Non-cyclic formylated dipyrrromethanes as phosphate anion receptors†

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New tetrakis- and hexakis(1H-pyrole-2-carbaldehyde) anion receptors are described. The anion binding properties of these receptors were studied in organic media and in the solid state. The receptors displayed good affinity for the dihydrogenphosphate and pyrophosphate anions (as the tetrabutylammonium salts) in chloroform even in the presence of a polar protic solvent, methanol. Solution phase spectroscopic analyses proved consistent with the binding mode seen in single crystal structural studies of the dihydrogenphosphate and pyrophosphate complexes and provided support for the contention that these receptors undergo conformational reorganization in order to accommodate the bound oxoanions both in chloroform solution and in the solid state.

Introduction

Inorganic phosphate and related anions are ubiquitous in nature. These anionic species play key roles in energy transduction and in the storage and expression of genetic information.1 Phosphates are also pervasive pollutants and may lead to eutrophication of lakes and waterways.2 The development of molecular constructs capable of binding the phosphate anion in its various protonated forms, as well as related oxoanion species is of critical interest since it may provide a first step towards solving various problems related to anion detection, extraction, and separation.3–6 In general, anions are larger than cations and therefore larger hosts are required to bind them.7 Fortunately, a number of binding motifs may be exploited to achieve anion recognition, including coulombic interactions, hydrogen bonding, halogen bonding, and anion–π interactions.8–12 Hydrogen bonding has proven particularly effective. However, certain anions, including oxoanions, such as dihydrogenphosphate (H2PO4−) and hydrogensulfate (HSO4−), are susceptible to proton transfer.13 As a result, ensuring anion binding rather than deprotonation (acid–base chemistry) has proven to be a challenging task.14,15 One hydrogen bond donor that is relatively less susceptible to deprotonation is pyrrole. An early example of a pyrrole-based system capable of binding the phosphate anion was the pentapyrrolic macrocycle, sapphyrin. This system, which is dicationic in its protonated form, proved capable of stabilizing a “sitting-atop” complex with the phosphate anion in the solid state and of binding DNA and other phosphate-containing species in solution.16 In 1996, Gale et al. reported that octamethylcalix[4]pyrrole (C[4]P; 1) may act as an anion receptor in organic media.17 Calix[4]pyrrole is a non-conjugated system long known in the literature.18 It was found to bind the fluoride and chloride anions in halogenated solvents, along with their counter cations in some cases.19,20 Halide anion recognition by C[4]P is accompanied by conformational change from a limiting 1,3-alternate conformation (the free form) to a cone conformation (halide-bonded form). This structural switching allows the formation of four NH–anion hydrogen bonds in the case of simple halide anions.17 Under most experimental conditions, C[4]P has proved less effective as a receptor for larger anions, including the dihydrogenphosphate anion. There thus remains a need for simple-to-prepare analogues of C[4]P that bind phosphate. Open-chain systems might allow this goal to be attained (Scheme 1).

![Scheme 1 Structures of compounds 1, 2a-b, and 3a-b.](image-url)
In the case of fluoride and chloride anion salts studied in organic media, compound 1 displayed association constants that were >2× greater than those recorded for the corresponding acyclic building block, dipyromethane (DPM) 2a. This difference was ascribed to the greater pre-organization of the cyclic systems, as well as to the increased number of pyrrolic NH hydrogen bond donors provided by 1 compared to 2a. Interestingly, compound 2a displayed a higher affinity towards H_{2}PO_{4}− (as its tetra-n-butylammonium (TBA′) salt) than did 1 in CD_{2}Cl_{2}. While this apparent dichotomy was not studied in detail, it might reflect the fact that the dihydrogenphosphate anion is too large to be accommodated well within the NH-rich calix[4]pyrrole pseudo cavity. In contrast, the presumably greater flexibility of 2a might allow the stabilization of a greater number of favourable contacts with this and other oxoanions than compound 1. While untested, this hypothesis provides an incentive to prepare and study new dipyromethane-based anion receptors. A goal of this effort would be to create phosphane anion receptors that rival C[4]P in terms of ease of preparation and which allow for good phosphate anion binding. In this study we focus on dipyromethane (DPM) receptors that have been subject to formylation in the so-called α-pyrrolic positions.

