



ELSEVIER

Contents lists available at ScienceDirect

Comptes Rendus Chimie

www.sciencedirect.com



Full paper/Mémoire

Towards the stability limit of cyclic diphosphonium bis-ylides

Mohammed Abdalilah^{a,b}, Yves Canac^{a,*,b}, Christine Lepetit^{a,b}, Remi Chauvin^{a,b,*}^a Laboratoire de chimie de coordination (LCC), CNRS, 205, route de Narbonne, 31077 Toulouse, France^b UPS, INP, LCC, université de Toulouse, 31077 Toulouse, France

ARTICLE INFO

Article history:

Received 5 February 2010

Accepted after revision 7 April 2010

Available online 31 May 2010

Keywords:

Phosphonium ylide
Carbodiphosphorane
Yldiide
Diylide
Carbon ligands

ABSTRACT

A typology of molecules containing two phosphonium ylide groups (bis-ylides) is proposed, *versus* the bridge length (fused, ω - and α -bisylides) and their relative topographical orientations (head-to-head, tail-to-tail or head-to-tail). The formal electrostatic constraint occurring in the head-to-head series is systematically addressed for cyclic representatives based on the *o*-bis(diphenylphosphonio)benzene framework. After a survey of previously reported results in the fused and β -bis-ylide series, emphasis is given to the cyclic α -bis-ylides. The non-substituted, non-stabilized version escaped isolation through spontaneous fragmentation to bis(diphenylphosphino)benzene and acetylene. Inspired by this result, the reverse process was employed for the generation of stabilized representatives with ethoxycarbonyl and benzoyl substituents at the ylidic carbon atoms. The stability and stereochemistry of the obtained head-to-head α -bis-ylides was investigated by NMR techniques and reproduced and analyzed by DFT calculations. The role of electrostatics in determining structural and reactivity features of cyclically constrained species is here illustrated.

© 2010 Académie des sciences. Published by Elsevier Masson SAS. All rights reserved.

1. Introduction

In a primary description, ylides can be represented as α -zwitterionic species where a positively charged heteroatom X with an octet electron count (such as P, S, N) is covalently bonded to a carbanionic center. In this category, phosphonium ylides \mathbf{F}_0 possessing an easily pyramidalized carbanion stabilized by an adjacent tetrahedral phosphonium center (Scheme 1), received a long-standing attention not only in the field of organic synthesis, but also of fundamental coordination chemistry. Beyond their ubiquitous role in the Wittig-type olefination of carbonyl compounds, phosphonium ylides are indeed intriguing stable electron-rich C- sp^3 carbyl

ligands of transition metals [1]. They are today attracting a renewed interest from the standpoints of stereochemistry and catalysis [1]. Depending on the charge delocalization exerted by the substituents, ylides are classified in three families constituted by the non-stabilized, the semi-stabilized and the stabilized versions. Due to their intrinsic stability and their ambidentate character (C- vs O- or N-coordinating modes), complexes of α -stabilized phosphonium ylide ligands have been naturally more exemplified [1].

Ideally C_2 -symmetric chelating ligands provide their complexes with particular kinetic stability (entropic factor) and stereochemical control (*cis vs trans*). Along the same lines as those governing the design of numerous diphosphines, diphosphonium bis-ylides deserve a particular typology. Three types of neutral bis-ylides can thus be distinguished (Scheme 1): the ω -type Ω_1 - Ω_3 , the α -type \mathbf{A}_1 - \mathbf{A}_3 and the fused type \mathbf{F}_1 - \mathbf{F}_3 . The ionic forms \mathbf{I}_0 - \mathbf{I}_3 of the respective types \mathbf{F}_0 - \mathbf{F}_3 , in particular yldiides \mathbf{I}_0 and

* Corresponding authors.

E-mail addresses: yves.canac@lcc-toulouse.fr (Y. Canac), chauvin@lcc-toulouse.fr (R. Chauvin).

diylides I_2 , should also be mentioned. Each type gathers three forms depending on the regiochemistry of the coupling between two ylide units F_0 , consisting each in a carbanionic head and a phosphonium tail: head-to-head, tail-to-tail or head-to-tail.

Among general ω -bis-ylides, the head-to-head form Ω_1 is exemplified by β -bis-ylides involving various bridging moieties (X: CO, PR, PR_2^+ , PYZ, P^+) [2]. When X = CO or when the ylides moieties are independently stabilized, the ambidentate abilities of such ligands are greatly enhanced. Several ω -bis-ylide ligands of the tail-to-tail form Ω_2 (X: $(CH_2)_n$, RPPR, ...) are also known, including β -representatives (X = RP) [3].

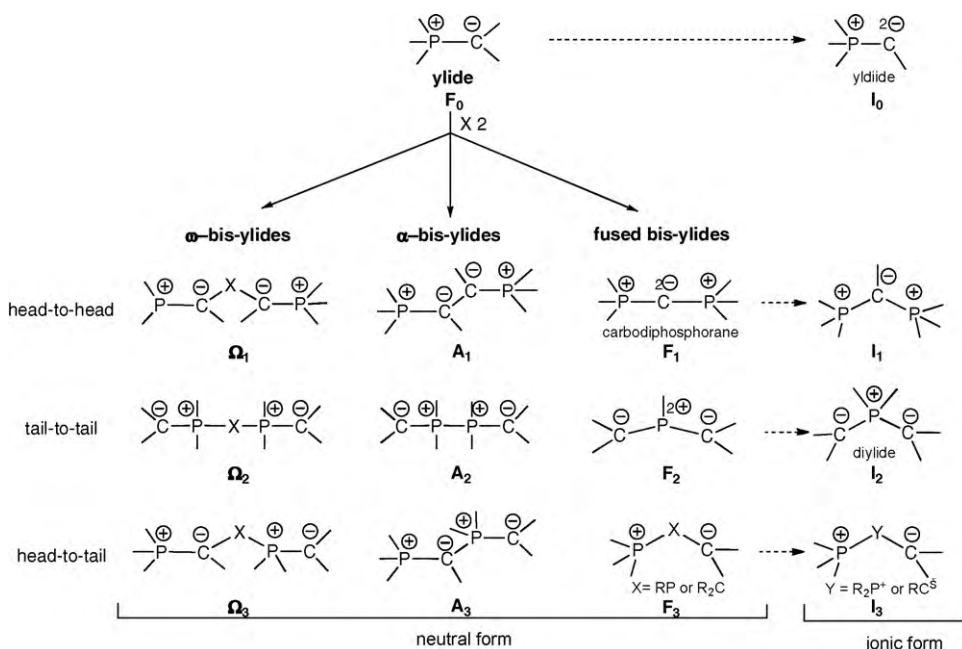
Fused bis-ylides are represented by their head-to-head (C-fused) form F_1 , corresponding to the carbodiphosphanes [4]. These species contain two cumulated ylide functions, with a formally divalent central carbon atom with two negative charges stabilized by two phosphonio groups. The presence of two electron pairs at the central carbon atom induces both a bent structure and a very strong nucleophilicity. Carbodiphosphanes can be regarded as neutral stabilized ylides of type I_0 , where a generic C-substituent is replaced by a phosphonium group [5]. Due to their strong nucleophilicity, carbodiphosphanes are poorly exemplified and undergo easy protonation or alkylation to the corresponding β -diphosphonium mono-ylides of type I_1 .

