Platinum phosphinothiolato hydride complexes: synthesis, structure and evaluation as tin-free hydroformylation catalysts†

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Ligand 2-diphenylphosphinothiophenol (Hsarp) reacted with Pt(PPh3)4 to yield trans-[PtH(sarp)(PPh3)]2, which undergoes fast exchange with free PPh3 on the NMR time scale and very slowly and reversibly formed some cis-[PtH(sarp)(PPh3)] over time in solution (11%, 24 h). Reaction of trans-[PtH(sarp)(PPh3)] with Hsarp in boiling toluene gave cis- and trans-[Pt(sarp)]2; the cis isomer being more stable. These complexes were characterized by 1H and 31P NMR and also analyzed by XRD in the case of trans-[PtH(sarp)(PPh3)], trans-[Pt(sarp)]2, and cis-[Pt(sarp)]. trans-[Pt(sarp)(PPh3)] was evaluated as a preformed, tin-free hydroformylation catalyst on styrene and found active at 100 °C, at pressures over 75 bar, yielding phenylpropanal (regioselectivities up to 83% in 2-phenylpropanal), with total conversions to aldehydes up to 100% at styrene/platinum ratios from 400/1 to 1000/1 and minimal hydrogenation products.

As part of a project on the synthesis of new amino- and phosphino-thiolates of group 10 metals, and on the analysis of the ligand-based stereoelectronic effects that are determinant in their properties and potential applications,10b,12 we now report on platinum 2-diphenylphosphinothiophenolato complexes, their synthesis and structural features and the first tests as homogeneous, tin halide free hydroformylation catalysts.13

Results and discussion

Synthetic aspects

Ligand Hsarp (2-diphenylphosphinothiophenol) reacted with Pt(PPh3)4 to yield modest amounts of 1 (30%) when used stoichiometrically (Scheme 1). When Hsarp was used in larger amounts the yield increased accordingly (up to 83%); this is interpreted as a need to displace three moles of PPh3 from the sphere of coordination of platinum(0) to maximize the amount of platinum hydride formed before the addition of Et2O to precipitate 1 and to keep PPh3 in solution (Scheme 1).

The yield of 1 is only dependent on the excess of Hsarp, and is independent of the temperature or the reaction time, within reasonable limits. Crude 1 contains basically trans-1 with small amounts of free PPh3 (ca. 0.5–1%). Crude 1 could be further purified by column chromatography on silica or by slow recrystallization to obtain pure trans-1. Solutions of trans-1 are stable to decomposition but evolve to an equilibrium mixture of trans-1 (89%) and cis-1 (11%) isomers, which is complete in 24 h.

Introduction

Phosphinothiolato ligands behave as soft–soft chelates with very different donors in terms of size, electronic properties and Bronsted basicity. This difference makes phosphinothiolato-species unique heterotopic ligands. Transition metal complexes of sulfur–phosphorus ligands have been studied for their potential applications.1 The areas of interest include carbonylation reactions,2 S-alkylation/S-dealkylation reactions,3 carbothiolations,4 and copolymerizations.5 These complexes are relevant in hydrodesulfurization (HDS),6 in the modeling of sulfur-containing metalloproteins7 and in imaging and radiotherapy.8

Olifin hydroformylation is a topic of continuing interest,9 including alternate metal catalysts (non-rhodium), which have been reviewed recently.10 The substitution of rhodium as the metal of choice for hydroformylation is not expected, but in some exceptional cases a third row metal (Ir) has been reported to outperform a second row metal (Rh) in catalytic carbonylation (see reports on the Cativa process).11

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The bischelate obtained in this way may contain small amounts of trans-[Pt(sarp)_2] which can be converted to cis-[Pt(sarp)_2] by further heating at 110 °C in toluene. The greater thermodynamic stability of the cis isomer is predicted by the greater trans influence of phosphorus over sulfur (or by the so-called antisymbiosis rule) while steric hindrance would favour the trans isomer. In the case of palladium, we have shown that trans and cis bischelates [Pd(sarp)_2] interconvert in solution and the equilibrium can be displaced towards the former or the latter by the use of polar or non-polar crystallization solvents respectively. In the case of platinum this procedure proved impractical, as trans-2 does not form in sufficient amounts in any solvent tested in reasonable reaction times. Bischelate cis-2 is also the identifiable platinum product of the thermal decomposition of 1 in boiling toluene.

