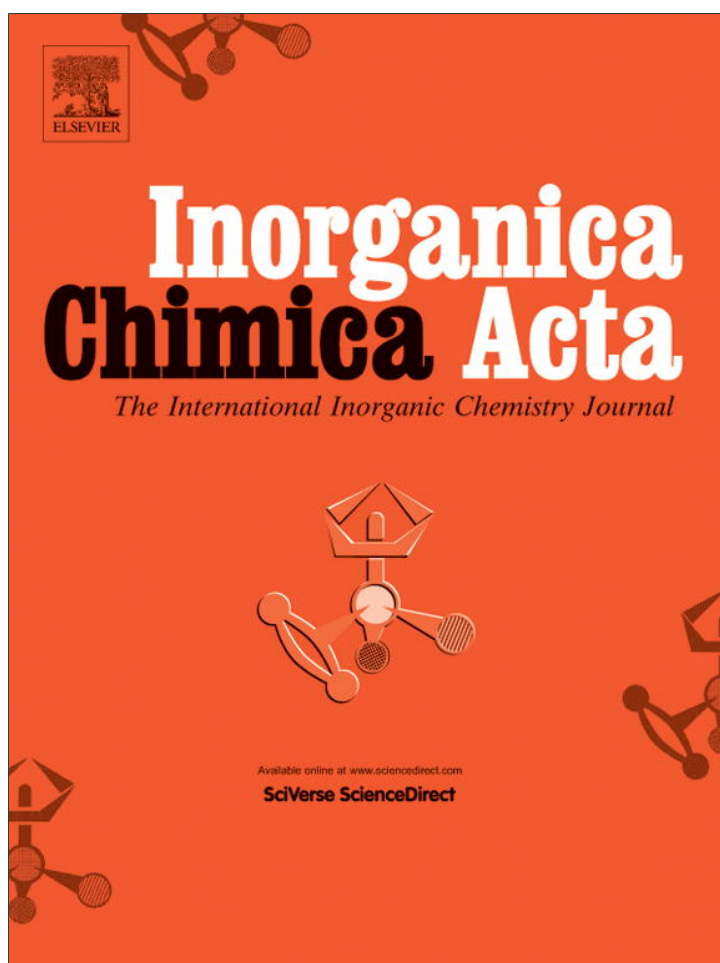


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The interaction between heteroditopic pyridine–nitrogen NHC with novel sulfur NHC ligands in palladium(0) derivatives: Synthesis and structural characterization of a bis-carbene palladium(0) olefin complex and formation in solution of an alkene–alkyne mixed intermediate as a consequence of the ligands hemilability

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ABSTRACT

We have synthesized a new class of heteroditopic chelating ligands bearing the phenylthiomethyl group and variously substituted imidazole derivatives and the corresponding complexes $[\text{Pd}(\eta^2\text{-ma})(\text{R-NHC-CH}_2\text{-SPh})]$ (ma = maleic anhydride; R = methyl, mesityl, di-*i*-propylphenyl). We have compared their chemical characteristics and reactivity with those of similar palladium(0) complexes bearing the R-NHC-CH₂-Py ligands.

The hemilability of the above-mentioned moieties is apparent when their palladium derivatives react with an excess of the same or different heteroditopic ligands. In these cases, we obtained complexes bearing maleic anhydride and two mono-coordinated heteroditopic ligands bound to the same metal center. However, at variance with the generally accepted opinion, precipitation of AgBr does not warrant the quantitative formation of the bis-heteroditopic Pd(0) derivative and in some cases, a reaction involving a heterogeneous equilibrium is observed.

The reaction of the palladium(0) carbene–sulfur or carbene–nitrogen complexes with the alkyne dimethyl-2-butyne dioate (dmbd) to give palladacyclometallate derivatives is not always warranted and formation of the intermediate $[\text{Pd}(\eta^2\text{-ma})(\eta^2\text{-dmbd})(\kappa^1\text{-Mes-NHC-CH}_2\text{-SPh})]$ is detected only in one case.

Finally, we have carried out a diffractometric study on the solid-state structures of two derivatives and in particular we describe the configurations of the complex $[\text{Pd}(\eta^2\text{-ma})(\kappa^1\text{-Me-NHC-CH}_2\text{-SPh})_2]$ and of $[\text{Pd}(\eta^2\text{-ma})(\text{Me-NHC-CH}_2\text{-SPh})]$.

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1. Introduction

Catalysts should be simultaneously characterized by a remarkable stability, a high reactivity and finally by selectivity toward the formation of the reaction products. These peculiarities which somehow might be considered chemically in conflict should all be present when metal complexes bearing hemilabile ligands [1] are used as catalysts. As a matter of fact, the introduction of a labile terminus into chelating ligands should promote the activation of a specific geometric site while the stability and the electronic requirements of the related complexes should be warranted by the pivot, which is the strong coordinating part of the ligands. The dangling wing bearing the labile nucleophile plays an impor-

tant role as well, since it will restore the starting substrates by re-coordinating the site made vacant as a consequence of the catalytic activity of the complexes [2].

Several classes of complexes of different metals and hemilabile ligands with a number of different scaffolds, different pivots and labile donors were synthesized and recently reviewed [2,3]. Furthermore, after the seminal article by Arduengo [4] a novel class of the performing donor ligands NHC was available for the chemists' community and hence several new heteroditopic and hemilabile NHC-E (E = P, N, O) ligands soon appeared in the literature [5]. Owing to our interest on chelating ligands bearing thioetheric sulfur [6] and the scarcity of heteroditopic NHC-S spectator ligands in the literature [7], we have synthesized some phenyl-thiomethyl-carbene (R-NHC-CH₂-SPh) (R = methyl, mesityl, di-*i*-propylphenyl) derivatives and the corresponding low valent palladium complexes and have compared their chemical characteristics and reactivity

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with those of the more studied R-NHC-CH₂-Py palladium(0) compounds [8].

Moreover, we have carried out a diffractometric study on the solid state structures of two olefin complexes and in particular we describe the configuration of a complex bearing maleic anhydride, Mes-NHC-CH₂-Py and Me-NHC-CH₂-SPh ligands coordinated to the same metal center.

2. Result and discussion

2.1. Synthesis of imidazolium bromide derivatives

The synthesis of the *N*-methylimidazolium (**1a** and **2a**) and *N*-arylimidazolium salts (**1b**, **1c**, **2b** and **2c**) was easily achieved in good yield (70–94%) by reacting 1-(2,4,6-trimethylphenyl)imidazole, 1-(2,6-di-*i*-propylphenyl)imidazole [9] or commercial 1-methylimidazole with an almost equimolecular amount of chloromethylphenylthioether or chloromethylpyridine in anhydrous acetonitrile in the presence of KBr, according to the literature procedure (Scheme 1): [10].

The formation of the imidazolium salts was easily detected by ¹H and ¹³C NMR technique owing to the very low field resonance of the acid imidazole proton between 10.0 and 10.5 and of the imidazole carbon between 136 and 139 ppm. Notably, compounds **2b** and **2c** which are known were synthesized by means of a slightly different procedure from that reported in the literature [11] (see Section 4).

2.2. R-NHC-CH₂-SPh and R-NHC-CH₂-Py silver(I) complexes

The transmetalation protocol with silver(I) NHC derivatives represents an advantageous and general synthetic procedure for the obtainment of transition metal carbene substrates [5g,5h,11]. As a matter of fact, the stability of silver carbene complexes does not require particular precautions and the synthesis of the corresponding transition metal carbene substrates has usually a remarkable yield. In particular, the transmetalation between silver(I) NHC derivatives and palladium(0) olefin precursors

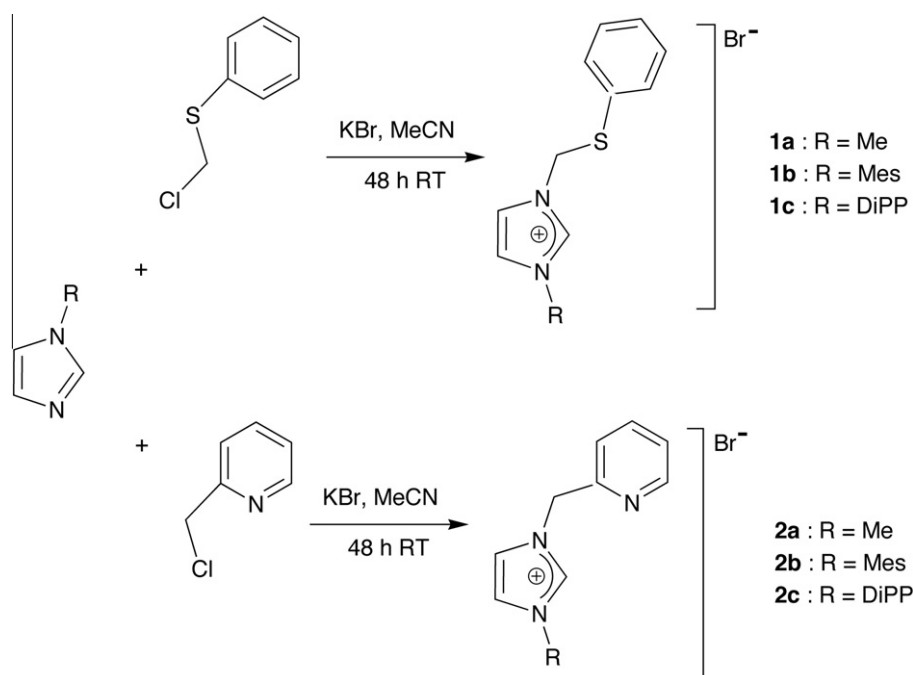
represents a very efficient synthetic approach to palladium(0) carbene complexes [5e,5j,7e,11b,11c,12]. Thus, we have synthesized the silver(I) NHC substrates according to the method reported in the literature [5g,5h] and described in Scheme 2.

The disappearance of the solid Ag₂O marks the reaction progress, whereas the ¹H and ¹³C NMR spectra of the products confirm the formation of the title complexes. In particular, the acid imidazolium proton is no more detectable in the NMR spectra of the silver derivatives and the coordinated imidazole carbon resonates at about 180 ppm. Remarkably, the methylene groups CH₂-Py or CH₂-S resonate as singlets thanks to the monocoordination of the R-NHC-CH₂-Py or R-NHC-CH₂-SPh and the consequent free rotation of the dangling uncoordinated wing bearing the thioether or pyridine fragment.

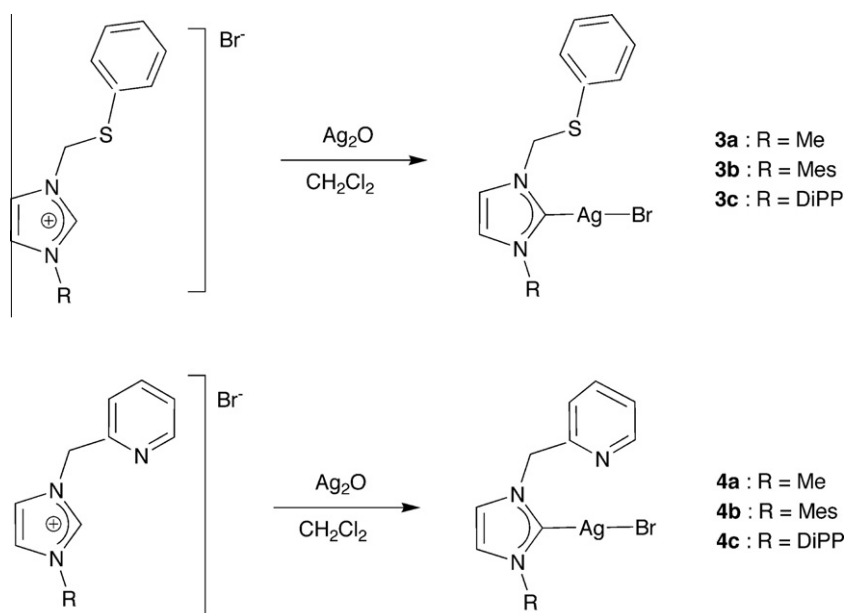
2.3. Synthesis of [Pd(η²-ol)(NHC)] complexes

Taking advantage of the lability of COD, Cavell and co-workers proposed the reaction between [Pd(η²-ma)(COD)] and free monodentate NHC ligands for the synthesis of some novel palladium(0) carbene derivatives [12]. However, such a direct approach is not possible with our carbene ligands since they are not available as free species. Therefore, we have reacted the same palladium(0) precursor with the silver derivatives R-NHC-CH₂-PyAgBr or R-NHC-CH₂-SPhAgBr. In order to minimize the decomposition of [Pd(η²-ma)(COD)], we have carried out the reaction by mixing the reactants at 253 K. The gradual temperature increase to 273 K promotes the reaction progress and the separation of the heteroditopic palladium(0) carbene complexes upon filtration of AgBr (Scheme 3).

