Linear Triphosphines as Ligands for Metal Complexes Immobilization in Ionic Liquids: Palladium-Catalyzed Methoxylation of Alkynes

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Abstract: Several novel palladium triphosphine complexes have been synthesized and tested as recyclable catalysts for the methoxylation of alkynes into acetals in ionic liquids. A complete conversion of phenylacetylene was achieved with only 0.2% of [Pd(Triphos)NCMe][(PF$_6$)$_2$] in a methanol/[BMIM][BF$_4$] mixture. We discovered that the addition of an ionic liquid to methanol allowed not only to increase the activity of the palladium catalyst but also to provide a recyclable catalyst which can be reused several times with a weaker drop of activity. To complete these catalytic studies, we describe the synthesis of the first poor $\pi$-electron-donating/strong $\pi$-electron-acceptor linear Triphosphine which, after palladium coordination, led to a better selectivity compared to its Triphos analogue. The performances of recovered ionic liquid reaction mixtures show for the first time that P-tridentate ligands efficiently immobilize palladium catalysts and lead to selective catalytic systems benign for environment.

Keywords: Polyphosphines, electron-poor ligands, palladium, ionic liquids, recyclable catalyst.

INTRODUCTION

The construction of C–O bonds by addition of alcohols to a C–C triple bond represents today one of the most useful methods to functionalize internal and terminal alkynes. As a consequence, a variety of catalysts have been studied for this reaction which leads essentially to acetals.

$$\text{R} = \text{C} = \text{C} \cdot \text{R'} + \text{R''OH} \rightarrow \begin{cases} \text{acetal} \\ \text{enol ether (traces)} \end{cases}$$


For instance, In(III), Zr(IV), Au(I) and Ir(I) complexes were found to be active in selective formation of spiroacetals from alkynes, [1-3] which are motifs or substructures of numerous fragrances, [4-6] insect pheromones, [7] polyketides natural products, [8,9] insecticidal drugs, [10] or biodegradable synthetic biomaterials. [11] Other reports indicate that the addition of methanol to non-activated alkynes can be promoted by rhodium, [12] platinum dihalides with sodium sulfate as co-catalyst, [13] phosphite-iridium complexes eventually combined with a Lewis acid, [14,15] or a phosphine-cationic gold complex in situ generated in the presence of a strong Brønsted acid. [16] We have focused our investigation on the activity of palladium complexes since, in addition to copper complexes, [17] they specifically catalyze one-pot three-component reactions using alkynes, alcohols and aryl iodides as starting material. [18] The control of metal-phosphine leaching in the mobile phase still remains in all cases a major problem in the alkyne methoxylation catalysis. To tackle this issue, two approaches can then be envisaged: i) either the use of imidazolium monophosphate salts as immobilizing ligands, similarly to studies devoted to palladium alkynylation, platinum hydrogenation and rhodium hydroformylation catalysis in ionic liquids, [19, 20] or ii) the use of flexible polyphosphine ligands to stabilize ionic palladium complexes. [21] We opted for the use of polyphosphine ligands with regards to their robustness, and their good metal-immobilizing abilities by coordination to cationic complexes in ionic liquids. They are also susceptible to functionalization by a dangling imidazolium (or imidazolidinium) ionic fragment to enhance their anchoring capacity into highly polar media. [22,23]

In several cases, the replacement of triphenylphosphine ligands by trifurylphosphine has led to enhanced activity in metal catalysis. For instance, better performances were found in palladium-catalysed aryl alkynylation, in platinum-catalysed hydrogenation of chloronitrobenzene, [19] for Negishi cross-coupling, [24] and for rhodium catalyzed reactions. [25] We thus postulated that better catalytic performances may also be achieved from an increased Lewis acidic character of the metal centre through the introduction of electron-withdrawing groups in a tridentate phosphine...
ligand. In the case of alkynes methoxylation, the nucleophilic addition of methanol to the palladium-coordinated alkyne may then be efficiently promoted. We focused our synthetic efforts on the preparation of the first example of a poor $\sigma$-electron-donating linear triphosphine. The resulting ligands and their corresponding palladium-complexes were fully characterized and tested in the methoxylation of phenylacetylene and 1-octyne in organic or ionic solvents. Recovery and recycling runs were successfully conducted in the ionic liquid [BMIM][BF$_4$].

RESULTS AND DISCUSSION

A. Synthesis of Triphosfurane Ligand

Our synthetic approach to get the modified triphosphine 2, named Triphosfurane, was based on the reaction of F$u'_{2}$PBr [bis(5-methyl-2-furyl)bromophosphine], prepared according to the literature, [26,27] with vinyl magnesium bromide via a Grignard reaction, similarly to its analogous vinyl diphenylphosphine Ph$_2$PCH=CH$_2$ [28] and followed by a radical-induced double hydrophosphorylation of phenylphosphine (See Scheme 1).

