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Isolation of singlet carbene derived 2-phospha-1,3-butadienes and their sequential one-electron oxidation to radical cations and dications†

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A synthetic strategy for the 2-phospha-1,3-butadiene derivatives [{(IPr)C(Ph)}P(cAAC^{Me})] (**3a**) and [{(IPr)C(Ph)} P(cAAC^{Cy})] (**3b**) (IPr = C{(NDipp)CH}$_2$, Dipp = 2,6-iPr$_2$C$_6$H$_3$; cAAC^{Me} = C{(NDipp)CMe$_2$CH$_2$CMe$_2$}; cAAC^{Cy} = C{(NDipp)CMe$_2$CH$_2$C(Cy)}, Cy = cyclohexyl) containing a C=C–P=C framework has been established. Compounds **3a** and **3b** have a remarkably small HOMO–LUMO energy gap (**3a**: 5.09; **3b**: 5.05 eV) with a very high-lying HOMO (−4.95 eV for each). Consequently, **3a** and **3b** readily undergo one-electron oxidation with the mild oxidizing agent GaCl$_3$ to afford radical cations [{(IPr)C(Ph)}P(cAAC^R)]GaCl$_4$ (R = Me **4a**, Cy **4b**) as crystalline solids. The main UV-vis absorption band for **4a** and **4b** is red-shifted with respect to that of **3a** and **3b**, which is associated with the SOMO related transitions. The EPR spectra of compounds **4a** and **4b** each exhibit a doublet due to coupling of the unpaired electron with the ^{31}P nucleus. Further one-electron removal from the radical cations **4a** and **4b** is also feasible with GaCl$_3$, affording the dications [{(IPr) C(Ph)}P(cAAC^R)](GaCl$_4$)$_2$ (R = Me **5a**, Cy **5b**) as yellow crystals. The molecular structures of compounds **3–5** have been determined by X-ray diffraction and analyzed by DFT calculations.

Introduction

Organic π-conjugated molecules are currently of great academic and significant technological interest due to their intriguing optoelectronic properties.[1] In this context, π-conjugated systems featuring heavier main-group elements[2] and systems exhibiting a considerable open-shell (radical-type) character[3] are particularly attractive as they display promising optical, electronic, and magnetic properties. Among heavier main-group elements, the choice to incorporate phosphorus into π-conjugated systems has been primarily driven by its semblance to the isoelectronic "CR" (R = H, alkyl or aryl group) unit, which is known as a diagonal relationship.[4] Moreover, while the calculated P=C π-bond strength (43 kcal mol^{-1}) is lower than the C=C π-bond of ethene (65 kcal mol^{-1})[5] the conjugative properties of both P=C and C=C bonds are comparable.[6]

Among stable main-group radicals,[7] various neutral,[8] cationic,[9] as well as anionic[10] phosphorus radicals have been also isolated

and structurally characterized, however, phosphorus radicals based on a π-conjugated framework remain scarce. 1,3-Butadiene **I** is the simplest molecule with conjugated π-bonds (Fig. 1) that has also been an important structural motif in phosphorus chemistry.[11] Indeed, unsubstituted as well as alkyl substituted phosphorus containing 1,3-butadiene derivatives were already reported by Appel,[11b] Regitz,[11f] and Denis,[11g] however, these compounds are unlikely to afford stable radical compounds on oxidation or reduction. In 2008, Robinson *et al.* reported a diphosphorus compound **II** containing a weak π-acceptor N-heterocyclic carbene (NHC).[12] Structural and theoretical data suggest that **II** should be better described as a base-stabilized diphosphinidene with C$_{(NHC)}$– P and P–P single bonds. Compound **III**, reported by Bertrand's group in 2010, features a strong π-acceptor cyclic alkyl amino carbene (cAAC^R) and exhibits short C–P bond lengths, thus it may be regarded as a genuine 2,3-diphospha-1,3-butadiene.[13] The same group also reported the 2-phospha-3-azabutadiene **IV** by an elegant choice of imine and cAAC precursors.[9c] Remarkably, these electron-rich species readily undergo one-electron oxidation to afford the corresponding radical cations (**II**)$^{•+}$, (**III**)$^{•+}$, and (**IV**)$^{•+}$.[9b,9c] We recently reported NHC-derived divinyldiphosphenes **V**[14] and isolated the corresponding radicals cations (**V**)$^{•+}$ by one-electron oxidation of **V**.[15] These and other early results[16] prompted us to reason that stable 2-phospha-1,3-butadienes **VI** as well as the corresponding radical cations (**VI**)$^{•+}$ and dications (**VI**)$^{++}$ should be synthetically accessible by a rational choice of substrates and reaction conditions.

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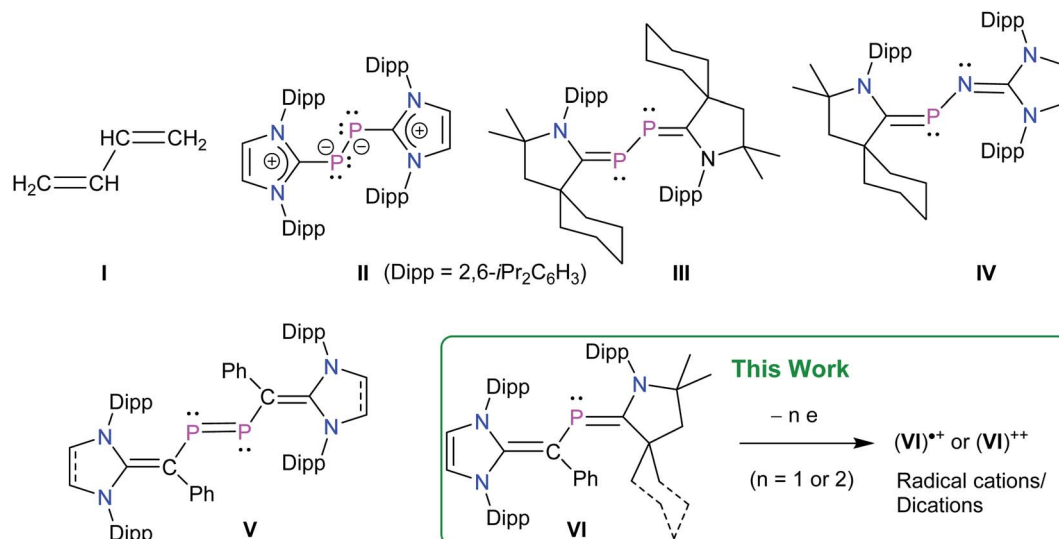
Fig. 1 1,3-Butadiene **I**. Selected examples of phosphorus containing derivatives **II–IV** and divinyldiphosphene **V** with singlet carbene frameworks.

Herein, we report the synthesis of 2-phospha-1,3-butadienes [{(IPr)C(Ph)}P(cAAC^{Me})] (**3a**) and [{(IPr)C(Ph)}P(cAAC^{Cy})] (**3b**) based on singlet carbene frameworks (IPr = C{(NDipp)CH}$_2$, Dipp = 2,6-iPr$_2$C$_6$H$_3$; cAAC^{Me} = C{(NDipp)CMe$_2$CH$_2$CMe$_2$}; cAAC^{Cy} = C{(NDipp)CMe$_2$CH$_2$C(Cy)}, Cy = cyclohexyl) as crystalline solids. Sequential one-electron oxidation of **3a** and **3b** leads to the formation of corresponding radical cations [{(IPr)C(Ph)}P(cAAC^{Me})](GaCl$_4$) (**4a**), [{(IPr)C(Ph)}P(cAAC^{Cy})](GaCl$_4$)

(**4b**) and dications [{(IPr)C(Ph)}P(cAAC^{Me})](GaCl$_4$)$_2$ (**5a**), [{(IPr)C(Ph)}P(cAAC^{Cy})] (GaCl$_4$)$_2$ (**5b**) as crystalline solids.