Tail-to-tail (P-fused) bis-ylides of the F_2 type, first exemplified by Appel et al. in 1982, correspond to $\sigma^3\lambda^5$ -bis-(methylene)phosphorane derivatives [6]. According to Schoeller's theoretical studies, these polarized π -systems, isoelectronic to allyl anions, possess a weakly pyramidalized phosphorus atom [7]. Their structural and

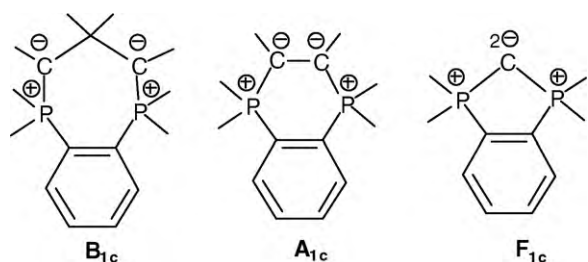
reactivity features are, however, basically similar to those of ylides I_0 . Ionic P-alkylated versions of bis-ylides F_2 are the so-called diylides I_2 , acting as stable ligands in transition metal complexes [8] possibly serving as catalyst precursors [8b]. The head-to-tail form F_3 , with a neutral sp^3 -C or -P center, is only a formal bis-ylide, i.e., a β -zwitterionic form where the ylidic character is no longer apparent.

In the α -bis-ylides series, the head-to-head (A_1) and tail-to-tail (A_2) forms exhibit strong C^-/C^- and P^+/P^+ electrostatic repulsion, respectively (Scheme 1). Head-to-head representatives A_1 may behave as genuine carbene transfer reagents upon coordination: cleavage of a P=C bond leads to an ylido-carbene and a phosphino group anchoring to the metal center [9]. The tail-to-tail form A_2 was first exemplified by Karsch et al., in 1988 [10]. In these species, both the repulsion of the adjacent cationic phosphorus atoms and the steric bulk of the carbanionic substituents (Me_3Si) may explain the observed length of the P^+-P^+ bond (2.278(1) Å), even longer than the P-P bond between five-coordinate P atoms in a bis-phosphorane (2.264(2) Å). To the best of our knowledge, the sole example of fully characterized head-to-tail α -bis-ylide form A_3 , with a triple charge alternance (Scheme 1), has been reported by Schmidbaur et al. in 1980 [11].

In contrast to linear diphosphonium bis-ylides (Scheme 1), their cyclic versions have attracted less attention [12]. The case of cyclic head-to-head β -, α - and fused bis-ylide types (B_{1c} , A_{1c} , F_{1c} , respectively) is hereafter systematically addressed (Scheme 2). The corresponding diphosphonium precursors based on the rigid, *cis*-spacing, insulating 1,2-phenylene bridge have been recently shown to be easily accessible [13].



Scheme 1. Typology of acyclic and cyclic phosphonium bis-ylides.



Scheme 2. Cyclic head-to-head β -, α - and fused bis-ylides on the *o*-phenylene framework.

2. Results and discussion

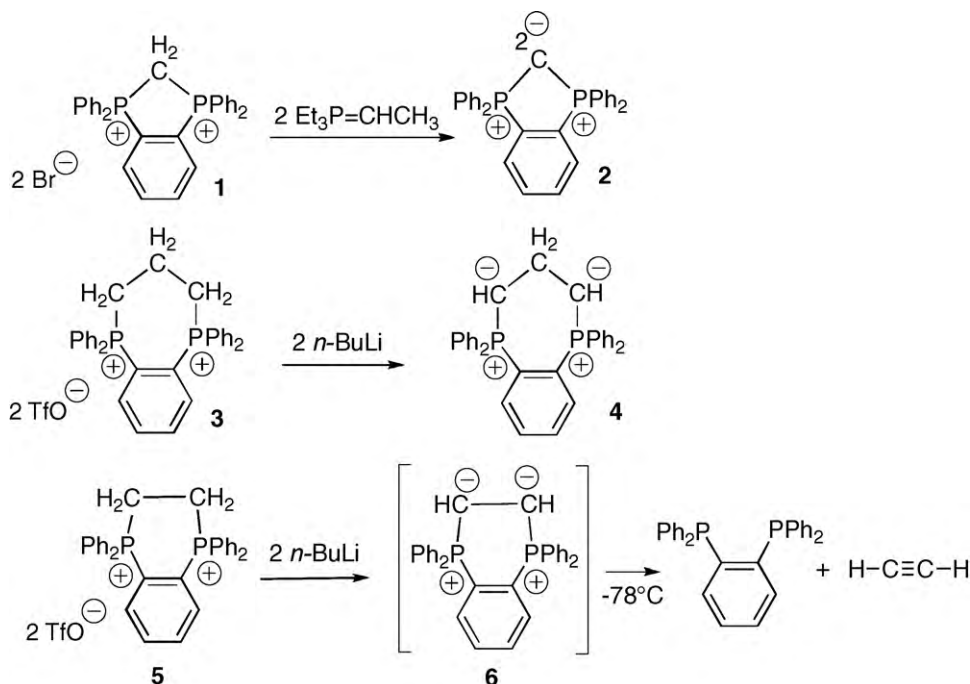
2.1. Non-stabilized head-to-head fused bis-ylides of cyclic diphosphoniums