Under such conditions, the vinyl difurylphosphine 1 was obtained selectively as shown by the presence in the $^{31}$P NMR spectrum of a single signal at −57 ppm. However, when the conditions described for the preparation of Triphosline [23] were applied to this vinyl difurylphosphine 1 in the presence of phenylphosphine and AIBN, several signals were observed in the corresponding spectrum (Fig 1).

![Scheme 1](image)

**Scheme.** (1).

The signals which appeared as a doublet and a triplet at −59 and at −17 ppm with a typical [22, 23] $^{3}J_{PP}$ coupling constant of 32 Hz were assigned to the expected Triphosfurane ligand 2a [PhP(CH$_2$CH$_2$PF$u'_{2}$)$_2$] respectively for the external and internal phosphorus atoms. The other signals appearing as a couple of doublet at −35, a triplet at −37 and a

![Fig. 1](image)

**Fig. (1).** Simulated and experimental $^{31}$P{${}^1$H} NMR spectra of Triphosfuranes 2.
doublet at \(-59\) ppm with same intensities were assigned to an AMX pattern corresponding to the diastereoisomers [Fu\(_2\)PCH\(_2\)CH\(_2\)PPhCH(CH\(_3\))PFu\(_2\)] \(2b\) and \(b'\), with an expected \(2b:2b'\) ratio equal to one, see Fig. (1). The \(2a\) and \(2b,b'\) isomers were present in similar amount (Scheme 1). The structure proposed for the Triphosfurane \(2b,b'\) was further proved by an additional electrospray analysis of the mixture which shows only one molecular peak for both Triphosfurane ligands (see experimental section for details). It is noteworthy that this kind of isomer \(2b,b'\) was never observed with the phenyl analogue. To explain the formation of this compound, we suggest that during the first addition reaction of vinyldifurylphosphine to the phenylphosphine radical \([\text{PhPH}]^*\) the presence of two electron-withdrawing furyl groups either renders the internal vinyl carbon more reactive than the external one, or destabilizes the secondary adduct radical \([\text{Fu}'_2\text{PCH}^*\text{-CH}_2\text{PHPh}]^*\). The regioselectivity of the second addition of vinyldifurylphosphine then follows a classical anti-Markovnikov addition which decreases the steric effects generated by the different bulky phosphorus fragments.

This poor regioselectivity during the formation of the expected Triphosfurane \(2a\) was improved by changing the reaction conditions. Consequently, \(\text{tBuOK}\) was tested as catalyst instead of AIBN, but the \(2a:2b, b'\) ratio remained unchanged. A better regioselectivity was observed when a photochemical irradiation was used to activate AIBN instead of heating. Regioselectivity became excellent by cooling of the reaction medium to 5 °C. The undesirable kinetic products Triphosfuranes \(2b,b'\) were then found to be less than 10%. A further crystallization allowed isolating pure Triphosfurane \(2a\) as colorless crystals suitable for X-ray structure analysis, as displayed in Fig. (2).

In the solid state molecular structure the backbone which incorporates the three phosphorus atoms and the methylene groups forms a semicircle arch, with the lone electron-pairs of each phosphorus atom alternatively pointing in opposite direction. The oxygen atoms from furyl fragments are found to be either on synperiplanar (O1 and O2, Fig. 2) or anti-periplanar position (O3 and O4, Fig. 2). Such arrangement induces a different chemical environment for methylene groups which is not detected at room temperature by \(^1\text{H}\) and \(^{13}\text{C}\) solution NMR spectroscopy in CDCl\(_3\), certainly due to free rotation of the P–CH\(_2\) bonds. This contrasts with the different chemical shifts observed for the methylene groups upon palladium coordination. The sum of P–C angles being identical for PFu'\(_2\)CH\(_2\) and PhP(\(\text{CH}_2\))\(_2\) moities means that the different phosphorus groups adopt the same pyramidal geometry (see Table 1 and 2). It is also interesting to compare the different P–C bond lengths. Indeed, they decrease continuously from P–C\(_{\text{alkyl}}\) (1.853 to 1.847 Å), P–C\(_{\text{Ph}}\) (1.835 Å) to P–C\(_{\text{Fu'}}\) bonds (1.811 to 1.804 Å) exactly like in the ethylbis(2-furyl)phenylphosphonium derivatives. [29] This observation is surprising because the lower electron density of the corresponding P–C bond due to the replacement of a phenyl group by an electron withdrawing substituent like furyl should increase the P–C bonds length. On the other hand, theoretical calculations performed on related phospho-
Table 1. Selected Bond Lengths [Å] and Bond Angles [°] for Compound 2a

