a solution of 1 mol of **TPPA** with 1.5 mol of HCl so that **TPPA** (with three protonable pyridine groups) would be protonated to one half. The IR spectrum of the resulting product (see Fig. S6c in the ESI†) is different from that of **TPPA** and corresponds to the spectrum of **MeTPPA**. Thus both the UV-Vis and IR spectra confirm that the **TPPA** guest in **ZrP·TPPA** and **ZrSPP·TPPA** is partially protonated.

Deconvolution of the longest-wavelength absorption maxima λ_{max} of **ZrP·TPPA** and **ZrSPP·TPPA** (see Fig. S7 in the ESI†) revealed two peaks appearing at ~370 and 450 nm that fit the positions of CT-peaks of **TPPA** and **MeTPPA** (Fig. 8). The observed bathochromic shift with $\Delta\lambda_{max}$ ~ 80 nm is similar to that observed in the solution (see above) and indicates enhanced ICT in both intercalates. In the case of **ZrP·TPPA** the shape of the spectrum suggests that the deprotonated form is present in the intercalate in a relatively higher amount than in **ZrSPP·TPPA**. This further implies that the interlayer environment of the **ZrSPP** host is more acidic than that of **ZrP**. The question arises whether this partially protonated **TPPA** in the intercalates might be further protonated by exposing the intercalates to an acidic environment. When the **ZrP·TPPA** and **ZrSPP·TPPA** intercalates are subjected to HCl vapors overnight, their UV-Vis spectra change distinctly and are identical to that of **MeTPPA** (see Fig. S8 in the ESI† for **ZrP·TPPA**). The powder XRD pattern of **ZrP·TPPA** after the exposure is identical to that before the exposure. It means that no deintercalation occurred in this case. On the other hand, the powder XRD pattern of **ZrSPP·TPPA** after the exposure is identical to that of **ZrSPP**; it means the **TPPA** guest molecules are deintercalated in an acidic environment.

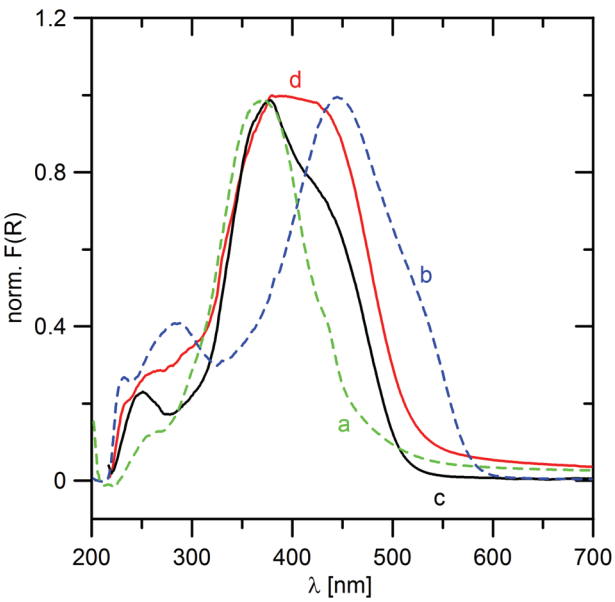

Fig. 8 UV-Vis spectra of **TPPA** (a), **MeTPPA** (b), **ZrP·TPPA** (c), and **ZrSPP·TPPA** (d).

Conclusion

Tris[4-(pyridin-4-yl)phenyl]amine was successfully intercalated into α-modification of zirconium phosphate and into zirconium 4-sulfophenylphosphonate. It was found that the guest amine is protonated during the intercalation on the peripheral pyridine-4-yl moieties. The UV-Vis spectra suggested an equilibrium between the protonated and non-protonated forms of the amine. However, a significant bathochromic shift and thus an ICT enhancement were revealed in the solution as well as in the solid state. This observation was further confirmed by the electrochemical measurements and calculations of further electronic properties for both limit **TPPA** and **MeTPPA** structures. Based on the amount of the intercalated species and the enlargement of the interlayer space caused by the intercalation the probable arrangement of the molecules of the guest in the intercalate was suggested. The triangular shape of the guest molecule leaves enough space between the layers of the host, so that water molecules can be accommodated in the interlayer space under ambient conditions. On heating, water is released from the intercalate and, as a consequence of this release, the interlayer distance in the intercalate decreases, which is caused by a reorientation (slanting) of the guest molecules with respect to the host layers. The prepared materials represent the first example of tripodal push–pull organic molecules introduced in a confined space of layered materials with the aim to influence their optical properties. In view of the current interest in novel inorganic–organic hybrid materials, this structure–property relationship study would serve as a useful guide for designing new intercalates with tunable optical properties.

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