The diphosphonium **1.Br₂** was first obtained by Schmidbauer et al. in 61% yield by refluxing *o*-bis(diphenylphosphino)benzene (*o*-dppb) and dibromomethane in toluene over 10 days [14]. We recently reported that replacing CH₂Br₂ by CH₂I₂ allowed for the isolation of the analogous diphosphonium **1.I₂** in 96% yield over 2 days only (Scheme 3) [13]. A rapid H/D exchange of the methylene protons was observed in the ¹H NMR spectrum of **1.I₂** in acidic deuterated solvents: the acidity of these protons ($\delta^1\text{H}(\text{P}^+\text{CH}_2\text{P}^+) = +5.38$ ppm, $t, {}^2J_{\text{PH}} = 11.7$ Hz) suggests the possible existence of the di-conjugated base, namely the cyclic carbodiphosphorane **2**. And indeed, according to Schmidbauer's results, the treatment of **1.Br₂** with an ylide base (Et₃P=CHMe) in THF at -78°C led to the

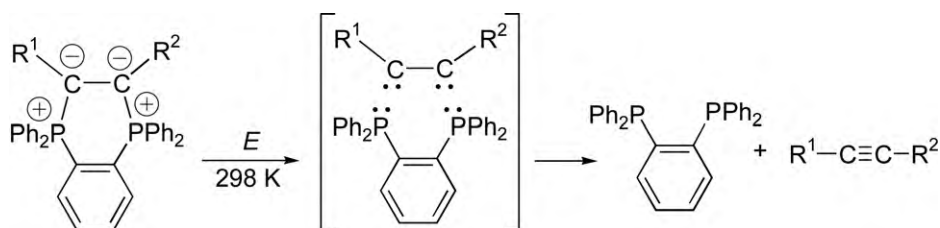
formation of the expected cyclic C-fused bis-ylide **2** [14]. This carbodiphosphorane was identified by ³¹P and ¹H NMR spectroscopy at -30°C , and by the regeneration of the starting diphosphonium **1.Br₂** upon quenching with one equivalent of HBr. The thermal instability of **2** is naturally explained by geometrical and electronic factors; the acute PCP angle in the five membered ring (*ca* 108°) indeed enforces the *sp*³ (vs *sp*) character of the central C atom, thus increasing the weight of the doubly zwitterionic resonance form, and finally a destabilizing charge separation.

2.2. Non-stabilized head-to-head β -bis-ylides of cyclic diphosphoniums

Elongation of the aliphatic carbon chain between the ylidic carbons and electron delocalization over two carbon atoms instead of one, should result in both geometrical and electronic relaxation. Following this principle, it was recently shown that addition of two equivalents of *n*-BuLi in THF at low temperature to the less strained diphosphonium **3.(TfO)₂** afforded the corresponding cyclic head-to-head β -bis-ylide **4** (Scheme 3) [15]. The structure of **4** was unambiguously established by multi-nuclear NMR spectroscopy, and in particular by the presence of a unique ³¹P signal at +24.4 ppm, which confirmed the onio character and the equivalence of the two phosphorus atoms. In the ¹H and ¹³C NMR spectra, the simultaneous presence of a triplet of triplets at $\delta_{\text{H}} = +0.59$ ppm ($J_{\text{HP}} = 26.8$, $J_{\text{HH}} = 3.3$ Hz) and of a triplet at $\delta_{\text{C}} = +21.9$ ppm ($J_{\text{CP}} = 4.4$ Hz) was assigned to the bridging methylene group of **4**. This bis-ylide was reported to act as a strongly donating C,C-chelating ligand of Rh(I) centers [15]. In spite of its strong



Scheme 3. Deprotonation of the cyclic diphosphoniums **1**, **3** and **5**.



Scheme 4. Fragmentation of cyclic head-to-head *o*-phenylene-diphosphonium α -bis-ylides, $R_1, R_2 = H, CO_2Me, CO_2Et$ (see calculated data in Table 1).

internal charge repulsion and its low stability ($t_{1/2} \approx 45$ min. at $+20^\circ C$), the β -bis-ylide **4** is stable enough to undergo selective transformations. Further bridge shortening from β to α was thus the next challenge.

2.3. Non-stabilized head-to-head α -bis-ylides of cyclic diphosphoniums

The cyclic *o*-phenylene-diphosphonium **5** with two carbon atoms in the aliphatic bridge, is easily prepared by reaction of *o*-dppb with 1,2-dichloro-ethane [13]. The corresponding head-to-head α -bis-ylide **6** was targeted by deprotonation of **5** with two equivalents of *n*-butyllithium in THF at $-78^\circ C$. The ^{31}P NMR spectrum indicated the disappearance of **5** ($\delta^{31}P = +12.6$ ppm) and the presence of a much more shielded new signal at $\delta^{31}P = -10.0$ ppm, which was attributed to *o*-dppb (Scheme 3). Multi-nuclear NMR monitoring in THF- d_8 at $-78^\circ C$ showed the immediate appearance of the *o*-dppb signal, along with a 1H signal attributable to acetylene, and this even when only a catalytic amount of *n*-butyllithium was added. Considering the chemical existence of the more strained fused bis-ylide **2** with two geminal negative charges, the non-detection of **6** even at low temperature, can be explained by electrostatic factors, in particular by the enhanced repulsion between the two vicinal negative charges: instantaneous fragmentation results in a neutral *o*-dppb molecule and a formal α -bis-carbene $H-C(:)-C(:)-H$ in resonance with acetylene (Scheme 4). Phosphonium ylides are indeed recognized as phosphino-carbenoid species [12].

This extrusion reaction is supported by the calculation of a strongly exergonic dissociation energy ($\mathbf{6} \rightarrow \textit{o}\text{-dppb} + \text{acetylene}$: $\Delta G(298\text{ K}) = -21.2$ kcal/mol) at the B3PW91/6-31G** level (Scheme 4, Table 1).

Upon substitution of one of the ylidic carbon atoms of **6** by an ester group (CO_2Et or CO_2Me), the calculated dissociation energy remains exergonic ($\Delta G(298\text{ K}) = -8.6$ kcal/mol), and it becomes slightly endergonic upon substitution of both ylidic carbons (Scheme

4). The stability of such α -bis-ylides, however, cancels if the alkoxycarbonyl groups are replaced by benzoyl groups (Table 1). These trends are in agreement with the relatively “stabilized” character of the corresponding mono-ylides in general. Although the positive Gibbs energies for the dissociation of ester-stabilized α -bis-ylides are quite small in the gas phase, solvent dielectric constant effects are expected to provide additional stabilization of these highly polar species. Experimental efforts in this direction are reported in the next section.