<table>
<thead>
<tr>
<th>Bond:</th>
<th>Angle:</th>
</tr>
</thead>
<tbody>
<tr>
<td>For Ph2P fragment</td>
<td></td>
</tr>
<tr>
<td>P(1)-C(1)Ph</td>
<td>1.8357(14)</td>
</tr>
<tr>
<td>C(1)-P(1)-C(19)</td>
<td>100.00(6)</td>
</tr>
<tr>
<td>P(1)-C(7)alkyl</td>
<td>1.8532(14)</td>
</tr>
<tr>
<td>C(1)-P(1)-C(7)</td>
<td>101.63(6)</td>
</tr>
<tr>
<td>P(1)-C(19)alkyl</td>
<td>1.8474(14)</td>
</tr>
<tr>
<td>C(19)-P(1)-C(7)</td>
<td>101.87(6)</td>
</tr>
<tr>
<td>Sum = 303.50(10)</td>
<td></td>
</tr>
<tr>
<td>For Fu'2P fragments</td>
<td></td>
</tr>
<tr>
<td>P(2)-C(8)alkyl</td>
<td>1.8486(14)</td>
</tr>
<tr>
<td>C(14)-P(2)-C(9)</td>
<td>101.91(7)</td>
</tr>
<tr>
<td>P(2)-C(9)Fu'</td>
<td>1.8080(15)</td>
</tr>
<tr>
<td>C(14)-P(2)-C(8)</td>
<td>101.75(6)</td>
</tr>
<tr>
<td>P(2)-C(14)Fu'</td>
<td>1.8040(14)</td>
</tr>
<tr>
<td>C(9)-P(2)-C(8)</td>
<td>97.45(6)</td>
</tr>
<tr>
<td>Sum = 301.11(11)</td>
<td></td>
</tr>
<tr>
<td>P(3)-C(20)alkyl</td>
<td>1.8474(14)</td>
</tr>
<tr>
<td>C(26)-P(3)-C(21)</td>
<td>101.68(7)</td>
</tr>
<tr>
<td>P(3)-C(21)Fu'</td>
<td>1.8114(15)</td>
</tr>
<tr>
<td>C(26)-P(3)-C(20)</td>
<td>100.34(7)</td>
</tr>
<tr>
<td>P(3)-C(26)Fu'</td>
<td>1.8052(15)</td>
</tr>
<tr>
<td>C(21)-P(3)-C(20)</td>
<td>99.97(6)</td>
</tr>
<tr>
<td>Sum = 301.99(12)</td>
<td></td>
</tr>
</tbody>
</table>

Nium cations rationalize this unusual short P-C bond length by a better orbital overlap in the case of furyl groups. [29]

B. Synthesis of Palladium-triphosphine Pre-catalysts

The well-known Triphos-palladium complex 4 was prepared by reaction between [PdCl2(NCPh)2] and the Triphos [[Ph2PCH2CH2)2PPh] ligand 3, according to literature reports. [24] The Triphosline-palladium complex 6 and 7 were similarly obtained starting from the lysidinyl- (or imidazolin-1-yl-) triphosphine 5, and the Triphosfurane ligand 2a respectively (Scheme 2). [23]

![Scheme 2](image)

Scheme 2.

The corresponding halide free complex 8 (see Scheme 3) was obtained from 4 by a further reaction with silver triflate. [30] In order to examine the effect of the counter-anion on the catalytic properties, we have also synthesized a new halide free Triphos-palladium complex 9 by a one-pot synthesis using TIPF6 as an halide abstracting reagent. The tridentate coordination mode in complex 9 was indicated by the existence in the 31P NMR spectrum of a triplet at 110.3 ppm and a doublet at 45.7 ppm. These chemical shifts correspond respectively to internal and external phosphorus groups, consistently with the analogous palladium complex bearing BF4- counter-anion. [31]

![Scheme 3](image)

Scheme 3.

In contrast to Triphos-palladium complex 8, the addition of AgOTf to 6 in similar conditions led in the 31P NMR spectrum to several multiplets around 97 ppm and between 56 and 50 ppm, which likely result from the decomposition of complex 10, see Scheme 4.

![Scheme 4](image)

Scheme 4.

Nonetheless, when TIPF6 in CH3CN/CH2Cl2 was used instead of the above mentioned silver salt, a single complex was formed and identified as complex 11 (see Scheme 4) with 31P NMR signals at 107.9, 30.7 and –144.3 ppm in a 1:2:2 ratio. Complex 11 could be very useful as precursor for
In contrast to complexes 4 and 6, complex 7 in the presence of TIPF₆ in a CH₂Cl₂/CH₃CN solvent mixture was partially converted to the corresponding halide free complex 12, which is characterized in its ³¹P NMR spectrum by the presence of a new doublet at 11.2 ppm and triplet at 117.0 ppm. However, in pure acetonitrile the reaction becomes almost complete.

Since the TICl formed during this halide abstraction reaction is slightly soluble by addition of dichloromethane to the reaction mixture, we suggest that the presence of the four furyl electron withdrawing groups in complex 12 significantly increase the metal electrophilic character which then reacts with traces of chloride anion. From a catalytic perspective the activity of complex 12 should not be altered by small amount of complex 7 since this one was found to be totally inactive in alkynes methoxylation.