In 2003, in the context of synthetic work aimed at producing a C[4]P-texaphyrin chimera, the pyrrolic α-positions of 2a were subject to formylation to produce 2b. The anion binding properties of 2b with TBAF were studied by isothermal calorimetry (ITC) in acetonitrile and via X-ray crystallography in the solid state. On this basis, it was concluded that 2b was not an effective anion receptor. On the other hand, we recently found that the bisdipyromethane (bisDPM) 3a and its formylated derivative 3b, act as effective and somewhat selective receptors for the TBA′ salts of dihydrogenphosphate (H_{2}PO_{4}−) and hydrogenpyrophosphate (HP_{2}O_{5}3−) in chloroform, as inferred from NMR spectroscopic and UV-Vis analyses. As compared to 1, the non-formylated system 3a proved to be a slightly better receptor for H_{2}PO_{4}− (K_{a} = 10^{5} vs. 10^{6} M^{−1} in CHCl_{3} and CH_{2}Cl_{2}, respectively). DPM derivatives, which do not bear substituents in the pyrrolic α-positions are notoriously unstable, being prone inter alia to oxidative coupling and electrophilic aromatic substitution. On the other hand, bis-formylation yields stable systems that we believe may have a role to play as anion receptors. In preliminary work, high phosphate anion affinities were noted for the four-fold formylated derivative of 3a, receptor 3b (K_{a} = (8 ± 2) × 10^{6} M^{−1} for 3b vs. (1.0 ± 0.1) × 10^{3} M^{−1} for 3a in CHCl_{3}; cf. Table 1). Presumably, this increased affinity reflects the favourable electronic changes, as well as the addition of hydrogen bond accepting sites, that result from formylation. To understand the underlying determinants and to improve on the phosphate binding affinity displayed by 3b, we have now prepared and studied several analogues bearing different spacer elements between the diformyl-DPM subunits, as well as a new hexakis(1H-pyrole-2-carbaldehyde) receptor, 7b. Compared to 3b, receptor 7b displays a phosphate anion affinity that is enhanced by roughly 66 fold when studied under identical conditions (i.e., CHCl_{3} containing 3% CH_{3}OH).

**Results and discussion**

**Syntheses and single crystal structures**

The synthesis and characterization of 3b–5b have been reported previously. In this paper, we present the new derivatives, 6b and 7b, obtained in moderate overall yields. The synthetic route to obtain 3b–7b is shown in Scheme 2. In the first step, commercially available diacetyl and triacetyl starting materials were converted into bisDPM 6a, and trisdipyrromethane (trisDPM) 7a, respectively, albeit in low yields (5–15%). The low yields stand in contrast to high yields observed for 3a and 5a, which precipitate from the reaction mixture and constitute the dominant reaction products. The unstable α-free bisDPM (6a) and trisDPM (7a) were then subject to Vilsmeier–Haack formylation to produce the stable formylated products 6b and 7b in relatively high yields (ca. 80%). Purification of 6b was carried out by recrystallization from N,N-dimethylformamide (DMF) at 80 °C, whereas 7b was purified by column chromatography over...
silica gel. The molecular structures were characterized by $^1$H and $^{13}$C NMR spectroscopy, high-resolution mass spectrometry (HRMS), and single crystal X-ray diffraction analyses.