Results and discussion

For the synthesis of desired 2-phospha-1,3-butadienes, N-heterocyclic vinyl (NHV)-substituted dichlorophosphine {(IPr)C(Ph)}PCl$_2$ (**1**)[14] and strong π-acceptor cAAC^R (ref. 17) were chosen as the appropriate precursors (Scheme 1).[18] Treatment of
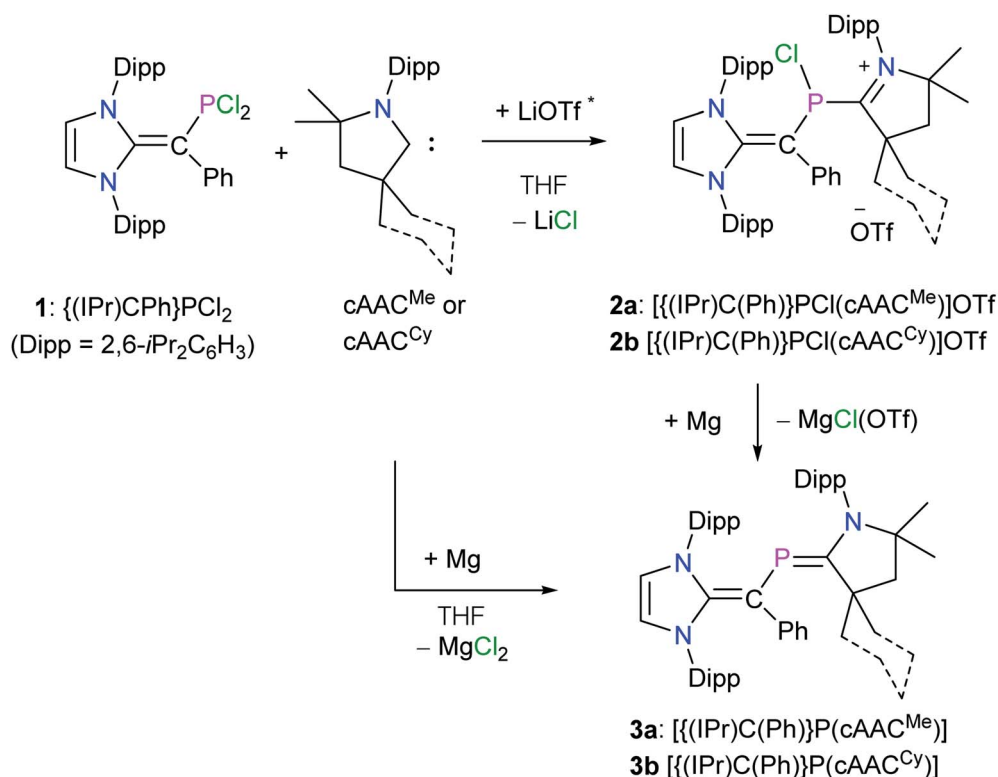
Scheme 1 Synthesis of 2-phospha-1,3-butadiene derivatives **3a** and **3b**. * cAACs were prepared by the deprotonation of their triflate salts with LDA and the side-product LiOTf was not separated.

a colorless THF solution of **1** with one equivalent of cAAC^{Me} or cAAC^{Cy} immediately resulted in the formation of dark blue solutions (Scheme 1). After workup, the ionic compounds [{(IPr)C(Ph)}P(Cl)(cAAC^R)](OTf) (R = Me **2a**, Cy **2b**) were isolated as violet crystalline solids. Compounds **2a** and **2b** are highly air sensitive solids and have been characterized by elemental analysis and NMR spectroscopy. The solid state molecular structure of a typical compound **2a** (Fig. S31†) was determined by X-ray diffraction. Reduction of **2a** and **2b** with magnesium turnings afforded the target compound **3a** and **3b**, respectively, as orange solids. Interestingly, **3a** and **3b** are also accessible in a one-pot reaction of **1** and cAAC^R with magnesium. Both **3a** and **3b** are soluble in common organic solvents (*n*-hexane, Et$_2$O, benzene, toluene, THF) and are stable under an inert gas atmosphere.

The ^{1}H NMR spectra of compounds **2a** and **2b** as well as **3a** and **3b** show expected resonances for the NHV and cAAC^R moieties. The ^{13}C{^{1}H} NMR spectrum of **2a** and **2b** as well as **3a** and **3b** each is consistent with the ^{1}H NMR resonances and exhibits expected doublets for the phosphorus bound carbon atoms (see the ESI†). The ^{31}P{^{1}H} NMR spectrum of **2a** (+100.9 ppm) and **2b** (+102.9 ppm) each shows a singlet, which is high-field shifted with respect to that of the **1** (+167 ppm). This is most likely due to the coordination of electron-rich cAAC^R to the phosphorus atoms in **2a** and **2b**. The ^{31}P{^{1}H} NMR signal for **3a** (+102.5 ppm) and **3b** (+108.6 ppm), respectively, appears at a higher field compared to that of **IV** (+134.0 ppm),[9c] which is expected because of the electronegativity difference between carbon and nitrogen.

The solid-state molecular structures of **3a** and **3b** (Fig. 2) adopt a *trans*-bent geometry along the C$_{NHV}$–P bond with the C2–P1 bond length of 1.818(1) and 1.820(1) Å, respectively. The C2–P1 bond length is larger compared to that in **1** (1.728(2) Å)[14] and **2a** (1.751(2) Å), but it is comparable with those of the diphosphenes **V** (1.785 to 1.797 Å).[15] The P1–C3

(**3a**: 1.735(1); **3b** 1.735(1) Å) and C1–C2 (**3a**: 1.386(1); **3b**: 1.384(2) Å) bonds are shorter compared to the same bonds in **2a** (1.860(2) and 1.437(2) Å, respectively). The P1–C3 bond lengths of **3a** and **3b** are nonetheless in line with those of the P=C double bonds in **III** (1.719(7) Å)[13] and **IV** (1.719(2) Å).[9c] The C1=C2, P1=C3, and P1–C2 bond lengths of **3a** and **3b** are in line with those of the literature known neutral 2-phospha-1,3-butadiene (Me$_3$SiO)*t*BuC=P–C(SiMe$_3$)=C(OSiiPr$_3$)*t*Bu (C=C: 1.356(4), P=C: 1.702(3), P–C: 1.846(3) Å).[19]

To shed light on the electronic structure of **3a** and **3b**, we performed DFT calculations at the M06-2X/def2-TZVPP//def2-SVP level of theory. The HOMO of **3a** and **3b** is a π-type orbital mainly located at the C$_{cAAC}$=P and C$_{IPr}$=C$_{Ph}$ bonds (Fig. 3). Remarkably, the HOMO energy of **3a** and **3b** (−4.95 eV) is quite high and comparable with those of related divinyldi-phosphenes **V** (−4.71 to −5.25 eV),[15] indicating the possibility of facile oxidation.
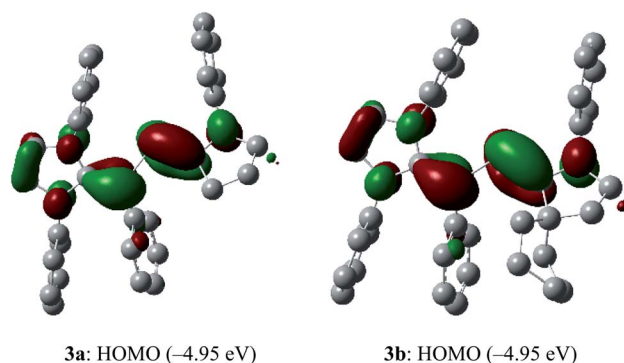

