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## Non-cyclic formylated dipyrromethanes as phosphate anion receptors<sup>†</sup>

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New tetrakis- and hexakis(1H-pyrrole-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.

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## 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.<sup>1</sup> 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.<sup>3-6</sup> 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- $\pi$  interactions.8-12 Hydrogen bonding has proven particularly effective. However, certain anions, including oxoanions, such as dihydrogenphosphate  $(H_2PO_4^{-})$  and hydrogensulfate  $(HSO_4^{-})$ , 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.<sup>16</sup> In 1996, Gale et al. reported that octamethylcalix[4]pyrrole (C[4]P; 1) may act as an anion receptor in organic media.<sup>17</sup> Calix[4]pyrrole is a nonconjugated system long known in the literature.<sup>18</sup> It was found to bind the fluoride and chloride anions in halogenated solvents, along with their counter cations in some cases.<sup>19,20</sup> 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 NHanion 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 simpleto-prepare analogues of C[4]P that bind phosphate. Open-chain systems might allow this goal to be attained (Scheme 1).

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**3b**, Y = -CHO **3a**, Y = -H;

Scheme 1 Structures of compounds 1, 2a-b, and 3a-b.

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In the case of fluoride and chloride anion salts studied in organic media, compound 1 displayed association constants that were  $>2\times$  greater than those recorded for the corresponding acyclic building block, dipyrromethane (DPM) 2a.21 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  $H_2PO_4^-$  (as its tetra-*n*-butylammonium (TBA<sup>+</sup>) salt) than did 1 in CD<sub>2</sub>Cl<sub>2</sub>. 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 dipyrromethane-based anion receptors. A goal of this effort would be to create phosphate 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 dipyrromethane (DPM) receptors that have been subject to formylation in the so-called  $\alpha$ -pyrrolic positions.

In 2003, in the context of synthetic work aimed at producing a C[4]P-texaphyrin chimera,<sup>22</sup> the pyrrolic  $\alpha$ -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.23 On this basis, it was concluded that 2b was not an effective anion receptor. On the other hand, we recently found<sup>24</sup> that the bisdipyyrromethane (bisDPM) 3a and its formylated derivative 3b, act as effective and somewhat selective receptors for the TBA<sup>+</sup> salts of dihydrogenphosphate  $(H_2PO_4^{-})$  and hydrogenpyrophosphate (HP<sub>2</sub>O<sub>7</sub><sup>3-</sup>) 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  $H_2PO_4^{-}$  ( $K_a = 10^3 vs. 10^2 M^{-1}$  in CHCl<sub>3</sub> and CH<sub>2</sub>Cl<sub>2</sub>, respectively). DPM derivatives, which do not bear substituents in the pyrrolic  $\alpha$ -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**  $(K_{\rm a} = (8 \pm 2) \times 10^{6} \,{\rm M}^{-1}$  for **3b** vs.  $(1.0 \pm 0.1) \times 10^{3} \,{\rm M}^{-1}$  for **3a** in CHCl<sub>3</sub>; cf. Table 1).<sup>24</sup> Presumably, this increased affinity reflects the favourable electronic changes, as well as the

Table 1 Calculated  $H_2PO_4^-$  binding affinities,  $K_a$  (M<sup>-1</sup>) of dipyrromethane derivatives before (**3a** and **5a**) and after formylation (**3b** and **5b**)

3a <sup><i>a</i>,<i>c</i></sup>	3 <b>b</b> <sup><i>b</i>,<i>c</i></sup>	5a <sup>a</sup>	$\mathbf{5b}^{b,c}$
$(1.0\pm0.1) imes10^3$	$(8\pm2) imes10^6$	$40.8\pm0.6$	$(4.6\pm0.9) imes10^5$
<sup>a</sup> Determined series	111 ND (D am a stre		

<sup>*a*</sup> Determined using <sup>1</sup>H NMR spectroscopy in CDCl<sub>3</sub>. <sup>*b*</sup> Determined using UV-Vis spectroscopy in CHCl<sub>3</sub>. <sup>*c*</sup> Values from ref. 24.

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*(1*H*-pyrrole-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<sub>3</sub> containing 3% CH<sub>3</sub>OH).