The bulkier ligand, 2-diphenylphosphino-6-trimethylsilyllithiophenol (Hsarp), reacted in the same way as Hsarp to yield the corresponding hydride [PtH(sarp')(PPh_3)] (Scheme 3). For comparison purposes, the ligand EtSCH_2CH_2SH was also prepared and used in the synthesis of [Pt(1:1:1:1EtSCH_2CH_2S,S,S,PPh_3),3964] which was made from Pt(PPh_3)_4 in the same way as 1 and 3, but by using a stoichiometric amount of the ligand. At room temperature, fast exchange was not observed in the NMR of 4 and isomerization was not detected (Scheme 3).

The reaction of Hsarp' with Pt(PPh_3)_4 in boiling toluene gave the bischelates 5, with the difference compared to 2, that a longer time (48 h) was needed for the bulkier ligand and the product was a mixture of cis and trans isomers, containing a slight excess of the latter (Scheme 3). Clearly, in the case of sarp' the trans isomer becomes more stable for steric reasons.

NMR spectroscopy

The hydride region of the ^1H NMR spectra of crude 1 consisted of a doublet with ^195Pt satellites and the ^31P NMR showed just noise at room temperature. This was attributed to the exchange phenomena, so the NMR were taken at low temperature (down to -60 °C) but the slow exchange limit was not achieved in this way. Purification of 1 by chromatography or recrystallization gave samples free of PPh_3 which exhibited non-exchanging, narrow lined ^1H and ^31P NMR spectra. The obvious conclusion is that exchange takes place between 1 and free PPh_3 and this is a bimolecular reaction that cannot proceed in the absence of PPh_3. The long accumulation of ^31P NMR spectra on samples of crude 1 at room temperature showed the NMR of Fig. 1, which consists of the very broad signals of trans-1, which exchanges with free PPh_3, and very small amounts of cis-1 and cis-2 (narrow lines).

Relevant NMR data are collected in Tables 1 and 2. Hydride signals are well shifted to low frequencies into negative delta values. Hydride coupling to platinum (J_{PtH}) is typically very large and it is observed in all cases. Coupling to the sarp...
phosphorus ($^{31}$P$_{PM}$, $P_A$) is also observed in fast exchange and no exchange conditions, while coupling to the PPh$_3$ phosphorus ($^{31}$P$_{PB}$, $P_B$) is only observed in the absence of free PPh$_3$, that is, in the absence of exchange. In this case the spectra have been obtained from samples that contained minute amounts of free PPh$_3$ (see Tables 1 and 2). Complex 4 showed no such exchange with added PPh$_3$ at room temperature, so that, we conclude that the exchange is directly related to the trans effect of the phosphorus of the sarp ligand.

$^{31}$P NMR mirrors in most ways the results observed in $^1$H NMR. In the absence of free PPh$_3$, the resonances of 1 are narrow and all $\delta$ and $J$ can be assigned. In the presence of PPh$_3$ (the fast exchange limit) only the chelate ligand phosphorus is observed, including its coupling to platinum ($J_{PM}$, $P_A$) but not to PPh$_3$ phosphorus ($P_B$).

The effect of free PPh$_3$ on the NMR of trans-1 was better understood by recording the $^{31}$P NMR spectra of a sample of trans-1 after adding known amounts of free PPh$_3$ in an NMR titration-like experiment (Fig. 2). The addition of just 0.5 mol% PPh$_3$ to pure trans-1 caused immediate collapse of the $^{31}$P signals; at 5% the signal corresponding to $P_A$ started to rise (broad), at about 50% the signal of $P_A$ (with $^{31}$P satellites) was narrower and the signal of free PPh$_3$ started to rise from the noise (broad), at 300% the exchange was fast enough for the PPh$_3$ to be quite narrow, but clearly displaced down-field from the known chemical shift of PPh$_3$. To see cis-1 in the NMR spectrum, solutions of trans-1 were allowed to age. After 24 h equilibrium was reached at room temperature; the equilibrium mixture in dichloromethane solution contained both trans-1 (89%) and cis-1 (11%), but crystallization from CH$_2$Cl$_2$/ Et$_2$O yielded only trans-1. The thermodynamic stability of the trans-P$_2$P isomer is in agreement with the high trans influence of hydride, in contrast with the low trans influence of chloride which gives cis-P$_{PM}$-P[PtCl(dppet-$^2$P$_S$)PPh$_3$]$_2$.  