Receptor 6b co-crystallised with two molecules of DMF. The single crystal structure revealed that two diformyl-DPM units residing on opposite cyclopentadienyl (Cp) rings of the ferrocene were locked at about 90° from one another, leading to the presence of a racemic mixture of conformational isomers within the single crystal. Fig. 1 shows one of the atropisomers of 6b. It crystallizes as a one-dimensional hydrogen bond interlocked ensemble, even in the presence of DMF. The formation of these assemblies can be rationalized on a geometric basis. The bridging ferrocene adopts a conformation such that the two constituent Cp rings, each bearing a DPM substituent, are offset from one another by 90°. This allows for the formation of four intermolecular hydrogen bonds between the two self-complementary diformyl-DPM subunits, as can be seen from an inspection of Fig. 1.

Receptor 7b, on the other hand, co-crystallised with CHCl$_3$. Each of the three diformyl-DPM units on the bridging benzene is oriented in a different direction (cf. Fig. 1). One of the three DPM subunits was found to resemble what is seen in the case of 3b wherein two diformyl-DPM units are seen to orient towards opposite π-faces of the intervening benzene ring. On the other hand, two of the diformyl-DPM units present in 7b adopt conformations in which the pyrrole NH groups point away from one another. As a result, 7b forms dimers stabilised by four hydrogen bonds that extend to create two-dimensional layers (cf. Fig. S2†). The assemblies formed by 6b and 7b may be viewed as being examples of Gale and co-worker’s “narcissistic dimer”† that are extended into one and two-dimensional space, respectively. The hydrogen bond interlocked aggregates seen in the solid state structures of 6b and 7b may account for the low solubility these species display in many solvents.

Anion binding studies

The anion selectivity for 3b–5b in CHCl$_3$ was found to be H$_2$PO$_4^-$ > HP$_2$O$_5^{3-}$ > HSO$_4^-$ > C$_6$H$_5$CO$_2^-$ ≫ NO$_3^-$, Cl$^-$ (as the commercially available TBA$^+$ salts). The observed selectivity was not found to correlate with the basicity of the anions (basicity order: HP$_2$O$_5^{3-}$ > C$_6$H$_5$CO$_2^-$ > H$_2$PO$_4^-$ > HSO$_4^-$, NO$_3^-$, Cl$^-$). Rather, the trend appears to correlate with an ability of the anion to interact with the receptor via hydrogen bond donation to the formyl groups (in addition to acting as a Lewis basic hydrogen bond acceptor for the pyrrolic NH protons). In fact, the H$_2$PO$_4^-$ anion, with two hydrogen bond donating sites (rather than one or none as in the other test anions), is bound with the highest relative affinity.

The effect of the formyl groups in abetting dihydrogenphosphate anion binding is substantial (i.e., several orders of magnitude difference in $K_a$ values). This was first noted for 3a vs. 3b in the context of our original report (vide supra). Further support for this conclusion came from a comparison of the α-free derivative 5a with its formylated congener 5b (cf. Table 1). Unfortunately, the other non-formylated DPM derivatives prepared in the context of this study proved too unstable to allow for analyses of their anion binding affinities.

Diffusion ordered NMR spectroscopy (DOSY) was used to examine the interactions between anions and receptor 3b. Receptor 3b was chosen for these studies for two reasons. First,
the low solubility of 3b in CHCl₃ means that an equilibrium was expected to exist between species free in solution and those tied up in the solid state. Secondly, on the ¹H NMR time scale, free receptor 3b and the H₃PO₄⁻ bound complex (3b·H₂PO₄⁻) were subject to slow-exchange, allowing both species to be observed concurrently. Fig. 2 shows the DOSY spectrum of a mixture of 3b at the half-equivalence point obtained by titrating 3b with TBAH₂PO₄ in CDCl₃. The one-dimensional ¹H NMR spectrum in the horizontal axis shows two sets of receptor signals; free 3b and 3b·H₂PO₄⁻, labelled with red and blue asterisks, respectively.