## Results and discussion

#### Syntheses and single crystal structures

The synthesis and characterization of **3b–5b** have been reported previously.<sup>24</sup> 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.<sup>24</sup> The unstable  $\alpha$ -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



Scheme 2 Synthesis of receptors **3b**-**7b**. \*Catalyst concentrations and reaction temperatures vary.

silica gel. The molecular structures were characterized by <sup>1</sup>H and <sup>13</sup>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<sub>3</sub>. 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<sup>24</sup> wherein two diformyl-DPM units are seen to orient towards opposite  $\pi$ -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<sup>†</sup>). The assemblies formed by **6b** and **7b** may be viewed as being examples of Gale and co-worker's "narcissistic dimer"15 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<sub>3</sub> was found to be  $H_2PO_4^- > HP_2O_7^{3-} > HSO_4^- > C_6H_5CO_2^- \gg NO_3^-$ , Cl<sup>-</sup> (as the



Fig. 1 Top left: Single crystal structure of **6b** (DMF solvent molecules are omitted for clarity); Top right: Single crystal structure of **7b** (CHCl<sub>3</sub> solvent molecules are omitted for clarity); Bottom: Linear assembly formed by one of the atropisomers of **6b**.

commercially available TBA<sup>+</sup> salts).<sup>24</sup> The observed selectivity was not found to correlate with the basicity of the anions (basicity order:  $HP_2O_7^{3-} > C_6H_5CO_2^{-} > H_2PO_4^{-} > HSO_4^{-}$ ,  $NO_3^{-}$ , Cl<sup>-</sup>). 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  $H_2PO_4^{-}$  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*).<sup>24</sup> Further support for this conclusion came from a comparison of the  $\alpha$ -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)<sup>25</sup> was used to examine the interactions between anions and receptor **3b**. Receptor **3b** was chosen for these studies for two reasons. First,



Fig. 2 DOSY spectrum of a 1 : 0.5 mixture of **3b** and TBAH<sub>2</sub>PO<sub>4</sub>. The mixture was produced by titrating a 0.8 mM CDCl<sub>3</sub> solution of TBAH<sub>2</sub>PO<sub>4</sub> into a turbid mixture of **3b** in CDCl<sub>3</sub> until the ratio of integration values corresponding to **3b** and **3b**  $\cdot$ H<sub>2</sub>PO<sub>4</sub><sup>-</sup> was equal. ^ shows signals ascribed to free **3b**; \* shows those assigned to **3b**  $\cdot$ H<sub>2</sub>PO<sub>4</sub><sup>-</sup>.

the low solubility of **3b** in CHCl<sub>3</sub> means that an equilibrium was expected to exist between species free in solution and those tied up in the solid state. Secondly, on the <sup>1</sup>H NMR time scale, free receptor **3b** and the H<sub>2</sub>PO<sub>4</sub><sup>-</sup> bound complex (**3b**·H<sub>2</sub>PO<sub>4</sub><sup>-</sup>) were subject to slow-exchange,<sup>24</sup> 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 TBAH<sub>2</sub>PO<sub>4</sub> in CDCl<sub>3</sub>. The one-dimensional <sup>1</sup>H NMR spectrum in the horizontal axis shows two sets of receptor signals; free **3b** and **3b**·H<sub>2</sub>PO<sub>4</sub><sup>-</sup>, 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  $3\mathbf{b} \cdot \mathrm{H_2PO_4^-}$  (blue) features a lower mobility than the free receptor  $3\mathbf{b}$  (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  $3\mathbf{b}$  and TBAH<sub>2</sub>PO<sub>4</sub> consist of only two receptor species, namely the free form and the bound complex,  $3\mathbf{b} \cdot \mathrm{H_2PO_4^-}$ . Support for the existence of two components in solution came from a UV-Vis study,<sup>24</sup> also carried out in CHCl<sub>3</sub>. When the total concentration of  $3\mathbf{b}$  was held constant, isosbestic behaviour was seen upon titration with TBAH<sub>2</sub>PO<sub>4</sub>. This is as expected for a receptor solution consisting of two interconverting species.<sup>24</sup>