2.4. Stabilized head-to-head α -bis-ylides of cyclic diphosphoniums

Preparation of head-to-head α -stabilized α -bis-ylides is envisaged through the reverse reaction of their fragmentation depicted in Scheme 4. Preparation of the corresponding diphosphonium precursors by adaptation of the previously used method [13] is indeed *a priori* tricky, and calculations suggest that the transformation is *a priori* not thermodynamically forbidden (Table 1). Moreover, arylphosphines are known to react with activated unsaturated substrates such as simple acetylenic Michael acceptors [9]. The envisioned process is formally a [2+1+1] cycloaddition which proceed in two steps, initiated by a Michael-type addition [16].

2.4.1. Ester group-stabilized head-to-head α -bis-ylides of cyclic diphosphoniums

As anticipated above, addition of one equivalent of diethyl acetylenedicarboxylate (DEAD) to *o*-dppb in toluene gave, after 10 min at room temperature, nearly quantitative yield of the targeted bis-ylide **7** as an apparent 70/30 mixture of two stereoisomers (Scheme 5). The occurrence of these two isomers likely results from a restricted rotation about the exocyclic $C-CO_2Et$ bonds. Strong delocalization of ylidic electron pairs is indeed classically observed in α -stabilized phosphonium ylides. According to this interpretation, the three *Z/Z*, *E/E* and *Z/E* stereoisomers could, however, be expected.

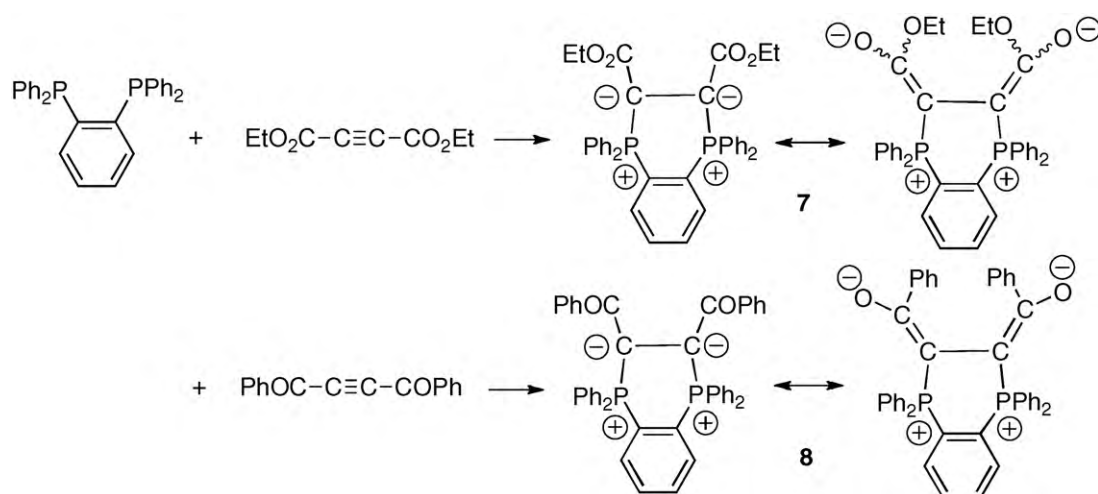
Table 1

Dissociation energies (in kcal/mol) of cyclic head-to-head α -bis-ylides into the corresponding diphosphine (*o*-dppb) and acetylene derivatives (Scheme 4), calculated at the B3PW91/6-31G** level^a.

	$R^1 = R^2 = H$	$R^1 = H, R^2 = CO_2Me$	$R^1 = R^2 = CO_2Et$	$R^1 = R^2 = CO_2Me$	$R^1 = R^2 = COPh$
ΔH (0 K)	-6.93	8.53	22.02	22.89	20.25
ΔH_{ZPE} (0 K) ^a	-9.47	6.14	19.39	20.14	17.56
ΔG (298.15 K)	-21.19	-8.60	2.00	2.60	-0.67

Dissociation energies were calculated from the most stable conformer of bis-ylides, diphosphines and acetylene derivatives

^a Zero-point energy corrected values.



Scheme 5. Synthetic approach of α -stabilized diphosphonium α -bis-ylides **7** and **8** from *o*-dpph.

The structure of **7** in CD_3CN solution was first assigned on the basis of the observed ^{31}P NMR chemical shifts of two broad singlets at $\delta_{\text{p}} = +0.52$ and $+2.21$ ppm, which are in the typical range for α -stabilized phosphonium ylides. The structure of **7** was then confirmed by ^1H and ^{13}C NMR spectroscopy and by mass spectrometry (electron spray $[\text{MH}]^+$: 617.2).

A variable temperature ^{31}P NMR experiment indicated a reversible coalescence of the two broad signals at $+60^\circ\text{C}$ corresponding to an activation barrier of *ca.* 15 kcal/mol, thus demonstrating the dynamic behavior of the different isomers of **7**.

By decreasing the temperature to 0°C , decoalescence in four broad singlets was observed, two of them having the same integral. These signals can thus be globally assigned to the expected three stereoisomers of bis-ylide **7** in a 41/22/37 ratio with respective chemical shifts $\delta_{\text{pa}} = +0.36$ and $\delta_{\text{pa}'} = +2.40$ ppm/ $\delta_{\text{pb}} = +0.60$ ppm/ $\delta_{\text{pc}} = +2.23$ ppm (Fig. 1). Due to dissymmetry, the *Z/E* isomer exhibits two non-equivalent phosphorus atoms and is therefore assigned to the set of the P_a and P_a' signals. In contrast, the symmetric *Z/Z* and *E/E* isomers exhibit only one type of phosphorus atom and are globally assigned to the P_b and P_c signals. Considering both attractive P^+/O^- and repulsive O^-/O^- electrostatic interactions, the *E/E* isomer is expected to be more stable than the *Z/Z* isomer: the major signal at $\delta_{\text{pc}} = +2.23$ ppm (37%) is therefore tentatively assigned to the *E/E* configuration, while the minor signal at $\delta_{\text{pb}} = +0.60$ ppm (22%) would correspond to the *Z/Z* configuration.