Coupling constants were useful in the characterization of these compounds. The $^{31}$P($^1$H) coupling constant in cis-1 is much larger (186 Hz) than any other $J_{PH}$ because this is the only case in which the hydride ligand is trans to phosphorus (Table 1). In the case of phosphorus to phosphorus, $^{3}$P$_{PH}$ is much larger in trans-1 (391 Hz) than in cis-1 (10.1 Hz). Coupling to platinum is strong both for $^1$H and $^{31}$P, but much larger $^{3}$P$_{PH}$ coupling constants are expected for cis-P$_2$P isomers than for trans-P$_2$P isomers, this criterion has been followed to assign the signals of trans-2 and cis-2 (Table 2), which was supported by the XRD studies (vide infra). However, coupling of the sarp phosphorus to platinum $^{3}$P$_{PH}$ is comparatively small in cis-1 (Table 2), this is most probably caused by the strong trans influence of the hydride ligand. There are numerous precursors of this effect: in cis-P[PtH$_2$($^2$PET$_3$)$_2$], $^{3}$P$_{PH}$ = 1984 Hz, while in trans-P[PtH$_2$($^2$PET$_3$)$_2$], $^{3}$P$_{PH}$ = 2764 Hz; and in cis-P[PtH$_2$(PMe$_3$)$_2$], $^{3}$P$_{PH}$ = 1875 Hz, while in trans-P[PtH$_2$(PMe$_3$)$_2$], $^{3}$P$_{PH}$ = 2394 Hz. cis-$^3$P$_{PH}$ Complexes cis-P([R$_2$P$_2$]), ($R$ = alkyl, $^1$H$_2$ aryly; $^{19}$L = PPh$_3$, PMe$_3$, PET$_3$) also feature comparatively small $^{3}$P$_{PH}$.

The behavior of 1 in solution is summarized in Scheme 4. The first reaction represents the fast exchange we observe in the NMR. Platinum(ii) will form a more stable trigonal bipyra-

![Fig. 1](image-url)
midal (tbp) intermediate with the bulky phosphorus ligands in equatorial (eq) positions and with the sarp chelate bonding in an axial-equatorial fashion. To form cis-1, the tbp intermediate must rearrange to a less favourable intermediate (in Scheme 4 this is depicted by L in an axial position), and this is presumably the reason why trans/cis isomerisation is such a slow process for 1.

In the exchange process of trans-1 we believe that sarp remains coordinated in the $S_{eq}$–$P_{eq}$ mode, supported by the fact that the chemical shift of $P_A$ is the same in the fast exchange limit and the no exchange (slow exchange) limit. The chemical shift of $^{31}P$ is strongly affected by the formation of cycles and any opening of the chelate should affect the $\delta_P$ of $P_A$. The coupling of $P_A$ to platinum is also retained in the fast exchange limit (Fig. 2).

**XRD crystal structures**

Slow crystallization of 1 gave yellow crystals of trans-1 exclusively; crystals of minor isomer cis-1 could not be obtained from normal laboratory solvents. Crystallization of 2 in CH$_2$Cl$_2$/Et$_2$O gave mainly chunky yellow crystals of cis-2-CH$_2$Cl$_2$, which is isostructural with the palladium complex cis [$Pd(sarp)_2$]-CH$_2$Cl$_2$.[16] However, careful inspection revealed the presence of a few smaller crystals of a more intense colour and an apparently different habit, which upon XRD analysis proved to be trans-2.

The structures exhibit slightly distorted square planar platinum(II) coordination. In the case of trans-1 (Fig. 3) it is worth noting the relatively long Pt–S distance (Table 3), which is ascribed to the strong trans influence of hydride. The chelate angle of sarp (Table 3) is always about 87°, as it is a rigid ligand. This angle makes sarp a well suited ligand both for cis coordination in sqp structures and for $S_{eq}$–$P_{eq}$ coordination in tbp intermediates, which are stabilized by the nature of sarp in terms of the bite angle and size of the coordinating groups (see above, Scheme 4).

