## Intra- and inter-molecular phosphoryl migration in phosphinothiazolines; precursors to polynuclear complexes and bimetallic coordination polymers<sup>†</sup>‡

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Two tautomers of the new phosphinoaminothiazoline Ph<sub>2</sub>PNHC=NCH<sub>2</sub>CH<sub>2</sub>S (1), obtained from the reaction of 2-amino-2-thiazoline (ATHZ) with Ph<sub>2</sub>PCl, have been structurally characterized and the intermediate formation of the diphosphine Ph<sub>2</sub>PN=CN(PPh<sub>2</sub>)CH<sub>2</sub>CH<sub>2</sub>S (2) has been demonstrated experimentally and by DFT calculations; 2 reacts with [AuCl(THT)] to give [(AuCl)<sub>2</sub>2] whereas the bidentate metalloligand *cis*-[Pt(1<sub>-H</sub>)<sub>2</sub>] reacts with AgOTf to form the Ag–Pt coordination polymer [Ag<sub>2</sub>[Pt(1<sub>-H</sub>)<sub>2</sub>].(OTf)<sub>∞</sub>.

The functionalization of phosphine ligands represents a major way to fine-tune their stereoelectronic characteristics and coordination properties as well as the dynamic behaviour and reactivity (stoichiometric and catalytic) of their metal complexes.<sup>1</sup> Increasing the diversity of chemical functions present on such ligands provides considerable opportunities for studying selectivity-related problems. We have now found that when 2-amino-2-thiazoline (ATHZ) is reacted with Ph2PCl in a 1 : 1 molar ratio, with the original aim of preparing the monophosphine ligand Ph<sub>2</sub>PNHC=NCH<sub>2</sub>CH<sub>2</sub>S (1a), corresponding to the most stable tautomer of ATHZ,<sup>2</sup> it is in fact the diphosphine  $Ph_2PN=CN(PPh_2)CH_2CH_2S$  (2) that is isolated in almost quantitative yield (based on phosphorus). This is rather surprising in view of recent results obtained with a closely related system.<sup>3</sup> However, when this reaction was performed in the presence of a base like Et<sub>3</sub>N or *n*-BuLi, 1 was isolated in two tautomeric forms, 1a (favoured in non-polar solvents) and 1b, and 2 was found to be a reaction intermediate (Scheme 1). It will be shown below that the role of the base is to scavenge the HCl liberated in preference to ATHZ. Another intermediate was formed prior to 2 and identified as 3 (as confirmed by <sup>31</sup>P NMR when reacting, at low temperature, Ph<sub>2</sub>PCl with lithiated ATHZ in large excess, see Supplementary Information), which results from nucleophilic attack of the endo-nitrogen atom of ATHZ on Ph2PCl. This kinetic product (+19.6 kJ/mol with respect to 1a), the formation of which is consistent with the endo-nitrogen atom being more nucleophilic than the exo-nitrogen,<sup>4</sup> was found to slowly and quantitatively isomerize to 1. DFT studies§ suggest for this 1,3-PPh<sub>2</sub> migration an intramolecular mechanism in which the PPh<sub>2</sub> group is transferred from the endocyclic to the exocyclic nitrogen via transition state TS1 (Fig. 1a). Attempts to identify an intermolecular Ph<sub>2</sub>P transfer were unsuccessful. The high energy computed for TS1 (+126 kJ/mol) disfavours this reaction path. In fact, when Ph<sub>2</sub>PCl is reacted with ATHZ in a 1 : 1 stoichiometry, an alternative path is preferred. The exocyclic nitrogen of 3, being more nucleophilic than the endo-nitrogen of ATHZ, rapidly attacks Ph2PCl, to form 2. Isolated 2 was shown independently to react with ATHZ to give first 1a, which rapidly equilibrates with 1b. Calculations showed that the reaction  $[2 + ATHZ \rightarrow 2 1a]$  is energetically favoured by 6.1 kJ/mol (Fig. 1b). We suggest a concerted mechanism in which the Ph2P group is transferred from the endocyclic nitrogen of 2 to the exocyclic nitrogen of the ATHZ imino-tautomer while a proton transfer occurs in the opposite direction (Fig. 1b). The calculated transition state TS2 is





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<sup>&</sup>lt;sup>†</sup> Dedicated to Prof. A. Tiripicchio (Parma) on the occasion of his 70th birthday, with our warmest wishes.