The structure of the three stereoisomers of bis-ylide **7** has been investigated at the B3PW91/6-31G** level of calculation. A crude conformational analysis on the CO_2Me -substituted bis-ylide model allowed to rule out C_s -symmetric conformers with planar or boat conformations of the aliphatic ring: these conformers are indeed more than 10 kcal/mol higher in energy than C_2 -symmetric conformers with a twisted aliphatic ring. Conformational analysis of the bis-ylide **7** itself restricted to C_2 symmetry yielded several equilibrium structures of quasi-degenerate

energies. As anticipated experimentally, the three most stable conformers rank in the energetic order: *Z/E* < *E/E* < *Z/Z* (Table 2). This confirms the assignment empirically proposed on the basis of elementary electrostatic considerations (see above): *Z/Z* isomer at $\delta_{\text{pb}} = +0.60$ ppm (22%); *E/E* isomer at $\delta_{\text{pc}} = +2.23$ ppm (37%).

The ^{31}P NMR chemical shifts of the three stereoisomers of **7** have been calculated at the B3PW91/6-31+G** level (Table 2). The chemical shift values obtained for the inequivalent phosphorus atoms of the *E/E* isomer in its frozen optimized C_1 -symmetry have been averaged to

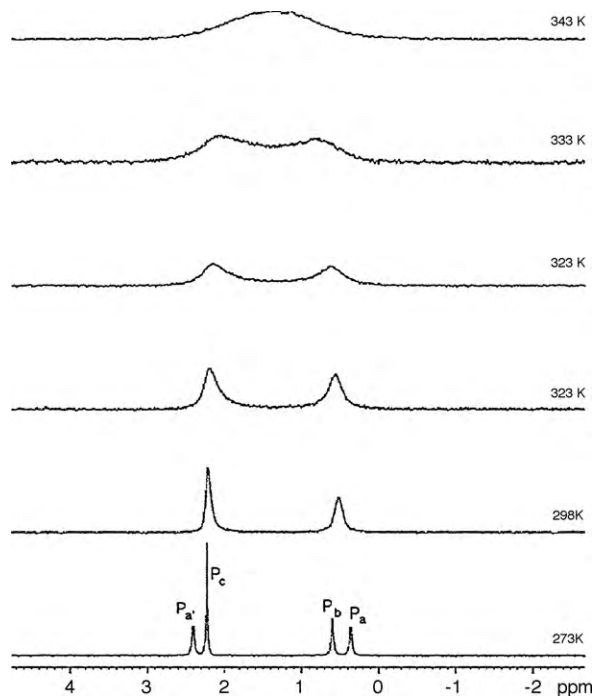


Fig. 1. Variable temperature ^{31}P NMR spectra of the bis-ylide **7** in CD_3CN at 202.53 MHz. For assignment, see text.

Table 2

Calculated structures of the three stereoisomers of the head-to-head α -bis-ylide **7** (B3PW91/6-31G**).^a ^{31}P NMR chemical shifts calculated at the B3PW91/6-31+G** level. For the *E/E* isomer, the given value is the mean of the calculated chemical shifts (in square brackets) for the frozen optimized C_1 -symmetric structure.

Geometry	Z/E	E/E	Z/Z
Symmetry	C_1	C_1	C_2
Relative energy (kcal/mol)	0.0	0.28	0.57
Calcd. ^{31}P δ (ppm) _a	-11.9	-8.57	-7.54
Exptl. ^{31}P δ (ppm)	3.9	[(-16.8 + 0.33)/2]	-7.54
	0.36	2.23	0.60
	2.40		

regenerate the apparent symmetry due their respective fluxionality at the NMR time scale. The calculated ^{31}P NMR chemical shifts are systematically more shielded than the experimental values. However, although the extreme chemical shifts calculated for the *Z/E* isomer (-11.9 and 3.9 ppm) correspond to the extreme chemical shifts observed for the same isomer (0.36 and 2.40 ppm), the rigid approximation is not sufficient for reproducing the relative experimental values with further accuracy. This is explained by the narrow range of chemical shifts observed in all the three isomers ($\Delta = 2.40 - 0.36 \approx 2$ ppm). Calculations of ^{31}P NMR chemical shifts of strained polar bis-ylides appear therefore less straightforward than those of related diphosponiums [13].

2.4.2. Keto group-stabilized head-to-head α -bis-ylides of cyclic diphosponiums

The procedure employed with the DEAD substrate was applied with another electron-poor symmetric alkyne, the 1,4-diphenylbut-2-yne-1,4-dione [17]. Reaction of this diketone with *o*-dppb in toluene at room temperature over 10 min thus afforded the corresponding cyclic α -bis-ylide **8** in nearly quantitative yield (Scheme 5). The bis-ylide was obtained as a single stereoisomer as indicated by the observation of a unique ^{31}P NMR signal at $\delta_{\text{p}} = -0.08$ ppm. A variable temperature ^{31}P NMR experiment ($-40 \rightarrow +50$ °C) did not indicate any significant change, except a slight sharpening of the signal at low temperatures which can be due to a higher rigidity of the structure of the diketone-bis-ylide **8** as compared to that of the bis(ethoxycarbonyl) bis-ylide **7** (section 2.5). On the basis of empirical geometrical and electrostatic considerations, the configuration of the observed stereoisomer of **8** is assigned to *Z/Z*, with two *cis*- $\text{Ph}_2\text{P}^+-\text{C}=\text{C}-\text{O}^-$ stereogenic units (Scheme 5).

The structure of the three stereoisomers of bis-ylide **8** has been calculated at the B3PW91/6-31G** level. In contrast to the bis-ylide **7**, but in agreement with the above-proposed assignment, the *Z/Z* stereoisomer of C_2

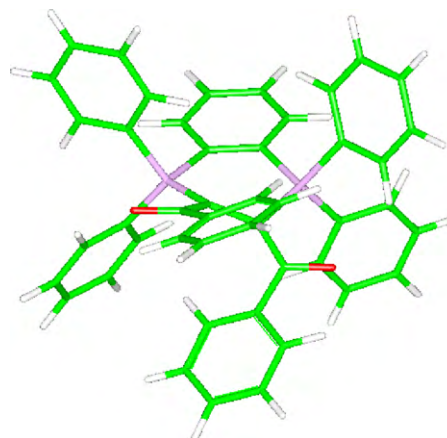


Fig. 2. Calculated structure of the *Z/Z* stereoisomer of the α -bis-ylide **8** (B3PW91/6-31G**).

symmetry is found much more stable than the *Z/E* and *E/E* isomers, lying 4.95 and 10.5 kcal/mol higher in energy, respectively (Fig. 2). The calculated ^{31}P NMR chemical shift of **8** (+6.6 ppm at the B3PW91/6-31+G** level) is quite shielded for a P(IV) nucleus, in qualitative agreement with the experimental value (-0.1 ppm).