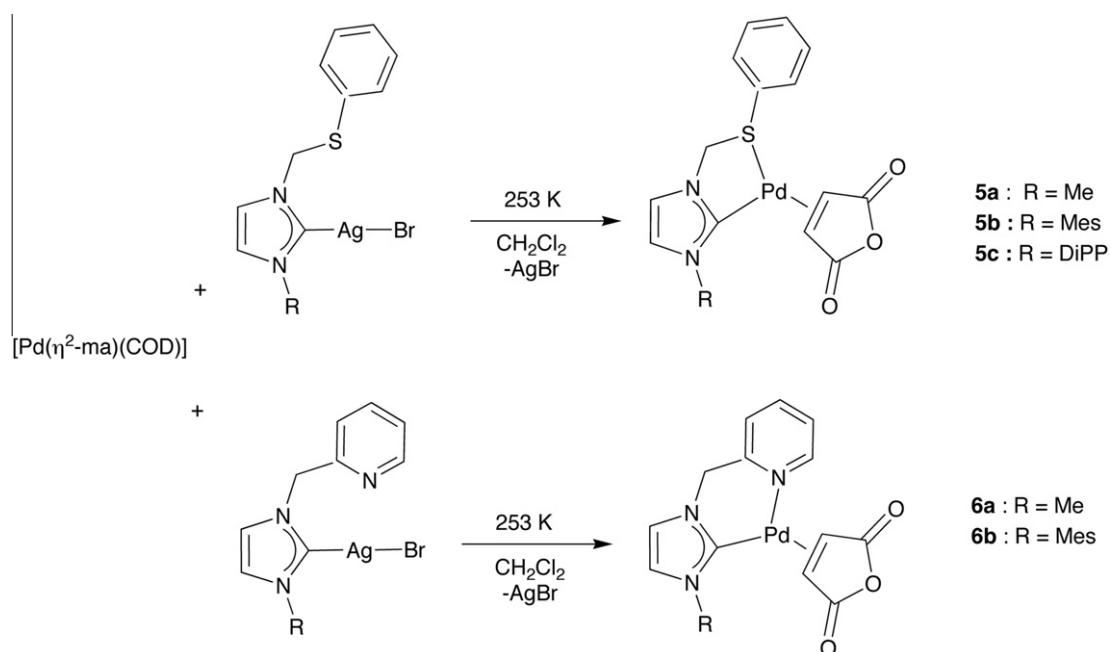
However, a limitation of such a method is represented by the fact that the ensuing carbene derivatives can only bear maleic anhydride (ma) or tetracyanoethene (tcne) since low valent palladium COD complexes with other, even slightly less stabilizing olefins are unstable and immediately decompose. The alternative approach to obtaining palladium(0) olefin complexes of the type [Pd(η²-ol)(L-L')] (ol = ma, fn, tcne; L-L' = NHC-Nitrogen) proposed by Elsevier and co-workers, consisting in the reaction between



Scheme 1. Synthetic methodology for the preparation of compounds **1a–1c** and **2a–2c**.



Scheme 2. Synthetic methodology for the preparation of compounds **3a–3c** and **4a–4c**.



Scheme 3. Synthetic methodology for the preparation of compounds **5a–5c** and **6a, 6b**.

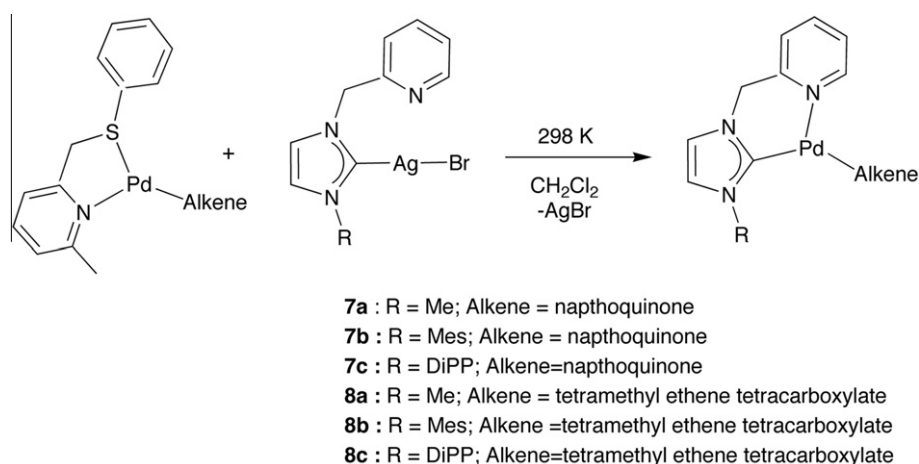
the complexes $[\text{Pd}(\eta^2\text{-ol})(t\text{-BuDAB})]$ (ol = ma, fn, tcne, *t*-BuDAB = *N,N'*-di-*t*-butyl-1,4-diaza-1,3-butadiene) and the silver carbene derivatives *NHC*- NMe_2AgBr [5g], *NHC*-pyridylAgBr, *NHC*-picolylAgBr (ol = ma) [11c] and *NHC*-pyrimidylAgBr (ol = ma) [13b], suggested to us an alternative protocol. Thus, we have reacted the *NHC* silver(I) derivatives with complexes of the type $[\text{Pd}(\eta^2\text{-ol})(\text{MeN-SPh})]$ (ol = ma, nq, tmetc; MeN-SPh = 2-methyl-6-sulfanylmethyl-pyridine) [14], the spectator ligand of which is displaced very easily by several nucleophiles [14b]. Thus, we were able to obtain the complexes of type $[\text{Pd}(\eta^2\text{-ol})(\text{R-NHC-CH}_2\text{-Py})]$ (ol = nq, tmetc) (Scheme 4):

Apparently, the complex $[\text{Pd}(\eta^2\text{-ol})(\text{MeN-SPh})]$ is more stable than $[\text{Pd}(\eta^2\text{-ma})(\text{COD})]$, thus the exchange reaction based on it

can be carried out at RT. However, addition of *R-NHC-CH}_2\text{-SPh-AgBr} to a solution of pyridylthioether palladium(0) derivatives leads to the formation of an equilibrium mixture of differently substituted species. This fact seems to suggest that the chelating ligands *R-NHC-CH}_2\text{-SPh} are less efficient than *R-NHC-CH}_2\text{-Py} in displacing the pyridylthioether fragment.***

The formation of the heteroditopic carbene olefin complexes can be clearly deduced from their ^1H and ^{13}C NMR spectra. In particular, the olefin protons and carbons resonate at higher fields than those of the uncoordinated ma ($\Delta\delta \sim 3\text{--}4$ and ~ 100 ppm, respectively).

In the low temperature (253 K) ^1H NMR spectra of the complexes bearing ma and *R-NHC-CH}_2\text{-SPh} (**5a–5c**), the olefin protons*



Scheme 4. Synthetic methodology for the preparation of compounds **7a–7c** and **8a–8c**.

always resonate as two doublets as a consequence of the asymmetry of the spectator ligands. Consistently, at the same temperature the olefin carbons show up as two separated signals.

At RT the olefin protons coalesce into two broad singlets. Such a phenomenon was often attributed to an apparent olefin rotation consisting in the de-coordination of the thioether wing followed by ligand rotation about the Pd–C bond and re-coordination at the opposite side (T-shape–Y-shape–T-shape mechanism). However, an alternative associative way triggered by the free electron pair on the sp^3 sulfur, cannot be ruled out [15].

Despite their diastereotopic nature at RT the CH_2 –S protons resonate as a broad singlet. This not unprecedented phenomenon is probably due to inversion of the sulfur absolute configuration due to fast exchange between the coordinated and the uncoordinated doublet of sp^3 sulfur [6a and refs. therein]. At low temperature (223 K), the broad singlet of complex **5c** only splits into an AB system.

The 1H NMR spectra of the complexes **5b** and **5c** at 253 K display the net separation of the signals ascribable to the *ortho* methyl protons of the mesityl and di-*i*-propylphenyl fragments, indicating that the rotation of those groups about their bond with the imidazole nitrogen is hampered.

In agreement with previous findings [11c] the carbenic carbons signal that is observed between 188.1 and 192.5 ppm exhibits a low field shift as compared to that of the Ag(I) complexes.

Finally, it is noteworthy that the resolved solid state structure of complex **5b** confirms the observation reported above (*vide post*).

Also in the case of the complexes bearing R-NHC- CH_2 -Py and naphthoquinone (**7a–7c**) the olefin protons are detected as broad singlet in the RT 1H NMR spectra. However, this signal splits into two separated signals at 253 K.

Owing to the ring inversion of the chelating ligand (olefin apparent rotation) the CH_2 -Py protons which are detected as an AB system at 253 K broaden at RT [11c].

The OCH_3 protons of the tetramethylethene tetracarboxylate derivatives (**8a–8c**) are detected at RT as two singlets. Probably, due to the steric hindrance in the complexes no olefin apparent rotation is operative in these cases.

2.4. Synthesis of bis-carbene palladium(0) olefin complexes

Taking advantage of the hemilability of these heteroditopic ligands, we have explored the possibility of introducing another carbene into the molecular scaffold of the palladium(0) olefin derivatives. Mc Guinness and Cavell [8a], Frøseth [16] and Fliedel and Braunstein [17] have obtained *trans* or *cis* bis-carbene

derivatives by reacting the Pd(II) cyclo-octadiene substrates [Pd(COD)(R)Cl], [Pd(COD)Cl₂] or [Pd(NCMe)₄](BF₄)₂ with the silver complexes of heteroditopic carbene-pyridine, carbene-imine or carbene-thioether respectively. Moreover, palladium(0) complexes bearing two monocoordinating carbene and maleic anhydride or tetracyanoethene have appeared so far in the literature [12,18]. However, attempts to synthesize complexes bearing two different heteroditopic carbenes simultaneously coordinated to low valent palladium were never carried out.

Therefore, we have reacted the palladium(0) derivative **5a** with the silver(I) complex **4b** in CH_2Cl_2 at RT. The ensuing fast reaction yields quantitatively the complex **9** represented in Chart 1. The structure of complex **9** in solution is deduced from its NMR spectra and confirmed by a diffractometric study. In particular, the concomitant coordination of both carbenes is testified by the presence of two distinct carbene carbon signals at 189.2 and 189.3 ppm in the ^{13}C NMR spectrum whereas the coordination of maleic anhydride is confirmed by the presence of two doublets of the olefin protons at 3.27 and 3.33 ppm in the 1H NMR spectrum. Curiously, the CH_2 -Py and CH_2 -S protons resonate as two AB systems although the sulfur and nitrogen are not bound to the metal center. This is probably due to the crowded molecular structure which hampers every rearrangement of the ligands, as can also be deduced from the hindered rotation of the mesityl ring about the C–N bond. As already stated, the mono-crystal diffractometric study of complex **9** confirms the proposed structure (*vide post*, Fig. 5a).

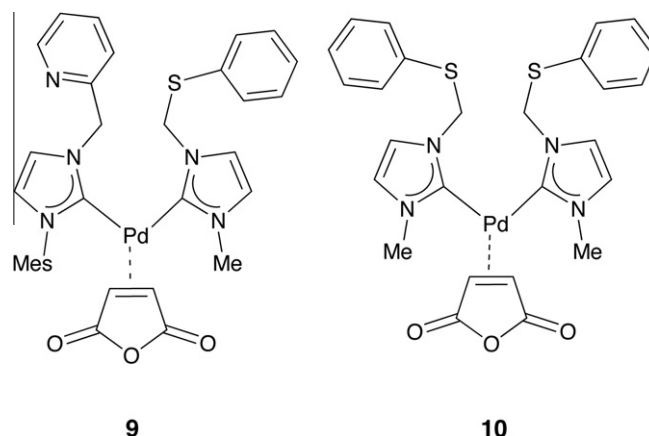


Chart 1. Topological representation of the complexes **9** and **10** bearing the two monocoordinated ditopic carbenes and the olefin coordinated to palladium.

In order to verify the general validity of such a protocol we have reacted palladium(0) and silver(I) complexes bearing the same carbene ligand. Thus, we have studied the reaction of the complex **5a** in CD_2Cl_2 with an equimolecular amount of the silver derivative **3a** by ^1H NMR at 298 K. Again, we have observed the fast and quantitative formation in solution of a new compound which was identified as complex **10** and isolated by addition of diethyl ether (Chart 1).

Surprisingly, the reaction between complexes **6b** and **4b** is not quantitative and is better represented by an equilibrium involving the incomplete displacement of the pyridine nitrogen of the coordinated ligand Mes-NHC-Py by the carbene carbon of the same ligand despite the commonly accepted idea that precipitation of insoluble AgBr should in any case drive this kind of reactions to completion. As a consequence, the contemporary presence in solution of **6b**, **4b** and **11** is clearly observed in the ensuing NMR spectrum whereas solid AgBr is detected as a cloudy suspension (Chart 2; Fig. 1).

Addition to the solution of further amounts of **4b** confirms the equilibrium nature of the reaction which can be easily driven to completion by addition of three equivalent of the silver derivative.

Similar equilibrium distributions can also be detected in the case of the reaction between complexes **6a** with **4a** and **6a** with **4b** (or **6b** with **4a**). It is apparent that the equilibrium position for these reactions is essentially determined by the insolubility of AgBr which dissolves in CH_2Cl_2 in an almost negligible extent, and by the efficiency of the carbene as strong σ -donor. These two concomitant phenomena support the widespread idea that these reactions are always quantitative. We have however shown that the chelating effect can efficiently contrast the reaction progress, and when two R-NHC- CH_2 -Py carbene ligands are involved at the same time, the reaction is not completely shifted to the right. At variance, these reactions are quantitative when at least one heteroditopic R-NHC- CH_2 -SPh carbene is involved. Evidently, the coordinative capability of the phenyl substituted sulfur appears to be less effective than that of the pyridine nitrogen, [6].

