Introduction

Much theoretical and synthetical effort has been devoted to lift non-ionic organic bases to the basicity level of common inorganic or metalorganic bases.\textsuperscript{1,2} With his famous phosphazenes Schwesinger established a widely used and commercially available class of (organo-)superbases.\textsuperscript{3,4} His homologization concept, the stepwise expansion of the molecular scaffold in order to better delocalize the positive charge formed upon protonation, was also applied to synthesize higher-order N-superbases of guanidines,\textsuperscript{5,6} imidazolidine amines\textsuperscript{7} and cyclopropenimines.\textsuperscript{8,9} However, such basicity enhancement is accompanied by an unwanted growth of the bases’ molecular weight. Therefore, other strategies for augmenting the intrinsic proton affinity have been investigated: in proton sponges, a second nitrogen basicity centre in close proximity to the proton a weight. \textsuperscript{10} Additional thermodynamic driving force comes from relief of strain of the aromatic backbone.\textsuperscript{11} Many derivatives of such proton sponges were designed by combining aforementioned superbasic functionalities with the 1,8-dia- minonaphthalene structural motif\textsuperscript{12} or as proton pincers with different backbones.\textsuperscript{13}

Atoms other than nitrogen as basicity centre were also applied, such as phosphorus.\textsuperscript{14,15} Recently, we demonstrated, that N-phosphazenyl substituted phosphines (PAPs) possess higher pK\textsubscript{BH}\textsuperscript{+} values as P\textsuperscript{III} bases than their corresponding phosphazene P\textsuperscript{V}N\textsubscript{Bu} counterparts as N bases.\textsuperscript{16} So far the limit of homologization is reached at the P\textsubscript{V} level both in phosphazenyl phosphazenes and phosphazenyl phosphines as both P\textsubscript{V} benchmark bases have only been isolated in their protonated form.\textsuperscript{16,17}

Non-ionic carbon is another contender to extend the basicity ladder to unmatched regions.\textsuperscript{18} In this respect phosphorus (mono-)ylides\textsuperscript{19,20} as well as bisylidic proton sponges\textsuperscript{21} were investigated on theoretical and experimental level. Although identified as potential superbases, the application of N-heterocyclic carbenes (NHCs),\textsuperscript{22} cyclic alkyl amino carbenes (CAACs),\textsuperscript{23} carbodicarbenes (CDCs),\textsuperscript{24} and carbodiphosphoranes (CDPs)\textsuperscript{25} has been exploited predominantly as strong Lewis bases towards transition and main group elements other than the proton.\textsuperscript{26}

The prototypic hexaphenyl carbodiphosphorane ([Ph]\textsubscript{6}-CDP) was first synthesized 1961 by Ramirez et al.\textsuperscript{27} Further compounds like the hexamethyl carbodiphosphorane ([Me]\textsubscript{6}-CDP),\textsuperscript{28} hexakis(dimethylamino) carbodiphosphorane ([dma]\textsubscript{6}-CDP),\textsuperscript{29} and mixed representatives followed.\textsuperscript{30–32}

Herein we promote carbodiphosphoranes with their electron-rich R\textsubscript{1}P–C–PR\textsubscript{3} functionality as exceptionally strong carbon Bronsted bases. As bisylides with a π-symmetric HOMO and σ-symmetric HOMO–1, both mainly located as lone pairs at the carbon, only slightly stabilized by back-bonding \textit{via} negative hyperconjugation,\textsuperscript{33} they provide outstanding pK\textsubscript{BH}\textsuperscript{+} values in particular for the first of two protonation steps. We present a synthesis for hexa(pyrrolidino) carbodiphosphorane ([prr]\textsubscript{6}-CDP) with its calculated first and second proton affinity (PA) of 287.6 and 188.9 kcal mol\textsuperscript{-1},\textsuperscript{34} which exceeds the PAs of (Ph)\textsubscript{6}-CDP (280.0 and 185.6 kcal mol\textsuperscript{-1})\textsuperscript{34} and (dma)\textsubscript{6}-CDP (279.9 and 174.9 kcal mol\textsuperscript{-1}).\textsuperscript{34} Furthermore we apply the homologization concept to CDPs by introducing PR\textsubscript{3} units bearing one intrinsically superbasic substituent R’ to access CDP...
superbases of second-order. We thereby focused on \(N\)-tetramethylguanidinyl (\(\text{tmg}\)) and \(N\)-tris(dimethylamino)phosphazenyldimethylamino (\(\text{dmap}_1\)) substituents targeting new carbodiphosphoranes \(\text{sym-}(\text{tmg})_2(\text{dmap})_4\)-CDP and \(\text{sym-}(\text{dmap})_2(\text{dmap})_4\)-CDP.