3. Conclusion

The synthesis and stability of diphosponiums head-to-head bis-ylides constrained in five-, six- and seven-membered rings have been systematically compared. In the non-stabilized series, it has been shown that while the fused bis-ylide **2** and the β -bisylide **4** are relatively stable, the α -bisylide **6** is unstable due to the repulsion of two vicinal negative charges and undergoes spontaneous fragmentation to *o*-dppb and acetylene. Introducing electron-withdrawing substituents at the two ylidic carbon atoms allowed for the isolation of stabilized α -bis-ylides **7** and **8**. The latter were obtained by a formal

[2+1+1] reaction of *o*-dppb with electron-poor alkynes following a stepwise mechanism. The stabilization effects and the stereochemical behaviour of the α -bis-ylides have been reproduced and analyzed at the theoretical level. In particular, it has been shown that, by contrast to unsubstituted parents, α -bis-ylides stabilized by ester groups exhibit endergonic dissociation. These results illustrate how electrostatic effects (e.g., repulsion between adjacent negative charges) may govern reactivity in geometrically strained frameworks [13].

4. Experimental section

4.1. General remarks

Toluene and diethyl ether were dried and distilled over sodium/benzophenone. All other reagents were used as commercially available. All reactions were carried out under argon atmosphere, using schlenk and vacuum line techniques. The following analytical instruments were used: Bruker ARX 250, DPX 300, AV 500. NMR chemical shifts δ are in ppm, with positive values to high frequency relative to the tetramethylsilane reference for ^1H and ^{13}C and to H_3PO_4 for ^{31}P .

7. To a solution of 1,2-bis(diphenylphosphino)benzene (*o*-dppb) (0.45 g, 1.0 mmol) in dry toluene (10.0 mL) at room temperature was added diethyl acetylenedicarboxylate (DEAD) (0.33 mL, 2.05 mmol). The reaction mixture was then stirred for 10 min. After evaporation of the solvent under vacuum, the reddish-yellow solid residue was washed by dry diethyl ether (2×20 mL) affording **7** as a yellow solid product (yield: 0.60 g, 98%).

^{31}P NMR (CD_3CN , 25 °C): $\delta = +0.52$ (s), $+2.21$ (s) ppm; ^{31}P NMR (CD_3CN , 0 °C): $\delta = +0.36$ (s) and $+2.40$ (s); $+0.60$ (s), $+2.23$ (s) ppm. ^1H NMR (CD_3CN , 0 °C): $\delta = 0.36$ (t, $J_{\text{HH}} = 7.1$ Hz, CH_3), 0.41 (broad t, $J_{\text{HH}} = 7.0$ Hz, CH_3), 1.07 (broad t, $J_{\text{HH}} = 7.0$ Hz, CH_3), 1.11 (broad t, $J_{\text{HH}} = 6.9$ Hz, CH_3), 3.33 (m, CH_2), 3.54 (m, CH_2), 3.78 (m, CH_2), 7.18–7.37 (m, H_{ar}), 7.44–7.69 (m, H_{ar}), 8.12–8.28 (m, H_{ar}). ^{13}C NMR (CD_3CN , 0 °C): $\delta = 13.3$ (s, CH_3), 13.4 (s, CH_3), 14.8 (s, CH_3), 14.9 (s, CH_3), 56.5 (s, CH_2), 56.6 (s, CH_2), 56.9 (s, CH_2), 128.1 (m, CH_{ar} , C_{ar}), 128.3 (d, $J_{\text{CP}} = 12.2$ Hz, CH_{ar}), 128.8 (d, $J_{\text{CP}} = 15.3$ Hz, CH_{ar}), 130.2 (d, $J_{\text{CP}} = 88.1$ Hz, C_{ar}), 130.9–131.8 (m, CH_{ar} , C_{ar}), 132.2 (s, CH_{ar}), 133.1 (d, $J_{\text{CP}} = 10.0$ Hz, CH_{ar}), 133.3 (d, $J_{\text{CP}} = 10.0$ Hz, CH_{ar}), 133.7 (m, CH_{ar}), 134.4 (pseudo ttt, $J_{\text{CP}} = 7.6$ and 24.0 Hz, CH_{ar}), 169.5 (d, $J_{\text{CP}} = 25.7$ Hz, CO), 170.1 (d, $J_{\text{CP}} = 28.4$ Hz, CO), 170.5 (d, $J_{\text{CP}} = 27.4$ Hz, CO), 170.6 (d, $J_{\text{CP}} = 27.0$ Hz, CO). MS (ES⁺) m/z 617.2 [MH]⁺. HRMS (ES⁺) calculated for $\text{C}_{38}\text{H}_{35}\text{O}_4\text{P}_2$: 617.2011; found 617.2037.

8. To a solution of 1,2-bis(diphenylphosphino)benzene (*o*-dppbo) (0.45 g, 1.0 mmol) in dry toluene (10.0 mL) at room temperature was added 1,4-diphenylbut-2-yne-1,4-dione (0.48 g, 2.05 mmol). The reaction mixture was then stirred for 10 min. After evaporation of the solvent under vacuum, the reddish-yellow solid residue was washed by dry diethyl ether (2×20 mL) giving **8** as a yellow solid product (yield: 0.61 g, 90%).