2.5. Reaction of the palladium(0) carbene derivatives with dimethyl-2-butyne dioate (dmbd)

We have also studied the attack of the activated alkyne dimethyl-2-butyne dioate (dmbd) on the complexes $[\text{Pd}(\eta^2\text{-ma})(\text{R-NHC-CH}_2\text{-Py})]$ and $[\text{Pd}(\eta^2\text{-ma})(\text{R-NHC-CH}_2\text{-SPh})]$. As can be deduced from the ^1H and ^{13}C NMR spectra, the complex **6a** reacts at RT with dmbd in CD_2Cl_2 giving quantitatively the palladium(II) cyclopentadienyl derivative $[\text{Pd}(\text{Me-NHC-CH}_2\text{-Py})(\text{C}_4(\text{COOMe})_4)]$

12. The presence of four distinct OCH_3 signals at 3.40, 3.42, 3.68 and 3.73 ppm and of the displaced maleic anhydride at about 7 ppm testifies the formation of the metallacyclopentadienyl derivative (Fig 2). Consistently, the carbene carbon signal shifts from 188.1 ppm of the starting complex **6a** to 180.5 ppm (see Section 4).

On the contrary, when complexes **5a**, **5b** and **6b** react with dmbd, an extensive decomposition with concomitant formation of metallic palladium was observed. In order to explain such a different behavior we resorted to an NMR study of the reaction between **5b** and the alkyne dmbd at low temperature (CD_2Cl_2 ; 253 K). The formation of a new organometallic species was immediately observed. The detailed investigation of its ^1H NMR spectrum strongly suggests that the derivative $[\text{Pd}(\eta^2\text{-ma})(\eta^2\text{-dmbd})(\kappa^1\text{-Mes-NHC-CH}_2\text{-SPh})]$ (**13**) bearing ma, dmbd and the monocoordinate carbene bound to palladium(0) is formed (Chart 3).

As a matter of fact, the olefin protons of the η^2 -coordinated ma resonate as a couple of doublets at 4.02 and 4.51 ppm ($J_{\text{HH}} = 4.5$ Hz), whereas the coordinated dmbd displays two singlets ascribable to the methyl protons of the non-equivalent methoxy fragments at 3.70 and 3.96 ppm, respectively. Interestingly, owing to the asymmetry of the intermediate, the $\text{CH}_2\text{-S}$ protons resonate as an AB system irrespective of the fact that the sulfur remains uncoordinated. This is probably due to the crowded molecular structure of **13** which hampers the free rotation of the wing bearing the thioether fragment, as was observed in the case of complex **9**, the open structure of which was unambiguously determined. Moreover, no uncoordinated ma is detectable in solution and the methyl groups of the mesityl fragment resonate as three distinct singlets since the free rotation of the latter is also hampered by steric hindrance (Fig. 3).

The HMQC spectrum confirms such an interpretation (see Supplementary material, Fig. 1S1); in particular, the coordination of the olefin ma is evident thanks to the cross-peaks between the olefin carbons (61.8, 62.8 ppm) and the olefin protons (4.02, 4.51 ppm) and to the signal resonance at lower field than that of the corresponding starting olefin complex.

Further confirmation comes from the presence of the olefin carbonyl carbon resonances at 168.4 and 168.7 ppm, the signal of the coordinated carbene carbon at 180.9 ppm and those of the non-equivalent ester carbons of the coordinated dmbd fragment at 159.8 and 160.8 ppm, respectively.

In some respect the compound **13** reminds similar species bearing two olefins and one NHC ligand coordinated to palladium, the existence of which was hitherto hypothesized by Cavell [19]. Moreover, beside the clear evidence displayed by the NMR spectra

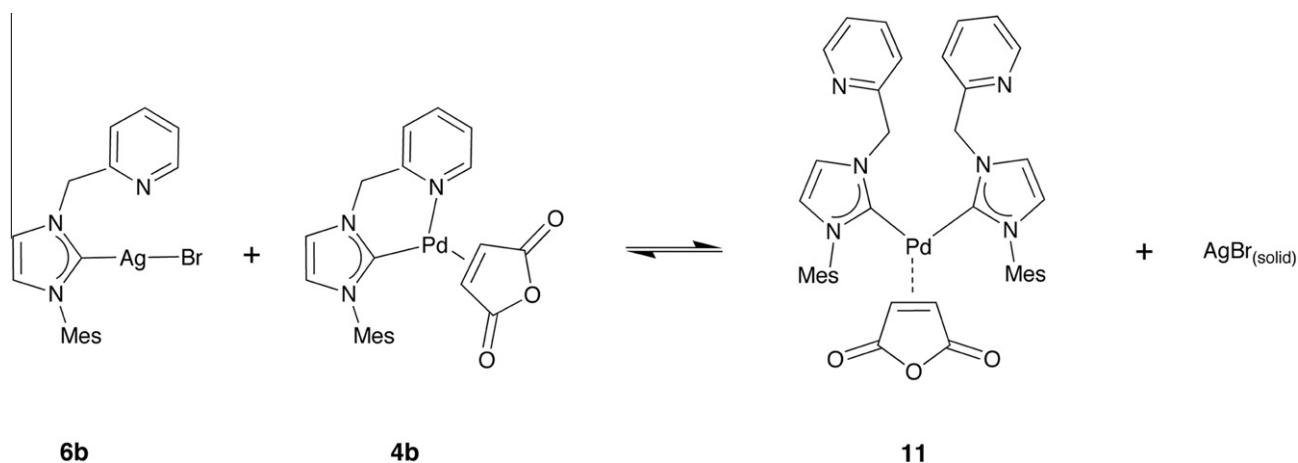


Chart 2. Equilibrium reaction between complexes **6b** and **4b** yielding complex **11**.

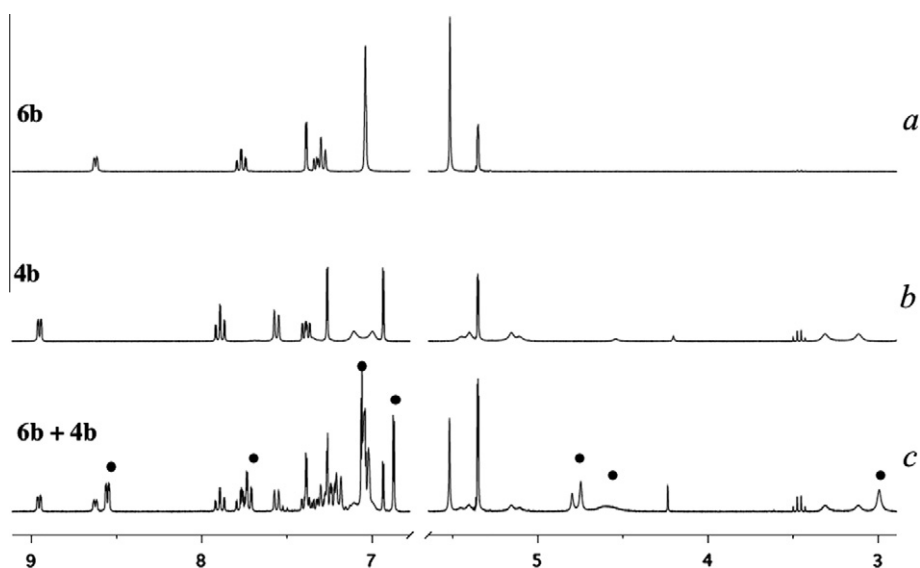


Fig. 1. Selected intervals of the ^1H NMR spectra at 298 K of CD_2Cl_2 solution of: (a) complex **6b**; $[\mathbf{6b}] = 1.5 \times 10^{-2}$ M; (b) complex **4b**; $[\mathbf{4b}] = 1.5 \times 10^{-2}$ M; (c) product **11** (●) from the equilibrium reaction between equimolecular amounts of complexes **4b** and **6b** (see Chart 2 and text).

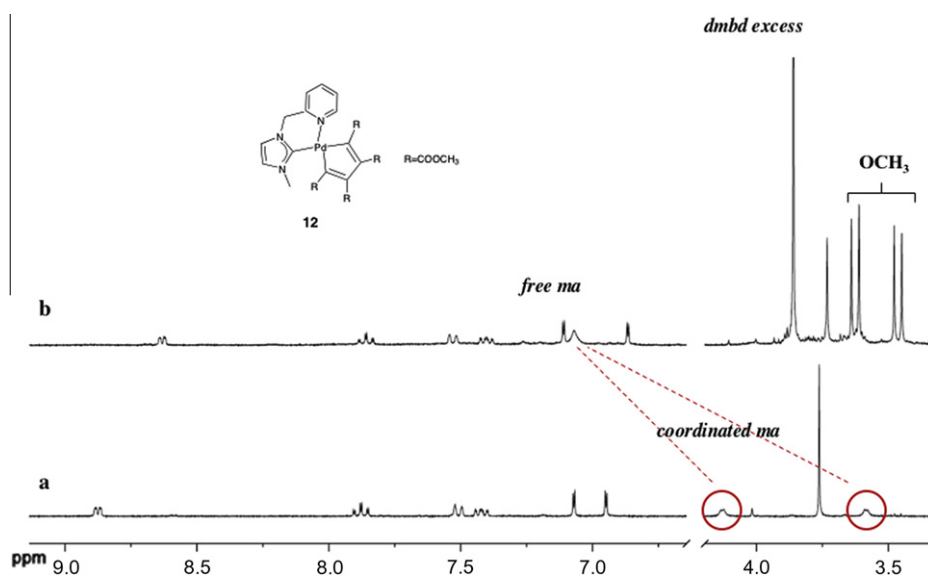


Fig. 2. Selected intervals of the ^1H NMR spectra at 298 K of CD_2Cl_2 solution of: (a) complex **6a**; $[\mathbf{6a}] = 1.5 \times 10^{-2}$ M; (b) complex **12**; $[\mathbf{12}] = 1.5 \times 10^{-2}$ M. (top insert) topological representation of complex **12**.

a theoretical investigation confirms the existence of a similar structure [20] (Chart 3b).

Remarkably, a palladium(0) complex bearing one olefin and one alkyne coordinated was hypothesised as a possible intermediate in the case of the attack of an alkyne to an olefin derivative bearing a pyridylthioether as spectator ligand, but in that case its presence in solution was not detectable [21a]. The stability of such an intermediate might probably be traced back to the donor nature of carbene and consequently to the high electron density on the Pd center which can be consequently easily shared between the two electron withdrawing unsaturated molecules.

Increasing the temperature of the solution induces a general decomposition with formation of metallic palladium and several unidentified species.

Similar behavior is detectable in the case of complexes **5a** and **6b** whereas, as already stated, only the complex **6a** yields the palladacyclopentadiene derivative $[\text{Pd}(\text{Me-NHC-CH}_2\text{-Py})(\text{C}_4(\text{COOMe})_4)]$

upon increasing the temperature. Interestingly, the decomposition of complex **5a** promotes the formation of the examethylmellitate derivative as can be deduced from the intense signal at 3.87 ppm ascribable to its eighteen protons [21]. The progress of the reaction yielding the palladacyclopentadiene product is probably controlled by small differences in the steric and electronic demands among the investigated complexes and in our case only the complex **6a** fulfills those steric requirements.

2.6. Crystal structure determination

ORTEP [22] views of complexes **5b** and **9** are shown in Figs. 4a and 5a. A selection of bond distances and angles is given in Table 1. In complex **5b** the palladium is bonded to a dissymmetric NHC bidentate ligand through the carbene carbon and the sulfur atom of the methylthiophenyl N-substituent and η^2 -coordinated to the C=C double bond of maleic anhydride. In complex **9** the palladium is

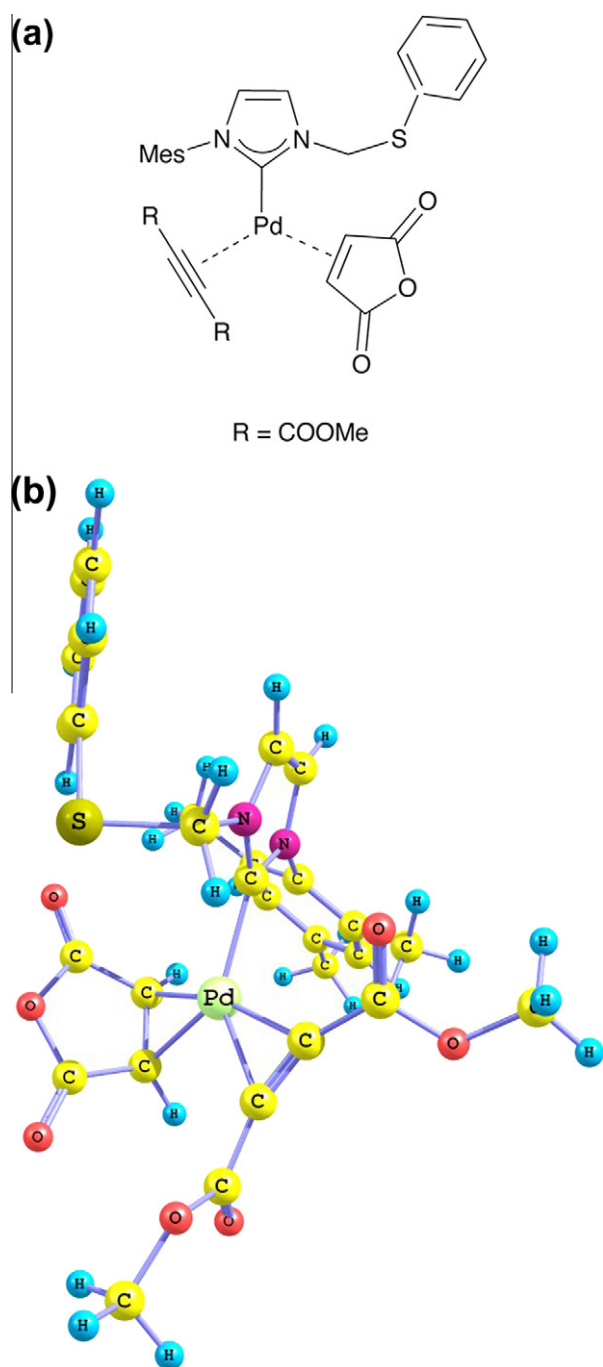


Chart 3. (a) Topological representation of the complex **13** bearing the monocoordinated ditopic carbene, the olefin ma and the alkyne dmbd coordinated and (b) structure of the complex **13** theoretically determined (see text).

bonded to two carbene carbons of two dissymmetric NHC ligands and η^2 -coordinated to the C=C double bond of maleic anhydride.