<sup>‡</sup> Electronic supplementary information (ESI) available: Computed atomic coordinates and distances; complete sets of crystallographic parameters; experimental procedures and spectroscopic characterizations. See DOI: 10.1039/b603495k



**Fig. 1** Relative energy profiles for the intramolecular  $1 \rightarrow 3$  tautomerism (a) and the intermolecular ATHZ +  $2 \rightarrow 21$  conversion (b).

+71.1 kJ/mol with respect to the reagents (ESI). Summarizing, two pathways are available to form 1: the aforementioned (disfavoured) direct 1,3-PPh<sub>2</sub> migration in 3 and the reaction of the rapidly formed 2 with the ATHZ present in the reaction mixture (Scheme 1). From the combined theoretical and experimental data, we conclude that the two step conversion  $3 \rightarrow 2 \rightarrow 1$  is preferred to the one step isomerization  $3 \rightarrow 1$ . The latter slowly occurs only if Ph<sub>2</sub>PCl is unavailable in the reaction mixture, which then precludes the intermediate formation of 2.

Alternative transition states for the ATHZ +  $2 \rightarrow 21$  reaction were computed (ESI). For instance, the conversion featuring the amino-tautomer of ATHZ was found to be energetically disfavoured (+181 kJ/mol), while an activated complex formed by two molecules of ATHZ and one of 2 was calculated to be energetically similar to TS2 but is entropically disfavoured.

Diphosphine 2 was isolated as the major product when ATHZ was reacted with  $Ph_2PCl$  in a 3 : 2 ratio, formation of ATHZ·HCl preventing reaction of the free amine with 2 to give 1. The molecular structures of 1 and  $2\P$  have been determined by X-ray diffraction (Fig. 2).

Tautomers **1a** and **1b** co-crystallized and adopt a head to tail intermolecular arrangement, forming pseudo-dimers through two N····H–N hydrogen bonds. Tautomer **1b** was computed to be only slightly more stable (0.95 kJ/mol) than **1a** and this is consistent with the *ca.* 1 : 1 mixture observed in the solid state. The



Fig. 2 ORTEP views of the molecular structures of compounds 1 (left) and 2 (right). Displacement parameters include 50% of the electron density. For 1 only one of the two tautomers (1a) is shown. Owing to crystallographic disorder, the bond parameters for this ligand are discussed in the Supplementary material. Bond distances (Å) and angles (°) for 2: S(1)-C(1) 1.7796(17), P(1)-N(1) 1.7511(15), P(2)-N(2) 1.7105(15), N(1)-C(1) 1.364(2), N(2)-C(1) 1.280(2); C(1)-N(1)-C(3) 115.53(14), C(1)-N(1)-P(1) 115.13(12), C(3)-N(1)-P(1) 129.32(12), C(1)-N(2)-P(2) 119.99(12).



Scheme 2

tautomerism should occur *via* a head to tail transition state (**TS3**). Although phosphine transamination represents a widely used strategy for aminophosphine synthesis (*e.g.*  $P(NR_2)_3$ , R = Me, Et as starting reagent), only a few examples concerning diaryl- or dialkyl-aminophosphines have been reported.<sup>5</sup> It is remarkable that ATHZ plays a dual role, both as a substrate for Ph<sub>2</sub>PCl attack and as a transamination reagent. To the best of our knowledge, this kind of reactivity has never been reported in P(III) chemistry. The Pl–N1 bond distance in **2** is significantly longer than P2–N2, consistent with the easier breaking of the endocyclic P–N bond. As expected, the C=N double bond is localized between C1 and the exocyclic N2 atom.

When 2 was reacted with two equivalents of [AuCl(THT)] in  $CH_2Cl_2$ , the dinuclear gold complex 4 was formed in quantitative yield (Scheme 2). A view of its molecular structure || is shown in Fig. 3.