Mass spectrometric (MS) analyses provided evidence that receptor **3b** would interact with the  $H_2PO_4^-$  anion in the gas phase. In these studies, an equimolar mixture of **3b** and TBAH<sub>2</sub>PO<sub>4</sub> in CHCl<sub>3</sub> was directly injected into the MS instrument without passing the species through a column. Negative ion electrospray ionization (ESI<sup>-</sup>) was used. The high-resolution MS analysis revealed two expected signals, namely, one at m/z505.1885 corresponding to [**3b**-H]<sup>-</sup> and one at m/z 603.1643 ascribable to the  $H_2PO_4^-$  complex ([**3b**·H<sub>2</sub>PO<sub>4</sub><sup>-</sup>]). (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.<sup>26</sup> Diffraction-grade single crystals of

the pyrophosphate complex,  $3\mathbf{b} \cdot (\text{TBA})_2 \text{H}_2 \text{P}_2 \text{O}_7$ , were obtained by layering a solution of  $3\mathbf{b} \cdot \text{H}_2 \text{P}_2 \text{O}_7^{2-}$  in  $\text{CH}_2 \text{Cl}_2$  with *n*-pentane. The complex  $3\mathbf{b} \cdot \text{H}_2 \text{P}_2 \text{O}_7^{2-}$  in  $\text{CH}_2 \text{Cl}_2$  was prepared by treating  $3\mathbf{b}$  with  $(\text{TBA})_3 \text{HP}_2 \text{O}_7$ . 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,  $[3\mathbf{b} \cdot \text{H}_2 \text{P}_2 \text{O}_7^{2-}]_2$  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  $[\mathbf{3b} \cdot \mathbf{H}_2 \mathbf{P}_2 \mathbf{O}_7^{2^-}]_2$  was deduced from the presence of two TBA<sup>+</sup> counter cations per pyrophosphate dianion. The H<sub>2</sub>P<sub>2</sub>O<sub>7</sub><sup>2-</sup> anion lies parallel to the  $\pi$ -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  $H_2P_2O_7^{2-}$  are favoured. The N-H···O distances between the NH protons of **3b** and the O atoms of  $H_2P_2O_7^{2-}$  range from 2.648 to 2.690 Å, separations that are short compared to typical N–H…O distances.<sup>27,28</sup> Each bound H<sub>2</sub>P<sub>2</sub>O<sub>7</sub><sup>2-</sup> 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.28-30 Because the hydrogen atoms on  $H_2P_2O_7^{2-}$  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  $H_2P_2O_7^{2-}$  guest.

Diffraction grade single crystals of the complex  $4b \cdot TBAH_2$ -PO<sub>4</sub> 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<sup>+</sup> counter cation and the bound dihydrogenphosphate guest, which precludes further inter-complex interactions. In the solid state, complex  $4b \cdot H_2PO_4^-$  displays the same conformation for the diformyl-DPM units as observed in  $3b \cdot H_2P_2O_7^{2-}$ . The four N–H…O distances that characterize the interaction between the NH protons of 4b and the O atoms of  $H_2PO_4^-$  range from 2.704 to 2.760 Å. The two O–H…O distances



**Fig. 3** Single crystal structure of the 2 : 2 complex  $[3b \cdot H_2P_2O_7^{2-}]_2$  viewed from the side relative to the bridging benzene rings. Two TBA<sup>+</sup> counter cations and the disordered CH<sub>2</sub>Cl<sub>2</sub> solvent molecules per pyrophosphate are omitted for clarity.



Fig. 4 Single crystal structure of the 1 : 1 complex 4b·H<sub>2</sub>PO<sub>4</sub><sup>-</sup> viewed from the top of the bound H<sub>2</sub>PO<sub>4</sub><sup>-</sup> molecular ion. One TBA<sup>+</sup> counter cation is associated with each complex. It and a CHCl<sub>3</sub> solvent molecule are omitted for clarity.