In both structures the palladium center is trigonal planar, as expected for zero-valent complexes of the type $[\text{PdL}'\text{L}'' (\eta^2\text{-alkene})]$. All bond distances involving the Pd(0) metal are in reasonable agreement with those in other complexes of Pd(0) with maleic anhydride [18,23]. In complex **5b** the Pd1–C2 length (2.125(3) Å) is considerably longer than the corresponding Pd1–C1 (2.060(3) Å) owing to the *trans* influence exerted by the carbene carbon of the ligand on the coordinated olefin, whereas the Pd–olefin distances in the case of complex **9** are similar in agreement with the findings of Lee et al. [18]. Remarkably, the Pd1–C5 and Pd1–C16

bonds in the bis-carbene derivative **9** are lengthened with respect to those of the Pd–C_{carbene} complex **5b** and of those of the chelate heteroditopic complexes described elsewhere [5g,11c,13b] but compare well with those of the similar complexes described in Ref. [18]. This fact is probably due to the increased electronic charge on the metal center. The C1=C2 alkene bond lengths of 1.430(4) and 1.444(4) Å, for structures **5b** and **9**, respectively, are about 0.1 Å longer than in the free alkene and are consistent with π -back bonding from Pd(0). The maleic anhydride rings are almost perpendicular to the basal coordination plane forming angles of 82.43(7)° with the Pd1, C1, C2, C5, S1 mean coordination plane of complex **5b** and of 79.5(8)° with the Pd1, C1, C2, C5, C16 mean coordination plane of **9** (Figs. 4b and 5b).

In the crystals the molecules are held together by means of C–H...O hydrogen bonds involving the oxygens of maleic anhydride as acceptors (Table 2).

3. Conclusion

We have synthesized new thioether derivatives of imidazolium salts **1a–1c**, the corresponding silver bromide derivatives **3a–3c** and the palladium(0) complexes **5a–5c**. We have compared the chemical characteristics and reactivity of the latter with those of complexes **6a** and **6b** bearing the carbene–pyridine R-NHC-CH₂-Py fragments as spectator ligands and resolved the solid state structure of complex **5b**.

Taking advantage of the hemilability of the above mentioned ligands we were able to obtain complexes with two mono-coordinated heteroditopic ligands and maleic anhydride concomitantly bound to the metal (complexes **9–11**) by adding one more equivalent of the silver carbenic precursor. From the NMR spectra of the reaction solutions it was clear that, despite the precipitation of AgBr an equilibrium mixture is often obtained in these cases. The structure of complex **9** bearing two different heteroditopic carbenes and an olefin concomitantly coordinated to a palladium(0) center, was determined by means of a diffractometric study. Moreover, in the attempt to obtain palladacyclometallate derivatives bearing R-NHC-CH₂-SPh or R-NHC-CH₂-Py as ancillary ligands we have reacted complexes **5a**, **5b**, **6a** and **6b** with the alkyne dimethyl-2-butyndioate. The expected complex was obtained only in the case of **6a**, whereas complexes **5a**, **5b** and **6b** undergo massive decomposition at RT. In the latter cases, the formation of an intermediate bearing ma, dmbd and the monodentate heteroditopic carbene coordinated to the same metal center was shown by means of low temperature NMR spectra.

4. Experimental

4.1. Materials

All solvents were purified by standard procedures and distilled under argon immediately before use. 1D- and 2D-NMR spectra were recorded using a Bruker 300 Avance spectrometer. Chemical shifts (ppm) are given related to TMS (¹H and ¹³C NMR). IR spectra were recorded on a Perkin-Elmer Spectrum One spectrophotometer.

4.2. Computational details

Theoretical calculations were performed with the GAUSSIAN 09 [20] package using the functional hybrid meta-GGA M06 [20c] and the Def2-TZVP basis set [20b]. The geometry optimization was performed without any symmetry constraint, followed by analytical frequency calculation to confirm that a minimum had been reached.

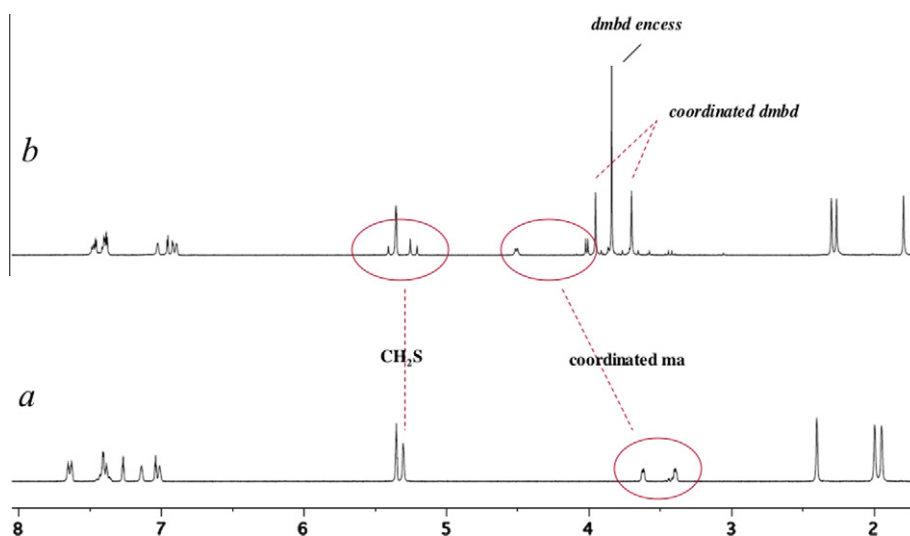


Fig. 3. Selected intervals of the ^1H NMR spectra at 298 K of a CD_2Cl_2 solution of: (a) complex **5b**; $[\mathbf{5a}] = 1.5 \times 10^{-2}$ M and (b) compound **13**.

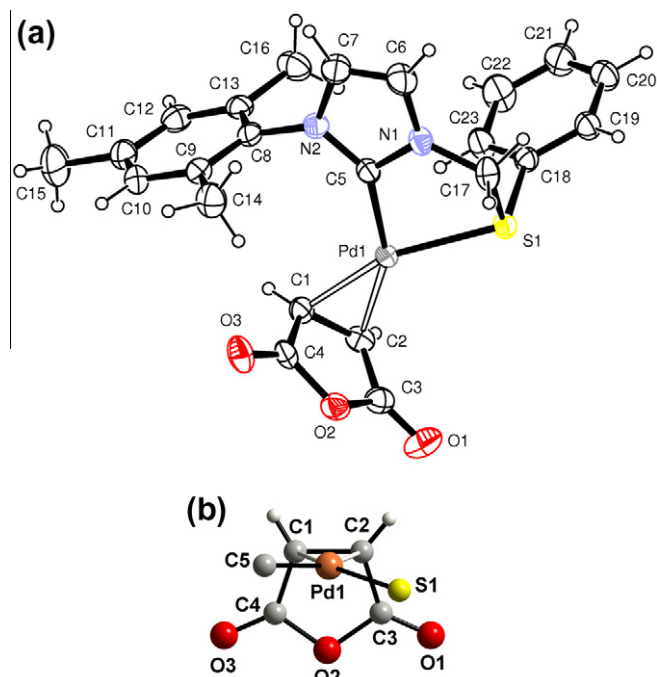


Fig. 4. (a) An ORTEP view of complex **5b** showing the thermal ellipsoids at 30% probability level. (b) The orientation of η^2 -maleic anhydride moiety with respect to the Pd coordination plane in complex **5b**.

Table 1
Selected bond distances and angles (Å and degrees).

	5b	9
<i>Distances</i>		
Pd1–C5	2.040(3)	2.067(3)
Pd1–S1	2.4157(8)	
Pd1–C16		2.061(2)
Pd1–C1	2.060(3)	2.101(3)
Pd1–C2	2.125(3)	2.121(3)
C1–C2	1.430(4)	1.444(4)
C5–N1	1.359(3)	1.354(4)
C5–N2	1.355(3)	1.357(4)
C16–N3		1.366(3)
C16–N4		1.362(3)
<i>Angles</i>		
C5–Pd1–S1	82.5(1)	
C5–Pd1–C16		98.6(1)
C5–Pd1–C1	116.2(1)	108.1(1)
C5–Pd1–C2	155.5(1)	147.9(1)
S1–Pd1–C1	161.1(1)	
S1–Pd1–C2	121.7(1)	
C16–Pd1–C1		152.8(1)
C16–Pd1–C2		113.4(1)
C1–Pd1–C2	39.9(1)	40.0(1)
<i>Dihedral Angles</i>		
Pd1,C1,C2,C5,S1 ^		
C1,C2,C3,C4,O2	82.43(7)	
Pd1,C1,C2,C5,C16^		
C1,C2,C3,C4,O2		79.54(8)

4.3. Crystal structure determinations

The crystal data of compounds **5b** and **9** were collected at room temperature using a Nonius Kappa CCD diffractometer with graphite monochromated $\text{Mo K}\alpha$ radiation. The data sets were integrated with the Denzo-SMN package [24] and corrected for Lorentz, polarization and absorption effects (SORTAV). [25] The structures were solved by direct methods using SIR97 [26] system of programs and refined using full-matrix least-squares with all non-hydrogen atoms anisotropically and hydrogens included on calculated positions, riding on their carrier atoms.

All calculations were performed using SHELXL-97 [27] and PARST [28] implemented in WINGX [29] system of programs. The crystal data are given in Table 1SI.

Crystallographic data (excluding structure factors) have been deposited at the Cambridge Crystallographic Data center and allocated the deposition numbers CCDC 848987 and CCDC 848988. These data can be obtained free of charge via www.ccdc.cam.ac.uk/conts/retrieving.html or on application to CCDC, Union Road, Cambridge, CB2 1EZ, UK (fax: +44 1223-336033, e-mail: deposit@ccdc.cam.ac.uk).

4.4. Synthesis of the precursors

The palladium precursors $\text{Pd}_2(\text{DBA})_3\cdot\text{CH}_2\text{Cl}$, [30] $[\text{Pd}(\eta^2\text{-ma-COD})]$, [31] $[(2\text{-methyl-6-(phenylthiomethyl)pyridine})\text{palladium}(0)(\text{naphthoquinone})]$ [23a] and $[(2\text{-methyl-6-(phenylthiomethyl)}$

Table 2
Hydrogen bond parameters (Å and degrees).