Equation. (2).

C. Catalytic Results in the Intermolecular Methoxylation of Alkynes

Since the above mentioned cationic Triphos-palladium complex 8 effectively catalyzes the conversion of phenylacetylene into dimethylacetal derivatives in methanol solvent, we thus started our catalytic studies using complex 8 under analogous conditions but in a 1:1 mixture of CH₃OH/[BMIM][BF₄] in view of performing catalyst recycling. ¹H NMR monitoring of the reaction in CD₃OD clearly showed the beneficial effect of the ionic liquid on the activity and chemoselectivity of alkoxylation reaction. Indeed, in this solvent mixture, a complete conversion of starting material is obtained after six days corresponding to a turnover number (TON) of 424. Five percent of acetophenone were detected in the products formed as a consequence of acetal hydrolysis. Conversely, seven days of reaction were required in pure methanol at 25 °C and the amount of acetophenone formed was doubled. In the next reactions the catalytic runs were thus stopped before complete conversion was achieved to better identify weak variations of activity in recycling experiments. For instance, in a first run after five days, 81% conversion of phenylacetylene into di methylacetal derivatives in methanol solvent, effectively catalyzes the conversion of phenylacetylene into dimethylacetal derivatives in methanol solvent. In the next reactions the catalytic runs were thus stopped before complete conversion was achieved to better identify weak variations of activity in recycling experiments. For instance, in a first run after five days, 81% conversion of starting material was obtained with a TON = 322. Catalytic results of methoxylation of phenylacetylene with the above starting material was obtained with a TON = 322. Catalytic results of methoxylation of phenylacetylene with the above complex 8 are summarized in Fig. 3. These results evidence an unexpected strong influence of the counteranion on the activity and stability of the palladium catalysts.

Complex 8 exhibits a slightly better activity than 9 in the first catalytic run (81% versus 75%). However, this activity dramatically decreases of practically 50% after each run. Although precursor 9 was less active than 8, the lifetime of its related catalytic species is greater and allows doubling the total turnover number compared to 8, with a value of about 

Phenylacetylene methoxylation with recovered Pd catalysts in [BMIM][BF₄]. Conversion of phenylacetylene in mol% (● black cylinder in the background), 1-octyne (◯ grained cylinder), 1-phenyl-1-propyne (■ grey square in the foreground) and performances of the recovered ionic liquid phase, determined by proton NMR in CDCl₃, with complexes 8, 9 and 11. Conditions: alkyne (4.55 mmol), pre-catalyst (0.2 mol %), [BMIM][BF₄] (0.5 ml), CH₃OH (0.5 ml), 5 days, 25 °C.

1.25x10³ mole of acetal per mole of 9 after five recycling runs. This clearly demonstrates the potential of catalytic systems incorporating 9 towards continuous-flow catalytic processes. Conversely, no activity was observed from Triphosline-palladium pre-catalyst 11. This could be a consequence of the lower electrophilic character of metal in complex 11 compared to 9 which is induced by the electron-donating dangling imidazole fragment and/or by its eventual coordination to the metallic species.

Although the drop of activity observed from recycling experiments with 9 is limited to seven percent after each catalytic run this might still remain too important for sustainable processes applications. We have subsequently performed additional experiments to determine the origin of this catalyst deactivation. While no ICP MS experiments were performed so far on the organic extracting solvent, the ligand-metal leaching is however excluded because it was not detected by proton NMR spectroscopy in extracting heptane phases. We then anticipated a slow decomposition of active species in the ionic liquid even at room temperature. So, the ionic liquid containing pre-catalyst 9 recovered after five successive catalytic experiments was left aside during five weeks at room temperature. After this inactivity period, phenylacetylene and methanol were then introduced and conversions of ca. 20 and 10 % were respectively found after the first and second recycling. These values are very close to those extrapolated at the 10th and 11th catalytic runs, if the recovered catalyst had been reused constantly during these five weeks. This experiment proves unambiguously that the drop of activity is consistent with a very slow deactivation process over time. Fig. (3) seems to indicate that catalytic systems based on complexes 8 and 9 deactivate in two different manners. In the case of complex 8, the decomposition rate is consistent with a first-order kinetic in catalyst concentration since the logarithm of the conversion changes in a linear manner with time. Conversely, this deactivation appears to be zero order in catalyst concentration in the case of complex 9, as shown in Fig. (3) from the linear decrease of conversion. The deactivation in this later case could be then correlated to physical or chemical properties of the catalytic medium which may evolve with time, either by slow introduction of air or moisture, or by the presence of residual methanol. In the case of complex 8, a slow reaction between traces of zerovalent palladium with the ionic liquid (palladium-carbene formation) may also indirectly decrease the catalyst concentration. Additional investigations are in progress to find out the origin of these deactivation processes.