associated with the hydrogen bonding between the formyl groups and the hydrogen atoms of  $H_2PO_4^-$  are 2.681 and 2.726 Å, respectively. Overall, derivative **4b** stabilizes six hydrogen bonding interactions with the  $H_2PO_4^-$  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  $7\mathbf{b} \cdot \mathbf{H}_2 \mathbf{PO}_4^-$  were obtained by using  $TBAH_2PO_4$  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  $4\mathbf{b} \cdot \mathbf{H}_2 \mathbf{PO}_4^{-}$  discussed above is seen as regard the diformyl DPMdihydrogenphosphate interactions, in  $7\mathbf{b} \cdot \mathbf{H}_2 \mathbf{PO}_4^-$ , the third diformyl-DPM of 7b is too distant to interact well with the bound H<sub>2</sub>PO<sub>4</sub><sup>-</sup> 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 NH-to-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 an ion- $\pi$  interactions, as well as pyrrole NH-anion hydrogen bonding interactions contribute to anion binding. To explore this possibility, the effect of adding TBAH<sub>2</sub>PO<sub>4</sub> to the most soluble receptor of the present series, namely the *meta*-derivative 4b, was investigated by <sup>1</sup>H NMR spectroscopy (Fig. 6). Upon addition of one molar equivalent of TBAH<sub>2</sub>PO<sub>4</sub> to a CDCl<sub>3</sub> 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 <sup>1</sup>H NMR spectroscopic studies of 5b carried out in CDCl<sub>3</sub>, 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<sup>†</sup>). Such shifts are consistent with the decrease in the deshielding effect of the intervening aromatic ring current as seen in previous studies of a,a,a,a-meso-



Fig. 5 Single crystal structure of the 2 : 2 complex  $[7b \cdot H_2PO_4^{-}]_2$ . One TBA<sup>+</sup> counter cation per  $H_2PO_4^{-}$  and 1.5 molecules of CHCl<sub>3</sub> per 7b have been omitted for clarity.



**Fig. 6** Partial <sup>1</sup>H NMR spectra of receptor **4b** before (bottom) and after (top) addition of one molar equivalent of  $TBAH_2PO_4$  in  $CDCl_3$ . Dashed lines show the changes in the chemical shifts of selected proton signals ascribable to receptor **4b**.

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

	٤	δ (ppm)	_	онс сно
Н	<b>4b</b> <sup><i>a</i></sup>	<b>4b</b> ·H₂PO₄ <sup>−b</sup>	$\Delta\delta$	NH HN
H <sup>1</sup>	7.25	7.10	- 0.15	
$H^2$	7.10	6.70	- 0.40	
H <sup>3</sup>	6.90	6.30	- 0.60	H <sup>1</sup> N CHO
$H^4$	6.80	6.85	+ 0.05	
$H^5$	6.00	6.30	+ 0.30	)=-{ Н⁴ СНО

<sup>*a*</sup> The solution of **4b** was prepared from diffraction grade single crystals of **4b**, which contained two DMF molecules of crystallization per equivalent of receptor. <sup>*b*</sup> Obtained by treating with the TBA<sup>+</sup> salt of  $H_2PO_4^-$ ; see text for details.

tetraaryl-C[4]P derivatives by Ballester and co-workers.<sup>31</sup> However, as noted by a reviewer, such an observation is not sufficient to confirm or rule out the presence of an anion– $\pi$ interaction. Intuitively, the interaction of anion H<sub>2</sub>PO<sub>4</sub><sup>-</sup> with the electron-rich intervening benzene ring of **4b** is expected to be repulsive, thus destabilizing the overall binding event. However, Deyà<sup>32</sup> has suggested that the polarization of a  $\pi$ 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– $\pi$  interactions (if any) are playing a substantial role in mediating the observed anion recognition behaviour.

Receptor 7**b** proved even less soluble in neat, non-hydrogen bonding solvents, such as CHCl<sub>3</sub>, than its congeners 3**b**-5**b**. This made it impossible to carry out binding studies that were directly comparable to those reported earlier.<sup>24</sup> However, in analogy to what was seen in the case of these other receptors, it was found that the solubility of 7**b** increased upon the addition of TBAH<sub>2</sub>PO<sub>4</sub>. Dilute solutions of 7**b**·H<sub>2</sub>PO<sub>4</sub><sup>-</sup> in CHCl<sub>3</sub> could thus be prepared by mixing single crystals of 7**b** with *ca.* 2.75 molar equivalents of TBAH<sub>2</sub>PO<sub>4</sub>. The UV-Vis spectrum of this solution was characterized by features similar to those seen in the reported spectrum of 3**b**·H<sub>2</sub>PO<sub>4</sub><sup>-</sup> at analogous concentrations. Thus, the observed spectral features were ascribed to the complex 7**b**·H<sub>2</sub>PO<sub>4</sub><sup>-</sup>. When the CHCl<sub>3</sub> solution made up from 7**b** and TBAH<sub>2</sub>PO<sub>4</sub> was titrated with CH<sub>3</sub>OH, spectral changes and isosbestic behaviour were observed. On this basis, we