Results and discussion

Synthesis

We experienced, that the established synthesis routes to CDPs are inappropriate for phosphines more electron-rich than \(\text{P}(\text{NMMe}_2)_3\): reactions between such phosphines \(\text{P}(\text{NR}_2)_2\text{R}\) and \(\text{CCl}_4\) did not follow the pattern outlined in ref. 32 and 35 but exclusively led to chlorination of the phosphine, whilst reactions with methylene bromide did not selectively follow the path outlined in ref. 30 and 36, but led to a 1:1-mixture of the methylated phosphonium bromide \([\text{R}(\text{NR}_2)_2\text{P-Me}]\text{Br}\) and the brominated species \([\text{R}(\text{NR}_2)_2\text{P-Br}]\text{Br}\). Therefore we further developed an alternative strategy laid out by Appel et al. for the synthesis of \(\text{dmap}_1\)-CDP.\(^{29}\) The doubly protonated precursors of the second-order carbodioporphane superbases, \(\text{sym-}(\text{tmg})_2(\text{dmap})_4\)-CDP (1) and \(\text{sym-}(\text{dmap})_2(\text{dmap})_4\)-CDP (2), were obtained in an oxidative imination sequence as shown in Scheme 1. Bis[\(\text{bis}(\text{dimethylamino})\text{phosphino})\text{methane} (3) was oxidized by \(\text{CCl}_4\) in presence of tetramethylguanidine (\(\text{Htmg}\)) or tris(dimethylamino)phosphazene \((\text{dmap}_1\cdot \text{H})\) instead of dimethylamine as nucleophile and auxiliary base. This reaction offers the advantage of preformed C-P-bonds avoiding the preparation of respective \(\text{P}^{\text{III}}\) nucleophiles.\(^{15,20,37}\) 3 is readily synthesized in two steps on a large scale\(^{18}\) and the selected superbasic building blocks oxidatively introduced as nucleophiles are either commercially available or easily accessible in few steps.\(^{4}\)

The synthesis of 4-\(\text{HBF}_4\), the precursor for (pyrr)\(_6\)-CDP 4, was accomplished in a one-pot synthesis (Scheme 2), since the intermediate bis[(pyrrolidino)phosphino)methane (5) turned out to decompose upon vacuum distillation. Starting from bis(dichlorophosphino)methane\(^{38}\) (6), 5 was prepared \(\text{in situ}\) with an excess of pyrrolidine (Fig. S1 in the ESI\(^+\)) and directly oxidized with \(\text{CCl}_4\).

In all three reactions the respective monoprotonated hydrochloride adducts were identified as products via \(\text{\textsuperscript{31}P}\) NMR spectroscopy. Therefore the second \(\text{pK}_\text{BHF}^+\) values in THF of these new CDPs are obviously lower than that of the auxiliary base pyrrolidine (13.5),\(^{39}\) tetramethylguanidine (15.5),\(^{40}\) or tris(dimethylamino)phosphazene 2a (19.7),\(^{41}\) respectively. For purification, the crude products were precipitated with \(\text{NaBF}_4\) from aqueous solution. These conditions lead to second protonation at the central carbon atom and a strongly alkaline solution. Therefore, even the monoprotonated CDPs can be considered as strong cationic bases in aqueous medium. Similar behaviour was found for \(\text{(Ph)}_6\)-CDP in water, although the latter is slowly hydrolysed under ambient conditions,\(^{27}\) which is not the case for peramated CDPs 1, 2 and 4 reported here.

The bis(tetrafluoridoborate) salts of 1, 2 and 4 were obtained in 50–60% yield as water and air stable, colourless solids, indefinitely storable. They are well soluble in polar organic solvents like methanol, acetonitrile or DMSO but insoluble in less polar solvents such as ethers and hydrocarbons.

For the liberation of the free CDPs different suitable bases were identified: for 4 potassium bis(trimethylsilyl)amide (\(\text{KHMD}\)) is of sufficient basicity, whilst for 1 the more basic sodium amide (\(\text{NaNH}_2\)) is necessary for full deprotonation. Both new bases 1 and 4 could be isolated in 70% and 60% yield, respectively, from \(\text{n}-\text{hexane}\) as pure colourless crystalline solids, indefinitely storable at room temperature under inert conditions. Contrastingly we were not able to isolate 2 as free CDP base form. Sodium amide in liquid ammonia or suspended in THF at room temperature selectively abstracts the first proton under formation of 2-\(\text{HBF}_4\) as colourless solid in 69% yield. At elevated temperature the central carbon atom is not further