Fig. 3 HOMOs (highest occupied molecular orbitals, isovalue 0.04) of **3a** and **3b** calculated at M06-2X/def2-TZVPP//def2-SVP level of theory. Hydrogen atoms, methyl as well as iso-propyl groups were omitted for clarity.
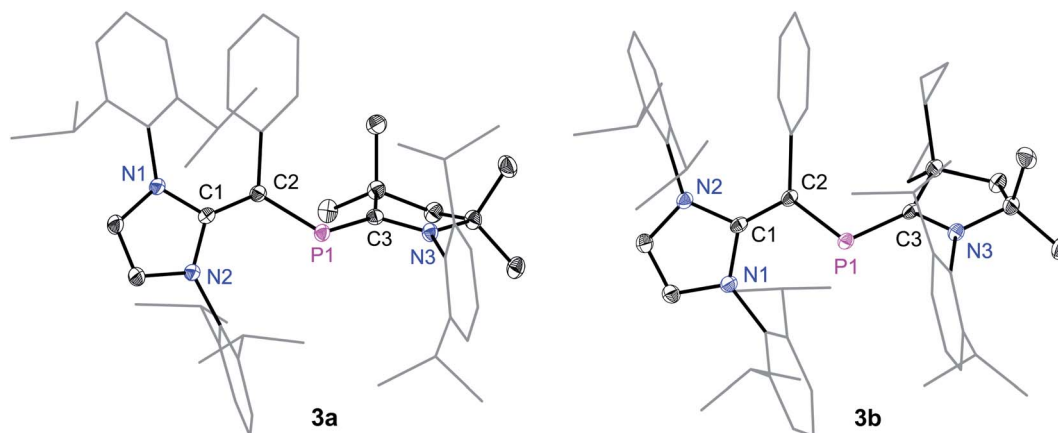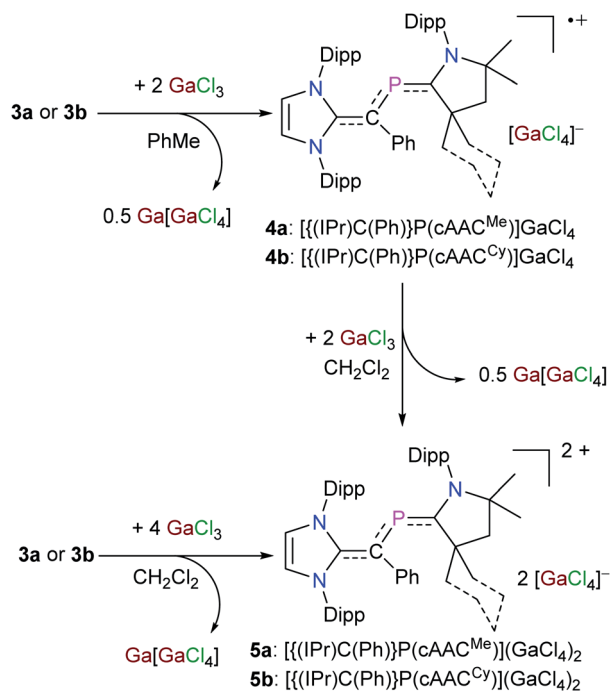

Fig. 2 Solid-state molecular structures of 2-phospha-1,3-butadienes **3a** and **3b**. Hydrogen atoms have been omitted for clarity. Selected bond lengths and bond angles are given in Table 1.

These preliminarily theoretical findings encouraged us to analyze the redox properties of **3a** and **3b** by electrochemical studies to gain an initial insight into the viability and stability of derived radicals. The cyclic voltammograms (CVs) of **3a** (Fig. S19†) and **3b** (Fig. S20†) show two main redox events in the −2.0 to 1.5 V region. The first reversible wave at $E_{1/2} = -1.06$ V

for **3a** and −1.08 V for **3b** may be assigned to the corresponding radical cation, whereas the second quasi-reversible wave at $E_{1/2} = -0.28$ V for **3a** and −0.24 V for **3b** may correspond to the dicationic species. Indeed, treatment of an orange toluene solution of **3a** or **3b** with GaCl₃ immediately led to the precipitation of a violet solid. After workup, the radical cation salts [{(IPr)C(Ph)}P(cAAC^Me)][GaCl₄] (**4a**) and [{(IPr)C(Ph)}P(cAAC^Cy)][GaCl₄] (**4b**) were isolated as violet crystals (Scheme 2). GaCl₃ acts as oxidizing agent and two molecules of GaCl₃ are required for one-electron oxidation.[15] Consistent with the CVs (Fig. S19 and S20†), the dication salts [{(IPr)C(Ph)}P(cAAC^Me)][GaCl₄]₂ (**5a**) and [{(IPr)C(Ph)}P(cAAC^Cy)][GaCl₄]₂ (**5b**) are selectively accessible on one-electron oxidation of **4a** and **4b** with GaCl₃ (Scheme 2). Alternatively, **5a** and **5b** can also be prepared directly from **3a** and **3b** with four equivalent of GaCl₃, respectively (Scheme 2). Compounds **4a**, **4b** and **5a**, **5b** are stable both in solutions as well as in the solid-state under an inert gas atmosphere, but decompose rapidly when exposed to air. The radicals **4a** and **4b** are NMR silent, while dicationic salts **5a** and **5b** are diamagnetic and exhibit well resolved ¹H and ¹³C{¹H} NMR signals for the NHV and cAAC^R units. The ³¹P{¹H} NMR signal for the dication salts **5a** (+244 ppm) and **5b** (+236 ppm) is downfield-shifted with respect to that of the 2-phospha-1,3-butadienes **3a** (+102 ppm) and **3b** (+108 ppm) but it is in the range expected for phosphaalkenes (200–300 ppm).[20]

Suitable single crystals for X-ray diffraction were obtained by a slow diffusion of *n*-hexane into a saturated THF or CH₂Cl₂ solution of each of radical cations **4a** and **4b** and dicationic salts **5a** and **5b**. The solid-state molecular structure of **4a** and **4b** (Fig. 4) as well as **5a** and **5b** (Fig. 5) each adopts a *trans*-bent geometry along the P–C_NHV bond and reveals an interesting bond length alteration trend with

Scheme 2 Sequential one-electron oxidation of 2-phospha-1,3-butadienes **3a** and **3b** with GaCl₃ to the corresponding radical cations **4a** and **4b** and dications **5a** and **5b**.
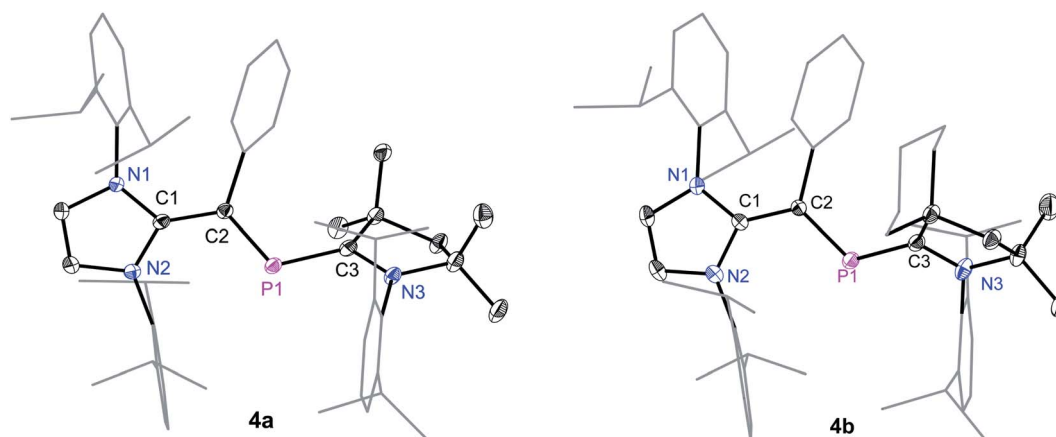

Fig. 4 Solid–state molecular structures of radical cation **4a** and **4b**. Hydrogen atoms, solvent molecules in **4b**, and the counter anions GaCl₄ have been omitted for clarity. Selected bond lengths and bond angles are given in Table 1.

