Formation and Structure of a Stable Monoradical Cation by Reduction of a Diphosphafulvenium Salt

BIASO, Frédéric, et al.

DOI: 10.1002/anie.200603009

Available at:
http://archive-ouverte.unige.ch/unige:3637

Disclaimer: layout of this document may differ from the published version.
The attention paid by physical chemists to organic radical ions has never ceased to increase over the last 20 years,[1,2] mainly because these species often act as reaction intermediates and because their structure can determine the result of many chemical processes.[3] Considerable efforts have been devoted to gaining a better knowledge of their electronic structure; unfortunately their high reactivity make their identification difficult, and reports on their crystal structures remain sparse.[4] Renewed interest in the behavior of organic ion radicals has recently appeared owing to the practical applications predicted for some of these compounds in materials science and because of their potential use as electronic devices, organic magnets, and organic conductors.[5,6] Fulvene derivatives (A, Scheme 1) play an important role in this field; for example, dithiafulvene (B) constitutes one of the elementary units of conducting, semiconducting, and superconducting molecular solids. In this context, a crucial property of tetrafulvalene lies in its capacity to easily release one electron to give, reversibly, a rather persistent radical cation.[7] The discovery of novel fulvene-centered systems that are able to reversibly stabilize a radical cation is of considerable interest. The results herein contribute to this project, and as will be seen further, phosphorus chemistry can open new and very attractive perspectives in this area.

On the basis of early reports on the chemistry of 1,4-diphosphoniacyclohexa-2,5-dienes,[8] as well as our recent EPR study on the reduction processes of monophospholium cations C+,[9] we postulated that pentavalent phosphorus atoms could behave as strong electron acceptors and thus allow the synthesis and stabilization of persistent radical cations through reduction processes. We validate this hypothesis herein in the case of a diphosphafulvenium dication D2+ whose reduction leads to a stable radical cation. To the best of our knowledge, diphosphafulvenium cations have never been reported so far.[10]

The trans-tetraphenyldiphosphafulvene derivative 1, whose synthesis was recently reported, was chosen as the starting precursor.[11] Reaction of 1 with excess methyl triflate (MeOTf) at 25 °C cleanly afforded dication 22+, which was isolated as a very stable white solid. The structure of 22+ was confirmed by NMR data and elemental analyses. Unfortunately, despite many attempts, 22+ proved to be reluctant towards crystallization, thus precluding the recording of an X-ray crystal structure. As expected, dication 22+ can be easily reduced, as indicated by cyclic voltammetry (CH3CN, room temperature), which revealed four waves at −0.54 V (reversible), −1.10 V (irreversible), −1.37 (partially reversible), and −1.72 V (irreversible) versus SCE. Importantly, the first reduction process was found to be reversible even at low scan rates (50 mV s−1), thereby suggesting that it would be possible to isolate a persistent radical cation. Reduction of 22+ was
carried out at low temperature by adding 1,2-dimethoxy-ethane (DME) to a solid mixture of $\mathbf{2}^+$ and [CoCp$_2$]. The solution immediately turned deep purple, and after warming to 25 °C, n-hexane was added to crystallize the radical monocation $\mathbf{2}^+$, which was then easily separated from cobaltocenium triflate (Scheme 2).

Scheme 2. Synthesis of compounds $\mathbf{2}^+$(TfO)$^-$ and $\mathbf{2}^{-}\text{TF}^-$.

The structure of $\mathbf{2}^+$ was confirmed by an X-ray crystallographic study. A view of one molecule of $\mathbf{2}^+$ is presented in Figure 1. The most significant piece of data concerns the two P1–C5 (1.713(3) Å) and P2–C5 (1.718(3) Å) bond lengths, which are very short compared to the P1–C1 and P2–C2 bond lengths (1.809(3) Å). On the other hand, the C5–C6 bond (1.473(4) Å) is also very long and has clearly lost its double-bond character. The geometry at C6 is planar ($\Sigma$angles = 358.8°). Furthermore, the CPh$_2$ group is twisted from the plane that is defined by the ring ($\Sigma$(P1-C5-C6-C31) = −70.5°) and participates in stabilizing the extra electron, as attested to by the short C6–C31 and C6–C37 bond lengths (av. 1.46 Å). All these data suggest that reduction has taken place at the P1–P2–C5–C6 segment. This observation is confirmed by the fact that the C1–C2 bond length remains close to what is expected for a classical carbon–carbon double bond (1.34 Å).

EPR experiments and DFT calculations were carried out to determine the precise electronic structure of $\mathbf{2}^+$. Electrolysis in the EPR cavity at a potential corresponding to the first reduction wave leads to the spectrum shown in Figure 2a.

This spectrum, measured at 300 K, is characterized by a $g$ value equal to 2.0029 and by a hyperfine interaction of about 30 G with two equivalent $^{31}$P nuclei. Additional structure resulting from coupling with several protons of two phenyl rings is clearly visible (Figure 2b). It is interesting to note that these couplings are quite similar to those measured by ENDOR on Ph$_3$C. A simulation was performed by using the following coupling constants (expressed in Gauss): $^{31}$P 30.1, H$_a$ 3.00, H$_b$ 2.83, H$_c$ 2.82, H$_d$ 1.35, H$_e$ 1.11. Note that the spectrum of $\mathbf{2}^+$, which was also recorded at 100 K, clearly shows that the $g$ and $^{31}$P coupling tensors do not exhibit any anisotropy. The reduced species is rather persistent, and its spectrum could be observed for several hours without voltage under a nitrogen atmosphere. The same spectrum was obtained by reducing a solution of $\mathbf{2}^+$ in THF with one equivalent of C$_8$H$_8$Na.