When the electron-rich unactivated alkyne 1-octyne was used instead of phenylacetylene, in the presence of palladium pre-catalysts 8, 9 and 11, we observed that only complex 8 effectively promotes the conversion of 1-octyne into the corresponding acetal with 66% yield in the first run. However, a rapid drop of conversion was observed, as previously obtained with phenylacetylene as substrate, see Fig. (3). The result concerning pre-catalyst 9 was surprising since it was active in methoxylation of phenylacetylene. We also noted that all the palladium pre-catalysts of our study remained inactive towards 1,2-disubstituted alkynes such as 1-phenyl-1-propyne.
To find out the reasons of these unexpected results, we started to investigate the mechanism of this reaction. The above-reported observations could be hardly correlated with the electrophilicity of the palladium metal centre which depends on the phosphine and counter-anion nature. To the best of our knowledge, mainly two different mechanisms have been proposed and discussed for this C–O bond formation. The first one is based on the addition of oxygenated nucleophiles, such as water or alcohols, to the coordinated alkyne inside [16] or outside [12] the metal coordination sphere. In our study, the fact that the rate of methanol addition onto the coordinated alkyne is lower when the phenyl group in Triphos is replaced by an electron-donating alkyl fragment in Triphosline is consistent with this mechanism. This assumption is confirmed by the higher conversion obtained with pre-catalyst 8 in the case of electron-poor alkenes such as phenylacetylene when compared to 1-octyne. Nevertheless, a second mechanism based on a σ-alkynyl-palladium intermediate [21] might also explain the inefficiveness of all our palladium pre-catalysts towards the activation of 1,2-disubstituted alkenes such as 1-phenyl-1-propyne. It is however dismissed because there are not yet reasonable catalytic pathways allowing the releasing of enol ether by an unlikely methanol addition to a covalent Pd–C bond. A third mechanism which involves as key step the protonation of a η^2-diyne Pd(0) complex should also be mentioned, [32, 33] but it was also excluded in the discussion because no acidic compound or zerovalent palladium species is present in our catalytic system.

The impact of electron-withdrawing groups in the Triphos ligand on activity and selectivity of palladium precatalysts was investigated. The catalytic properties of 12 were examined and compared to those found with the cationic Triphos-palladium complex 9. After five days in neat methanol at room temperature, we found rather similar conversions of phenylacetylene with 51% for 12 and 57% for 9. This result was disappointing and did not fit our previsions in terms of activity enhancement. We discovered that the selectivity is, conversely, much more sensitive to slight variations of electrophilic character at the metal center. Indeed, the bis-addition [PhC(OMe)_2CH_3]/mono-addition [PhC(OMe)_2CH_3] products ratio was found to be 2.4 with complex 12, while under identical conditions it was of 6.1 with complex 9.

In view of getting more mechanistic insights, a phosphorus NMR spectroscopy monitoring was performed with complex 12 in the presence of either five equivalents of phenylacetylene or methanol in CDCl₃. In both cases, signals of complex 12 at 117.1 and 11.2 ppm were fully replaced by new triplets at 116.2 and 10.6 ppm and new doublets at 115.2 and 9.9 ppm, respectively. This attests that methanol and phenylacetylene can both easily coordinate palladium centers. Since the Δδ₀ for the palladium species coordinating methanol is higher than the Δδ₀ for palladium species coordinating phenylacetylene, a stronger ligand-metal interaction is attributed to methanol-palladium. Therefore, a favored thermodynamic intermediate for the first coordination to the metal is assumed to incorporate the alcohol preferentially instead of the alkyne, leading to complex (A) which is similar to those reported in the methoxycarbonylation of ethane, [33] see Scheme 5. This is perfectly consistent with a cationic Triphosfurane-palladium catalysts behaving similarly to the Lewis acid monophosphine gold catalyst. [16] Additional NMR kinetic experiments mixing phenylacetylene and methanol resulted in very quick modification of complexes at the NMR time-scale and did not allow any easy interpretation. Consistently with the NMR spectroscopic observations and above mentioned catalytic cycles, we depict the concomitant formation of mono- and bis-addition products in Scheme 5.

After replacement of one coordinated methanol by a phenylacetylene molecule, an inner nucleophilic attack of the alcohol to the C–C triple bond takes place to form the carbopalladated complex. A further internal protonation of the Pd–C bond as previously described with analogous gold complexes [16] would lead to complex (B). The active species (A) could be regenerated by direct releasing of the mono-addition product or by a second methanol nucleophilic addition followed by the bis-addition product release.
It is interesting to note that the different enol ether/acetal ratios found of 2.4 and 6.1 with complexes 9 and 12 respectively are correlated to the electrophilic character of the palladium center which depends on the P-substituents. Indeed, in the case where the triphosphine ligand bears the four electron-withdrawing furyl groups, the strength of the methanol-palladium interaction is enhanced. Subsequently, the enol ether elimination and its replacement by methanol could be favored against the second C–O bond formation. This effect would annihilate the expected beneficial effect of the presence of the electron-poor triphosphine ligand, as shown by the slightly decreased activity observed with the related Triphosfurane-palladium catalyst.