A–H...B	A–H	A...B	Symm. Op.	H...B	A–H...B
5b					
C6–H6...O3	0.93	3.167(4)	$3/2 - x, y - 1/2, z - 1/2$	2.24	172
C16–H16A...O1	0.96	3.449(5)	$x - 1/2, 3/2 - y, z$	2.58	151
C22–H22...O2	0.93	3.409(4)	$x, y - 1, z$	2.52	159
C19–H19...O2	0.93	3.527(3)	$2 - x, 2 - y, z - 1/2$	2.65	158
9					
C31–H31...O3	0.93	3.364(5)	$1/2 + x, 1/2 - y, z - 1/2$	2.48	159
C17–H17...O3	0.93	3.258(3)	$1/2 - x, y - 1/2, 1/2 - z$	2.40	153
C28–H28B...O1	0.97	3.439(3)	$-x, -y, -z$	2.55	152

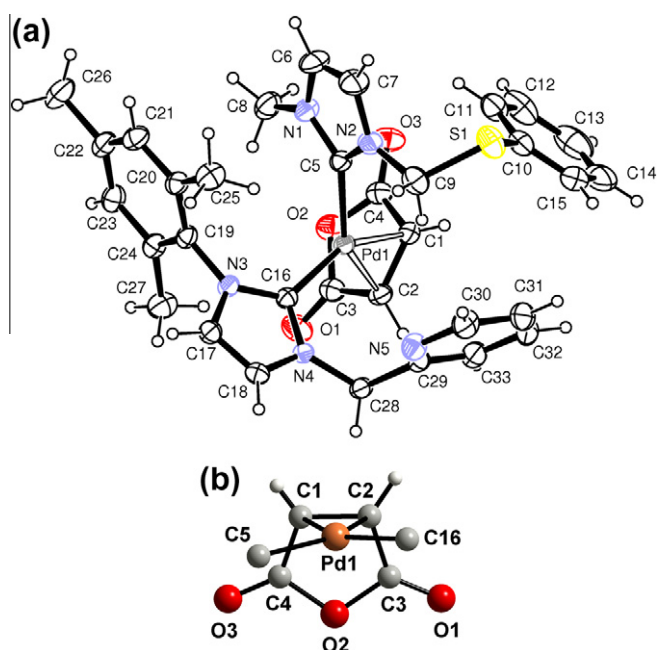


Fig. 5. (a) An ORTEP view of complex **9** showing the thermal ellipsoids at 30% probability level. (b) The orientation of η^2 -maleic anhydride moiety with respect to the Pd coordination plane in complex **9**.

pyridine)palladium(0)(tetramethyl ethene tetracarboxylate)] [**23e**] were synthesized according to published procedures.

4.5. Synthesis of the *N*-arylimidazoles

The syntheses of the *N*-mesityl-imidazole and *N*-2,6-diisopropylphenyl-imidazole were carried out according to the optimized published procedure. [9] The nucleophilic attack of the appropriate arylamine to oxalaldehyde in MeOH at RT was followed by the cyclization reaction of the cheto-imino derivatives in the presence of ammonium chloride, formaldehyde and phosphoric acid. The reaction products were extracted in diethylether and re-crystallized. The *N*-methyl-imidazole was a commercial product and used as purchased.

4.6. Synthesis of the *N*-arylimidazolium salts

[1-(2-Pyridyl)methylene-3-mesityl]imidazolium bromide (**2b**), [1-(2-pyridyl)methylene-3-(2,6-di-*i*-propylphenyl)]imidazolium bromide (**2c**) {[1-(2-pyridyl)methylene-3-mesityl]imidazolyl-2-

ene}silver bromide (**4b**) {[1-(2-pyridyl)methylene-3-(2,6-diisopropylphenyl)]imidazolyl-2-ene}silver bromide (**4c**) were prepared following literature procedures [11c].

4.7. [1-(Phenylthio)methylene-3-methyl]imidazolium bromide (**1a**)

To a solution of 0.93 g (13.8 mmol) of *N*-methyl-imidazole in anhydrous MeCN (40 ml), 2 g (12.6 mmol) of chloro-methyl-phenyl-sulfide and 1.65 g (13.9 mmol) of KBr were added under inert atmosphere (Ar). The resulting suspension was stirred for 48 h, treated with 60 ml of CH₂Cl₂ and filtered on a Celite filter. The clear solution was dried under vacuum and the residue was re-dissolved in CH₂Cl₂ (2–3 ml) and re-precipitated with diethylether as a sticky white solid. 2.68 g (yield 94%) of the title dry product were obtained upon decantation and evaporation of the solvent.

¹H NMR (CDCl₃, *T* = 298 K, ppm) δ : 3.99 (s, 3H, NCH₃), 6.28 (s, 2H, SCH₂), 5.86 (s, 2H, *m*-mesityl), 7.33 (m, 3H, SPh), 7.40 (t, *J* = 1.8 Hz, 1H, CH=CH Im), 7.44 (m, 2H, SPh), 7.51 (t, *J* = 1.7 Hz, 1H, CH=CH Im), 9.99 (bs, 1H, NCHN). ¹³C {¹H} NMR (CDCl₃, *T* = 298 K, ppm) δ : 36.8 (CH₃, NCH₃), 53.4 (CH₂, SCH₂), 121.7 (CH, CH=CH Im), 123.8 (CH, CH=CH Im), 128.8 (CH, SPh-CH), 129.7 (CH, SPh-CH), 130.4 (C, SPh-C), 132.2 (CH, CPh-CH), 136.5 (CH, NCHN). *Anal. Calc.* for C₁₁H₁₃BrN₂S: C, 46.32; H, 4.59; N, 9.82. Found: C, 46.21; H, 4.62; N, 9.90%.

The following derivatives were prepared in a similar way using the appropriate reactants.

4.8. [1-(Phenylthio)methylene-3-mesityl]imidazolium bromide (**1b**)

White micro crystals: Yield 77%. ¹H NMR (CDCl₃, *T* = 298 K, ppm) δ : 1.87 (s, 6H, *o*-mesityl-CH₃), 2.33 (s, 3H, *p*-mesityl-CH₃), 6.28 (s, 2H, SCH₂), 6.95 (s, 2H, *m*-mesityl-H), 7.04 (t, *J* = 1.8 Hz, 1H, CH=CH Im), 7.35 (m, 3H, SPh), 7.62 (m, 2H, SPh), 7.69 (t, *J* = 1.7 Hz, 1H, CH=CH Im), 10.22 (bs, 1H, NCHN).

¹³C {¹H} NMR (CDCl₃, *T* = 298 K, ppm) δ : 17.3 (CH₃, *o*-mesityl-CH₃), 20.9 (CH₃, *p*-mesityl-CH₃), 53.4 (CH₂, SCH₂), 121.7 (CH, CH=CH Im), 122.8 (CH, CH=CH Im), 129.0 (CH, SPh-CH), 129.5 (C, SPh-C), 129.6 (CH, SPh-CH), 129.7 (CH, *m*-mesityl-CH), 130.2 (C, *o*-mesityl-C), 132.7 (CH, SPh-CH), 134.0 (C, SPh-C), 138.4 (CH, NCHN), 141.4 (C, *p*-mesityl-C). *Anal. Calc.* for C₁₉H₂₁BrN₂S: C, 58.61; H, 5.44; N, 7.19. Found: C, 58.59; H, 5.47; N, 7.12%.

4.9. [1-(Phenylthio)methyl-3-(2,6-di-*i*-propylphenyl)]imidazolium bromide (**1c**)

White micro crystals: Yield 68%. ¹H NMR (CD₂Cl₂, *T* = 298 K, ppm) δ : 1.06 (d, *J* = 6.8 Hz, 6H, *i*Pr-CH₃), 1.13 (d, *J* = 6.8 Hz, 6H, *i*Pr-CH₃), 2.01 (sept, *J* = 6.8 Hz, 2H, *i*Pr-CH), 6.42 (s, 2H, SCH₂), 7.07 (d, *J* = 1.8 Hz, 1H, CH=CH Im), 7.26 (d, *J* = 7.9 Hz, 2H, *m*-aryl-H), 7.30–7.40 (m, 3H, SPh), 7.50 (t, *J* = 7.9 Hz, 2H, *p*-aryl-H), 7.65–7.68 (m, 2H, SPh), 7.89 (d, *J* = 1.8 Hz, 1H, CH=CH Im), 10.37 (bm, 1H, NCHN). ¹³C {¹H} NMR (CD₂Cl₂, *T* = 298 K, ppm) δ : 23.8 (CH₃, *i*Pr-CH₃), 22.9 (CH₃, *i*Pr-CH₃), 28.4 (CH, *i*Pr-CH), 53.6 (CH₂, SCH₂), 122.0 (CH, CH=CH Im), 124.2 (CH, CH=CH Im), 124.5 (CH, *m*-aryl-CH), 128.8 (CH, SPh-CH), 129.6 (CH, SPh-CH), 129.7 (C, *i*-aryl-C), 129.8 (C, SPh-C), 131.7 (CH, *p*-aryl-CH), 132.3 (CH, SPh-CH), 138.2 (CH, NCHN), 128.7 (C, *o*-aryl-C). *Anal. Calc.* for C₂₂H₂₇BrN₂S: C, 61.25; H, 6.31; N, 6.49. Found: C, 61.31; H, 6.28; N, 6.53%.

4.10. [1-(2-Pyridyl)methylene-3-methyl]imidazolium bromide (**2a**)

To an anhydrous MeCN solution (70 ml) of 3 g (23.5 mmol) of 2-chloromethylpyridine (obtained by neutralization with Na₂CO₃ of the corresponding hydrochloride), 4.13 g (34.5 mmol) of KBr and 2.1 ml (25.5 mmol) of *N*-methylimidazole were added under inert

atmosphere (Ar). The red-orange mixture was stirred for 48 h and the solvent reduced to 3–4 ml under vacuum. An equivalent volume of CH_2Cl_2 (70 ml) was then added and the resulting suspension was filtered on a Celite filter. The clear solution was evaporated to dryness and the reddish sticky residue was re-dissolved in CH_2Cl_2 (15 ml) and precipitated with diethylether (40 ml). 5.08 g (yield 87%) of red hygroscopic micro crystals were obtained.

^1H NMR (CDCl_3 , $T = 298$ K, ppm) δ : 4.06 (s, 3H, NCH_3) 5.76 (s, 2H, NCH_2), 7.27 (d, $J = 1.6$ Hz, 1H, $\text{CH}=\text{CH}$ Im), 7.30 (m, 1H, 5-Pyr), 7.63 (d, $J = 1.6$ Hz, 1H, $\text{CH}=\text{CH}$ Im), 7.77 (td, $J = 7.4$, 1.6 Hz, 1H, 4-Pyr), 7.84 (d, $J = 7.4$ Hz, 1H, 3-Pyr), 8.55 (d, $J = 4.9$ Hz, 1H, 6-Pyr), 10.71 (bs, 1H, NCHN). ^{13}C $\{^1\text{H}\}$ NMR (CDCl_3 , $T = 298$ K, ppm) δ : 36.6 (CH_3 , NCH_3), 53.7 (CH_2 , NCH_2), 122.7 (CH, 5-Py), 123.3 (CH, $\text{CH}=\text{CH}$ Im), 123.6 (CH, $\text{CH}=\text{CH}$ Im), 123.8 (CH, 3-Py), 137.3 (CH, 4-Py), 137.5 (CH, NHC), 149.7 (CH, 6-Py), 152.3 (C, 2-Py). *Anal. Calc.* for $\text{C}_{10}\text{H}_{12}\text{BrN}_2\text{S}$: C, 47.26; H, 4.76; N, 16.54. Found: C, 47.19; H, 4.73; N, 16.59%.

4.11. *{[1-(Phenylthio)methylene-3-methyl]imidazolyl-2-ene}silver bromide (3a)*

To a solution of 0.322 g (1.13 mmol) of $[\text{Me-NHC-CH}_2\text{-SPh}]\text{Br}$ in anhydrous CH_2Cl_2 (20 ml) under inert atmosphere (Ar) 0.144 g (0.621 mmol) of Ag_2O was added. The mixture was stirred in the dark for 12 h and filtered on a Celite filter. The clear solution was evaporated to small volume under vacuum and the title product was precipitated by addition of diethylether filtered on gouch and washed with diethylether and pentane. 0.378 g (yield 85%) of the complex as a whitish solid was obtained. ^1H NMR (CD_2Cl_2 , $T = 298$ K, ppm) δ : 3.76 (s, 3H, NCH_3), 5.42 (s, 2H, SCH_2), 7.00 (d, $J = 1.8$ Hz, 1H, $\text{CH}=\text{CH}$ Im), 7.11 (d, $J = 1.8$ Hz, 1H, $\text{CH}=\text{CH}$ Im), 7.38 (m, 5H, SPh). ^{13}C $\{^1\text{H}\}$ NMR (CDCl_3 , $T = 253$ K, ppm) δ : 38.7 (CH_3 , NCH_3), 56.2 (CH_2 , SCH_2), 120.4 (CH, $\text{CH}=\text{CH}$ Im), 122.6 (CH, $\text{CH}=\text{CH}$ Im), 129.7 (CH, SPh-CH), 129.0 (CH, SPh-CH), 130.9 (C, SPh-C), 133.3 (CH, SPh-CH), 182.2 (C, NCN). *Anal. Calc.* for $\text{C}_{11}\text{H}_{12}\text{AgBrN}_2\text{S}$: C, 33.70; H, 3.09; N, 7.15. Found: C, 33.73; H, 2.99; N, 7.21%.

The following derivatives were prepared in a similar way using the appropriate reactants.

4.12. *{[1-(Phenylthio)methylene-3-mesityl]imidazolyl-2-ene}silver bromide (3b)*

Light brown microcrystals: Yield 83%. ^1H NMR (CDCl_3 , $T = 298$ K, ppm) δ : 1.80 (s, 6H, *o*-mesityl CH_3), 2.31 (s, 3H, *p*-mesityl CH_3), 5.51 (s, 2H, SCH_2), 6.91 (m, 3H, $\text{CH}=\text{CH}$ Im, *m*-mesityl-H), 7.34 (m, 3H, SPh), 7.37 (d, $J = 1.8$ Hz, 1H, $\text{CH}=\text{CH}$ Im), 7.45 (m, 2H, SPh), ^{13}C $\{^1\text{H}\}$ NMR (CDCl_3 , $T = 298$ K, ppm) δ : 17.5 (CH_3 , *o*-mesityl- CH_3), 20.9 (CH_3 , *p*-mesityl- CH_3), 56.4 (CH_2 , SCH_2), 120.2 (CH, $\text{CH}=\text{CH}$ Im), 123.4 (CH, $\text{CH}=\text{CH}$ Im), 129.2 (CH, *m*-mesityl-CH), 129.3 (C, *o*-mesityl-C), 129.8 (CH, SPh-CH), 130.01 (C, SPh-C), 133.8 (CH, SPh-CH), 134.4 (CH, SPh-CH), 134.9 (C, *i*-mesityl-C), 139.5 (C, *p*-mesityl-C), 183.1 (C, NCN). *Anal. Calc.* for $\text{C}_{19}\text{H}_{20}\text{AgBrN}_2\text{S}$: C, 45.99; H, 4.06; N, 5.65. Found: C, 46.03; H, 4.11; N, 5.67%.