Fig. 7 Left: Stacked UV-Vis spectra for titration of 7b ( $1.1 \times 10^{-5}$  M) with TBAH<sub>2</sub>PO<sub>4</sub> ( $2.9 \times 10^{-4}$  M) in CHCl<sub>3</sub> containing 3% (v/v) CH<sub>3</sub>OH. Right: Binding curve fitting plotted from titration data at 302 nm.  $K_a = (1.4 \pm 0.2) \times 10^5$  M<sup>-1</sup>.

conclude that the  $H_2PO_4^-$  affinity of **7b** decreases in the presence of increasing quantities of CH<sub>3</sub>OH (Fig. S15<sup>†</sup>). However, it was also apparent that large quantities of methanol were required to disrupt the complex completely. It was thus decided that a solution of 3% (v/v) CH<sub>3</sub>OH in CHCl<sub>3</sub> would provide a good balance between solubility and binding affinity and allow for quantitative analyses.

In a first set of experiments, an 11  $\mu$ M solution of 7b in 3% (v/ v) CH<sub>3</sub>OH in CHCl<sub>3</sub> was prepared and then subjected to titration with TBAH<sub>2</sub>PO<sub>4</sub> (*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 H<sub>2</sub>PO<sub>4</sub><sup>-</sup> binding in accord with the methods described previously.<sup>24</sup> Similar UV-Vis titrations were carried out with a series of anions in the form of their respective TBA salts. It was found that titration with either the H<sub>2</sub>PO<sub>4</sub><sup>-</sup> or HP<sub>2</sub>O<sub>7</sub><sup>3-</sup> anions gave rise to significant spectral changes with saturation behaviour being observed in the presence of fewer than ten molar equivalents of these two anions. On the other hand, no discernable spectral changes were seen upon titration with HSO<sub>4</sub><sup>-</sup>, Cl<sup>-</sup>, or NO<sub>3</sub><sup>-</sup>.

In the initial studies of **3b–5b**,  $CHCl_3$  was used as the solvent. To allow for a comparison between the original *tetrakis*-(1*H*-pyrrole-2-carbaldehyde) anion receptors and the newer and *hex-akis* analogue **7b**, compound **3b** was subject to titration with  $H_2PQ_4^-$  in 3% (v/v) CH<sub>3</sub>OH–CHCl<sub>3</sub>. The results from this comparison are summarised in Table 3. Briefly, **7b** is characterized by a *ca.* 2 orders of magnitude higher dihydrogenphosphate anion affinity. This increase is in accord with the design expectations underlying **7b**, namely that the presence of a third DPM

Table 3 Summary of calculated binding affinities,  $K_a$  (M<sup>-1</sup>) as determined by UV-Vis spectroscopy in two separate solvent systems<sup>*a*</sup>

$\begin{array}{cccc} & & & \\ H_2 P O_4^{} & & (8 \pm 2) \times 10^6 & & (2.1 \pm 0.3) \times 10^3 & (1.4 \pm 0.2) \times 1 \\ H P_2 O_7^{3-} & & (1.4 \pm 0.1) \times 10^5 & & (3.8 \pm 0.6) \times 10^3 & & (3.6 \pm 0.2) \times 1 \\ H S O_4^{-} & & (4.1 \pm 0.2) \times 10^3 & \text{nd} & & \text{nd} \end{array}$	$\mathbf{3b}^{c}$	$3\mathbf{b}^b$	ns <sup>a</sup> 3	Anions <sup>a</sup>
$NO_3^-$ nd nd nd	$ \begin{array}{ccc} 6 & (2.1\pm0.3)\times10^3 \\ 4 & 10^5 & (3.8\pm0.6)\times10^3 \\ 4 & 10^3 & \text{nd} \\ & \text{nd} \\ & & \text{nd} \end{array} $	$egin{aligned} (8\pm2) imes10^6\ (1.4\pm0.1) imes10^5\ (4.1\pm0.2) imes10^3\ nd \end{aligned}$	$D_4^-$ ( $D_7^{3-}$ ( $4^-$ ( - r	$H_2PO_4^{-}$ $HP_2O_7^{3-}$ $HSO_4^{-}$ $NO_3^{-}$