To summarize, the electron-withdrawing ligand does not alter the activity of palladium catalyst in the methoxylation of alkynes but enhances the productivity in mono-addition product. Further studies are under investigation to confirm these proposals and in particular theoretical calculations could help in this purpose.

CONCLUSION

Cationic palladium complexes containing the linear triphosphine ligands Triphos, Triphosline, Triphosfurane were studied as catalytic precursors in the methoxylation of aromatic and aliphatic-substituted alkynes. In the case of the Triphos ligand, a complete conversion of phenylacetylene was achieved with only 0.2% of pre-catalyst [Pd(Triphos)NCMe][(PF6)2] in a methanol/[BMIM][BF4] mixture. We discovered that the addition of an ionic liquid to pure methanol allowed not only to increase the activity of the palladium catalyst but also to provide a recyclable catalyst which can be reused more than a dozen times with a drop of activity of only 1% per day. This activity decreasing being zero-order in catalyst concentration we believe that it is due to evolution of physical/chemical properties of the catalytic medium in the course of recycling runs.

In an effort to further improve these catalytic performances, the first example of poor electron-donating/strong π-electron-acceptor linear trisphosphine, named Triphosfurane, [Ph[P(CH2CH2PFu'2)2]], has been prepared and fully characterized by NMR spectroscopy and X-ray diffraction. It has been obtained with high selectivity at low temperature from a novel vinyldifurylphosphine reagent and phenylphosphine in the presence of AIBN under irradiation conditions. Such specific conditions are required because the introduction of furyl groups onto vinylfurlyphosphine affects strongly its reactivity towards radical species. Upon palladium coordination, we found that the presence of electron-withdrawing groups in Triphos plays an important role on the selectivity of the phenylacetylene methoxylation without altering the activity. The performances of recovered ionic liquid described in the present study show for the first time that P-tridentate ligands exhibit excellent metal-immobilizing abilities in ionic liquids, and may lead to efficient catalytic systems, selective and benign for environment.

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EXPERIMENTAL SECTION

General Procedures. All reactions were performed in Schlenk-type flasks under an argon atmosphere. Solvents were purified and dried by conventional methods and distilled under argon. All analysis were performed at the "Plateforme d'Analyses Chimiques et de Synthèse Moléculaire de l'Université de Bourgogne" (PACSMUB). 1H, 31P{1H} and 13C{1H} NMR spectra were recorded at 298 K on a Bruker 300 Avance spectrometer. All chemical shifts are relative to SiMe4 (for 1H and 13C NMR spectroscopy) and 85% H3PO4 (for 31P NMR spectroscopy) and are given in ppm. Mass spectra were obtained on a Bruker microTOF-Q ESI-MS spectrometer. The elemental analyses were performed on a Fisons EA 1108 CHNS-O apparatus. The Triphos 3, AgOTf, TIPF6, acetonitrile, phenylacetylene, and 1-octyne were commercial products and were used as received. The bis(5-methyl-2-furyl) bromophosphine (Fu2PBr), ionic liquid [BMIM][BF4] were prepared according to the literature [26, 34]. The solids (catalyst and reagents) and the ionic liquid were degassed under vacuum before use. Ligand 5 and complexes [PdCl2(NCPh)2], 4, 6 and 8 were prepared according to the literature. [35, 21, 23]

Synthesis of Vinyldifurylphosphine 1

A solution of Fu2PBr (3.17 g, 11.6 mmol) in 10 ml of THF was slowly added to vinyl magnesium chloride (8.71 ml, 1.6 mol.l\(^{-1}\) in THF) at room temperature. The resulting yellow green solution was stirred for 3.5 hours. The reaction was hydrolyzed by 10 ml of degassed water and the solvent was removed under vacuum. The product I was extracted by 10 x 10 mL of CH2Cl2. The resulting solution was dried over MgSO4 and filtered over Celite®. Evaporation of solvent led to an orange viscous oil (2.35 g, 92 %). 1H NMR δ (CDCl3) = 6.67 (dd, 2H, J\(_{HH}\) = J\(_{HP}\) = 3Hz, H\(^a\)), 6.65 (dd, 1H, H\(^f\), J\(_{HP}\) = 12Hz, J\(_{HH}\) = 15Hz, J\(_{HP}\) = 24Hz), 6.00 (m, 2H, H\(^h\)), 5.82 (dd, H, H\(^i\), J\(_{HP}\) = 12Hz, J\(_{HH}\) = 2Hz, J\(_{HP}\) = 33Hz), 5.64 (ddd, H, H\(^i\), J\(_{HP}\) = 15Hz, J\(_{HH}\) = 2Hz, J\(_{HP}\) = 18Hz), 2.34 (s, 6H, CH\(_3\)). 13C\(^{13}\)C{1H} NMR δ (CDCl3) = 157.24 (d, 2C, J\(_{PC}\) = 2 Hz, C\(^5\)), 148.50 (d, 2C, J\(_{PC}\) = 8 Hz, C\(^6\)), 133.52 (d, 1C, J\(_{PC}\) = 3Hz, CH\(_{3vinyl}\)), 127.86 (d, 1C, J\(_{PC}\) = 22Hz, CH\(_{3vinyl}\)), 121.73 (d, 2C, J\(_{PC}\) = 24Hz, C\(^7\)), 107.05 (d, 2C, J\(_{PC}\) = 6Hz, C\(^8\)), 13.85 (s, 2C, CH\(_3\)). 31P{1H} NMR δ (CDCl3) = -57.03(s). Satisfactory elemental analysis could not be obtained due to its waxy nature and high air-sensitivity.