4.13. *{[1-(Phenylthio)methylene-3-(2,6-di-*i*-propylphenyl)]imidazolyl-2-ene}silver bromide (3c)*

White microcrystals: Yield 92%. ^1H NMR (300 MHz, CDCl_3) δ : 1.06 (d, $J = 6.9$ Hz, 6H, *iPr-CH}_3), 1.11 (t, $J = 6.8$ Hz, 6H, *iPr-CH}_3), 2.12 (sept, $J = 6.8$ Hz, 2H, *iPr-CH}), 5.54 (s, 2H, $\text{CH}_2\text{-S}$), 6.95 (s, 1H, $\text{CH}=\text{CH}$ Im), 7.19 (d, $J = 7.8$ Hz, 2H, *CH o-Ph*), 7.33–7.38 (m, 3H, CH mesityl), 7.39–7.43 (m, 2H, $\text{CH}=\text{CH}$ Im, *CH p-Ph*), 7.44–7.50 (m, 2H, CH m-Ph). ^{13}C NMR (75 MHz, CDCl_3) δ : 24.2 (CH_3 , *iPr-CH}_3), 24.3 (CH_3 , *iPr-CH}_3), 28.0 (CH, *iPr-CH}), 56.3 (CH_2 , $\text{CH}_2\text{-S}$),******

119.9 (CH, $\text{CH}=\text{CH}$ Im), 124.1 (CH *o-Ph*), 124.7 (CH, $\text{CH}=\text{CH}$ Im), 129.2 (CH, *p*-DiPP), 129.8 (CH, *m-Ph*), 130.5 (CH, *p-Ph*), 133.5 (C, *i*-DiPP), 145.4 (C, *o*-DiPP-C), 183.6 (C, NHC). *Anal. Calc.* for $\text{C}_{22}\text{H}_{26}\text{AgBrN}_2\text{S}$: C, 49.09; H, 4.87; N, 5.20. Found: C, 49.18; H, 4.75; N, 5.29%.

4.14. *{[1-(2-Pyridyl)methylene-3-methyl]imidazolyl-2-ene}silver bromide (4a)*

White microcrystals: Yield 71%. ^1H NMR (CDCl_3 , $T = 298$ K, ppm) δ : 3.87 (d, $J = 0.7$ Hz, 3H, NCH_3), 5.44 (s, 2H, NCH_2), 7.00 (d, $J = 1.8$ Hz, 1H, $\text{CH}=\text{CH}$ Im), 7.18 (d, $J = 1.8$ Hz, 1H, $\text{CH}=\text{CH}$ Im), 7.24 (dd, $J = 7.6$, 4.9 Hz, 1H, 5-Pyr), 7.39 (d, $J = 7.6$ Hz, 1H, 3-Pyr), 7.69 (td, $J = 7.6$, 1.6 Hz, 1H, 4-Pyr), 8.56 (d, $J = 4.9$ Hz, 1H, 6-Pyr). ^{13}C $\{^1\text{H}\}$ NMR (CDCl_3 , $T = 253$ K, ppm) δ : 39.2 (CH_3 , NCH_3), 57.4 (CH_2 , NCH_2), 122.3 (CH, $\text{CH}=\text{CH}$ Im), 122.7 (CH, $\text{CH}=\text{CH}$ Im), 123.1 (CH, 5-Pyr), 123.8 (CH, 3-Pyr), 137.8 (CH, 4-Pyr), 150.3 (CH, 6-Pyr), 155.0 (C, 2-Pyr), 182.4 (C, NCN). *Anal. Calc.* for $\text{C}_{10}\text{H}_{11}\text{AgBrN}_3$: C, 33.27; H, 3.07; N, 11.64. Found: C, 33.35; H, 3.21; N, 11.51%.

4.15. *{[1-(Phenylthio)methylene-3-mesityl]imidazolyl-2-ene}palladium(0) (maleic anhydride) (5b)*

To the yellowish solution of 0.05 g (0.16 mmol) of $[\text{Pd}(\eta^2\text{-ma})(\text{COD})]$ under inert atmosphere (Ar) in 15 ml of anhydrous CH_2Cl_2 at 253 K (liquid N_2 /acetone) 0.0793 g (0.16 mmol) of the silver(I) complex (**3b**) dissolved in 10 ml of anhydrous CH_2Cl_2 previously cooled at 253 K was added in two successive aliquots (5 ml each). The resulting brownish mixture was stirred for 1 h and the bath temperature was gradually increased to 270 K. The precipitated AgBr and traces of metallic palladium were removed by Millipore filtration and the resulting yellow solution evaporated to dryness. Addition of diethylether (20 ml) induced the precipitation of 0.0675 g (yield 83%) of the title complex which was filtered off, washed with diethylether and stored at 253 K.

^1H NMR (CD_2Cl_2 , $T = 298$ K, ppm) δ : 2.00 (s, 6H, *o*-mesityl- CH_3), 2.43 (s, 3H, *p*-mesityl CH_3), 3.55 (bs, 2H, $\text{CH}=\text{CH}$ ma), 5.32 (s, 2H, SCH_2), 7.04 (d, $J = 1.9$ Hz, 1H, $\text{CH}=\text{CH}$ Im), 7.09 (s, 2H, *m*-mesityl-H), 7.27 (d, $J = 1.9$ Hz, 1H, $\text{CH}=\text{CH}$ Im), 7.41 (m, 3H, SPh), 7.65 (m, 2H, SPh). ^1H NMR (CD_2Cl_2 , $T = 253$ K, ppm) δ : 1.95 (s, 3H, *o*-mesityl- CH_3), 2.00 (s, 3H, *o*-mesityl- CH_3), 2.40 (s, 3H, *p*-mesityl CH_3), 3.40 (d, $J = 3.3$ Hz, 1H, $\text{CH}=\text{CH}$ ma), 3.62 (d, $J = 3.3$ Hz, 1H, $\text{CH}=\text{CH}$ ma), 5.30 (bs, 2H, SCH_2), 7.01 (s, 1H, *m*-mesityl-H), 7.04 (d, $J = 1.9$, 1H, $\text{CH}=\text{CH}$ Im), 7.14 (s, 1H, *m*-mesityl-H), 7.26 (d, $J = 1.9$, 1H, $\text{CH}=\text{CH}$ Im), 7.40 (m, 3H, SPh), 7.64 (m, 2H, SPh). ^{13}C $\{^1\text{H}\}$ NMR (CD_2Cl_2 , $T = 253$ K, ppm) δ : 17.5 (CH_3 , *o*-mesityl- CH_3), 17.8 (CH_3 , *p*-mesityl- CH_3), 21.0 (CH_3 , *o*-mesityl- CH_3), 42.4 (CH, $\text{CH}=\text{CH}$ ma), 47.1 (CH, $\text{CH}=\text{CH}$ ma), 59.9 (CH_2 , SCH_2), 118.8 (CH, $\text{CH}=\text{CH}$ Im), 123.6 (CH, $\text{CH}=\text{CH}$ Im), 128.2 (CH, *m*-mesityl-CH), 129.2 (CH, *m*-mesityl-CH), 129.5 (CH, SPh-CH), 130.0 (CH, SPh-CH), 131.7 (C, SPh-C), 132.7 (CH, SPh-CH), 134.4 (C, *o*-mesityl-C), 135.6 (C, *o*-mesityl-C), 136.4 (C, *i*-mesityl-C), 138.7 (C, *p*-mesityl-C), 173.0 (C, $\text{C}=\text{O}$ ma), 173.2 (C, $\text{C}=\text{O}$ ma), 190.3 (C, NCN), IR (KBr pellet, cm^{-1}) ν_{CO} : 1791, 1724. *Anal. Calc.* for $\text{C}_{24}\text{H}_{24}\text{N}_2\text{O}_2\text{PdS}$: C, 56.42; H, 4.73; N, 5.48. Found: C, 56.31; H, 4.83; N, 5.36%.

The following derivatives were prepared in a similar way using the appropriate reactants.

4.16. *{[1-(Phenylthio)methylene-3-methyl]imidazolyl-2-ene}palladium(0) (maleic anhydride) (5a)*

Yellow microcrystals: yield 70%. ^1H NMR (CD_2Cl_2 , $T = 298$ K, ppm) δ : 3.92 (s, 3H, NCH_3), 3.95 (bs, 2H, $\text{CH}=\text{CH}$ ma), 5.18 (s, 2H, SCH_2), 7.09 (bm, 2H, $\text{CH}=\text{CH}$ Im), 7.41 (m, 3H, SPh), 7.61 (m, 2H, SPh). ^1H NMR (CD_2Cl_2 , $T = 253$ K, ppm) δ : 3.82 (bs, 1H, $\text{CH}=\text{CH}$

ma), 3.90 (s, 3H, NCH₃), 4.47 (bs, 1H, CH=CH ma), 5.17 (s, 2H, SCH₂), 7.10 (bs, 2H, CH=CH Im), 7.39 (m, 3H, SPh), 7.59 (m, 2H, SPh). ¹³C {¹H} NMR (CDCl₃, T = 253 K, ppm) δ: 39.0 (CH₃, NCH₃), 42.0 (CH, CH=CH ma), 47.6 (CH, CH=CH ma), 59.2 (CH₂, SCH₂), 118.4 (CH, CH=CH Im), 123.4 (CH, CH=CH Im), 129.6 (CH, SPh-CH), 129.9 (CH, SPh-CH), 131.7 (C, SPh-C), 132.7 (CH, SPh-CH), 172.7 (C, C=O ma), 172.8 (C, C=O ma), 188.1 (C, NCN), IR (KBr pellet, cm⁻¹) ν_{CO}: 1785, 1719. Anal. Calc. for C₁₆H₁₆N₂O₂PdS: C, 47.24; H, 3.96; N, 6.89. Found: C, 47.36; H, 3.83; N, 6.73%.

4.17. {[1-(Phenylthio)methylene-3-(2,6-di-i-propylphenyl)imidazolyl-2-ene]palladium(0) (maleic anhydride) (**5c**)}

Yellow microcrystals: yield 92%. ¹H NMR (CD₂Cl₂, T = 298 K, ppm) δ: 1.10 (d, J = 6.9 Hz, 6H, iPr-CH₃), 1.23 (d, J = 6.9 Hz, 6H, iPr-CH₃), 2.47 (bm, 2H, iPr-CH), 3.61 (bs, 2H, CH=CH ma), 5.33 (s, 2H, SCH₂), 7.10 (d, J = 1.9 Hz, 1H, CH=CH Im), 7.27 (d, J = 1.9 Hz, 1H, CH=CH Im), 7.36 (d, J = 7.7 Hz, 2H, m-aryl-H), 7.43 (m, 3H, SPh-CH), 7.56 (t, J = 7.7 Hz, 1H, p-aryl-H), 7.64 (m, 2H, SPh-CH). ¹³C {¹H} NMR (CDCl₃, T = 298 K, ppm) δ: 23.2 (CH₃, iPr-CH₃), 24.7 (CH₃, iPr-CH₃), 28.4 (CH₃, iPr-CH), 59.1 (CH₂, SCH₂), 118.2 (CH, CH=CH Im), 124.8 (CH, CH=CH Im), 124.0 (CH, m-aryl-CH), 129.6 (CH, SPh-CH), 129.8 (CH, p-aryl-CH), 129.9 (CH, SPh-CH), 131.54 (C, SPh-C), 132.0 (CH, SPh-CH), 135.9 (C, i-aryl-C), 144.9 (C, o-aryl-C), 192.5 (C, NCN). ¹H NMR (CD₂Cl₂, T = 253 K, ppm) δ: 1.07 (bs, 6H, iPr-CH₃), 1.18 (bd, J = 6.5 Hz, 3H, iPr-CH₃), 1.26 (bd, J = 7.4 Hz, 3H, iPr-CH₃), 2.28 (bm, 1H, iPr-CH), 2.50 (bm, 1H, iPr-CH), 3.58 (bd, J = 3.1 Hz, 1H, CH ma), 3.77 (bd, J = 3.1 Hz, 1H, CH ma), 5.24 (d, J = 12.1 Hz, 1H, CH₂S), 5.31 (d, J = 12.6 Hz, 1H, CH₂S), 7.07 (d, J = 1.7 Hz, 1H CH=CH Im), 7.26 (d, J = 1.8 Hz, 1H CH=CH Im), 7.32–7.41 (m, 5H, CH Ph, DiPPH), 7.55–7.62 (m, 3H, CH Ph, DiPPH). ¹³C NMR (CD₂Cl₂, T = 253 K, ppm) δ: 23.2 (CH₃, iPr-CH₃), 23.5 (CH₃, iPr-CH₃), 24.8 (CH₃, iPr-CH₃), 25.0 (CH₃, iPr-CH₃), 28.3 (CH, iPr-CH), 28.4 (CH, iPr-CH), 43.3 (CH, Ma), 47.9 (CH, Ma), 59.5 (CH₂, CH₂S), 118.6 (CH, CH=CH Im), 123.5 (CH, CH=CH Im), 124.5 (CH, m-DiPP), 124.8 (CH, m-DiPP), 129.7 (CH, o-Ph-C), 130.0 (CH, p-DiPP), 130.1 (CH, p-Ph-C), 131.4 (C, i-Ph), 132.5 (CH, m-Ph-C), 135.8 (C, i-DiPP), 144.5 (C, o-DiPP-C), 145.3 (C, o-DiPP-C), 171.2 (CO, Ma), 173.7 (CO, Ma), 191.3 (C, NHC). IR (KBr pellet, cm⁻¹) ν_{CO}: 1790, 1720. Anal. Calc. for C₂₇H₃₀N₂O₂PdS: C, 58.64; H, 5.47; N, 5.07. Found: C, 58.77; H, 5.59; N, 5.21%.