**Table 3** Relevant distances (Å) and angles (°) for trans-1, trans-2 and cis-2, obtained in this work

|               | trans-1  | trans-2  | cis-2-CH$_2$Cl$_2$ |
|---------------|----------|----------|--------------------|
| Pt–S          | 2.357(3) | 2.315(2) | 2.313(5)           |
| Pt–P1         | 2.243(3) | 2.293(2) | 2.236(4)           |
| Pt–P2         | 2.278(3) | 2.263(4) | 2.249(4)           |
| S–Pt–P1       | 98.6(1)  | 96.9(7)  | 87.7(2)            |
| S–Pt–Pt–P2    | 171.6(1) | 107.5(2) | 84.7(2)            |
| P1–Pt–P2      | 104.6(3) | 107.5(2) | 172.0(2)           |
| P1–C1         | 109.1(3) |          | 171.7(2)           |

$^a$ Chelate angle. $^b$ Open angle, platinum is at the inversion center, $S'$ is related to S by $-x$, $-y$, $-z$. $^c$ Chelate angle, S1 and P1 belong to one ligand, and S2 and P2 belong to the other.

The Pt–P2 distance in trans-1, corresponding to PPh$_3$ is somewhat longer than the Pt–P1 distance corresponding to sarp (Fig. 3). The P–Pt–P angle is deformed ($ca.$ 172°) from linearity in trans-1 and cis-2, compared to symmetric trans-2 (Fig. 4, center of symmetry at Pt), owing to steric reasons. In trans-1, the hydride ligand has not been located but it must be placed trans-1.

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**Fig. 2** The addition of free PPh$_3$ to a sample of pure trans-1 (uppermost trace) caused a dramatic change in its $^{31}$P(II) NMR spectrum (101 MHz, CD$_2$Cl$_2$, rt), the added amounts are indicated by the traces in mol% of added PPh$_3$. A small amount of cis-[$Pt(sarp)_2$] was added for reference (*).
to sulfur (NMR, IR), thus the PPh₃ is somewhat displaced towards the smaller ligand. In the structures of 2 the sarp ligands are symmetrically disposed only in the trans isomer (Fig. 4), while in the cis isomer the P–Pt–P angle increases to 100.3(2)° and the S–Pt–S angle decreases to 84.7(2)° owing to the repulsion of the −PPh₂ groups. The crystal structure of cis-1 could not be determined by XRD, but considering the structures of trans-1, trans-2 and cis-2, together with the spectroscopic data, leaves little doubt about its structure.

**Catalytic hydroformylation**

Complex trans-1 exhibits desirable characteristics for a homogenous catalyst: it undergoes fast ligand exchange, it contains a hydride ligand susceptible to olefin insertion reactions and also triaryl phosphines which usually favor small molecule activation, and addition and elimination processes associated with homogeneous catalysis. However, ligand sarp is a sulfur donor and these ligands are often associated with catalyst deactivation rather than catalyst activation. The thiophilicity of late transition metals, coupled with the ability of thiolates to give one, two or even three electron pairs in terminal or bridged compounds respectively, may block catalytic activity in homogeneous systems, but this should not be the case here because of the presence of good competing ligands.

Nevertheless, trans-1 was tested as a hydroformylation catalyst on styrene in the absence of tin, chloride or any other extra ligands. The first results were disappointing: no activity was detected with syngas pressures below 75 bar. At 75 bar and higher pressures the results were positive (Table 4), using styrene/platinum ratios of 400/1 conversions were essentially complete in 24 h, with very low hydrogenation to ethylbenzene and a regioselectivity towards branched aldehyde of ca. 70%. Lowering the styrene/platinum ratio it can be seen that average TON numbers are about 400 and TOF numbers around 18 h⁻¹ (Table 4, entries 3 and 4). Selectivity increased somewhat upon increasing the hydrogen partial pressure (Table 4, entry 4). Complex cis-2 was identified in the reaction mixtures as one of the platinum products formed after the reaction was over (after all, it is the thermal decomposition product of trans-1). However, testing showed that cis-2 was not responsible for the previous results, as it turned out to perform very poorly, with residual activity and low selectivity (Table 4, entry 5). Instead of preformed trans-1, the mixture of Pt(PPh₃)₄ plus Hsarp was also tested with the above olefins, and found inactive; one reason for this could be the inherent presence of excess PPh₃.
Table 4  Hydroformylation of styrene with trans-1

| Catalyst | [S]/[Pt] | Pt \( t \) | \( \eta^2 \text{C}_8\text{H}_8 \) | \( \eta^2 \text{C}_8\text{H}_8 \) | TON | \( R^+ \) (%) | HP (%) |
|----------|---------|----------|----------------|----------------|-----|----------------|--------|
| trans-1  | 400     | 75       | 24             | 100            | 400 | 70             | 0.4    |
| trans-1  | 400     | 125      | 97             | 98             | 400 | 76             | 0.6    |
| trans-1  | 900     | 125      | 24             | 48             | 432 | 66             | 1.2    |
| trans-1  | 1000    | 75       | 24             | 5              | 20  | 58             | 0.2    |
| cis-2    | 400     | 75       | 24             | 5              | 20  | 58             | 0.2    |

* Conditions: 19 mmol styrene (S) in 20 mL toluene at 100 °C. *Total pressure (bar), H₂/CO = 1. *Reaction time. *Conversion of styrene to products. *Average turnover number. *Regioselectivity in iso-aldehyde. *Hydrogenation product (ethylbenzene). *In this experiment, the pressure ratio H₂/CO = 1.3.