^{31}P NMR (CD_3CN , 25 °C): $\delta = -0.08$ ppm; ^{31}P NMR (CD_3CN , -40 °C): $\delta = +0.24$ ppm. ^1H NMR (CD_3CN , -40 °C): $\delta = 6.42$ (d, $J_{\text{HH}} = 6.7$ Hz, 4H, H_{ar}), 6.68 (m, 4H,

H_{ar}), 7.00 (m, 2H, H_{ar}), 7.17–7.29 (m, 4H, H_{ar}), 7.40 (m, 3H, H_{ar}), 7.50 (m, 1H, H_{ar}), 7.58 (m, 2H, H_{ar}), 7.66 (m, 1H, H_{ar}), 7.77–7.90 (m, 9H, H_{ar}), 8.11 (m, 4H, H_{ar}). ^{13}C NMR (CD_3CN , -40 °C): $\delta = 124.4$ (d, $J_{\text{CP}} = 73.0$ Hz, C_{ar}), 126.5 (brs, CH_{ar}), 128.1 (brs, CH_{ar}), 128.8–129.7 (m, CH_{ar} , C_{ar}), 130.9 (brs, CH_{ar}), 131.7 (brs, CH_{ar}), 131.9 (brs, CH_{ar}), 132.7 (brs, CH_{ar}), 133.5 (m, CH_{ar}), 136.1 (brs, CH_{ar}), 141.9 (brs, C_{ar}), 185.7 (pseudo t, $J_{\text{CP}} = 9.4$ Hz, CO). MS (ES⁺): m/z 681.2 [MH]⁺; HRMS (ES⁺) calculated for $\text{C}_{46}\text{H}_{35}\text{O}_2\text{P}_2$ 681.2112; found, 681.2115.

4.2. Computational details

Geometries were fully optimized at the B3PW91/6-31G** level of calculation using Gaussian 03 [18]. Vibrational analysis was performed at the same level of calculation as the geometry optimization. The magnetic shielding tensor was calculated at the B3PW91/6-31+G** level using the gauge-independent atomic orbital (GIAO) method implemented in Gaussian 03 [18]. The ^{31}P NMR chemical shifts were estimated with respect to the usual H_3PO_4 reference.

Acknowledgements

The authors thank the French *ministère de l'Enseignement supérieur et de la recherche*, the French *Centre national de la recherche scientifique*, the ANR program (ANR-08-JCJC-0137-01) for financial support. The authors also thank *Calcul intensif en Midi-Pyrénées* (CALMIP, Toulouse, France), *institut du développement et des ressources en informatique scientifique* (IDRIS, Orsay, France) and *centre informatique de l'enseignement supérieur* (CINES, Montpellier, France) for computing facilities. M.A. thanks UNESCO, the French Consulate in East Jerusalem, and the *conseil régional de Midi-Pyrénées* for financial support.

Thanks are also due to one of the referees of this article who draw our attention to the point mentioned in reference [16].

Appendix A. Supplementary data

Supplementary data associated with this article can be found, in the online version, at doi:10.1016/j.crci.2010.04.005.

References

- [1] (a) E.P. Urriolabeitia, Top. Organomet. Chem. 'Late transition metal complexes of neutral η^1 -carbon ligands: R. Chauvin, Y. Canac, (Eds.), Springer, 30, 2010, (chapter 2), p. 15; (b) E.P. Urriolabeitia, Dalton Trans. 42 (2008) 5673; (c) L.R. Falvello, J.C. Ginés, J.J. Carbó, A. Lledos, R. Navarro, T. Soler, E.P. Urriolabeitia, Inorg. Chem. 45 (2006) 6803; (d) J. Vicente, M.T. Chicote, Coord. Chem. Rev. 193–195 (1999) 1143; (e) O.I. Kolodiazhnyi, Tetrahedron 52 (1996) 1855; (f) W.C. Kaska, K.A. Ostojka Starzewski, in: A.W. Johnson (Ed.), Ylides and Imines of Phosphorus, John Wiley & Sons: New York, 1993, (Chapter 14); (g) H. Schmidbaur, Angew. Chem. Int. Ed. Engl. 22 (1983), 22 907.
- [2] (a) J. Vicente, M.T. Chicote, I. Saura-Llamas, P.G. Jones, K. Meyer-Bäse, C. Freire Erdbrügger, Organometallics 7 (1988) 997; (b) A. Schmidpeter, G. Jochem, C. Klinger, C. Robl, H. Nöth, J. Organomet. Chem. 529 (1997) 87;