Scheme 1 Preparation of CDP precursors 1-\(\text{HBF}_4\) and 2-\(\text{HBF}_4\) together with subsequent deprotonation to 1 (one exemplary mesomeric structure displayed) and 7, respectively. Numbering schemes refer to assigned NMR signals in the experimental section.
deprotonated, even though it is the thermodynamically most acidic site (see Theoretical Calculations). Instead NaNH₂ deprotonates selectively one of the dimethylamino groups at the terminal phosphazene moiety which results in the irreversible elimination of N-methylmethanimine and reduction of the phosphazene to a phosphine (Scheme 1). A related deprotonation and reduction of tetrakis(dimethylamino)phosphonium bromide under the action of NaNH₂ was described by Pinchuk et al.⁴⁵ In case of this reaction is slow but highly selective and 2 could be obtained as sole product as pale yellow highly viscous oil. The proposed configuration was confirmed via ¹H, ¹³C, and ³¹P NMR spectroscopy and by HR mass spectrometry. 7 can be considered as a hybrid between mixed valence phosphazyl phosphines⁵⁻⁶ and ylide PIII/PV compounds of the type (Me₂N)₃P=CH(P₂) (ref. 42) or other ylide-functionalized phosphines.⁴⁶ Further attempts to deprotonate 2·2HBF₄ with other bases or reducing agents resulted either in only single deprotonation (benzyl potassium in THF), in an unselective disintegration (nBuLi) or in the same deprotonation of the P-NMe₂ group (potassium in liquid ammonia, ethylene diamine, THF, or DME or an excess of benzyl potassium in THF). The reaction of potassium hydride in THF gave a mixture of 7 as minor component and presumably free CDP 2 as major product by means of ³¹P NMR spectroscopy (Fig. S29 in the ESI†). Clearly the acidity of P³⁻attached NMe₂ groups limits the accessibility of 2. Under the action of excess of strong inorganic bases at elevated temperatures the stability limit of these phosphazene moieties seems to have been reached.

For analytical reasons the monoprotonated forms of 1 and 4 were prepared on NMR scale either via commutation between the free CDP and its bisprotonated form or by protonating the free CDPs with one equivalent triflimidic acid (HTFSI).

**Structural features**

For X-ray structure determination suitable single crystals were obtained from n-hexane for both presented CDPs 4 and 1. They crystallize solvent-free in space group P2₁/c or Pbnm, respectively, with one complete molecule per asymmetric unit (Fig. 1). Contrary to the parent compound (dma)₆-CDP, one of the hitherto two reported linear CDPs,²⁸,⁴⁴ a bent structure with P-C-P angles of 155.9(2)° and 147.3(9)°, respectively is found. Since the potential for bending at the central P-C-P carbon atom in polymorphic (Ph)₆-CDP is very flat⁴⁴ and reveals high dependence of the crystallization method,⁴⁵ the obtained crystals of (dma)₆-CDP from the melt are maybe the reason for its linearity.²⁹ The P–C_central distances are with 1.606 Å (4) and 1.618 Å (1) in the for CDPs reported range: (dma)₆-CDP: 1.584(1) Å,²⁹ (Me)₆-CDP: 1.594(3) Å,⁴⁶ (Ph)₆-CDP: 1.601–1.635 Å.⁴⁴,⁴⁷ On average, pyrrolidine N–P distances in 4 are 1.68 Å while those of dma and tmg groups in 1 are 1.70 Å and 1.66 Å respectively.

Single crystals obtained from reaction control samples for the synthesis of 4·2HBF₄ turned out to be a cococrystallate of 4·2HCl and pyrrolidinium chloride (Fig. 2). Cations and anions form a C•••H–Cl•••H–N hydrogen bond network with C–H–Cl distances of 3.600(2) Å and N–Cl distances of 3.018(2) Å and

![Image](http://example.com/image1.png)

**Fig. 1** Molecular structure of 4 (top) and 1 (bottom). Hydrogen atoms omitted for clarity, ellipsoids at 50% probability. Selected bond length/Å and angles/°: 4 P₁–C₁ 1.605(2), P₁–N₁ 1.672(2), P₁–N₂ 1.678(2), P₁–N₃ 1.694(2), P₂–C₁ 1.606(2), P₂–N₁ 1.669(2), P₂–N₂ 1.671(2), P₁–C₁–P₂ 159.5(9), C₁–P₁–N₁ 110.2(1), C₁–P₁–N₂ 115.1(1), C₁–P₁–N₃ 121.8(1), C₁–P₂–N₁ 118.4(1), C₁–P₂–N₂ 115.1(1), C₁–P₂–N₃ 111.3(1), C₃–P₁–C₁–P₂ 168.0(4), N₄–P₂–C₁–P₁ 130.6(4), P₁–C₁–N₁ 1.619(1), P₁–N₁ 1.680(1), P₃–N₁ 1.671(1), C₁–P₁–N₁ 1.714(1), P₁–N₁ 1.665(1), N₁–C₂ 1.298(2), N₂–C₂ 1.377(2), N₃–C₂ 1.382(2), P₂–C₁ 1.671(1), P₂–N₉ 1.719(1), P₂–N₁ 1.680(1), P₂–N₆ 1.664(1), N₆–C₁–N₁ 1.299(2), N₆–C₁ 1.376(2), N₈–C₁–N₁ 1.379(2), P₂–C₁–P₁ 147.3(9), C₁–P₁–N₁ 109.5(2), C₁–P₁–N₂ 109.5(2), C₁–P₁–N₃ 119.8(5), C₂–N₁–P₁ 128.1(1), C₁–P₂–N₉ 120.76(6), C₁–P₂–N₁ 110.08(6), C₁–P₂–N₆ 119.47(6), C₁–N₆–P₂ 127.3(1), N₄–P₁–C₁–P₂ 162.2(2), N₁₀–P₂–C₁–P₁ 155.8(2).
Peripheral hydrogen atoms and BF$_4$-anions omitted for clarity, ellipsoids independent molecules depicted, structure factors given for both).