The two-dimensional spectrum is characterized by the presence of two sets of receptor signals corresponding to slightly different diffusion rates. The complex 3b·H₂PO₄⁻ (blue) features a lower mobility than the free receptor 3b (red). No evidence of higher order (i.e., aggregated) species is seen. Based on these observations, we conclude that in the absence of complete conversion to the bound form, samples of TBAH₂PO₄ consist of only two receptor species, namely the free receptor and the bound complex, 3b·H₂PO₄⁻. Support for the existence of two components in solution came from a UV-Vis study, also carried out in CHCl₃. When the total concentration of 3b was held constant, isobestic behaviour was seen upon titration with TBAH₂PO₄. This is as expected for a receptor solution consisting of two interconverting species.²⁴

Mass spectrometric (MS) analyses provided evidence that receptor 3b would interact with the H₃PO₄⁻ anion in the gas phase. In these studies, an equimolar mixture of 3b and TBAH₂PO₄ in CHCl₃ was directly injected into the MS instrument without passing the species through a column. Negative ion electrospray ionization (ESI⁻) was used. The high-resolution MS analysis revealed two expected signals, namely, one at m/z 505.1885 corresponding to [3b·H]⁻ and one at m/z 603.1643 ascribable to the H₃PO₄⁻ complex ([3b·H₃PO₄⁻]). (cf. Fig. S4†).

Further insights into the binding modes operative in the case of the present series of receptors came from single crystal X-ray diffraction analyses.²⁶ Diffraction-grade single crystals of the pyrophosphate complex, 3b·(TBA)₂H₂P₂O₇, were obtained by layering a solution of 3b·H₃P₂O₇⁻ in CHCl₃ with n-pentane. The complex 3b·H₂P₂O₇⁻ in CH₂Cl₂ was prepared by treating 3b with [(TBA)₃HP₂O₇]. This served to improve the net solubility by converting the poorly soluble receptor (vide supra) to the corresponding pyrophosphate complex. In the solid state, a 2 : 2 complex, [3b·H₂P₂O₇⁻]₂ is seen. The overall complex consists of a hydrogen-bonded dimer that lies across a crystallographic inversion centre (cf. Fig. 3).

The protonation state of the bound pyrophosphate in the complex [3b·H₂P₂O₇⁻], was deduced from the presence of two TBA⁺ counter cations per pyrophosphate dianion. The H₃P₂O₇⁻ anion lies parallel to the π-surface of the bridging benzene ring. Both diformyl-DPM arms are oriented such that four hydrogen bonds between the diformyl-DPM units in 3b and the bound H₃P₂O₇⁻ are favoured. The N–H···O distances between the NH protons of 3b and the O atoms of H₃P₂O₇⁻ range from 2.648 to 2.690 Å, separations that are short compared to typical N–H···O distances.²⁷,²⁸ Each bound H₃P₂O₇⁻ interacts with another pyrophosphate anion with the result that a hydrogen bonded dimer is formed. In the dimer, four protons are shared between eight oxygen atoms. The observed O···O distances of 2.508–2.518 Å are within the range expected for this type of proton bridged O–H···O type interaction.²⁷,²⁸ Because the hydrogen atoms on H₃P₂O₇⁻ are involved in dimer formation, the formyl groups do not participate in the hydrogen bonds that serve to bind the anion. Indeed, the formyl groups point away from the bound H₃P₂O₇⁻ guest.

Diffraction grade single crystals of the complex 4b·TBAH₂PO₄ were obtained in a similar way. The resulting structure is shown in Fig. 4. There is no evidence of dimerization in this case. The lack of dimerization is ascribed to the relatively tight binding between the TBA⁺ counter cation and the bound dihydrogenphosphat e guest, which precludes further inter-complex interactions. In the solid state, complex 4b·H₃PO₄⁻ displays the same conformation for the diformyl-DPM units as observed in 3b·H₂P₂O₇⁻. The four N–H···O distances that characterize the interaction between the NH protons of 4b and the O atoms of H₃PO₄⁻ range from 2.704 to 2.760 Å. The two O–H···O distances...
associated with the hydrogen bonding between the formyl groups and the hydrogen atoms of H$_2$PO$_4^-$ are 2.681 and 2.726 Å, respectively. Overall, derivative 4b stabilizes six hydrogen bonding interactions with the H$_2$PO$_4^-$ anion. Specifically, it acts as a four-fold donor via the pyrrolic NH protons and acts as a two-fold acceptor via the two formyl moieties.