Fig. 5 Solid-state molecular structures of **5a** and **5b**. Hydrogen atoms, solvent molecules in **5b**, and the counter anions $GaCl_4$ have been omitted for clarity. Selected bond lengths and bond angles are given in Table 1.
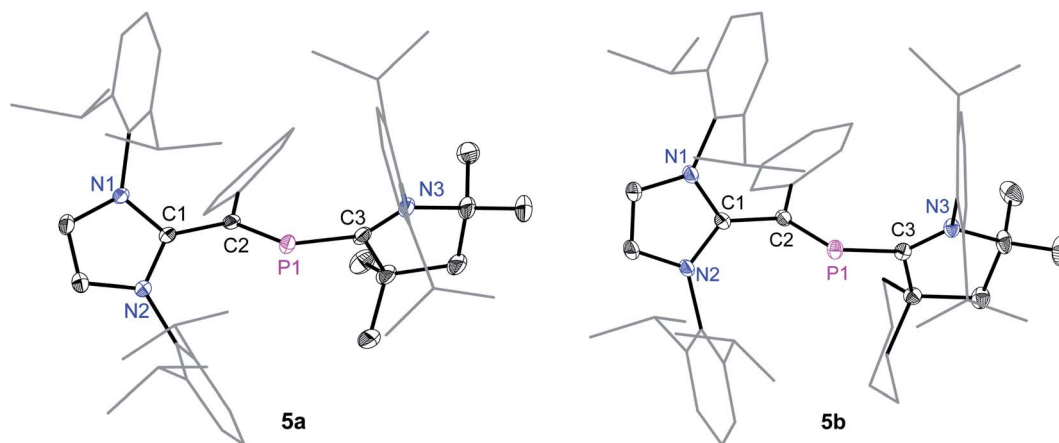
respect to the precursors **3a** and **3b** (Table 1). The C2–P1 bond length of **4a** (1.758(2) Å) and **4b** (1.761(2) Å) is smaller compared to that of the respective 2-phospha-1,3-butadienes **3a** (1.818(1) Å) and **3b** (1.820(1) Å) but longer with respect to that of the dications **5a** (1.692(3) Å) and **5b** (1.692(2) Å) Å. The P1–C3 bond, however, steadily stretches on going from neutral to radical cations and to dications: **3a**: 1.735(1), **3b**: 1.735(1); **4a**: 1.785(2), **4b**: 1.790(2); **5a**: 1.865(2), **5b**: 1.853(2) Å. A similar trend in the C1–C2 bond length stretching can also be seen in **3a**: 1.386(1), **3b**: 1.384(2); **4a**: 1.432(2), **4b**: 1.433(2); and **5a**: 1.479(3), **5b**: 1.478(2) Å. In conclusion, the formally C1═C2 and P1═C3 double bonds in **3a** and **3b** become comparable to C_{sp^2}–C_{sp^2} (ca. 1.47 Å) and P–C_{sp^2} (1.85 Å) single bond lengths[21] in **5a** and **5b**, whereas the C2–P1 single bond in **3a** and **3b** adopts double bond lengths (1.60–1.70 Å) in **5a** and **5b** as expected for phosphaalkenes.[22]

Table 1 Selected bond lengths (Å) and angles (°) of 2-phospha-1,3-butadienes (**3a** and **3b**), radical cations (**4a** and **4b**) and dication salts (**5a** and **5b**)

	3a	3b	4a	4b	5a	5b
C1–C2	1.386(1)	1.384(2)	1.432(2)	1.433(2)	1.479(3)	1.478(2)
C2–P1	1.818(1)	1.820(1)	1.758(2)	1.761(2)	1.692(2)	1.692(2)
P1–C3	1.735(1)	1.735(1)	1.785(2)	1.790(2)	1.865(2)	1.853(2)
C1–N1	1.401(1)	1.403(1)	1.375(2)	1.375(2)	1.351(3)	1.344(2)
C1–N2	1.408(1)	1.412(1)	1.382(2)	1.385(2)	1.355(3)	1.356(2)
C3–N3	1.387(1)	1.388(1)	1.349(2)	1.349(2)	1.295(3)	1.295(3)
C1–C2–P1	117.7(1)	118.0(1)	117.3(1)	117.4(1)	114.9(1)	115.3(1)
C2–P1–C3	109.7(1)	110.9(1)	108.8(1)	110.3(1)	106.1(1)	105.2(1)
P1–C3–N3	117.6(1)	116.4(1)	116.0(1)	115.1(1)	118.1(2)	121.5(1)
N1–C1–N2	103.0(1)	103.3(1)	104.8(1)	104.9(1)	107.2(2)	107.3(2)

DFT calculated geometries of **3–5** are found to be fully in agreement with their solid-state molecular structures determined by X-ray diffraction (Table S5†). An increasing value of the WBIs (Wiberg Bond Indices) for the C_{NHV}–P bond of **3a** (0.95), **3b** (0.95), **4a** (1.22), **4b** (1.22), **5a** (1.63), and **5b** (1.63) is consistent with the experimental C2–P1 bond lengths (Table 1). Similarly, the WBIs for the P–C_{cAAC} (**3a**: 1.57, **3b**: 1.56; **4a**: 1.19, **4b**: 1.18; **5a**: 0.92, **5b**: 0.91) as well as C–C_{Ph} (**3a**: 1.49, **3b**: 1.49; **4a**: 1.20, **4b**: 1.20; **5a**: 1.06, **5b**: 1.06) bonds also exhibit the expected trend. The NPA (Natural Population Analysis) atomic partial charges (Table S5†) calculated using the NBO (Natural Bond Orbital) method indicate that the phosphorous atom in **3a** (0.49e), **3b** (0.49e), **4a** (0.65e), **4b** (0.65e), **5a** (0.85e), and **5b** (0.85e) carries a positive charge. The former IPr carbene carbon atom (C1) also bears a positive charge (**3a** 0.45e, **3b** 0.44e, **4a** 0.47e, **4b** 0.47e, **5a** 0.43e and **5b** 0.43e), which is larger compared to that of the cAAC carbene carbon atom C3 (**3a** −0.08e, **3b** −0.08e, **4a** 0.04e, **4b** 0.04e, **5a** 0.23e and **5b** 0.24e). This is most likely because of the greater π-acceptor property of cAACs compared to the IPr.

The SOMO (singly occupied molecular orbital) of **4a** and **4b** (Fig. 6) is the π-orbitals of the C_{IPr}═C_{vinyl} and P═C_{cAAC} bonds with a small contribution from the nitrogen atoms of the pyrrolidine and imidazole ring, whereas the LUMO (lowest unoccupied molecular orbital) is the π*-orbitals of the P atom along with the π*-orbitals of C_{IPr}═C_{vinyl} and C_{cAAC}–N bonds of NHV and cAAC, respectively. The localized Mulliken atomic spin density (Fig. 7) and the plot of the SOMO (Fig. 6) for **4a** and **4b** reveal that the unpaired electron is mainly delocalized over the CPCN moiety with an almost equal spin density distribution on the phosphorus (**4a**: 17%, **4b**: 18%), carbene C_{cAAC} (**4a**: 19%, **4b**: 20%), and vinylic C_{Ph} (**4a**: 26%, **4b**: 25%) atoms. The spin-density at the imidazole-ring nitrogen atoms (6% at each of N) of **4a** and **4b** is rather small compared to that at the pyrrolidine nitrogen atom (17%).