1.799(1), P1

$soids at 50% probability. # marked atoms generated from $2HBF_4$ as well. All three bisprotonated CDPs exhibit a strong influence of charge delocalization as the reason for their extraordinary basicity: upon protonation the P–C bonds elongate from 1.606 Å (4) and 1.618 Å (1) to 1.799 Å in $2HCl$ and 1.821 Å in $2HBF_4$ and $2HBF_4$, whilst the P–N bonds become shorter to average 1.62 Å for pyrrolidine and 1.64 Å for dimethylamine substituents. This complies with distances found in protonated phosphazenes$^{23}$ and phosphorus ylids$^{22}$ and proves the electron donating effect of the amino substituents. The P–N bonds to the tmg groups in $2HBF_4$ exhibits with 1.58 Å (1.66 Å in 1) clearly double-bond character. The P–N=C angles are expanded from 127° to 128° to 132° and 136°. A diminishing difference of formal N=C single and double bonds in the tmg group indicates the conjugation within the CN$_2$ moiety. The formal P–N single and double bonds of the phosphazene substituents in $2HBF_4$ equalize at 1.57–1.59 Å with P–N angles between 134° and 142°. Similar influence of negative hyperconjugation for charge delocalization was found in superbasic PAPs$^{26}$ and protonated diphosphazenes.$^{51}$ The P–C–P angles in the bisprotonated forms (4: 120°, 1: 113°, 2: 121°) are more acute than in the free CDPs (4: 156°, 1: 147°). The difference to ideal tetrahedral geometry presumably arise from the bulkiness of the PR$_3$ moieties.

**NMR spectroscopic features**

All six presented compounds were characterized by $^1$H, $^{13}$C, and $^{31}$P NMR spectroscopy. Selected chemical shifts and couplings are collected in Table 1. Proton shifts of bis- and monoprotonated CDPs lie around 3 ppm for CH$_2$ and below 1 ppm for CH groups, both decreasing with increasing basicity of the parent CDP indicating less polarized C–H bonds. This shielding trend is not observed in the $^{13}$C NMR shifts of the carbon nuclei: the most basic CDP 1 exhibits a triplet at 9.5 ppm compared to $–1.6$ ppm (4) and $–6.8$ ppm ([dma]$_n$CDP)$_2$. Surprisingly the $^{13}$C chemical shift for 1 is even higher than for its monoprotonated form (1–HTFSI: 9.3 ppm) contrasting the typical trend 1.579(4), P4–N10/P8–N16 1.644(4)/1.652(4), P4–N11/P8–N17 1.635(4)/1.648(4), P4–N12/P8–N18 1.655(4)/1.637(4), P3–C1–P1/P5–C22 120.9(2)/121(2), N1–P1–C1/N19–P5–C22 110.8(2)/109.8(2), N2–P1–C1/N20–P6–C22 103.8(2)/104.0(2), N3–P1–C1/N21–P5–C22 107.9(2)/108.4(2), P1–N3–P2/P5–N21–P6 138.2(3)/135.7(3), N7–P3–C1/N14–P7–C22 111.2(2)/112.4(2), N8–P3–C1/N13–P7–C22 105.0(2)/103.2(2), N9–P3–C1/N15–P7–C22 107.9(2)/107.4(2), P3–N9–P4–P7–N15–P8 133.6(3)/141.6(3), N2–P1–C1/P3–N20–P5–C22–P7 164.8(3)/166.8(3), N8–P3–C1/P1/N13–P7–C22–P5 165.6(3)/164.3(3).
Quantumchemical calculations