To get further insights into the electronic structure of $\mathbf{2}^+$, DFT calculations were carried out on the actual molecules $\mathbf{2}^+$ and $\mathbf{2}^-$ at the quantum level. A good agreement was found between the experimental and theoretical structures of $\mathbf{2}^+$. Only the dihedral angle of the CPh$_2$ group differs. An examination of the molecular orbitals of the two species furnishes important information. In $\mathbf{2}^+$ the LUMO is antibonding between C5 and C6 and bonding between the phosphorus atoms and C5 (Figure 3), thus explaining why the P–C5 bonds are shortened upon reduction and why the C5–C6 bond is significantly elongated. Upon reduction, one electron is added into the LUMO of $\mathbf{2}^-$ as shown in the SOMO of $\mathbf{2}^-$ (Figure 3). Calculations of Wiberg bond indices...
confirm that the order of the C5–C6 bond has been reduced on going from 2+ to 2+ (1.58 to 1.11).

Isotropic and anisotropic hyperfine interactions with 1H ($A_{iso} = -2.63$ and $-2.84$ G for $H_{iso}$, 1.16 and 1.43 G for $H_{max}$, and $-3.05$ G for $H_{min}$) and $^{31}$P nuclei ($A_{iso} = 32.9$ and 33.2 G; $\tau_{min} = -4$, $-3$, and $+7$) were obtained from the DFT calculations and agree with the experimental values. In accord with frozen-solution spectra, the anisotropy of the $^{31}$P coupling was also found to be rather small. All these data suggest that in diphosphafulvenium systems the two pentavalent phosphorus atoms behave as strong acceptors and prevent delocalization over the ring. Such systems are therefore different from their sulfur or carbon counterparts. The excess of electron density gained upon reduction mainly resides on the exocyclic double bond and is strongly stabilized by negative hyperconjugation with the two pentavalent phosphorus atoms. Therefore, two ylidic forms can be proposed to rationalize the electronic structure of 2+ (Scheme 3).

Scheme 3. Ylidic forms involved in 2+.

From this study we can conclude that 1,1-bis(phosphonio) olefins can act as suitable precursors for the synthesis of persistent monoradical cations upon reduction at low potential. Further studies aimed at expanding this simple idea to the elaboration of highly conjugated paramagnetic structures are currently underway in our laboratories.

**Experimental Section**

All experiments were carried out in a dry argon or nitrogen atmosphere by using distilled and degassed solvents. For the synthesis of 2+ (TIO)2, see the Supporting Information. 2+ (TIO)2: DME (25 mL) was condensed onto a mixture of 2+ (TIO)2 (0.24 g, 0.27 mmol) and [CoCp2] (51 mg, 0.27 mmol) at $-78^\circ$C. The solution was warmed to room temperature and stirred for two hours. A mixture of hexanes was then condensed onto the crude mixture at $-78^\circ$C, which resulted in the formation of deep purple crystals of 2+ (TIO)2 within 12 h. The resulting crystals were isolated and washed with diethyl ether (3 x 5 mL) to afford 2+ (TIO)2 in 93% yield (0.19 g, 0.25 mmol). No NMR spectrum is available for this compound owing to its paramagnetic nature.

Received: July 26, 2006
Published online: October 2, 2006

**Keywords:** density functional calculations - EPR spectroscopy - phosphorus - radicals - reduction

[12] Crystal data for 2+ (TIO)2: C20H20P2C2O2S, monoclinic, space group P21/n, a = 10.5080(10), b = 17.3590(10), c = 19.5300(10) Å, $\beta = 90.3500(10)^\circ$, $V = 3685.54(1) Å^3$, $Z = 4$, $\mu_{tot} = 1.35$ g cm$^{-3}$, $F(000) = 1564$, MoK$\alpha$ radiation ($\lambda = 0.71069$ Å), $\mu = 0.230$ cm$^{-1}$, crystal dimensions 0.20 x 0.20 x 0.18 mm. Data collection was performed on a Nonius Kappa CCD single-crystal diffractometer at $T = 150 K$; the crystal structure was solved with SIR97[16] refinement against $F^2$ (SHELXL-97) with anisotropic thermal parameters for all nonhydrogen atoms, calculated hydrogen positions with riding isotropic thermal parameters, $hkl$ ranges: $-13/13$, $-20/22$ - 24/24; 13478 data collected, 7774 unique data ($R_{int} = 0.0203$), 5796 data with $I > 2\sigma(I)$, 427 parameters refined, GoF($F^2$) = 1.022, final R indices ($R1 = \sum||F_o||-|F_c||/\sum||F_o||$, $wR2 = \sum||F_o^2 - F_c^2||/\sum||F_o^2||^{1/2}$), $R1 = 0.0584$, $wR2 = 0.1643$, max/min residual electron density 0.664(0.055)/ -0.504(0.055) e Å$^{-3}$. CCDC-612414 contains the supplementary crystallographic data for this paper. These data can be obtained.
A search of the Cambridge Structural Database afforded 15 structures of triarylmethyl radicals. We found an average C–Cipso bond length of 1.46 Å. For comparison, an average corresponding bond length of 1.53 Å is found for Ar3CH derivatives.


See the Supporting Information for further details.

ζ(P1-C5-C6-C31) = 53.3° for the optimized structure versus −70.5° for the X-ray structure.

P–C5 bonds are shortened from 1.819 Å in 2+ to 1.756 Å in 2+ (calculated values).

C5–C6 bond is elongated from 1.397 Å in 2+ to 1.469 Å in 2+ (calculated values).