4.18. {[1-(2-Pyridyl)methylene-3-methylimidazolyl-2-ene]palladium(0) (maleic anhydride) (**6a**)}

Yellow microcrystals: yield 95%. ¹H NMR (CD₂Cl₂, T = 298 K, ppm) δ: 3.58 (d, J = 3.2 Hz, 1H, CH=CH ma), 3.76 (s, 3H, NCH₃), 4.13 (d, J = 3.2 Hz, 1H, CH=CH ma), 4.98, 5.22 (AB system, 1H, J = 14.4 Hz, SCH₂), 6.95 (d, J = 1.8 Hz, 1H, CH=CH Im), 7.07 (d, J = 1.8 Hz, 1H, CH=CH Im), 7.42 (ddd, J = 7.6, 5.3, 1.7 Hz, 1H, 5-Pyr), 7.51 (d, J = 7.6 Hz, 1H, 3-Pyr), 7.87 (td, J = 7.6, 1.7 Hz, 1H, 4-Pyr), 8.88 (dd, J = 5.3, 0.9 Hz, 1H, 6-Pyr). ¹³C {¹H} NMR (CD₂Cl₂, T = 298 K, ppm) δ: 37.7 (CH₃, NCH₃), 40.1 (CH, CH=CH ma), 41.3 (CH, CH=CH ma), 55.3 (CH₂, NCH₂), 120.7 (CH, CH=CH Im), 120.8 (CH, CH=CH Im), 124.4 (CH, 3-Pyr), 124.7 (CH, 5-Pyr), 138.4 (CH, 4-Pyr), 153.1 (CH, 6-Pyr), 153.2 (C, 2-Pyr), 174.0 (C, C=O ma), 186.0 (C, NCN). IR (KBr pellet, cm⁻¹) ν_{CO}: 1779, 1714. Anal. Calc. for C₁₅H₁₅N₃O₂Pd: C, 47.95; H, 4.02; N, 11.18. Found: C, 47.84; H, 4.18; N, 11.01%.

4.19. {[1-(2-Pyridyl)methylene-3-mesityl]imidazolyl-2-ene]palladium(0) (maleic anhydride) (**6b**)}

Yellow microcrystals: yield 95%. ¹H NMR (CD₂Cl₂, T = 298 K, ppm) δ: 1.94 (bs, 3H, o-mesityl-CH₃), 2.07 (bs, 3H, o-mesityl-

CH₃), 2.40 (s, 3H, p-mesityl-CH₃), 3.12 (bs, 1H, CH=CH ma), 3.31 (bs, 1H, CH=CH ma), 5.13, 5.43 (AB system, J = 14.8 Hz, 1H, NCH₂), 6.94 (d, J = 1.8 Hz, 1H, CH=CH Im), 7.03 (m, 2H, m-mesityl-H), 7.26 (d, J = 1.8 Hz, 1H, CH=CH Im), 7.39 (ddd, J = 7.6, 5.3, 1.7 Hz, 1H, 5-Pyr), 7.56 (d, J = 7.6 Hz, 1H, 5-Pyr), 7.89 (td, J = 7.6, 1.7 Hz, 1H, 4-Pyr), 8.96 (dd, J = 1.0, 5.3 Hz, 1H, 6-Pyr). IR (KBr pellet, cm⁻¹) ν_{CO}: 1785, 1715. Anal. Calc. for C₂₃H₂₃N₃O₂Pd: C, 57.57; H, 4.83; N, 8.76. Found: C, 57.42; H, 4.95; N, 8.64%.

4.20. {[1-(2-Pyridyl)methylene-3-mesityl]imidazolyl-2-ene]palladium(0)(naphthoquinone) (**7b**)}

To a solution of [2-methyl-6-(phenylthiomethyl)pyridine]palladium(0)(naphthoquinone) (0.050 g, 0.10 mmol) in dichloromethane (10 ml) was added a solution of {[1-(2-pyridyl)methylene-3-mesityl]imidazolyl-2-ene}silver bromide (**4b**) (0.046 g, 0.10 mmol) in 10 ml in the same solvent. Immediate precipitation of silver bromide was observed. After a half hour the reaction mixture was filtered through Celite and concentrated to a small volume at reduced pressure. The slow addition of diethylether gave the precipitation of the product as microcrystalline red-orange solid (0.044 g, 82%).

¹H NMR (CD₂Cl₂, T = 253 K, ppm) δ: 1.46 (s, 3H, o-mesityl-CH₃), 1.99 (s, 3H, o-mesityl-CH₃), 2.47 (s, 3H, p-mesityl-CH₃), 4.18, 4.36 (AB system, J = 6.7 Hz, 2H, CH=CH nq), 5.04, 5.24 (AB system, J = 14.6 Hz, 2H, NCH₂), 6.76 (d, J = 1.8 Hz, 1H, CH=CH Im), 6.87 (bs, 1H, m-mesityl-H), 7.07 (bs, 1H, m-mesityl-H), 7.21 (d, J = 1.8 Hz, 1H, CH=CH Im), 7.33–7.50 (m, 5H, 3-Pyr, 5-Pyr, CH-nq), 7.84 (td, J = 7.7, 1.7 Hz, 1H, 4-Pyr), 7.92 (d, J = 7.4 Hz, 1H, CH-nq), 8.47 (d, J = 4.5 Hz, 1H, 6-Pyr).

¹³C NMR (CD₂Cl₂, T = 253 K, ppm) δ: 17.3 (CH₃, o-mesityl), 18.0 (CH₃, o-mesityl), 21.4 (CH₃, p-mesityl), 55.5 (CH₂-Py), 58.7 (CH=CH nq), 65.5 (CH=CH nq), 121.1 (CH=CH Im), 122.0 (CH=CH Im), 123.6 (CH, 3-Py), 124.0 (CH, 5-Py), 125.3 (CH, Nq), 125.5 (CH, Nq), 128.8 (CH, m-mesityl), 129.4 (C, p-mesityl), 130.6 (CH, m-mesityl), 134.9 (C, o-mesityl), 135.2 (CH, Nq), 135.4 (C, o-mesityl), 135.8 (CH, Nq), 137.5 (C, i-mesityl), 138.1 (CH, 4-Py), 138.5 (C, Nq), 150.7 (CH, 6-Py), 152.3 (C, 2-Py), 174.1 (CO), 183.8 (CO), 187.0 (C, NHC). IR (KBr pellet, cm⁻¹) ν_{CO}: 1790, 1720. Anal. Calc. for C₂₈H₂₅N₃O₂Pd: C, 62.06; H, 4.65; N, 7.75. Found: C, 62.14; H, 4.51; N, 7.88%.

The following derivatives were prepared in a similar way using the appropriate silver complex.

4.21. {[1-(2-Pyridyl)methylene-3-methylimidazolyl-2-ene]palladium(0)(naphthoquinone) (**7a**)}

Red-orange microcrystals: yield 64%. ¹H NMR (CD₂Cl₂, T = 253 K, ppm) δ: 3.54 (s, 3H, NCH₃), 4.88, 4.54 (AB system, J = 6.7 Hz, 2H, CH=CH nq), 4.89, 5.17 (AB system, J = 14.5 Hz, 2H, NCH₂), 6.83 (d, J = 1.7 Hz, 1H, CH=CH Im), 7.02 (d, J = 1.7 Hz, 1H, CH=CH Im), 7.43–7.53 (m, 4H, 3-Pyr, 5-Pyr, CH-nq), 7.81 (td, J = 7.6, 1.6 Hz, 1H, 4-Pyr), 8.01–8.04 (m, 2H, CH-nq), 8.08–8.11 (m, 2H, CH-nq), 8.39 (d, J = 5.1 Hz, 1H, 6-Pyr). ¹³C {¹H} NMR (CDCl₃, T = 298 K, ppm) δ: 37.0 (CH₃, NCH₃), 55.3 (CH₂, NCH₂), 61.0 (CH, CH=CH nq), 63.8 (CH, CH=CH nq), 120.4 (CH, CH=CH Im), 121.4 (CH, CH=CH Im), 123.8 (CH, 5-Pyr), 124.0 (CH, CH-nq), 125.3 (CH, 3-Pyr), 130.3 (CH, CH-nq), 130.8 (CH, CH-nq), 137.9 (CH, 4-Pyr), 149.6 (CH, 2-Pyr), 150.9 (CH, 6-Pyr), 180.8 (C, CO), 188.7 (C, NCN). IR (KBr pellet, cm⁻¹) ν_{CO}: 1633. Anal. Calc. for C₂₀H₁₇N₃O₂Pd: C, 54.87; H, 3.91; N, 9.60. Found: C, 54.72; H, 3.79; N, 9.74%.

4.22. {[1-(2-Pyridyl)methylene-3-(2,6-di-i-propylphenyl)imidazolyl-2-ene]palladium(0) (naphthoquinone) (**7c**)}

Red-orange microcrystals, yield 68%. ¹H NMR (CD₂Cl₂, T = 253 K, ppm) δ: 0.20 (d, J = 6.8 Hz, 3H, iPr-CH₃), 0.75 (d, J = 6.8 Hz, 3H,

*i*Pr-CH₃), 1.10 (d, *J* = 6.8 Hz, 3H, *i*Pr-CH₃), 1.50 (d, *J* = 6.8 Hz, 3H, *i*Pr-CH₃), 1.54 (sept, *J* = 6.8 Hz, 1H, *i*Pr-CH), 2.60 (sept, *J* = 6.8 Hz, 1H, *i*Pr-CH), 4.42, 4.46 (AB system, *J* = 6.8 Hz, 2H, CH=CH nq), 5.42, 5.02 (AB system, *J* = 14.4 Hz, 2H, NCH₂), 6.84 (d, *J* = 1.7 Hz, 1H, CH=CH Im), 6.98 (d, *J* = 7.7 Hz, 2H, *m*-aryl-H), 7.19 (d, *J* = 1.7 Hz, 1H, CH=CH Im), 7.31–7.49 (m, 6H, *m*-aryl-H, 3-pyr, 5-Pyr, CH-nq), 7.59 (t, *J* = 7.8 Hz, 1H, *p*-aryl-H), 7.81 (td, *J* = 7.6, 1.5 Hz, 1H, 4-Pyr), 8.02 (d, *J* = 6.9 Hz, 1H, CH-nq), 8.37 (d, *J* = 4.4 Hz, 1H, 6-Pyr), ¹³C {¹H} NMR (CD₂Cl₂, *T* = 253 K, ppm) δ: 22.2 (CH₃, *i*Pr-CH₃), 22.7 (CH₃, *i*Pr-CH₃), 22.2 (CH₃, *i*Pr-CH₃), 24.0 (CH₃, *i*Pr-CH₃), 27.8 (CH, *i*Pr-CH), 28.6 (CH, *i*Pr-CH), 55.6 (CH₂, NCH₂), 57.2 (CH, CH=CH nq), 64.2 (CH, CH=CH nq), 120.8 (CH, CH=CH Im), 123.1 (CH, CH=CH Im), 123.9 (CH, *m*-aryl-CH), 124.0 (CH, CH-nq) 125.4 (CH, 3-Pyr), 125.9 (CH, 5-pyr), 129.4 (CH, *p*-aryl-CH), 130.0 (CH, CH-nq), 130.8 (CH, CH-nq), 135.1 (C, *i*-aryl-C), 135.5 (C, C-nq), 137.2 (C, C-nq), 138.0 (CH, 4-Pyr), 144.9 (C, *o*-aryl-C), 145.2 (C, *o*-aryl-C), 150.1 (C, 6-Pyr), 152.2 (CH, 2-Pyr), 176.7 (C, C=O), 185.4 (C, C=O), 189.1 (C, CN). IR (KBr pellet, cm⁻¹) ν_{CO}: 1641. *Anal.* Calc. for C₃₁H₃₁N₃O₂Pd: C, 63.75; H, 5.35; N, 7.20. Found: C, 63.88; H, 5.49; N, 7.07%.