First and second proton affinity (PA) and gas-phase basicity (GB) of carbodiphosphoranes 1, 2, 4 and phosphine 7 are calculated utilizing M06-2X/6-31+G(d) theoretical model. pK_BH⁺ values in THF are obtained using the same functional and basis set whereas solvent is treated as dielectric continuum utilizing the SMD solvation model. pK_BH⁺ values are calculated as relative values using an isodesmic reaction approach where Schwesingers (dma)P₂-tBu phosphazene with pK_BH⁺ of 33.9 (ref. 20) has served as a reference base. Calculated values for protonation at central carbon atom, and in case of 7 protonation at the PIII atom as well, are presented in Table 2. It appears that the first proton affinity as well as pK_BH⁺ values of 1 and 2 are higher than in Schwesingers (dma)P₂-tBu phosphazene which has PA of 293.3 kcal mol⁻¹ calculated at the same level of theory. Interestingly first GB of 1 is slightly lower than the GB of (dma)P₂-tBu (GB = 288.2 kcal mol⁻¹) implying that the higher pK_BH⁺ value of 1 relative to (dma)P₂-tBu is a result of a more pronounced solvation effect in the carbodiphosphorane. This is unexpected considering that the N–H bond in a protonated phosphazene has a higher polarity than the C–H bond in protonated CDP as a result of lower electronegativity of carbon relative to nitrogen. The calculated pK_BH⁺ (THF) 39.1 of 2 would be far higher than the pK_BH⁺ (THF) 33.9 of (dma)P₂-tBu, the strongest commercially available superbase. As described isolation of neutral base 2 is not achieved experimentally as other C–H bonds in the precursor 2–H⁺ seemed to have a higher kinetic and thermodynamic acidity. In order to understand the deprotonation path of 2–H⁺ under the action of NaN₃, the reaction profile is calculated and presented in Fig. S36 in the ESI.† It appears, that the deprotonation of peripheral NMe₂ group in combination with the irreversible elimination of N-methylmethanimine is thermodynamically feasible (exergonic), however, kinetically hindered by a high barrier (ΔG² = 32.8 kcal mol⁻¹). This explains, that deprotonation induced degradation is competitive to deprotonation of central carbon atom at elevated temperatures, though the central carbon atom in 2–H⁺ is the thermodynamically most acidic site. It appears that decomposition product – phosphine 7 – has a gas-phase

Experimental values in parentheses.

Table 2. Calculated first and second proton affinity (PA) and gas phase basicity (GB) together with pK_BH⁺ values in THF

<table>
<thead>
<tr>
<th>Species</th>
<th>PA/kcal mol⁻¹</th>
<th>GB/kcal mol⁻¹</th>
<th>pK_BH⁺ in THF</th>
</tr>
</thead>
<tbody>
<tr>
<td>4</td>
<td>291.1</td>
<td>282.2</td>
<td>32.8 (30.1–32.9)</td>
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<tr>
<td>2</td>
<td>191.6</td>
<td>184.0</td>
<td>—</td>
</tr>
<tr>
<td>1</td>
<td>294.4</td>
<td>287.2</td>
<td>34.9 (35.8 ± 1)</td>
</tr>
<tr>
<td>2</td>
<td>202.0</td>
<td>194.1</td>
<td>—</td>
</tr>
<tr>
<td>2</td>
<td>305.3</td>
<td>299.7</td>
<td>39.1</td>
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<tr>
<td>2</td>
<td>212.1</td>
<td>202.2</td>
<td>—</td>
</tr>
<tr>
<td>7</td>
<td>275.9</td>
<td>268.7</td>
<td>24.4</td>
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<tr>
<td>At carbon</td>
<td>276.2</td>
<td>268.8</td>
<td>21.1</td>
</tr>
<tr>
<td>At phosphorus</td>
<td>276.2</td>
<td>268.8</td>
<td>21.1</td>
</tr>
</tbody>
</table>

Experimental values in parentheses.
basicity (30.9 kcal mol\(^{-1}\)) much lower than CDP 2. Interestingly, GB value for protonation at central carbon and \(\text{P}^{\text{III}}\) phosphorus of 7 is almost the same, whereas \(\text{pK}_{\text{BH}^+}\) in THF for protonation at \(\text{P}^{\text{III}}\) is by 3.3 orders of magnitude lower than \(\text{pK}_{\text{BH}^+}\) for protonation at carbon, which again indicates a more pronounced solvation effect in C-protonated CDP.

Conclusions

In this work we presented the most basic uncharged carbon bases known so far. A convenient synthesis for first- and second-order carbodiophosphorane superbases was presented. The CDPs \((\text{pyrr})_6\)-CDP 4 and sym-(tmg)\(_2\)(dma)\(_4\)-CDP 1 were synthesized as free base as well as in their mono- and bisprotonated forms. In our attempt to synthesize the even more outstanding base sym-(dmaP\(_1\))\(_2\)(dma)\(_4\)-CDP 2 an unexpected, but highly selective deprotonation at peripheral PNH\(_3\) bonds induced an irreversible elimination path towards phosphine 7. This reaction is indicating a potential basicity limit for phosphazene containing superbases. Structural as well as spectroscopic features were investigated and the basicity was quantified by theoretical and experimental means. Remarkable \(\text{pK}_{\text{BH}^+}\) values for 4 and 1 confirm them as benchmark breakers for non-ionic carbon bases on the THF basicity scale. Compared to the top Schenwiesi bases, this basicity is even more outstanding, if their molecular weight below 500 g mol\(^{-1}\) is considered. We expect, that such simply synthesized carbodi-phosphoranes with water stable protonated forms will enter the field of organic superstarbase catalysis.\(^1\)

Experimental section

General

All Reactions with air or moisture sensitive substances were carried out under inert atmosphere using standard Schlenk techniques. Air or moisture sensitive substances were stored in a nitrogen-flushed glovebox. Solvents were purified according to common literature procedures and stored under an inert atmosphere over molsieve (3 Å or 4 Å).\(^4\) Pyrrolidine and tetramethylguanidine were distilled from CaH\(_2\), trichloromific acid was purified by sublimation under argon. Bis(dichlorophosphino)methane\(^8\) (6), bis[bis(dimethylamino)phosphino]methane\(^8\) (3), tris(dimethylamino)phosphazene\(^4\) and \((\text{pyrr})_6\)-tBu\(^4\) were prepared according to literature-known procedures. \((\text{dma})\_4\)-tBu was purchased as 1 M solution in hexane and dried in high vacuum. All other reagents were used as provided.