 $^a$  The anions were studied as their TBA+ salts.  $^b$  In CHCl<sub>3</sub>; data from ref. 24.  $^c$  In CHCl<sub>3</sub> containing 3% (v/v) CH<sub>3</sub>OH.

subunit would force this system to adopt a preorganised conformation in a way that isn't possible for **3b**.

To exploit the binding characteristics of the present diformyl-DPM receptors for the purposes of anion sensing, analogue 6b was prepared. It incorporates a ferrocene spacer and was designed to perform as an electro-active analogue of the tetrakis(1H-pyrrole-2-carbaldehyde) receptors 3b-5b. This new derivative proved poorly soluble in typical organic solvents used for electrochemical studies (e.g., CH<sub>3</sub>CN, DMF, CH<sub>2</sub>Cl<sub>2</sub>). However, as above, an increase in its solubility was seen upon the addition of TBAH<sub>2</sub>PO<sub>4</sub>. This was seen as a "useful property"33 that could be used to obtain insights into the H<sub>2</sub>PO<sub>4</sub><sup>-</sup> binding events under conditions of electrochemical analysis. It was found, for instance, that initial turbid mixtures of free receptor 6b in CH<sub>3</sub>CN 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 TBAH2-PO<sub>4</sub> 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  $H_2PO_4^-$ , 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**  $\cdot H_2PO_4^-$ . 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**  $\cdot H_2PO_4^-$ . When the increase in the anodic peak current is plotted against the added  $H_2PO_4^-$  concentration, a binding isotherm could be



**Fig. 8** Stacked cyclic voltammogram of **6b** in CH<sub>3</sub>CN 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 TBAH<sub>2</sub>PO<sub>4</sub> until the TBAH<sub>2</sub>-PO<sub>4</sub> concentration reached 2.8 mM. TBAPF<sub>6</sub> (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.

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Fig. 9 Views of the single crystal structures of 3b, 4b, and 7b in the absence and presence of bound phosphate derivatives.

obtained (cf. Fig. S20†) from which an approximate  $H_2PO_4^-$  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**<sup>+</sup>) and  $H_2PO_4^-$ .

### 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 <sup>1</sup>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  $H_2PO_4^-$  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.