Single crystals of complex 7b-H$_2$PO$_4^-$ were obtained by using TBAH$_2$PO$_4$ to solubilize receptor 7b in CHCl$_3$ (by conversion to the corresponding dihydrogenphosphate complex) and then layering with n-pentane. The resulting structure is shown in Fig. 5. Although some analogy to that of 4b–H$_2$PO$_4^-$ discussed above is seen as regard the diformal DPM-dihydrogenphosphate interactions, in 7b–H$_2$PO$_4^-$, the third diformal-DPM of 7b is too distant to interact well with the bound H$_2$PO$_4^-$ anion. It thus interacts in an intermolecular sense with a dihydrogenphosphate anion bound to a second receptor. The result is a dimeric complex with overall 2 : 2 stoichiometry. The four pyrrole N-H-deprotonated phosphate oxygen N–H···O distances range from 2.730 to 2.827 Å. The intermolecular hydrogen bond that is presumed to play a role in stabilizing the 2 : 2 complex is characterized by an O–H···O distance of 2.598 Å.

Based on the binding mode observed in the solid state, we considered it likely that anion–π interactions, as well as pyrrolic NH–anion hydrogen bonding interactions contribute to anion binding. To explore this possibility, the effect of adding TBAH$_2$PO$_4$ to the most soluble receptor of the present series, namely the meta-derivative 4b, was investigated by $^1$H NMR spectroscopy (Fig. 6). Upon addition of one molar equivalent of TBAH$_2$PO$_4$ to a CDCl$_3$ solution of 4b, discernible shifts in the proton resonances of 4b were observed (cf. Table 2). Of particular note were the 0.6 ppm upfield shifts seen for the proton signals of the linking benzene ring. Similar $^1$H NMR spectroscopic studies of 5b carried out in CDCl$_3$, 7b in a more competitive solvent mixture, CDCl$_3$:DMSO-d$_6$ (8 : 1), likewise revealed upfield shifts in the intervening benzene resonances (cf. Fig. S8 and S10†). Such shifts are consistent with the decrease in the deshielding effect of the intervening aromatic ring current as seen in previous studies of $\alpha,\alpha,\alpha,\alpha$-meso-tetraaryl-C[4]P derivatives by Ballester and co-workers.\textsuperscript{14}

However, as noted by a reviewer, such an observation is not sufficient to confirm or rule out the presence of an anion–π interaction. Intuitively, the interaction of anion H$_2$PO$_4^-$ with the electron-rich intervening benzene ring of 4b is expected to be repulsive, thus destabilizing the overall binding event. However, Dey\textsuperscript{32} has suggested that the polarization of a π-electron system by an anion induces a dipole moment that may provide a contribution to the anion–receptor interaction. In the case of the present system, further studies will be necessary to determine whether anion–π interactions (if any) are playing a substantial role in mediating the observed anion recognition behaviour.

Receptor 7b proved even less soluble in neat, non-hydrogen bonding solvents, such as CHCl$_3$, than its congeners 3b–5b. This made it impossible to carry out binding studies that were directly comparable to those reported earlier.\textsuperscript{24} However, in analogy to what was seen in the case of these other receptors, it was found that the solubility of 7b increased upon the addition of TBAH$_2$PO$_4$. Dilute solutions of 7b–H$_2$PO$_4^-$ in CHCl$_3$ could thus be prepared by mixing single crystals of 7b with ca. 2.75 molar equivalents of TBAH$_2$PO$_4$. The UV-Vis spectrum of this solution was characterized by features similar to those seen in the reported spectrum of 3b–H$_2$PO$_4^-$ at analogous concentrations. Thus, the observed spectral features were ascribed to the complex 7b–H$_2$PO$_4^-$. When the CHCl$_3$ solution made up from 7b and TBAH$_2$PO$_4$ was titrated with CH$_3$OH, spectral changes and isosbestic behaviour were observed. On this basis, we