\(^1\)H, \(^13\)C, and \(^31\)P NMR spectra were recorded on a Bruker Avance III HD 250, Avance II 300, Avance III HD 300 or Avance III HD 500 spectrometer. Chemical shift \(\delta\) is denoted relatively to SiMe\(_4\) \((\text{H}, \text{C})\) or 85% H\(_2\)PO\(_4\) \((\text{P})\). \(^1\)H and \(^13\)C NMR spectra were recorded on a Bruker Alpha ATR-FT-IR. CCDC 1903830 (4·2HCl + PyrrCl\(_4\)), 1903833 (1·2HBF\(_4\)), 1903838 (2·2HBF\(_4\)), 1903840 (1), 1903841 (4·2HBF\(_4\)), and 1903843 (4) contain the supplementary crystallographic data for this paper.\(^\dagger\)

General procedure for the precipitation of BF\(_4\)-salts

The crude product was dissolved in a minimum amount of water and a concentrated aqueous sodium tetrafluoroborate solution (2.0 eq.) was added. The resulting precipitate was filtered off, rinsed three times with small portions of cold water, washed with THF and dried in high vacuum.

\((\text{pyrr})_6\)-CDP 2HBF\(_4\) (4·2HBF\(_4\))

6 (3.60 g, 16.5 mmol, 1.00 eq.) was dissolved in THF (60 mL), cooled to \(-78 ^\circ\)C and pyrrolidine (17.7 mL, 216 mmol, 13.1 eq.) was added dropwise. Afterwards the cooling bath was removed and the mixture stirred for additional 6 h. Carbon tetrachloride (3.12 mL, 32.3 mmol, 1.96 eq.) was added at \(-78 ^\circ\)C and the mixture allowed to warm to room temperature overnight. The suspension was filtered under air and the filter cake extracted with THF (3 × 60 mL). The solvent was removed under reduced pressure and the residue dried in high vacuum. The crude product was converted to its tetrafluoroborate salt as described in the general procedure and recrystallized from ethanol/ethanol. 4·2HBF\(_4\) (6.38 g, 9.52 mmol, 58%) was obtained as colourless solid.

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[\text{C}_{25}\text{H}_{50}\text{B}_{2}\text{F}_{8}\text{N}_{6}\text{P}_{2}] (670.27 \text{ g mol}^{-1}) \quad \text{H NMR (500.2 MHz, CD}_{2}\text{CN)}: \delta (\text{ppm}) = 3.43 (t, 3_{\text{J}_{\text{PH}}} = 19 \text{ Hz}, \text{H}, \text{CH}_{2}), 3.25–3.22 (m, 24\text{H}, \text{H}1), 1.97–1.95 (m, 24\text{H}, \text{H}2, \text{overlapped with the solvent signal}). \quad \text{13}^1\text{C}[^1\text{H}] \text{ NMR (125.8 MHz, CD}_{2}\text{CN)}: \delta (\text{ppm}) = 48.7 (s, \text{C}1), 26.9–26.8 (m, \text{C}2), 26.4 (t, \text{J}_{\text{P}_{\text{C}}} = 110 \text{ Hz}, \text{CH}_{2}). \quad \text{31}^1\text{P}[^1\text{H}] \text{ NMR (121.5 MHz, CD}_{2}\text{CN)}: \delta (\text{ppm}) = 32.7. \quad \text{ESI}^{+}(\text{MS (MeOH): m/z (\%)) = 495.6 (100) [M – H – 2BF\(_4\)]^+, 583.2 (5) [M – BF\(_4\)]^+. \quad \text{ESI}^{+}(\text{HRMS: m/z [M – H – 2BF\(_4\)]\_c} = 495.3488, \text{found 495.3505; [M – BF\(_4\)]\_c} = 583.3600, \text{found 583.3611. Elemental analysis: calced C 44.80%, H 7.52%, N 12.54%; found C 44.49%, H 7.50%, N 12.46%. IR (neat): \(\tilde{\nu} (\text{cm}^{-1})\) = 2970 (w), 2879 (w), 1458 (w), 1251 (w), 1210 (m), 1134 (m), 1047 (vs), 1021 (vs), 918 (m), 870 (m), 824 (m), 779 (m), 699 (m), 581 (w), 549 (w), 517 (m) 484 (s). \quad \text{XRD: for single crystal X-ray structure determination suitable single crystals were obtained by slow evaporation of a concentrated solution in chloroform.}

sym-(tmg)\(_2\)(dma)\(_4\)-CDP 2HBF\(_4\) (1·2HBF\(_4\))