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## Notes and references

- 1 J. L. Sessler, P. A. Gale and W.-S. Cho, *Anion Receptor Chemistry*, Royal Society of Chemistry, Cambridge, 2006.
- 2 J. H. Ryther and W. M. Dunstan, *Science*, 1971, **171**, 1008–1013.
- 3 A. K. H. Hirsch, F. R. Fischer and F. Diederich, *Angew. Chem.*, *Int. Ed.*, 2007, **46**, 338–352.
- 4 Y. Zhang, E. Desmidt, A. Van Looveren, L. Pinoy,
  B. Meesschaert and B. Van Der Bruggen, *Environ. Sci. Technol.*, 2013, 47, 5888–5895.
- 5 E. A. Katayev, Y. A. Ustynyuk and J. L. Sessler, *Coord. Chem. Rev.*, 2006, **250**, 3004–3037.
- 6 A. E. Hargrove, S. Nieto, T. Zhang, J. L. Sessler and E. V. Anslyn, *Chem. Rev.*, 2011, **111**, 6603–6782.
- 7 J. W. Steed, D. R. Turner and K. Wallace, *Core Concepts in Supramolecular Chemistry and Nanochemistry*, John Wiley, Hoboken, NJ, 2007.
- 8 P. A. Gale, Coord. Chem. Rev., 2003, 240, 191-221.
- 9 S. L. Tobey and E. V. Anslyn, J. Am. Chem. Soc., 2003, 125, 14807–14815.
- 10 J. L. Sessler, M. J. Cyr, V. Lynch, E. McGhee and J. A. Ibers, J. Am. Chem. Soc., 1990, 112, 2810–2813.
- 11 G. Cavallo, P. Metrangolo, T. Pilati, G. Resnati, M. Sansotera and G. Terraneo, *Chem. Soc. Rev.*, 2010, **39**, 3772–3783.
- 12 P. Gamez, T. J. Mooibroek, S. J. Teat and J. Reedijk, Acc. Chem. Res., 2007, 40, 435-444.
- 13 G. A. Jeffrey, *An Introduction to Hydrogen Bonding*, Oxford University Press, Oxford, 1997.
- 14 Note: Under certain conditions, deprotonation, rather than anion binding, has been observed with pyrrolic receptors.
- 15 S. Camiolo, P. A. Gale, M. B. Hursthouse, M. E. Light and A. J. Shi, *Chem. Commun.*, 2002, **3**, 758–759.
- 16 B. L. Iverson, K. Shreder, V. Král and J. L. Sessler, J. Am. Chem. Soc., 1993, 115, 11022–11023.
- 17 P. A. Gale, J. L. Sessler, V. Král and V. Lynch, *J. Am. Chem. Soc.*, 1996, **118**, 5140–5141.
- 18 A. Baeyer, Ber. Dtsch. Chem. Ges., 1886, 19, 2184-2185.
- 19 R. Custelcean, L. H. Delmau, B. A. Moyer, J. L. Sessler, W.-S. Cho, D. Gross, G. W. Bates, S. J. Brooks, M. E. Light and P. A. Gale, *Angew. Chem., Int. Ed.*, 2005, 44, 2537–2542.
- 20 D. E. Gross, F. P. Schmidtchen, W. Antonius, P. A. Gale, V. M. Lynch and J. L. Sessler, *Chem.-Eur. J.*, 2008, 14, 7822–7827.

- 21 W.-S. Cho, PhD Dissertation, The University of Texas at Austin, 2005.
- 22 J. L. Sessler, W.-S. Cho, S. P. Dudek, L. Hicks, V. M. Lynch and M. T. Huggins, *J. Porphyrins Phthalocyanines*, 2003, **07**, 97–104.
- 23 J. L. Sessler, S. Camiolo and P. A. Gale, *Coord. Chem. Rev.*, 2003, **240**, 17–55.
- 24 M. K. Deliomeroglu, V. M. Lynch and J. L. Sessler, *Chem. Commun.*, 2014, **50**, 11863–11866.
- 25 R. Neufeld and D. Stalke, Chem. Sci., 2015, 6, 3354-3364.
- 26 R. Custelcean, Chem. Soc. Rev., 2010, 39, 3675-3685.
- 27 N. L. Bill, D.-S. Kim, S. K. Kim, J. S. Park, V. M. Lynch, N. J. Young, B. P. Hay, Y. Yang, E. V. Anslyn and J. L. Sessler, *Supramol. Chem.*, 2012, 24, 72–76.

- 28 J. Cai, B. P. Hay, N. J. Young, X. Yang and J. L. Sessler, *Chem. Sci.*, 2013, **4**, 1560–1567.
- 29 V. Král, H. Furuta, K. Shreder, V. Lynch and J. L. Sessler, *J. Am. Chem. Soc.*, 1996, **118**, 1595–1607.
- 30 M. Wei, B. Wu, L. Zhao, H. Zhang, S. Li, Y. Zhao and X.-J. Yang, *Org. Biomol. Chem.*, 2012, **10**, 8758–8761.
- 31 G. Gil-Ramírez, E. C. Escudero-Adán, J. Benet-Buchholz and P. Ballester, *Angew. Chem., Int. Ed.*, 2008, 47, 4114–4118.
- 32 D. Quiñonero, A. Frontera and P. M. Deyà, in *Anion Coordination Chemistry*, ed. K. Bowman-James, A. Bianchi and E. García-Espana, Wiley-VCH Verlag & Co. KGaA, Weinheim, 2012, ch. 6, pp. 321–362.
- 33 K. A. Connors, *Binding constants: The Measurement of Molecular Complex Stability*, Wiley, New York, 1987.