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**Table 2** Summary of chemical shifts of selected proton resonances of 4b before and after addition of one molar equivalent of TBAH$_2$PO$_4$ in CDCl$_3$

<table>
<thead>
<tr>
<th>H</th>
<th>4b$^a$</th>
<th>4b–H$_2$PO$_4^-$</th>
<th>$\Delta$δ</th>
</tr>
</thead>
<tbody>
<tr>
<td>$H^1$</td>
<td>7.25</td>
<td>7.10</td>
<td>-0.15</td>
</tr>
<tr>
<td>$H^2$</td>
<td>7.10</td>
<td>6.70</td>
<td>-0.40</td>
</tr>
<tr>
<td>$H^3$</td>
<td>6.90</td>
<td>6.30</td>
<td>-0.60</td>
</tr>
<tr>
<td>$H^4$</td>
<td>6.80</td>
<td>6.85</td>
<td>+0.05</td>
</tr>
<tr>
<td>$H^5$</td>
<td>6.00</td>
<td>6.30</td>
<td>+0.30</td>
</tr>
</tbody>
</table>

$^a$ The solution of 4b was prepared from diffraction grade single crystals of 4b, which contained two DMF molecules of crystallization per equivalent of receptor. $^b$ Obtained by treating with the TBA$^+$ salt of H$_2$PO$_4^-$; see text for details.
CH3OH in CHCl3 was prepared and then subjected to titration. It was also apparent that large quantities of methanol were present, and that this would allow for quantitative analyses.

In a first set of experiments, an 11 μM solution of 7b in 3% (v/v) CH3OH in CHCl3 was prepared and then subjected to titration with TBAH2PO4 (cf. Fig. 7). The induced spectral changes were then fit to a 1:1 binding profile and used to calculate the binding constant corresponding to H2PO4⁻ binding in accord with the methods described previously. Similar UV-Vis titrations were carried out with a series of anions in the form of their TBA+ salts. To allow for a comparison between the original tetrakis-(1H-pyrrole-2-carbaldehyde) receptors 3b-5b, this new derivative proved poorly soluble in typical organic solvents used for electrochemical studies (e.g., CH2CN, DMF, CH2Cl2). However, as above, an increase in its solubility was seen upon the addition of TBAH2PO4. This was seen as a “useful property” that could be used to obtain insights into the H2PO4⁻ binding events under conditions of electrochemical analysis. It was found, for instance, that initial turbid mixtures of free receptor 6b in CH2CN containing 10% (v/v) DMF give rise to a quasi-reversible cyclic voltammogram (CV) (black line in Fig. 8). The oxidation wave for 6b becomes increasingly irreversible as the relative concentration of TBAH2PO4 increases. Eventually, the CV becomes characterized by a relatively intense anodic peak and a near-absence of a corresponding cathodic feature.

The observed increase in the anodic peak current is ascribed to the increase in the concentration of 6b (both free and complexed) that occurs as the titration runs its course. Since the 6b is not completely soluble in the solvent mixture in the absence of H2PO4⁻, the concentration of free 6b in the titration mixture should be constant up until the point where all traces of solid 6b are dissolved, presumably by conversion to the corresponding dihydrogenphosphate complex, 6b·H2PO4⁻. To the extent such a supposition is correct, it would account for the observed increase in the peak current, which could be attributed directly to the formation of the readily oxidizable species, 6b·H2PO4⁻.

When the increase in the anodic peak current is plotted against the added H2PO4⁻ concentration, a binding isotherm could be determined by UV-Vis spectroscopy in two separate solvent systems as described in Table 3.

**Table 3** Summary of calculated binding affinities, $K_a$ (M⁻¹) as determined by UV-Vis spectroscopy in two separate solvent systems.