4a: SOMO (−8.08/−9.66 eV)   **4a**: LUMO (−3.01/−4.59 eV)

4b: SOMO (−8.04/−9.63 eV)   **4b**: LUMO (−3.05/−4.55 eV)
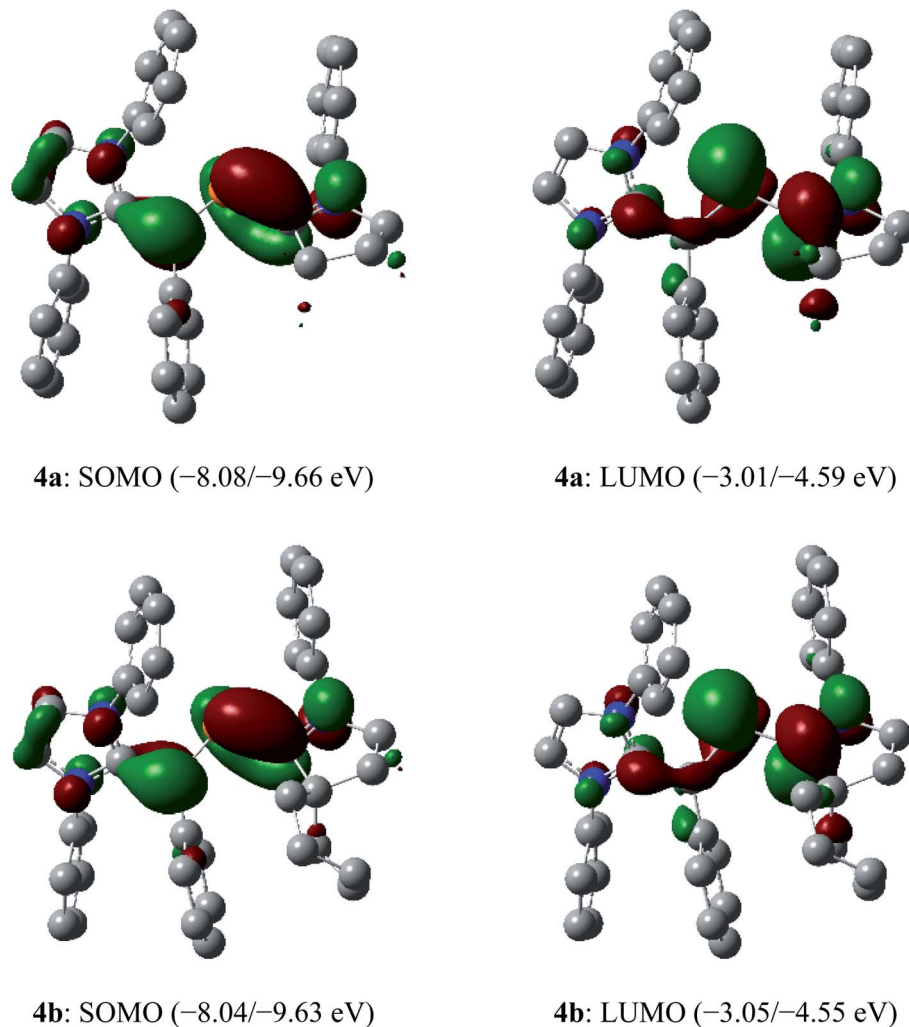
Fig. 6 Selected molecular orbitals (isovalue 0.04) of the radical cations **4a** and **4b** (α/β spin orbital energy) calculated at the M06-2X/def2-TZVPP//def2-SVP level of theory. Hydrogen atoms, methyl groups as well as iso-propyl groups were omitted for clarity.
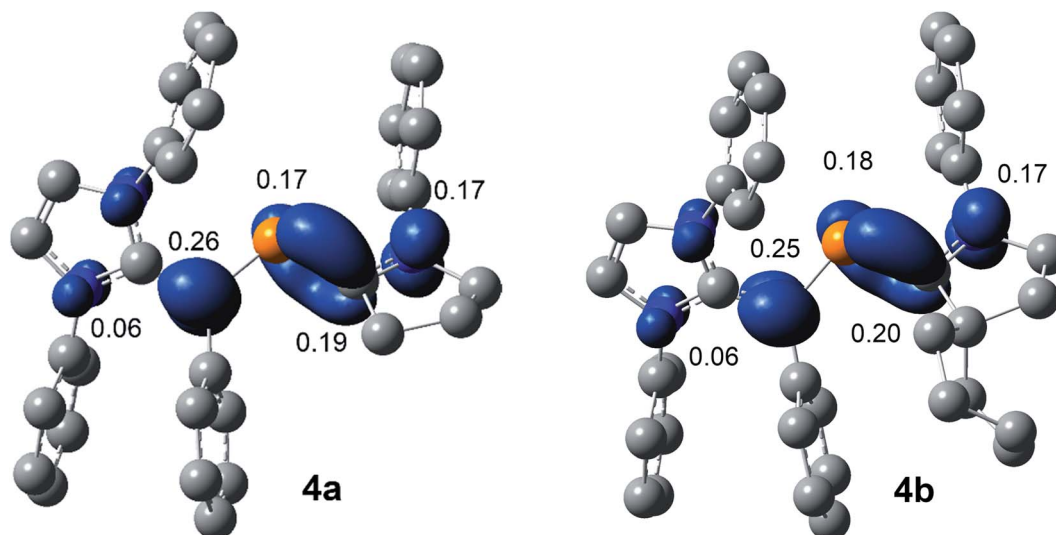
Fig. 7 Calculated Mulliken spin densities (isovalue 0.004 a.u.) of radical cations **4a** and **4b** at M06-2X/def2-TZVPP//def2-SVP level of theory.
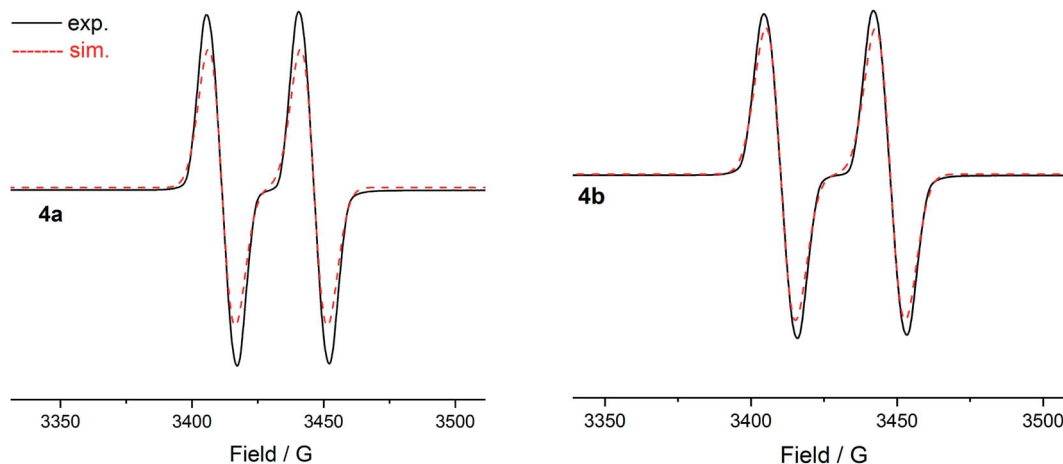
Fig. 8 X-band EPR spectra of radical cations **4a** and **4b** in THF at 298 K.

The room temperature X-band EPR spectra of **4a** ($g = 2.0064$) and **4b** ($g = 2.0064$) in THF exhibit a doublet (Fig. 8) owing to the coupling of the unpaired electron with the phosphorous nucleus ($A_{iso}(^{31}P) = 35$ G **4a**; 38 G **4b**). The magnitude of the hyperfine coupling constant (hfc) of **4a** and **4b** is comparable to those observed for radical cations $(\mathbf{II})^{\bullet+}$ ($A_{iso}(^{31}P) = 44$ G)[9b] $(\mathbf{III})^{\bullet+}$ ($A_{iso}(^{31}P) = 42$ G),[9b] and $(\mathbf{IV})^{\bullet+}$ ($A_{iso}(^{31}P) = 44$ G),[9c] but that is larger than those of $(\mathbf{V})^{\bullet+}$ ($A_{iso}(^{31}P) = 12$–20 G) featuring a longer π-conjugated system (Fig. 1).[15] This is, however, considerably smaller than that of the phosphinyl radical cation $[(cAAC^{Cy})P(R)]^{\bullet+}$ ($A_{iso}(^{31}P) = 99$ G) (R = 2,2,6,6-tetramethylpiperidino)[9a] as well as those observed for phosphinyl radicals $R_2P^{\bullet}$ ($A_{iso}(^{31}P) = 92$–96 G) (R = HC(SiMe$_3$)$_2$ or N(SiMe$_3$)$_2$),[8a,23] for which, particularly for the latter, the unpaired electron resides predominantly in a 3p(P) valence orbital. The value of coupling constants is in good agreement with the computed values (Table S12†). These hfcs corroborate with the delocalization of the spin-density along the CPCN moiety of **4a** and **4b** (Fig. 7). The measured EPR spectra of **4a** and **4b** were simulated by employing the g values, the hyperfine coupling for the phosphorus atom, and two linewidth parameters (Table S13†). The EPR spectra of **4a** (Fig. S29†) and **4b** (Fig. S30†) measured in a frozen THF solution at 80 K show an anisotropic pattern. The g-factors (**4a**: $g_{\parallel} = 2.0062$, $g_{\perp} = 2.0082$; **4b**: $g_{\parallel} = 2.0069$, $g_{\perp} = 2.0079$) and hfc tensors (**4a**: $A_z(^{31}P) = 149$ MHz, $A_x(^{31}P) = -83$ MHz, $A_y(^{31}P) = 39$ MHz; **4b**: $A_z(^{31}P) = 179$ MHz, $A_x(^{31}P) = -71$ MHz, $A_y(^{31}P) = 49$ MHz) were determined. The analysis of hfc tensors reveals the major contribution of phosphorus 3p (**4a**: 11.8%; **4b**: 12.8%) orbital to the SOMO, whereas the contribution of the 3s (**4a**: 0.45%; **4b**: 0.46%) orbital is small.