We have drawn (Scheme 5) a working mechanism that could rationalize the activity of complex 1 (before more data available), which also considers the special characteristics of ligand sarp. The insertion of the olefin into the Pt-H bond gives rise to an alkyl complex, which can be linear or branched, thus determining the regioselectivity of the reaction. It should be noted that for the branched alkyl, \( \eta^2 \text{C}_8\text{H}_8 \) coordination is possible, thus favoring both activity towards styrene and regioselectivity towards a branched product. It is very difficult to ascertain if \( L = \text{PPH}_3 \) or CO at this point (besides, the presence of CO in the coordination sphere of platinum does not even seem necessary until later). Insertion of CO causes the formation of the acyl complex which must react with hydrogen to yield the aldehyde and to regenerate the starting platinum species. For the activation of hydrogen by transition metal complexes, the first identified mechanism was oxidative addition, which would yield a platinum(IV), d⁸, species (central path in Scheme 5, in blue). After the elimination of the aldehyde, the cycle would be complete. However, given the nature of sarp (containing a proton-basic moiety not present in other ligands) we suggest that a different form of H₂ activation can be considered (the lower path in Scheme 5, in green), with no need for oxidation of Pt(II) to Pt(IV) in a reducing medium. This would be a heterolytic cleavage across the Pt-S bond, or hydrogenolysis-type activation. A similar activation has been described for rhodium and iridium arylthiolates.14 Moreover, the activation of H₂ to Pt-H and H₂O⁺ (i.e. platinum mediated disproportionation of H₂ to H⁺ and H⁺) has been implicitly proposed in the context of the platinum catalyzed hydroformylation of styrene.23 In this mechanism the elimination of aldehyde gives a platinum(0) species that was already invoked in the synthetic part of this paper, and which we assume to readily react to form a platinum hydride and follow the cycle.

**Experimental**

Reactions were performed under nitrogen using standard techniques.24 The isolated products are fairly air stable, and hydride complexes were kept refrigerated. Solvents were dried, distilled under nitrogen, kept over molecular sieves and deaerated prior to use.25 Tetrakis(triphenylphosphine)platinum(0) was prepared by established procedures.26 The synthesis of the ligands Hsarp and Hsarp’ have been described.27 NMR data are given on the \( \delta \) scale and referenced in the usual manner. Catalysis experiments were performed in a stainless steel Autoclave Engineers EZE seal reactor of a standard vace (ca. 80 mL) and cover design, mechanical stirring (magnetic drive), heating and temperature control through a thermocouple, and sample port. GC analyses were performed using capillary columns HP-5, 0.25 mm diameter × 30 m length, using FID and MS detectors. Elemental analyses (C, H and S) were performed with a Carlo Erba CHNS EA-1108. The platinum content was determined by repeated cycles of mineralization of a weighed sample with conc. HNO₃ in a heated crucible (fume hood), followed by ignition in an electric furnace at temperatures over 900 °C (open crucible, 24 h), this gave platinum metal that was weighed as such.

**X-ray diffraction structure determination**

Data were collected using Mo Kα radiation. For compounds trans-1 and cis-2-CH₂Cl₂ an Enraf-Nonius CAD4 diffractometer \( (2θ_{\text{max}} = 50^\circ) \) was used; for compound trans-2, a Kuma KM4CCD \( (2θ_{\text{max}} = 56^\circ) \) was used. Empirical absorption corrections were applied. Structures were solved by direct methods (SHELXS-86) and refined by full-matrix least-squares methods on \( F^2 \) for all reflections (SHELXL-97). Non-hydrogen atoms were refined anisotropically. The hydride atoms were placed in calculated positions with isotropic displacement parameters fixed at 1.2 times the \( U_{eq} \) of the corresponding carbon atoms. In trans-1, the hydrogen bonded to the platinum atom was not localized in the difference Fourier map, and it has not been included in the refinement.
Crystal data and further refinement details are presented as ESL.† Compound cis-2-CH2Cl2 is a solvate and some decay (caused by solvent loss) was observed during data collection; this may be the reason why in this structure high residual peaks (5.43 and –4.43 e Å−3) were found in the final difference density map, near to the Pt atom (1.05 Å and 1.12 Å respectively).