<table>
<thead>
<tr>
<th>Anions</th>
<th>3b</th>
<th>3b</th>
<th>7b</th>
</tr>
</thead>
<tbody>
<tr>
<td>H2PO4⁻</td>
<td>$\left(8 \pm 2\right) \times 10^6$</td>
<td>$\left(2.1 \pm 0.3\right) \times 10^3$</td>
<td>$\left(1.4 \pm 0.2\right) \times 10^5$</td>
</tr>
<tr>
<td>HP2O7⁻</td>
<td>$\left(1.4 \pm 0.1\right) \times 10^3$</td>
<td>$\left(3.8 \pm 0.6\right) \times 10^3$</td>
<td>$\left(3.6 \pm 0.2\right) \times 10^4$</td>
</tr>
<tr>
<td>HSO4⁻</td>
<td>$\left(4.1 \pm 0.2\right) \times 10^3$</td>
<td>nd</td>
<td>nd</td>
</tr>
<tr>
<td>NO3⁻</td>
<td>nd</td>
<td>nd</td>
<td>nd</td>
</tr>
<tr>
<td>Cl⁻</td>
<td>nd</td>
<td>nd</td>
<td>nd</td>
</tr>
</tbody>
</table>

* The anions were studied as their TBA⁺ salts. In CHCl3; data from ref. 24. In CHCl3 containing 3% (v/v) CH3OH.

**Fig. 8** Stacked cyclic voltammogram of 6b in CH2CN containing 10% (v/v) DMF. The initial turbid mixture of 6b contained a quantity of material sufficient to produce a 1.0 mM solution once completely dissolved. This mixture was titrated with TBAH2PO4 until the TBAH2PO4 concentration reached 2.8 mM. TBAFP6 (0.1 M) was used as the supporting electrolyte. Glassy carbon was used as the working electrode, a Pt wire as the counter electrode, and a Ag/AgCl couple as the reference electrode.
obtained (cf. Fig. S20†) from which an approximate $\text{H}_2\text{PO}_4^-$ affinity, $(1.0 \pm 0.1) \times 10^3 \text{M}^{-1}$, could be calculated (cf. Table ESI-1†). The loss in reduction signal intensity is believed to reflect the strong interaction between the oxidized ferrocenium receptor ($6b^+$) and $\text{H}_2\text{PO}_4^-$. 

Conclusions

In summary, we have extended the tetrakis(1H-pyrrole-2-carbaldehyde) receptor family to include a new electroactive derivative, $6b$, incorporating a ferrocene linker. Also reported here is a hexaformyl trisDPM derivative, $7b$. The anion binding properties of these new systems were studied by monitoring the increase in solubility electrochemically, through high-resolution mass spectrometric analyses, 1D $^1\text{H}$ NMR and 2D DOSY spectroscopic experiments, single crystal X-ray diffraction analyses, and UV-Vis spectroscopic titrations. Conformational switching from trans-like conformations as seen in the free tetrakis- ($3b$,$4b$) and hexakis- ($7b$) (1H-pyrrole-2-carbaldehyde) receptors to the corresponding cis-like conformations upon interacting with phosphate derivatives was revealed by X-ray diffraction analyses in the solid state (Fig. 9). In mixed methanol–chloroform solution, the new hexakis-carbaldehyde derivative $7b$ was found to exhibit an affinity for $\text{H}_2\text{PO}_4^-$ that is roughly 100 times higher than that displayed by the tetrakis analogue $3b$. This increase in binding affinity is rationalized in terms of a receptor design that overcomes in part the conformational penalty that needs to be paid in order to achieve substrate binding. As such, the present findings provide experimental insights that might prove useful in the design of anion receptors based on rather simple binding motifs, such as pyrroles and dipyrromethane subunits.

Acknowledgements

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Notes and references

14 Note: Under certain conditions, deprotonation, rather than anion binding, has been observed with pyrrolic receptors.