3 (831 mg, 3.29 mmol, 1.00 eq.) and tetramethylguanidine (1.14 g, 9.88 mmol, 3.00 eq.) were dissolved in THF (60 mL). Carbon tetrachloride (640 \mu L, 6.62 mmol, 2.01 eq.) was added at \(-78 ^\circ\)C and the mixture allowed to warm to room temperature overnight. The suspension was filtered under air and the filter cake extracted with THF (3 × 20 mL). The solvent was removed under reduced pressure and the residue dried in high vacuum. The crude product was converted to its tetrafluoroborate salt as described in the general procedure and recrystallized from ethanol. 1·2HBF\(_4\) (1.08 g, 1.66 mmol, 50%) was isolated as colourless solid.
A solution of potassium bis(trimethylsilyl)amide (558 mg, 2.80 mmol, 2.09 eq.) in THF (15 mL) was added to a suspension of \( \text{[C}_{10}\text{H}_{50}\text{B}_{2}\text{F}_{6}\text{N}_{10}\text{P}_{2}] \) (654.24 g mol\(^{-1}\)) \(^1\)H NMR (500.2 MHz, CD\(_2\)CN): \( \delta \) (ppm) = 3.16 (d, \( ^1J_{F\text{H}} = 17 \text{ Hz}, \text{CH}_2 \)), 2.91 (s, 24H, \( \text{H} \)), 2.53 (d, \( ^1J_{F\text{H}} = 10 \text{ Hz}, \text{H} \)), 2.16 (t, \( ^1J_{F\text{H}} = 11 \text{ Hz}, \text{N} \)). \(^{13}\)C\(^{1}\)H NMR (125.8 MHz, CD\(_2\)CN): \( \delta \) (ppm) = 161.6 (dd, 2\( \times \)^1J\(_{PC} = 2 \text{ Hz}, \text{CN} \)), 40.9 (s, C1), 37.1 (dd, 2\( \times \)^1J\(_{PC} = 2 \text{ Hz}, \text{C} \)), 25.2 (t, \( ^1J_{PC} = 112 \text{ Hz}, \text{CH} \)). \(^{31}\)P\(^{1}\)H NMR (205.25 MHz, CD\(_2\)OH): \( \delta \) (ppm) = 20.8 (s, \( ^1J_{PC} = 113 \text{ Hz} \) (satellites)). ESI(+) MS (MeOH): \( m/z \) \( \% \) = 479.5 (100) [M - H - 2BF\(_4\)^+]. ESI(+) HRMS: \( m/z \) [M - H - 2BF\(_4\)]\(^+\) calcd. 479.3622, found 479.3625. Elemental analysis: calc C 34.38\%, H 7.70\%, N 21.41\%; found C 34.98\%, H 7.84\%, N 21.39\%. IR (neat): \( \tilde{\nu} \) (cm\(^{-1}\)) = 2911 (br. w.), 1539 (s), 1486 (m), 1429 (m), 1401 (m), 1356 (m), 1238 (m), 1186 (m), 1161 (m), 1046 (vs.), 1034 (vs.), 979 (vs.), 933 (vs.), 784 (s), 771 (s), 739 (m), 716 (m), 690 (w), 672 (w), 618 (w), 572 (m), 519 (m), 459 (m), 437 (m). XRD: for single crystal X-ray structure determination suitable single crystals were obtained from ethanol/n-hexane at \(-25^\circ\)C.

**sym-(dmpa)\(^2\)(dmpa)\(_4\)CDP (2-2HBF\(_4\))**

A mixture of \( \text{1-}2\text{HBF}_4 \) (190 mg, 290 \( \mu \)mol, 1.00 eq.) and sodium amide (113 mg, 2.90 mmol, 10.0 eq.) was stirred in THF (15 mL) for 16 h at room temperature. The suspension was filtered over Celite and the filter cake extracted with THF (3 \x3c; 5 mL). All volatiles were removed in \textit{vacuo}, n-hexane (10 mL) added to the residue, filtered again over Celite and extracted with n-hexane (3 \x3c; 4 mL). Evaporation of the solvent and drying in high vacuum yielded 1 (86 mg, 0.17 mmol, 60%) as colourless solid. \[^1\]H NMR (500.2 MHz, CD\(_2\)CN): \( \delta \) (ppm) = 2.88 (dd, 2\( \times \)^1J\(_{F\text{H}} = 5 \text{ Hz}, 24H, H)), 2.73 (s, 24H, \( \text{H} \)), \(^{13}\)C\(^{1}\)H NMR (125.8 MHz, CD\(_2\)CN): \( \delta \) (ppm) = 156.0 (s, CN), 40.1 (s, C1), 38.3 (s, C2), 9.5 (t, \( ^1J_{PC} = 209 \text{ Hz}, \text{PC} \)). \(^{31}\)P\(^{1}\)H NMR (205.25 MHz, CD\(_2\)OH): \( \delta \) (ppm) = 18.2. LIFDI(+) HRMS: \( m/z \) [M + H\(^+\)]\(^+\) calcd. 479.3619, found 479.3629. Elemental analysis: calc C 47.68\%, H 10.11\%, N 29.27\%; found C 47.54\%, H 9.96\%, N 29.47\%. IR (neat): \( \tilde{\nu} \) (cm\(^{-1}\)) = 3006 (w), 2847 (m), 2810 (m), 2778 (s), 1766 (vs.), 1496 (s), 1472 (m), 1453 (m), 1440 (m), 1241 (m), 1358 (vs.), 1281 (m), 1251 (m), 1235 (m), 1173 (m), 1128 (s), 1052 (m), 971 (s), 949 (vs.), 917 (m), 860 (vs.), 796 (m), 748 (m), 685 (s), 652 (s), 629 (vs.), 568 (m), 527 (s), 452 (s). XRD: for single crystal X-ray structure determination suitable single crystals were obtained from n-hexane at \(-25^\circ\)C.