Preparation of trans-Pt[PPh3][sarp-κ2-P,S][PPPh3] (trans-1). A solution of Pt[PPh3]4 (500 mg, 0.40 mmol) was dissolved in dichloromethane (ca. 30 mL) to give a clear yellow solution, which was cooled down to 0 °C. Upon slow addition of Hsarp (2-diphenylphosphinothiophenol, 235 mg, 0.80 mmol dissolved in 5 mL of ethyl ether) to the ice-cold Pt[PPh3]4 solution, only partial loss of color was observed. The pale yellow solution was allowed to react for 10 min at room temperature and then its volume was reduced to ca. 1 mL under vacuum. Addition of ethyl ether (3 mL) and cooling to 0 °C caused the precipitation of a yellow microcrystalline solid. The solution was kept cold for 1 h to allow for the full precipitation of the product. The solid was collected by filtration, washed with ether (3 × 5 mL), and dried under vacuum. Yield: 248 mg (83%) of crude 1, a microcrystalline and yellow in color. Data for crude 1: 1H NMR (250 MHz, CDCl3, 298 K): δ = –7.58 ppm (s, with 195Pt satellites, 1H, H–Pt, JHPt = 1023 Hz); 6.8–7.4 ppm (m, 29H arom.), unchanged down to 213 °C. This product was qualified as “crude” because, as prepared, it contained some PPh3 (from 0.5 to 1% depending on the sample) and some [Pt(sarp)3] (from 0.1 to 1%). Owing to the content in PPh3, the NMR spectra of crude 1 were always broad (195Pt) or consistent with the fast-exchange limit (1H). High purity trans-1 was obtained by column chromatography (over silica 60 A, 3 cm diameter, 40 cm length, diethyl ether/dichloromethane 90/10 eluent) or by slow crystallization. Procedure for crystallization: crude 1 (300 mg) was dissolved in dichloromethane (60 mL), the clear solution was filtered through a short column of Celite (1 cm diameter, 3 cm length) and poured into a clean, scratch-free flatbed crystallizer. The yellow solution was layered with ethyl ether (10 mL) and allowed to rest after being covered with a watch glass (under air). After 48 h, the loss of volume caused by evaporation was made up with fresh ether (ca. 30 mL). After a further 72 h the solvent had evaporated completely. The residue consisted of large (ca. 1/16–1/32) yellow pseudo-cubes of pure trans-1, a very small amount of small prismatic crystals of [Pt(sarp)3], over a small amount of a fine white powder (PPh3 and OPPh3), the recovery of pure trans-1 was above 90%. Anal. Found: C, 57.01; H, 3.93; S, 4.42; Pt, 26.27. Calc. C, 57.52; H, 4.02; S, 4.27; Pt, 25.95. trans-1 was the only isomer present in the solid state or in freshly prepared solutions, cis-1 arose only with time and could not be isolated. The trans/cis equilibrium was reached in 24 h, with a trans/cis ratio of 89/11 in CD2Cl2, as measured by 1H NMR. Data for pure 1: 1H NMR (250 MHz, CDCl3, 298 K) trans-Pt,P-isomer (major): δ = –7.61 ppm (d, with 195Pt satellites, 1H, H–Pt, JHPt = 1151 Hz, JPPt = 340 Hz); 6.8–7.4 ppm (m, 29H arom.). cis-Pt,P isomer (minor): δ = –4.77 ppm (dd, with 195Pt satellites, H–Pt, JHPt = 950 Hz, JPPt = 186 Hz, JPP = 10 Hz); 6.8–7.4 ppm (m, 29H arom.). 31P{1H} NMR (100 MHz, CDCl3, 298 K) trans-Pt,P isomer (major): δ = 31.9 (d, with 195Pt satellites, Pt–PPh3, JPPt = 2915.2 Hz, JPP = 391.2 Hz); 50.47 (d, with 195Pt satellites, Pt–PPPh3Ar, JPPt = 2818.6 Hz, JPP = 391.2 Hz), cis-Pt,P complex (minor): δ = 24.3 (d, with 195Pt satellites, Pt–PPh3, JPPt = 3187 Hz, JPP = 10.1 Hz); 56.6 (d, with 195Pt satellites, Pt–PPPh3Ar, JPPt = 1922 Hz, JPP = 10.1 Hz). IR (KBr) ν(Pt–H): 2100 cm−1 (st). It was seen that the intensity of the (Pt–H) stretching, relative to the more intense ligand-originated vibration at 1569 cm−1, is a good qualitative measure of the contamination by PPh3 and [Pt(sarp)3], for pure 1 the intensity ratio was ca. 0.87 on the transmittance scale.