Synthesis of Triphosfurane 2a and Isomers 2b,b’

A mixture of 1 (303 mg, 1.37 mmol), phenylphosphine (83 µL, 0.75 mmol) and AIBN (12.4 mg, 0.075 mmol) was cooled to 4 °C and irradiated at this temperature with quartz jacketed high-pressure mercury lamp during 18 hours under stirring. The ligand 2a was obtained as white oil contaminated by traces of 2b,b’ (304 mg, 90 %). After several days...
at room temperature, pure ligand 2a was obtained as colorless crystals suitable for X-ray analysis. Data for 2a: 1H NMR δ (CDCl₃) = 7.42 (m, 2H, H₈), 7.12 (m, 3H, H₆), 6.72 (m, 4H, H₇), 5.80 (m, 4H, H₃), 2.43 (m, 4H, CH₂P₆H₅), 2.03 (s, 6H, CH₃), 2.01 (s, 6H, CH₃), 1.91 (m, 4H, CH₂P₆H₅), 1.31 (m, 2H, PH₇). 31P{1H} NMR δ (CDCl₃) = 139.64 (t, 1P, P, C = 4 Hz, C₄), 139.84 (t, 1P, P, C = 4 Hz, C₄), 123.57 (m, 2H, H₂), 119.32 (m, 2H, H₂), 121.51 (s, 2C, CH₃). F NMR δ (CDCl₃) = -60.32 (t, 2F, J₈,F = 13 Hz, F₈) and -60.94 (t, 2F, J₈,F = 13 Hz, F₈). For ligand 3, 1H NMR δ (CDCl₃) = 7.81 (m, 2H, H₈), 7.41 (m, 3H, H₆), 6.85 (m, 2H, H₇), 5.78 (m, 4H, H₃), 2.43 (m, 4H, CH₂PO₂), 2.03 (s, 6H, CH₃), 2.01 (s, 6H, CH₃), 1.91 (m, 4H, CH₂P₆H₅), 1.31 (m, 2H, PH₇). 31P{1H} NMR δ (CDCl₃) = -61.32 (t, 1P, P, C = 4 Hz, C₄), -61.34 (t, 1P, P, C = 4 Hz, C₄). The complexes were analyzed by ESI-MS (positive mode, CH₂Cl₂/MEOH) for C₃₆H₃₆N₅P₅PdF₁₂ (971.04) m/z = 699.09 [M+F - H]⁺, simulated: 699.101. Anal. Calc. for C₃₆H₃₆N₅P₅PdF₁₂ (M=971.04): C, 38.66, N, 1.74.

Synthesis of Complex 11

A mixture of complex PdCl₂(NCPh₂) (145 mg, 0.378 mmol) and ligand 5 (298 mg, 0.523 mmol) were added 10 ml of CH₂Cl₂. A red solution was obtained after a few minutes. After stirring overnight, the solution of complex PdCl₂(NCPh₂) was filtered and dried under vacuum for 3 hours. TlPF₆ (0.319 g, 0.913 mmol), 15 ml of CH₂Cl₂ and few drops of CH₃CN were then added and the resulting mixture became a red suspension after 3 hours stirring. After filtration, the solvent was partially removed and a beige powder was formed by addition of 10 ml of Et₂O, which was washed with Et₂O and dried under vacuum (243 mg, 64 %). 1H NMR δ (CDCl₃) = 7.83-7.11 (m, 20 H, H aromatics), 3.31-2.32 (m, 16 H, N(CH₂)₅, 5 PCH₂), 1.88 (s, 3 H, CH₃), 1H NMR δ (CDCl₃) = 163.36 (s, 1 C, CNC), 133.52-128.28 (m, 24 C, C aromatics), 51.42 (s, 1 C, CH₂(N=C)), 50.38 (s, 1 C, CH₂(N-C)), 41.52 (s, 1 C, CH₃N), 29.88 (s, 1 C, P(CH₂)), 27.42 (s, 2 C, 2 CH₂PPh₂), 15.53 (s, 1 C, CH₃). Found: C, 44.95, H, 4.45. Anal. Calc. for C₃₆H₄₂N₅P₅PdF₁₂ (M=1005.09): C, 42.98; H, 4.21; N, 4.18. Found: C, 42.66, H, 4.35, N, 4.45.