- (c) L.R. Falvello, S. Fernández, R. Navarro, A. Rueda, E.P. Urriolabeitia, *Organometallics* 17 (1998) 5887;
 (d) S. Fernández, R. Navarro, E.P. Urriolabeitia, *J. Organomet. Chem.* 602 (2000) 151;
 (e) E. Serrano, C. Vallés, J.J. Carbó, A. Lledós, T. Soler, R. Navarro, E.P. Urriolabeitia, *Organometallics* 25 (2006) 4653;
 (f) E. Serrano, R. Navarro, T. Soler, J.J. Carbó, A. Lledós, R. Navarro, E.P. Urriolabeitia, *Inorg. Chem.* 48 (2009) 6823.
- [3] (a) A. Spannenberg, W. Baumann, U. Rosenthal, *Organometallics* 19 (2000) 3991;
 (b) L.R. Falvello, M.E. Margalejo, R. Navarro, E.P. Urriolabeitia, *Inorg. Chim. Acta* 347 (2003) 75;
 (c) H.H. Karsch, E. Witt, A. Schneider, E. Herdtweck, M. Heckel, *Angew. Chem. Int. Ed.* 34 (1995) 557.
- [4] (a) W. Petz, G. Frenking, in: R. Chauvin, Y. Canac (Eds.), *Late transition metal complexes of neutral η^1 -carbon ligands*, Springer, 30, 2010, (Chapter 3), p. 49.
 (b) R. Tonner, F. Öxler, B. Neumüller, W. Petz, G. Frenking, *Angew. Chem. Int. Ed.* 45 (2006) 8038;
 (c) R. Tonner, G. Frenking, *Angew. Chem. Int. Ed.* 46 (2007) 8695;
 (d) R. Tonner, G. Frenking, *Chem. Eur. J.* 14 (2008) 3260;
 (e) S. Marrot, T. Kato, H. Gornitzka, A. Baceiredo, *Angew. Chem. Int. Ed.* 45 (2006) 2598.
- [5] (a) M. Schlosser, T. Kadibelban, G. Steinhoff, *Angew. Chem. Int. Ed.* 11 (1966) 968;
 (b) E.J. Corey, J. Kang, *J. Am. Chem. Soc.* 104 (1982) 4724;
 (c) E.J. Corey, J. Kang, K. Kyler, *Tett. Lett.* 26 (1985) 555;
 (d) H.J. Bestmann, M. Schmidt, *Angew. Chem. Int. Ed.* 26 (1987) 79;
 (e) M.I. Khalil, *Main Group Met. Chem.* 21 (1998) 193;
 (f) B. Schaub, T. Jenny, M. Schlosser, M. Tett. Lett. 25 (1985) 4097;
 (g) B. Schaub, M. Schlosser, *Tett. Lett.* 26 (1985) 1623;
 (h) K. Korth, J. Sundermeyer, *Tett. Lett.* 41 (2000) 5461;
 (i) S. Goumri-Magnet, H. Gornitzka, A. Baceiredo, G. Bertrand, *Angew. Chem. Int. Ed.* 38 (1999) 678;
 (j) R. Zurawinski, C. Lepetit, Y. Canac, M. Mikolajczyk, R. Chauvin, *Inorg. Chem.* 48 (2009) 2147.
- [6] (a) R. Appel, J. Peters, A. Westerhaus, *Angew. Chem. Int. Ed.* 21 (1982) 80;
 (b) R. Appel, A. Westerhaus, *Tett. Lett.* (1982) 2017;
 (c) A.R. Barron, A.H. Cowley, *J. Chem. Soc., Chem. Commun.* (1987) 1092;
 (d) H.J. Metternich, E. Niecke, *Angew. Chem. Int. Ed.* 30 (1991) 312;
 (e) R. Appel, in: M. Regitz, O. J. Scherer (Eds.), *Multiple Bonding and Low Coordination in Phosphorus Chemistry*, Thieme (1990) 367.
- [7] W.W. Schoeller, J. Niemann, *J. Am. Chem. Soc.* 108 (1986) 22.
- [8] (a) G. Wittig, M. Rieber, *Justus Liebigs Ann. Chem.* 562 (1949) 177;
 (b) R.A. Grey, L.R. Anderson, *Inorg. Chem.* 16 (1977) 3187;
 (c) H.J. Cristau, *Chem. Rev.* 94 (1994) 1299;
 (d) M. Taillefer, H.J. Cristau, *Top. Curr. Chem.* 229 (2003) 41.
- [9] (a) M.A. Shaw, J.C. Tebby, R.S. Ward, D.H. Williams, *J. Chem. Soc. (C)* (1967) 2442;
 (b) M.A. Shaw, J.C. Tebby, *J. Chem. Soc. (C)* (1970) 5;
 (c) J.C. Tebby, I.F. Wilson, D. Vaughan Griffiths, *J. Chem. Soc. Perkin I.* (1979) 2133;
 (d) L.R. Falvello, R. Luser, M.E. Margalejo, R. Navarro, E.P. Urriolabeitia, *Organometallics* 22 (2003) 1132.
- [10] (a) H.H. Karsch, B. Deubelly, G. Müller, *J. Chem. Soc., Chem. Commun.* (1988) 517;
 (b) H.H. Karsch, B. Deubelly, G. Grauvogl, G. Müller, *J. Organomet. Chem.* 459 (1993) 95;
 (c) H.H. Karsch, E. Witt, *J. Organomet. Chem.* 529 (1997) 151.
- [11] H. Schmidbaur, U. Deschler, B. Zimmer-Gasser, D. Neugebauer, U. Schubert, *Chem. Ber.* 113 (1980) 902.
- [12] (a) Y. Canac, S. Conejero, M. Soleilhavoup, B. Donnadiou, G. Bertrand, *J. Am. Chem. Soc.* 128 (2006) 459;
 (b) S. Conejero, M. Song, D. Martin, Y. Canac, M. Soleilhavoup, G. Bertrand, *Chem. Asian. J.* 1–2 (2006) 155.
- [13] M. Abdalilah, R. Zurawinski, Y. Canac, B. Laleu, J. Lacour, C. Lepetit, G. Magro, G. Bernardinelli, B. Donnadiou, C. Duhayon, M. Mikolajczyk, R. Chauvin, *Dalton Trans.* (2009) 8493.
- [14] G.A. Bowmaker, R. Herr, H. Schmidbaur, *Chem. Ber.* 116 (1983) 3567.
- [15] (a) Y. Canac, C. Lepetit, M. Abdalilah, C. Duhayon, R. Chauvin, *J. Am. Chem. Soc.* 130 (2008) 8406;
 (b) Y. Canac, C. Duhayon, R. Chauvin, *Angew. Chem. Int. Ed.* 46 (2007) 6313.
- [16] (a) M.A. Shaw, J.C. Tebby, R.S. Ward, D.H. Williams, *J. Chem. Soc. (C)* (1970) 504;
 (b) M. Davies, A.N. Hughes, S.W.S. Jafry, *Can. J. Chem.* 50 (1972) 3625.
- [17] Z.Z. Zhang, G.B. Schuster, *J. Am. Chem. Soc.* 111 (1989) 7149.
- [18] M.J. Frisch, G.W. Trucks, H.B. Schlegel, G.E. Scuseria, M.A. Robb, J.R. Cheeseman, J.A. Montgomery, T. Vreven, K.N. Kudin, J.C. Burant, J.M. Millam, S.S. Iyengar, J. Tomasi, V. Barone, B. Mennucci, M. Cossi, G. Scalmani, N. Rega, G.A. Petersson, H. Nakatsuji, M. Hada, M. Ehara, K. Toyota, R. Fukuda, J. Hasegawa, M. Ishida, T. Nakajima, Y. Honda, O. Kitao, H. Nakai, M. Klene, X. Li, J.E. Knox, H.P. Hratchian, J.B. Cross, C. Adamo, J. Jaramillo, R. Gomperts, R.E. Stratmann, O. Yazyev, A.J. Austin, R. Cammi, C. Pomelli, J.W. Ochterski, P.Y. Ayala, K. Morokuma, G.A. Voth, P. Salvador, J.J. Dannenberg, V.G. Zakrzewski, S. Dapprich, A.D. Daniels, M.C. Strain, O. Farkas, D.K. Malick, A.D. Rabuck, K. Raghavachari, J.B. Foresman, J.V. Ortiz, Q. Cui, A.G. Baboul, S. Clifford, J. Ciołowski, B.B. Stefanov, G. Liu, A. Liashenko, P. Piskorz, I. Komaromi, R. L. Martin, D.J. Fox, T. Keith, M.A. Al-Laham, C.Y. Peng, A. Nanayakkara, M. Challacombe, P.M.W. Gill, B.W. Johnson, W. Wong, C. Gonzalez, and J.A. Pople, *Gaussian 03, Revision B.05*, Gaussian, Inc., Pittsburgh PA, 2003.