Preparation of [PtH(sarp-κ2-P,S)][PPPh3] (3). A solution of Pt[PPh3]4 (300 mg, 0.24 mmol) was dissolved in dichloromethane (ca. 20 mL) to give a clear yellow solution, which was cooled down to 0 °C. Upon the slow addition of Hsarp (134 mg, 0.36 mmol dissolved in ca. 10 mL of ethyl ether) to the ice-cold Pt[PPh3]4 solution, only partial loss of color was observed. The pale yellow solution was allowed to react for 20 min. at room temperature. The solvents were removed under vacuum and the resulting oily residue was stirred with 2 mL of cold ethyl ether (ice bath). This resulted in the formation of a pale yellow precipitate which was isolated by filtration and washed with a small amount (ca. 1 mL) of diethyl ether, and dried under vacuum. Yield: 120 mg (60%) of crude 3, yellow in color, complex 3 is much more soluble in common organic solvents than 1. Complex 3 thus obtained was contaminated with trace amounts of PPh3. Anal. Found: C, 56.49; H, 4.93; S, 3.53; Pt, 23.37. Calc. C, 56.85; H, 4.65; S, 3.89; Pt, 23.68. 1H NMR (250 MHz, CDCl3, 298 K): δ = –7.08 ppm (d, with 195Pt satellites, 1H, H–Pt, JHPt = 1151 Hz, JPPt = 6.6 Hz); 0.03 (s, 9H, Si(CH3)3); 7.3 ppm (m, 28H, arom.). 31P{1H} NMR (101.3 MHz, CDCl3, 298 K): δ = 48.2 ppm (s, with 195Pt satellites, sarp‘, JPPt = 2990 Hz). IR (KBr) ν(Pt–H): 2068 cm−1 (m).

Preparation of EtSCH2CH2SH, 2-ethylthioethanethiol. A 100 mL side-arm flask was loaded with ethylene sulfide (C2H4S, 3.0 g, 50 mmol) dissolved in 60 mL of dry methanol and placed in an ice bath. To this, excess solution of potassium ethane thiolate (prepared by mixing 9.3 g of ethanethiol, 150 mmol, and 9.0 g of 85% KOH, 137 mmol, in 60 mL of dry methanol) was slowly added. The mixture was allowed to warm up to room temperature and stirred for 1 h. A solution of ammonium chloride (5.0 g, 93 mmol, in 40 mL of water) was added to quench the reaction, followed by drops of conc. HCl until acid to litmus. The precipitation of potassium chloride was observed at this point. The organic solvents were removed under vacuum. The KCl precipitate re-dissolved and oil separated, this oil was extracted with dichloromethane (4 × 10 mL). The combined organic phases were dried with anhydrous magnesium sulfate. The solvent was removed at reduced pressure and the resulting oil was subjected to microdistillation at 3 mm Hg, the fraction was collected at 88–92 °C. Yield: 3.78 g (62%) of a colorless, malodorous, viscous liquid, 98%+ by GC.
Preparation of [Pt(sarp′-κ²P, S)S][Pt(PPh₃)₄] (5), bis[(2-diphenylphosphino)-6-trimethylsilylthiophenolato-κ²P, S]triphenyolphosphineplatinum(ii). The platinum(0) complex Pt[PPh₃]₄ (400 mg, 0.30 mmol) was dissolved in toluene (ca. 20 mL) to give a yellow solution and Hsarp′ (234 mg, 0.60 mmol, dissolved in 10 mL of toluene) was added. The pale yellow solution was heated to the reflux temperature (ca. 110 °C) and allowed to react until the Pt-H stretching absorption (2068 cm⁻¹) was not detectable in the IR and the hydride signal was not seen in the H1 NMR, which took a minimum of 28 h. The solvent was removed under vacuum and the oily residue was stirred with 20 mL of ethyl ether at 0 °C (ice bath). The solid that formed was isolated by filtration, washed with cold ethyl ether (2 × 4 mL) and vacuum dried. Yield: 162 mg (55%) of 5, yellow in color. This material was found to be a mixture of cis-PP and trans-PP isomers (ca. 45/55). Anal. Found: C, 54.56; H, 4.81; S, 6.63. Calc. C, 54.47; H, 4.79; S, 6.92. Data for cis-5: H1 NMR (250 MHz, CDCl₃, 298 K) δ 0.24 [s, 9H, Si(CH₃)₃]; 6.6–7.7 (m, 13H, arom). 31P{H} NMR (101.3 MHz, CDCl₃, 298 K) δ 41.90 [s, with 195Pt satellites, JPPt = 2867 Hz]. Data for trans-5: H1 NMR (250 MHz, CDCl₃, 298 K) δ 0.39 [s, 9H, Si(CH₃)₃]; 6.6–7.7 (m, 13H, arom). 31P{H} NMR (101.3 MHz, CDCl₃, 298 K) δ 48.90 [s, with 195Pt satellites, JPPt = 2629 Hz].