**Attempted synthesis of sym-(dmpa)\(^2\)(dmpa)\(_4\)CDP (2)**

A mixture of 2-2HBF\(_4\) (136 mg, 174 \( \mu \)mol, 1.0 eq.) and freshly ground sodium amide (75 mg, 1.99 mmol, 11 eq.) was suspended in THF (15 mL) and stirred for 72 h at 60 \( ^\circ\)C. The solid was removed by filtration over Celite and the filtrate evaporated to dryness. The residue was dissolved in n-pentane (20 mL), cleared via syringe filtration, the solvent removed and the residue dried in high vacuum to give 7 as pale yellow high viscous oil.

\[^{1}\]H NMR (300.3 MHz, CD\(_2\)CN): \( \delta \) (ppm) = 2.99 (d, \( ^3J_{F\text{H}} = 9 \text{ Hz}, 12\text{H}, \text{H} \)), 2.88 (d, \( ^3J_{F\text{H}} = 10 \text{ Hz}, \text{H} \)), 2.83 (d, \( ^3J_{F\text{H}} = 11 \text{ Hz}, \text{H} \)), 2.32 (d, \( ^3J_{F\text{H}} = 18\text{H}, \text{H} \)). \(^{13}\)C\(^{1}\)H NMR (75.5 MHz, CD\(_2\)CN): \( \delta \) (ppm) = 38.5 (dd, \( ^3J_{PC} = 4 \text{ Hz}, \text{PC} \)), 38.4 (dd, \( ^3J_{PC} = 16 \text{ Hz}, \text{C} \)), 38.1 (dd, \( ^3J_{PC} = 4 \text{ Hz}, \text{C} \)), 37.1 (d, \( ^3J_{PC} = 4 \text{ Hz}, \text{C} \)), 13.0 (dd, \( ^3J_{PC} = 3 \text{ Hz}, \text{C} \)), 10.9 (s, \( ^1J_{PC} = 1 \text{ Hz}, \text{C} \)), 10.8 (s, \( ^1J_{PC} = 1 \text{ Hz}, \text{C} \)).
BF4]+. LIFDI(+) HRMS: (m/z \[M}\] + calcd 495.34939, found 495.35146.

A mixture of 2-HBF4 (600 mg, 769 \mu mol, 1.00 eq.) and finely ground sodium amide (321 mg, 8.23 mmol, 10.7 eq.) was suspended in THF-d8 (0.5 mL) and used for analytics.

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\text{[C_2H_4F_6N_4O_3P_8S_2]} (759.75 \text{ g mol}^{-1}) \text{H NMR (300.3 MHz, THF-d8):} \delta (\text{ppm}) = 2.90 (24\text{H, H1}), 1.26–2.64 (24\text{H, H2}), 0.55 (t, \text{J}_{\text{PH}} = 4\text{ Hz}, 1\text{H, CH}). \text{13C}{\text{^1H}} \text{NMR (75.5 MHz, THF-d8):} \delta (\text{ppm}) = 161.1 (2\text{C, CN}), 121.1 (q, J_{\text{FC}} = 322\text{ Hz, CF}), 40.3 (8\text{C, C1}), 37.7 (dd, 2\text{C, J}_{\text{FC}} = 2\text{ Hz, C2}), 9.3 (t, J_{\text{FC}} = 185\text{ Hz, CH}). \text{31P}{\text{^1H}} \text{NMR (121.5 MHz, C6D6):} \delta (\text{ppm}) = 37.1. \text{LIFDI(+) MS (THF):} m/z = 353.4 \text{([M - TFSI]^-). LIFDI(+) HRMS:} m/z = 353.4. \text{calcd 353.4, found 353.4.}

Notes and references


Chemical Science


51 (a) S. Ulrich, CCDC 1903847: Experimental Crystal Structure Determination; (b) S. Ulrich, CCDC 1903849: Experimental Crystal Structure Determination.