The UV-vis spectra of both 2-phospha-1,3-butadienes **3a** (277, 331, 427 nm) and **3b** (271, 322, 430 nm) exhibit three main absorptions (Fig. 9) which, based on TD-DFT
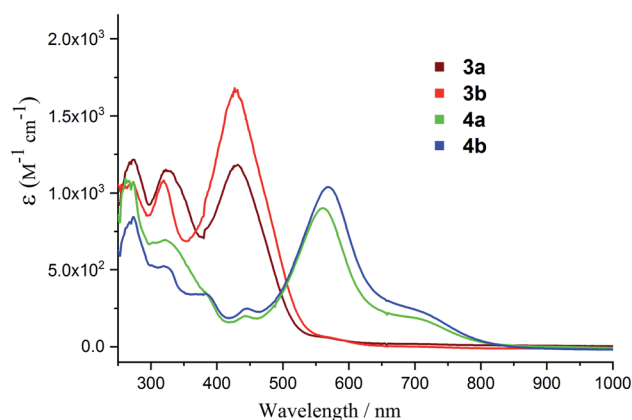

Fig. 9 UV-vis spectra of **3a** and **3b** and their radical cations **4a** and **4b** in THF.

calculations at TD-PCM(thf)/M06-2X/def2-SVP level of theory, comprise dominant contributions of the H−1 → L, H → L+2, and H → L transitions, respectively (Fig. S32 and S33, Tables S6 and S7†). The UV-vis spectra of the radical cations **4a** and **4b**, respectively show a broad absorption at 563 and 571 nm along with a shoulder at ca. 700 nm (Fig. 9) that corresponds to the SOMO-related transitions S → L and S−1 → S (Tables S8 and S9†).

In the dications **5a** and **5b**, the HOMO-2 is a π-type orbital with contributions from the aryl groups and C_{vinyl}–P bond (Fig. 10). The LUMO of **5a** and **5b** is the π-orbital of the C_{IPr}=C_{vinyl} and P=C_{cAAC} bonds with a small contribution from the nitrogen atoms of pyrrolidine and imidazole rings. Upon removal of one and two electrons from the π-type orbital of **3a** and **3b**, the HOMO of **3a** and **3b** (Fig. 3) becomes SOMO in the radical cations **4a** and **4b** (Fig. 6) and LUMO in the dications **5a** and **5b** (Fig. 10).
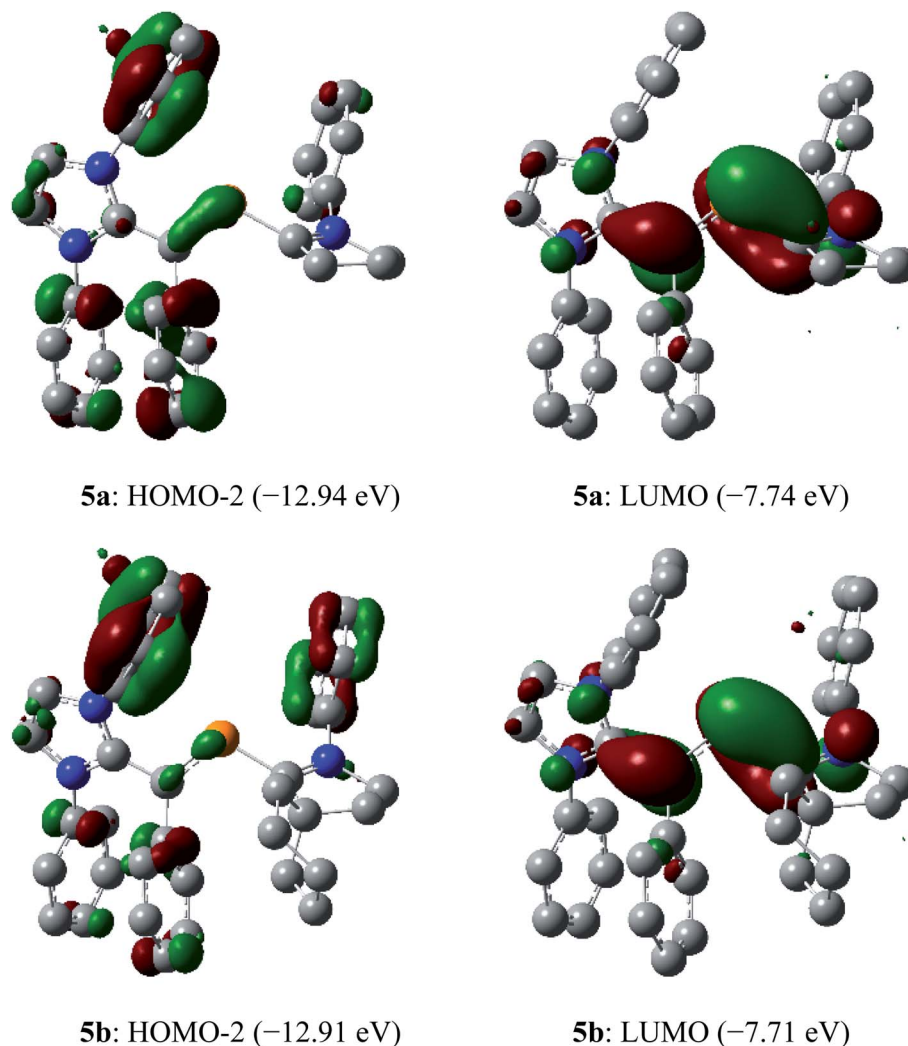
5a: HOMO-2 (−12.94 eV)

5a: LUMO (−7.74 eV)

5b: HOMO-2 (−12.91 eV)

5b: LUMO (−7.71 eV)

Fig. 10 Selected molecular orbitals (isovalue 0.04) of the dications **5a** and **5b** calculated at M06-2X/def2-TZVPP//def2-SVP level of theory. Hydrogen atoms, methyl groups as well as iso-propyl groups were omitted for clarity.