Each metal centre is coordinated, in a slightly distorted linear geometry, by a chlorine and a phosphorus atom. The coordinated ligand retains a geometry similar to that found in **2**. On the other hand, both P–N distances shorten upon coordination. Reactions of **2** with  $[PtCl_2(NCMe)_2]$  or  $[PdCl_2(COD)]$  led to the formation of poorly soluble precipitates, which are likely to be oligomeric



Fig. 3 ORTEP view of the crystal structure of compound 4 in  $4 \cdot CHCl_3$ . Displacement parameters include 50% of the electron density. Selected bond distances (Å) and angles (°): Au(1)–P(1) 2.229(2), Au(2)–P(2) 2.2212(17), Au(2)–Cl(2) 2.2834(19), Au(1)–Cl(1) 2.299(2), P(1)–N(1) 1.675(6), P(2)–N(2) 1.713(6), N(2)–C(1) 1.379(8), N(1)–C(1) 1.294(8); P(1)–Au(1)–Cl(1) 174.99(7), P(2)–Au(2)–Cl(2) 177.44(7), N(1)–P(1)–Au(1) 117.2(2), N(2)–P(2)–Au(2) 113.4(2), C(1)–N(1)–P(1) 124.8(5), C(1)–N(2)–P(2) 117.7(4), C(1)–N(2)–C(2) 115.1(5), C(2)–N(2)–P(2) 126.3(5), N(1)–C(1)–S(1) 128.7(5), N(2)–C(1)–S(1) 111.2(5), N(1)–C(1)–N(2) 120.1(6).



**Fig. 4** Views of the crystal structure of compound **6** in  $6\cdot 3CH_2Cl_2$ . Solvent molecules and OTf anions have been omitted for clarity. a) Asymmetric unit (displacement parameters include 50% of the electron density); b) two orthogonal projections of the polymer chain. Selected bond distances (Å) and angles (°): Pt–N(4) 2.084(4), Pt–N(2) 2.096(4), Pt–P(2) 2.2309(14), Pt–P(1) 2.2314(14), Ag(1)–N(3) 2.078(4), Ag(1)–N(1) 2.096(5), P(1)–N(1) 1.686(5), P(2)–N(3) 1.687(5), N(1)–C(1) 1.324(7), N(2)–C(1) 1.307(7), N(4)–C(4) 1.319(7), N(3)–C(4) 1.339(7); N(4)–Pt–N(2) 100.04(18), N(4)–Pt–P(2) 80.95(13), N(2)–Pt–P(1) 80.49(13), P(2)–Pt–P(1) 98.51(5), N(3)–Ag–N(1) 172.08(19).

coordination compounds in which the diphosphine **2** would act as a bridging ligand.

Ligand 1 reacted with *t*-BuLi and  $[PtCl_2(NCPh)_2]$  in a 2 : 2 : 1 ratio to afford the bidentate metalloligand *cis*- $[Pt(1_{-H})_2]$  (5) which, upon reaction with AgOTf, yielded quantitatively the new bimetallic, cationic coordination polymer  $[Ag_{\infty}[Pt(1_{-H})_2]_{\infty}](OTf)_{\infty}$  (6, Scheme 2). Views of the molecular structure of  $6.3CH_2Cl_2^{**}$  are depicted in Fig. 4.

The platinum coordination geometry is square planar, the four L–Pt–L angles summing to  $359.9(2)^{\circ}$ , while the Ag atoms, which are bound to the exocyclic N atoms of **5**, adopt a slightly bent geometry [N3–Ag–N1 172.08(19)°]. The C=N double bonds are delocalized over the N1,C1,N2 and N3,C4,N4 atoms, respectively, the C–N distances ranging from 1.307(7) [N2–C1] to 1.339(7) Å [N3–C4].

The resulting infinite chains form a zig-zag and wave-like arrangement (Fig. 4b); the silver atoms occupy the maximum and the minimum points of the sinusoid, and are separated by 9.899(3) Å. To the best of our knowledge, only two examples of structurally characterized Pt–Ag coordination polymers in which linear Ag centres connect Pt metalloligands have been reported before.<sup>6</sup>

The full characterization of tautomers 1a,b and the intermediate formation of the diphosphine ligand 2 in the course of their synthesis emphasize the subtlety of the bonding in these systems and are relevant to their use in coordination chemistry where ligand interactions with one or many metal centres may profoundly affect the nature of the reactive sites. Further studies with Cu(I), Ag(I) and Au(I) are in progress to delineate the conditions for the formation of new bimetallic coordination polymers.