Synthesis of Complex 12

A mixture of complex 7 (150 mg, 0.206 mmol) and thallium hexafluorophosphate (250 mg, 0.715 mmol) were added 7 ml of acetonitrile. The yellow suspension was stirred for 3 hours at room temperature and then filtered over Celite. After evaporation of solvent, a yellow viscous residue was obtained which was washed with Et₂O and dried for 2 hours under vacuum (200 mg, 85%). 1H NMR δ (CDCl₃) = 8.30 (m, 2H, H₈), 7.65 (m, 3H, H₆), 7.48 (m, 2H, H₇), 6.27 (m, 2H, H₉), 5.78 (m, 2H, H₇), 5.61 (m, 2H, H₈), 5.58 (m, 2H, H₆), 5.50 (m, 2H, H₉), 5.37 (m, 2H, H₅), 4.52 (m, 2H, H₄), 4.23 (m, 2H, H₆), 4.09 (m, 2H, H₈), 3.86 (m, 2H, H₇), 3.65 (m, 2H, H₉), 3.05 (m, 2H, H₆), 2.93 (m, 2H, H₇), 2.84 (m, 2H, H₈), 2.64 (m, 2H, H₉), 2.43 (m, 2H, H₅), 2.25 (m, 2H, H₄), 2.17 (m, 2H, H₃), 1.67 (m, 2H, H₂). Found: C, 44.10; H, 4.46.

Synthesis of Complex 9

To a mixture of PdCl₂(NCPh₂) (0.101 g, 264 mmol) and Triphos (0.143 g, 267 mmol) were added 6 ml of CH₂Cl₂. After stirring overnight, addition of 50 ml of Et₂O allowed to precipitate the intermediate [PdCl(Triphos)]Cl which was filtered and dried under vacuum. TIPF₆ (0.187 g, 0.536 mmol), 7 ml of CH₂Cl₂ and few drops of CH₃CN were then added to this complex and the resulting suspension was stirred for 4 hours. After filtration, the solution was concentrated to the half volume under vacuum. Addition of 10 ml of Et₂O afforded the expected complex 6 as a beige powder which was dried under vacuum (139 mg, 91 %). 1H NMR δ (CDCl₃) = 7.85-7.20 (m, 25 H, H aromatics), 3.81-2.74 (m, 8 H, 4 PCH₂), 2.17 (s, 3 H, CH₃). No 13C¹{¹H}NMR data due to the lower solubility of 6 in common polar solvents. 31P¹{¹H} NMR δ (CDCl₃) = 110.38 (t, 1P, J₈,F = 10.6 Hz), 45.70 (d, 2 P, 3P₈,F = 10.6 Hz), -144.25 (hept., 1 P, PF₆, J₈,F = 571 Hz). ESI-MS (positive mode, CH₂Cl₂/MEOH) found for C₃₆H₃₆N₅P₅PdF₁₂ (971.04) m/z = 699.09 [M+F - H]⁺, simulated: 699.101. Anal. Calc. for C₃₆H₃₆N₅P₅PdF₁₂ (M=971.04): C, 44.49; H, 3.74; N, 1.44. Found: C, 44.95, H, 3.86, N, 1.74.

General Procedure for Methoxylation of Alkynes in an Ionic Liquid

Example given with complex 9: (10.3 mg, 0.011 mmol, 0.2 mol %) of 9 was dissolved in a mixture of 0.5 ml of [BMIM]BF₄ and 0.5 ml of CH₃OH (12.3 mmol). After addition of phenylacetylene (0.5 ml, 4.55 mmol), the solu-
Crystalline Structure Determination for Compound 2a

Crystal data and refinement details are reported in Table 2. Data sets were collected on an Enraf-Nonius Kappa CCD diffractometer at 115 K using MoKα radiation. The structure was solved using Direct Methods (SIR 92) [36] and refined with full-matrix least-squares methods based on [2] on the data of the WINGX [38] program. All non-hydrogen atoms were refined on anisotropic thermal parameters. Hydrogen atoms were included in their calculated positions and refined as riding atoms. CCDC-808224 for 2a contains the supplementary crystallographic database for this paper. This data can be obtained free of charge from www.ccdc.cam.ac.uk/conts/retrieving.html [or from the Cambridge Crystallographic Data Centre, 12, Union Road, Cambridge CB2 1EZ, UK; fax: (internat.) +44-1223/336-8082 for the supplementary crystallographic database].

Abbreviation

Triphos = [Ph2P(CH2)2PPh(CH2)2PPh3]

Conflict of Interest

None declared.

References

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