Conclusions

In conclusion, we have isolated the crystalline 2-phospha-1,3-butadiene derivatives **3a** and **3b** by a rational choice of combining a weak π-acceptor (IPr) and a strong π-acceptor (cAAC^R) singlet carbene scaffolds. Sequential one-electron oxidation of **3a** and **3b** affords the radical cations **4a** and **4b** and the dications **5a** and **5b**. The isolation of **4a**, **4b** and **5a**, **5b** as crystalline solids is consistent with the redox properties of **3a** and **3b** analyzed by electrochemical studies. Molecular structures of all compounds in the solid-state were established by single crystal X-ray diffraction. Computational and EPR spectroscopic data indicate that the unpaired electron in **4a** and **4b** is delocalized over the CPCN π-conjugated framework. The study emphasizes the advantage of merging singlet carbenes with dissimilar donor–acceptor properties in accessing stable open-shell π-conjugated systems. As a variety of stable singlet carbenes with adaptable properties are readily accessible, it is very likely that many π-conjugated systems with other heavier main-group elements, which are hitherto believed to be synthetically challenging targets, may be isolated.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

References

1 (a) C. Wang, H. Dong, W. Hu, Y. Liu and D. Zhu, *Chem. Rev.*, 2012, **112**, 2208–2267; (b) W. Wu, Y. Liu and D. Zhu, *Chem.*

Soc. Rev., 2010, **39**, 1489–1502; (*c*) Y. Wang, L. Sun, C. Wang, F. Yang, X. Ren, X. Zhang, H. Dong and W. Hu, *Chem. Soc. Rev.*, 2019, **48**, 1492–1530; (*d*) J. Yang, D. Yan and T. S. Jones, *Chem. Rev.*, 2015, **115**, 5570–5603; (*e*) C. Lee, S. Lee, G.-U. Kim, W. Lee and B. J. Kim, *Chem. Rev.*, 2019, **119**, 8028–8086.

2 (*a*) F. Vidal and F. Jäkle, *Angew. Chem., Int. Ed.*, 2019, **58**, 5846–5870; (*b*) F. Pop, N. Zigon and N. Avarvari, *Chem. Rev.*, 2019, **119**, 8435–8478; (*c*) M. Stolar and T. Baumgartner, *Chem. Commun.*, 2018, **54**, 3311–3322; (*d*) M. Hirai, N. Tanaka, M. Sakai and S. Yamaguchi, *Chem. Rev.*, 2019, **119**, 8291–8331; (*e*) K. Dhbaibi, L. Favereau and J. Crassous, *Chem. Rev.*, 2019, **119**, 8846–8953; (*f*) T. Baumgartner and R. Réau, *Chem. Rev.*, 2006, **106**, 4681–4727; (*g*) S. M. Parke, M. P. Boone and E. Rivard, *Chem. Commun.*, 2016, **52**, 9485–9505; (*h*) A. M. Priegert, B. W. Rawe, S. C. Serin and D. P. Gates, *Chem. Soc. Rev.*, 2016, **45**, 922–953.

3 (*a*) Z. Zeng, X. Shi, C. Chi, J. T. Lopez Navarrete, J. Casado and J. Wu, *Chem. Soc. Rev.*, 2015, **44**, 6578–6596; (*b*) T. Y. Gopalakrishna, W. Zeng, X. Lu and J. Wu, *Chem. Commun.*, 2018, **54**, 2186–2199; (*c*) M. Nakano, K. Fukuda, S. Ito, H. Matsui, T. Nagami, S. Takamuku, Y. Kitagawa and B. Champagne, *J. Phys. Chem. A*, 2017, **121**, 861–873; (*d*) M. Nakano, *Chem. Rec.*, 2017, **17**, 27–62; (*e*) Y. Yang, E. R. Davidson and W. Yang, *Proc. Natl. Acad. Sci. U. S. A.*, 2016, **113**, E5098–E5107; (*f*) Y. Su, X. Wang, X. Zheng, Z. Zhang, Y. Song, Y. Sui, Y. Li and X. Wang, *Angew. Chem., Int. Ed.*, 2014, **53**, 2857–2861; (*g*) J. Casado, *Top. Curr. Chem.*, 2017, **375**, 73.

4 (*a*) M. C. Simpson and J. D. Protasiewicz, *Pure Appl. Chem.*, 2013, **85**, 801–815; (*b*) D. A. Pantazis, J. E. McGrady, J. M. Lynam, C. A. Russell and M. Green, *Dalton Trans.*, 2004, 2080–2086; (*c*) K. B. Dillon, F. Mathey and J. F. Nixon, *Phosphorus: The Carbon Copy*, John Wiley & Sons, New York 1998.

5 (*a*) M. W. Schmidt, P. N. Truong and M. S. Gordon, *J. Am. Chem. Soc.*, 1987, **109**, 5217–5227; (*b*) P. v. R. Schleyer and D. Kost, *J. Am. Chem. Soc.*, 1988, **110**, 2105–2109.

6 (*a*) R. C. Fischer and P. P. Power, *Chem. Rev.*, 2010, **110**, 3877–3923; (*b*) L. Nyulaszi, T. Veszpremi and J. Reffy, *J. Phys. Chem.*, 1993, **97**, 4011–4015; (*c*) R. Szűcs, P.-A. Bouit, L. Nyulászi and M. Hissler, *ChemPhysChem*, 2017, **18**, 2618–2630; (*d*) L. Nyulászi, *Chem. Rev.*, 2001, **101**, 1229–1246.

7 (*a*) R. G. Hicks, *Stable Radicals: Fundamentals and Applied Aspects of Odd-Electron Compounds*, John Wiley & Sons Ltd, Chichester, 2010; (*b*) S. Kundu, S. Sinhababu, V. Chandrasekhar and H. W. Roesky, *Chem. Sci.*, 2019, **10**, 4727–4741; (*c*) G. W. Tan and X. P. Wang, *Chin. J. Chem.*, 2018, **36**, 573–586; (*d*) Y. Kim and E. Lee, *Chem.–Eur. J.*, 2018, **24**, 19110–19121; (*e*) Y. Su and R. Kinjo, *Coord. Chem. Rev.*, 2017, **352**, 346–378; (*f*) M. Soleilhavoup and G. Bertrand, *Acc. Chem. Res.*, 2015, **48**, 256–266; (*g*) C. D. Martin, M. Soleilhavoup and G. Bertrand, *Chem. Sci.*, 2013, **4**, 3020–3030; (*h*) T. Chivers and J. Konu, in *Comprehensive Inorganic Chemistry II: From Elements to Applications*, ed. J. Reedijk and K. Poeppelmeier, Elsevier, Amsterdam, 2013, vol. I, pp. 349–373; (*i*) J. Konu and T. Chivers, in *Stable Radicals: Fundamental and Applied Aspects of Odd-Electron Compounds*, ed. R. G. Hicks, John Wiley & Sons, Ltd, 2010, pp. 381–406; (*j*) F. Breher, *Coord. Chem. Rev.*, 2007, **251**, 1007–1043; (*k*) P. P. Power, *Chem. Rev.*, 2003, **103**, 789–809; (*l*) H. Grützmacher and F. Breher, *Angew. Chem., Int. Ed.*, 2002, **41**, 4006–4011; (*m*) K. C. Mondal, S. Roy and H. W. Roesky, *Chem. Soc. Rev.*, 2016, **45**, 1080–1111.

8 (*a*) S. L. Hinchley, C. A. Morrison, D. W. H. Rankin, C. L. B. Macdonald, R. J. Wiacek, A. H. Cowley, M. F. Lappert, G. Gundersen, J. A. C. Clyburne and P. P. Power, *Chem. Commun.*, 2000, 2045–2046; (*b*) A. Armstrong, T. Chivers, M. Parvez and R. T. Boeré, *Angew. Chem., Int. Ed.*, 2004, **43**, 502–505; (*c*) S. Ito, M. Kikuchi, M. Yoshifuji, A. J. Arduengo III, T. A. Konovalova and L. D. Kispert, *Angew. Chem., Int. Ed.*, 2006, **45**, 4341–4345; (*d*) P. Agarwal, N. A. Piro, K. Meyer, P. Müller and C. C. Cummins, *Angew. Chem., Int. Ed.*, 2007, **46**, 3111–3114; (*e*) O. Back, B. Donnadieu, M. von Hopffgarten, S. Klein, R. Tonner, G. Frenking and G. Bertrand, *Chem. Sci.*, 2011, **2**, 858–861; (*f*) T. Beweries, R. Kuzora, U. Rosenthal, A. Schulz and A. Villinger, *Angew. Chem., Int. Ed.*, 2011, **50**, 8974–8978; (*g*) A. Hinz, A. Schulz and A. Villinger, *Angew. Chem., Int. Ed.*, 2015, **54**, 668–672; (*h*) L. Gu, Y. Zheng, E. Haldón, R. Goddard, E. Bill, W. Thiel and M. Alcarazo, *Angew. Chem., Int. Ed.*, 2017, **56**, 8790–8794; (*i*) X. Chen, A. Hinz, J. R. Harmer and Z. Li, *Dalton Trans.*, 2019, **48**, 2549–2553; (*j*) Z. Li, Y. Hou, Y. Li, A. Hinz, J. R. Harmer, C.-Y. Su, G. Bertrand and H. Grützmacher, *Angew. Chem., Int. Ed.*, 2018, **57**, 198–202; (*k*) A. M. Tondreau, Z. Benko, J. R. Harmer and H. Grützmacher, *Chem. Sci.*, 2014, **5**, 1545–1554.