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## Notes and references

§ All calculations were done using the program system TURBOMOLE.<sup>8a</sup> If not mentioned explicitly otherwise, we used the DFT method with the Becke–Perdew functional (BP86).<sup>8b</sup> The Coulomb terms were treated by the RI-J approximation.<sup>8c</sup> For the structure optimizations we used SV(P) basis sets<sup>8d</sup> (single zeta for core orbitals, double zeta for the valence shells and one set of polarization functions for all centres except hydrogen). Single point energies were calculated with larger triple zeta valence plus polarization basis sets (TZVP).<sup>8e</sup>

Crystal data for 1:  $C_{15}H_{15}N_2PS$ , M = 286.32, monoclinic  $P2_1/c$ , a =15.5330(3), b = 11.8340(2), c = 16.1740(3) Å,  $\beta = 105.3850(3)^{\circ}$ , V = 2866.52(9) Å<sup>3</sup>, Z = 8,  $D_{c} = 1.327$  g cm<sup>-3</sup>,  $\mu$ (Mo-K $\alpha$ ) = 0.325 mm<sup>-1</sup>, F(000) = 1200. A Nonius Kappa CCD diffractometer was employed for the collection of 8350 unique reflections (Mo-K $\alpha$ ,  $\lambda = 0.71073$  Å, T =173 K). The structure was solved by direct methods and refined by a full matrix least-squares technique based on  $F^2$  (SHELX97<sup>7</sup>). The final  $R_1$  and  $wR_2$  are 0.046 and 0.112, respectively, for 359 parameters and 5517 reflections  $[I > 2\sigma(I)]$ . CCDC 601675. The same data collection and refinement procedures were used for 2, 4·CHCl<sub>3</sub> and 6·3CH<sub>2</sub>Cl<sub>2</sub>. Crystal data for 2:  $C_{27}H_{24}N_2P_2S$ , M = 470.48, triclinic P-1, a = 9.1420(2), b = 10.0760(2), c = 14.4330(4) Å, α = 92.1400(9), β = 105.8700(9), γ = 107.9900(12)°, V = 1205.47(5) Å<sup>3</sup>, Z = 2, D<sub>c</sub> = 1.296 g cm<sup>-3</sup>, μ(Mo-Kα) =  $0.285 \text{ mm}^{-1}$ , F(000) = 492.  $R_1$  and  $wR_2$  are 0.048 and 0.123, respectively, for 289 parameters and 4950 reflections  $[I > 2\sigma(I)]$  (7015 unique). CCDC 601676. For crystallographic data in CIF or other electronic format see DOI: 10.1039/b603495k

|| Crystal data for **4**·CHCl<sub>3</sub>: C<sub>27</sub>H<sub>24</sub>Au<sub>2</sub>Cl<sub>2</sub>N<sub>2</sub>P<sub>2</sub>S·CHCl<sub>3</sub>, M = 1054.68, triclinic P-1, a = 9.610(1), b = 13.281(2), c = 13.769(2) Å,  $\alpha = 98.04(5)$ ,  $\beta = 101.87(5)$ ,  $\gamma = 104.52(5)^{\circ}$ , V = 1630.5(4) Å<sup>3</sup>, Z = 2,  $D_c = 2.148$  g cm<sup>-3</sup>,  $\mu$ (Mo-K $\alpha$ ) = 9.582 mm<sup>-1</sup>, F(000) = 992. The final  $R_1$  and  $wR_2$  are 0.041 and 0.11, respectively, for 361 parameters and 7031 reflections  $[I > 2\sigma(I)]$ (9536 unique). CCDC 601677. For crystallographic data in CIF or other electronic format see DOI: 10.1039/b603495k

\*\* Crystal data for 6·3CH<sub>2</sub>Cl<sub>2</sub>: C<sub>30</sub>H<sub>28</sub>AgN<sub>4</sub>P<sub>2</sub>PtS·3CH<sub>2</sub>Cl<sub>2</sub>:CF<sub>3</sub>O<sub>3</sub>S, M = 1277.43, monoclinic P2<sub>1</sub>/n, a = 9.185(3), b = 19.488(6), c = 25.892(9) Å,  $\beta = 99.74(2)^{\circ}$ , V = 4568(3) Å<sup>3</sup>, Z = 4,  $D_c = 1.858$  g cm<sup>-3</sup>,  $\mu$ (Mo-K $\alpha$ ) = 4.097 mm<sup>-1</sup>, F(000) = 2488.  $R_1$  and  $wR_2$  are 0.050 and 0.112, respectively, for 514 parameters and 10225 reflections [ $I > 2\sigma(I)$ ] (13265 unique). CCDC 601678. For crystallographic data in CIF or other electronic format see DOI: 10.1039/b603495k

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