Hydroformylation experiments

The reactor was conditioned by vacuum and syngas replenishment cycles. The reaction mixture was injected into the reactor through an inlet valve (followed by some fresh solvent in order to clean the valve) at room temperature and the pressure was set to about 30 bar. The temperature was raised to working temperature using the temperature regulator and then the pressure was adjusted to working pressure through the gas supply system. The mechanical stirring was initiated (400 s⁻¹) and this was considered time zero. The reaction mixture consisted of, typically, the alkene (in the case of styrene, 2.00 g, 0.0192 mol), complex 1 (0.0364 g, 4.75 × 10⁻³ mol; this value depends on the entry in Table 4) and toluene (20 mL total solvent). The organic products were analyzed by GC and NMR. Blank experiments under the conditions of Table 4 run without complex 1 added or with PPh₃ added instead of complex 1 (0.02–0.20 g) gave no conversion. Out of the reactor all solutions were homogeneous and yellow in color with no black platinum or other precipitates. After some time, the formation of some crystals was generally observed; they were identified as cis-2.

Conclusions

Ligands Hsarp and Hsarp′ readily reacted with Pt(PPh₃)₄ to give hydride complexes trans-[PTH(sarp)[PPh₃]]). The yield of 1 depends mainly on the stoichiometry employed, an excess of Hsarp being necessary for good yields. This does not seem to be a kinetic effect as yields are not influenced by the reaction time or temperature. This dependence is ascribed to an equilibrium in the chelate-assisted oxidative addition of ArS–H to Pt.
The crude also contains very small amounts of cis-[Pt- Hsarp(PPh₃)] and cis-[Pt(sarp)] and PPh₃. Under these conditions the trans-P₃P hydride undergoes a very fast exchange of coordinated PPh₃ with free PPh₃ and has very broad signals in the NMR, while the cis-P₃P hydride exhibits narrow signals at the same temperature, which means that it does not undergo fast PPh₃ exchange. These hydrides are remarkably stable towards oxygen and moisture, which allowed purification by column chromatography. Pure trans-1 does not show any broadness in the NMR; the addition of free PPh₃ shows that the exchange of PPh₃ is bimolecular. Only trans-1 is isolated as solid crystals upon recrystallisation, fresh solutions show the signals of trans-1 only, but with time the cis isomer becomes apparent. Equilibrium was reached in 24 h and the solution contained 89% trans-1 and 11% cis-1. Fast exchange and trans/cis equilibrium was not observed in other non-sarp containing complexes such as 4. The reaction of Pt[PPh₃]₄ or 1 with excess Hsarp (at 110 °C, 24 h) yields bischelates [Pt(sarp)]₂, the cis isomer being more stable thermodynamically. In ³¹P NMR, cis-2 shows a signal at δ 42.80 with ¹⁹⁵Pt satellites (¹J_PtP = 2912 Hz) and trans-2 shows a signal at δ 50.80 (¹J_PtP = 2645 Hz) which is in agreement with XRD results and with the general observation that direct Pt–P coupling constants are larger for cis-P₃P isomers.

Hydride complex trans-1 was found worth evaluating as a hydroformylation catalyst but was found inactive at low pressures. At higher pressures, however, it was found active on styrene. The activity of trans-1 is comparable to platinum/diphosphine/SnCl₃ systems published recently, but its selectivity is different. This difference in selectivity would support trans-1 as a new catalytic system.

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