9 (*a*) O. Back, M. A. Celik, G. Frenking, M. Melaimi, B. Donnadieu and G. Bertrand, *J. Am. Chem. Soc.*, 2010, **132**, 10262–10263; (*b*) O. Back, B. Donnadieu, P. Parameswaran, G. Frenking and G. Bertrand, *Nat. Chem.*, 2010, **2**, 369–373; (*c*) R. Kinjo, B. Donnadieu and G. Bertrand, *Angew. Chem., Int. Ed.*, 2010, **49**, 5930–5933; (*d*) G. Ménard, J. A. Hatnean, H. J. Cowley, A. J. Lough, J. M. Rawson and D. W. Stephan, *J. Am. Chem. Soc.*, 2013, **135**, 6446–6449; (*e*) X. Pan, Y. Su, X. Chen, Y. Zhao, Y. Li, J. Zuo and X. Wang, *J. Am. Chem. Soc.*, 2013, **135**, 5561; (*f*) X. Pan, X. Chen, T. Li, Y. Li and X. Wang, *J. Am. Chem. Soc.*, 2013, **135**, 3414–3417; (*g*) Y. Su, X. Zheng, X. Wang, X. Zhang, Y. Sui and X. Wang, *J. Am. Chem. Soc.*, 2014, **136**, 6251–6254; (*h*) A. Brückner, A. Hinz, J. B. Priebe, A. Schulz and A. Villinger, *Angew. Chem., Int. Ed.*, 2015, **54**, 7426–7430; (*i*) X. Pan, X. Wang, Z. Zhang and X. Wang, *Dalton Trans.*, 2015, **44**, 15099–15102.

10 (*a*) T. Cantat, F. Biaso, A. Momin, L. Ricard, M. Geoffroy, N. Mézailles and P. Le Floch, *Chem. Commun.*, 2008, 874–876; (*b*) C. Pi, Y. Wang, W. Zheng, L. Wan, H. Wu, L. Weng, L. Wu, Q. Li and P. v. R. Schleyer, *Angew. Chem., Int. Ed.*, 2010, **49**, 1842–1845; (*c*) X. Pan, X. Wang, Y. Zhao, Y. Sui and X. Wang, *J. Am. Chem. Soc.*, 2014, **136**, 9834–9837; (*d*) G. Tan, S. Li, S. Chen, Y. Sui, Y. Zhao and

X. Wang, *J. Am. Chem. Soc.*, 2016, **138**, 6735–6738; (*e*) S.-s. Asami, S. Ishida, T. Iwamoto, K. Suzuki and M. Yamashita, *Angew. Chem., Int. Ed.*, 2017, **56**, 1658–1662; (*f*) M. K. Mondal, L. Zhang, Z. Feng, S. Tang, R. Feng, Y. Zhao, G. Tan, H. Ruan and X. Wang, *Angew. Chem., Int. Ed.*, 2019, **58**, 15829–15833.

11 (*a*) R. Appel, U. Kündgen and F. Knoch, *Chem. Ber.*, 1985, **118**, 1352–1370; (*b*) R. Appel, F. Knoch and H. Kunze, *Chem. Ber.*, 1984, **117**, 3151–3159; (*c*) R. Appel, V. Barth and F. Knoch, *Chem. Ber.*, 1983, **116**, 938–950; (*d*) B. Breit, H. Memmesheimer, R. Boese and M. Regitz, *Chem. Ber.*, 1992, **125**, 729–732; (*e*) M. Seidl, M. Stubenhofer, A. Y. Timoshkin and M. Scheer, *Angew. Chem., Int. Ed.*, 2016, **55**, 14037–14040; (*f*) T. W. Mackewitz, D. Ullrich, U. Bergsträßer, S. Leininger and M. Regitz, *Liebigs Ann. Chem.*, 1997, **1997**, 1827–1839; (*g*) J. C. Guillemin, J. L. Cabioch, X. Morise, J. M. Denis, S. Lacombe, D. Gonbeau, G. Pfister-Guillouzo, P. Guenot and P. Savignac, *Inorg. Chem.*, 1993, **32**, 5021–5028.

12 Y. Wang, Y. Xie, P. Wei, R. B. King, H. F. Schaefer III, P. v. R. Schleyer and G. H. Robinson, *J. Am. Chem. Soc.*, 2008, **130**, 14970–14971.

13 O. Back, G. Kuchenbeiser, B. Donnadieu and G. Bertrand, *Angew. Chem., Int. Ed.*, 2009, **48**, 5530–5533.

14 D. Rottschäfer, M. K. Sharma, B. Neumann, H.-G. Stammler, D. M. Andrada and R. S. Ghadwal, *Chem.–Eur. J.*, 2019, **25**, 8127–8134.

15 M. K. Sharma, D. Rottschäfer, S. Blomeyer, B. Neumann, H.-G. Stammler, M. van Gastel, A. Hinz and R. S. Ghadwal, *Chem. Commun.*, 2019, **55**, 10408–10411.

16 (*a*) D. Rottschäfer, B. Neumann, H.-G. Stammler, R. Kishi, M. Nakano and R. S. Ghadwal, *Chem.–Eur. J.*, 2019, **25**, 3244–3247; (*b*) M. K. Sharma, S. Blomeyer, B. Neumann, H. G. Stammler and R. S. Ghadwal, *Chem.–Eur. J.*, 2019, **25**, 8249–8253; (*c*) M. K. Sharma, S. Blomeyer, B. Neumann, H.-G. Stammler, M. van Gastel, A. Hinz and R. S. Ghadwal, *Angew. Chem., Int. Ed.*, 2019, **58**, 17599–17603.

17 V. Lavallo, Y. Canac, C. Präsang, B. Donnadieu and G. Bertrand, *Angew. Chem., Int. Ed.*, 2005, **44**, 5705–5709.

18 No reaction between {(IPr)C(Ph)PCl$_2$} (**1**) and a weak π-acceptor NHC (IPr or SIPr) was observed under similar experimental conditions.

19 B. Manz, U. Bergsträßer, J. Kerth and G. Maas, *Chem. Ber.*, 1997, **130**, 779–788.

20 (*a*) O. Kühl, *Phosphorus-31 NMR Spectroscopy*, Springer-Verlag, Berlin Heidelberg, 2008; (*b*) V. B. Gudimetla, L. Ma, M. P. Washington, J. L. Payton, M. Cather Simpson and J. D. Protasiewicz, *Eur. J. Inorg. Chem.*, 2010, 854–865.

21 P. Pyykkö and M. Atsumi, *Chem.–Eur. J.*, 2009, **15**, 186–197.

22 F. Mathey, *Angew. Chem., Int. Ed.*, 2003, **42**, 1578–1604.

23 S. L. Hinchley, C. A. Morrison, D. W. H. Rankin, C. L. B. Macdonald, R. J. Wiacek, A. Voigt, A. H. Cowley, M. F. Lappert, G. Gundersen, J. A. C. Clyburne and P. P. Power, *J. Am. Chem. Soc.*, 2001, **123**, 9045–9053.