4.23. {[1-(2-Pyridyl)methylene-3-mesityl]imidazolyl-2-ene}palladium(0)(tetramethyl ethene tetracarboxylate) (**8b**)

To a solution of [2-methyl-6-(phenylthiomethyl)pyridine]palladium(0)(tetramethyl ethene tetracarboxylate) (0.050 g, 0.09 mmol) in dichloromethane (10 ml) was added a solution of {[1-(2-pyridyl)methylene-3-mesityl]imidazolyl-2-ene}silver bromide (**4b**) (0.040 g, 0.09 mmol) in 10 ml of the same solvent. Immediate precipitation of silver bromide was observed. After a half hour the reaction mixture was filtered through a pad of Celite and concentrated to a small volume at reduced pressure. The slow addition of diethyl ether gave the precipitation of the product as microcrystalline yellow solid (0.041 g, 74%). ¹H NMR (CD₂Cl₂, *T* = 298 K, ppm) δ: 2.01 (s, 6H, *o*-mesityl-CH₃), 2.34 (s, 3H, *p*-mesityl-CH₃), 3.29 (s, 6H, OCH₃), 3.49 (s, 6H, OCH₃), 5.29 (s, 2H, NCH₂), 6.77 (d, *J* = 1.8 Hz, 1H, CH=CH Im), 6.95 (s, 2H, *m*-mesityl-H), 7.25 (d, *J* = 1.8 Hz, 1H, CH=CH Im), 7.40 (ddd, *J* = 7.6, 4.3, 1.3 Hz, 1H, 5-Pyr), 7.52 (d, *J* = 7.6 Hz, 1H, 3-Pyr), 7.84 (td, *J* = 7.6, 1.7 Hz, 1H, 4-Pyr), 9.60 (d, *J* = 4.3 Hz, 1H, 6-Pyr), ¹³C {¹H} NMR (CD₂Cl₂, *T* = 298 K, ppm) δ: 17.4 (CH₃, *o*-mesityl-CH₃), 20.6 (CH₃, *p*-mesityl-CH₃), 51.1 (CH₃, OCH₃), 55.6 (CH₂, NCH₂), 120.8 (CH, CH=CH Im), 121.6 (CH, CH=CH Im), 123.7 (CH, 3-Pyr), 124.1 (CH, 5-Pyr), 128.6 (CH, *m*-mesityl-CH), 135.3 (C, *o*-mesityl-C), 135.9 (C, *i*-mesityl-C), 137.8 (CH, 4-Pyr), 137.9 (C, *p*-mesityl-C), 153.4 (C, 2-Pyr), 155.3 (CH, 6-Pyr), 170.5 (C, C=O), 170.6 (C, C=O), 188.6 (C, CN).

IR (KBr pellet, cm⁻¹) ν_{CO} = 1724, 1682. *Anal.* Calc. for C₂₈H₂₃N₃O₈Pd: C, 52.22; H, 4.85; N, 6.53. Found: C, 52.08; H, 4.71; N, 6.41%.

The following derivatives were prepared in a similar way using the appropriate silver complex.

4.24. {[1-(2-Pyridyl)methylene-3-methyl]imidazolyl-2-ene}palladium(0)(tetramethyl ethene tetracarboxylate) (**8a**)

Yellow microcrystals: yield 64%. ¹H NMR (CD₂Cl₂, *T* = 298 K, ppm) δ: 3.60 (s, 6H, OCH₃), 3.63 (s, 6H, OCH₃), 3.76 (s, 3H, NCH₃), 5.12 (s, 2H, NCH₂), 6.92 (d, *J* = 1.8 Hz, 1H, CH=CH Im), 7.06 (d, *J* = 1.8 Hz, 1H, CH=CH Im), 7.41 (ddd, *J* = 7.6, 4.6, 1.0 Hz, 1H, 5-Pyr), 7.47 (d, *J* = 7.6 Hz, 1H, 3-Pyr), 7.84 (td, *J* = 7.6, 1.6 Hz, 1H, 4-Pyr), 9.63 (d, *J* = 4.6 Hz, 1H, 6-Pyr), ¹³C {¹H} NMR (CD₂Cl₂, *T* = 298 K, ppm) δ: 37.4 (CH₃, NCH₃), 51.3 (CH₃, OCH₃), 51.5 (CH₃, OCH₃), 55.2 (CH₂, NCH₂), 120.6 (CH, CH=CH Im), 120.9 (CH, CH=CH Im), 124.0 (CH, 5-Pyr), 124.4 (CH, 3-Pyr), 138.1 (CH, 4-

Pyr), 153.4 (C, 2-Pyr), 155.7 (CH, 6-Pyr), 170.5 (C, C=O), 171.1 (C, C=O), 185.2 (C, CN). IR (KBr pellet, cm⁻¹) ν_{CO}: 1707, 1681. *Anal.* Calc. for C₂₀H₂₃N₃O₈Pd: C, 44.50; H, 4.29; N, 7.78. Found: C, 44.63; H, 4.37; N, 7.65%.

4.25. {[1-(2-Pyridyl)methylene-3-(2,6-di-*i*-propylphenyl)]imidazolyl-2-ene}palladium(0)(tetramethyl ethene tetracarboxylate) (**8c**)

Yellow microcrystals: yield 71%. ¹H NMR (CD₂Cl₂, *T* = 298 K, ppm) δ: 1.01 (d, *J* = 6.8 Hz, 6H, *i*Pr-CH₃), 1.31 (d, *J* = 6.8 Hz, 6H, *i*Pr-CH₃), 2.43 (bm, 2H, *i*Pr-CH), 3.21 (bs, 6H, OCH₃), 3.45 (s, 6H, OCH₃), 5.34 (s, 2H, NCH₂), 6.83 (d, *J* = 1.6 Hz, 1H, CH=CH Im), 7.21 (d, *J* = 1.8 Hz, 1H, CH=CH Im), 7.22 (d, *J* = 7.7 Hz, 2H, *m*-aryl-H), 7.37 (m, 2H, *p*-aryl-H, 5-Pyr), 7.48 (d, *J* = 7.7 Hz, 1H, 3-Pyr), 7.85 (td, *J* = 7.6, 1.5 Hz, 1H, 4-Pyr), 9.48 (d, *J* = 4.4 Hz, 1H, 6-Pyr), ¹³C {¹H} NMR (CD₂Cl₂, *T* = 298 K, ppm) δ: 23.4 (CH₃, *i*Pr-CH₃), 24.7 (CH₃, *i*Pr-CH₃), 28.2 (CH, *i*Pr-CH), 51.3 (CH₃, OCH₃), 51.5 (CH₃, OCH₃), 55.8 (CH₂, NCH₂), 120.2 (CH, CH=CH Im), 123.3 (CH, CH=CH Im), 123.7 (CH, 3-Pyr), 123.7 (CH, *m*-aryl-CH), 124.1 (CH, 5-Pyr), 129.1 (CH, *p*-aryl-CH), 135.5 (C, *i*-aryl-C), 137.5 (CH, 4-Pyr), 146.0 (C, *o*-aryl-C), 153.5 (C, 2-Pyr), 155.3 (CH, 6-Pyr), 170.3 (C, C=O), 170.8 (C, C=O), 191.1 (C, CN). ν_{CO}: 1729, 1699, 1680.

Anal. Calc. for C₃₁H₃₇N₃O₈Pd: C, 54.27; H, 5.44; N, 6.12. Found: C, 54.38; H, 5.38; N, 6.19%.

4.26. {κ¹-C-[1-(Phenylthio)methylene-3-methyl]imidazolyl-2-ene}{κ¹-C-[1-(2-pyridyl)methylene-3-mesityl]imidazolyl-2-ene}palladium(0)(maleic anhydride) (**9**)

To a solution of {[1-(2-pyridyl)methylene-3-mesityl]imidazolyl-2-ene}palladium(0)(maleic anhydride) (**6b**) (0.064 g, 0.133 mmol) in dichloromethane (10 ml) was added a solution of {[1-(phenylthio)methylene-3-methyl]imidazolyl-2-ene}silver bromide (**2a**) (0.052 g, 0.133 mmol) in 10 ml of the same solvent. Immediate precipitation of silver bromide was observed. After a half hour the reaction mixture was filtered through a Millipore filter yielding a clear solution. Reduction to small volume (~3 ml) and addition of diethylether gave the products as microcrystalline yellow solid after cooling to 0 °C (0.062 g, 67%). The same product was obtained starting from {[1-(2-pyridyl)methylene-3-mesityl]imidazolyl-2-ene}silver bromide (**4b**) and {[1-(phenylthio)methylene-3-methyl]imidazolyl-2-ene} (**5a**). ¹H NMR (CD₂Cl₂, *T* = 298 K, ppm) δ: 1.62 (s, 3H, *p*-mesityl-CH₃), 1.85 (s, 3H, *o*-mesityl-CH₃), 2.34 (s, 3H, *o*-mesityl-CH₃), 2.89 (s, 3H, NCH₃), 3.29, 3.37 (AB system, *J* = 3.7 Hz, 2H, CH=CH ma) 4.81, 5.06 (AB system, *J* = 14.1 Hz, 2H, CH₂-Pyr), 5.59, 5.84 (AB system, *J* = 15.5 Hz, 2H, SCH₂) 6.7 (d, *J* = 1.9 Hz, 1H, CH=CH Im), 6.81 (d, *J* = 1.9 Hz, 1H, CH=CH Im), 6.83 (s, 1H, *m*-mesityl-H), 6.87 (d, *J* = 1.9 Hz, 1H, CH=CH Im), 6.95 (s, 1H, *m*-mesityl-H), 7.25 (d, *J* = 1.9 Hz, 1H, CH=CH Im), 7.27–7.34 (m, 6H, S-Ph 5Pyr) 7.41 (d, *J* = 7.9 Hz, 1H, 3-Pyr), 7.79 (td, *J* = 7.7, 1.9 Hz, 1H, 4-Pyr), 8.63 (d, *J* = 4.0 Hz, 1H, 6-Pyr). ¹³C {¹H} NMR (CD₂Cl₂, *T* = 298 K, ppm) δ: 16.7 (CH₃, *p*-mesityl-CH₃), 16.8 (CH₃, *p*-mesityl-CH₃), 20.57 (CH₃, *o*-mesityl-CH₃), 36.4 (CH₃, NCH₃), 38.2 (CH, CH=CH ma), 38.7 (CH, CH=CH ma), 54.1 (CH₂, CH₂-Pyr), 56.5 (CH₂, SCH₂), 119.1 (CH, CH=CH Im), 121.3 (CH, 5-Pyr), 121.9 (CH, CH=CH Im), 122.4 (CH, CH=CH Im), 122.8 (CH, 3-Pyr), 123.0 (CH, CH=CH Im), 128.3 (CH, *m*-mesityl), 128.8 (CH, *p*-Ph), 129.1 (C, *p*-mesityl-C), 129.2 (CH, *o*-Ph), 131.6 (C, *i*-Ph), 133.4 (CH, *m*-Ph), 136.0 (C, *o*-mesityl-C), 136.2 (C, *o*-mesityl-C), 137.1 (CH, 4-Pyr), 138.4 (C, *i*-mesityl-C), 149.4 (CH, 6-Pyr), 156.5 (C, 2-Pyr), 174.0 (C, C=O), 174.4 (C, C=O), 189.2 (C, CN), 189.3 (C, CN). IR (KBr pellet, cm⁻¹) ν_{CO}: 1765, 1704. *Anal.* Calc. for C₄₂H₄₃N₅O₂PdS: C, 63.99; H, 5.50; N, 8.88. Found: C, 63.89; H, 5.58; N, 8.81%.

4.27. Bis- $\{\kappa^1\text{-C-[1-(phenylthio)methylene-3-methylimidazolyl-2-ene]\}_2$ palladium(0)(maleic anhydride) (**10**)

To a solution of $\{[1-(phenylthio)methylene-3-methylimidazolyl-2-ene]\text{palladium(0)(maleic anhydride)}\}$ (**5a**) (0.025 g, 0.061 mmol) in dichloromethane (10 ml) was added a solution of $\{[1-(phenylthio)methylene-3-methylimidazolyl-2-ene]\}$ silver bromide (**2a**) (0.024 g, 0.061 mmol) in 10 ml of the same solvent. Immediate precipitation of silver bromide was observed. After a half hour the reaction mixture was filtered through a Millipore filter yielding a clear solution. Reduction to small volume (~ 3 mL) and addition of diethylether gave the product as microcrystalline yellow solids after cooling to 0 °C. (0.031 g, 82%). $^1\text{H NMR}$ (CD_2Cl_2 , $T = 298$ K, ppm) δ : 3.54 (s, 2H, HC=CH ma), 3.69 (s, 6H, NCH₃), 5.46, 5.33 (AB system, $J = 14.3$ Hz, 2H, SCH₂), 6.90 (bs, 4H, HC=CH Im), 7.32–7.40 (m, 10H, SPh-CH). $^{13}\text{C}\{^1\text{H}\}$ NMR (CD_2Cl_2 , $T = 298$ K, ppm) δ : 37.9 (CH₃, NCH₃), 38.7 (CH, CH=CH ma), 55.0 (CH₂, NCH₂), 120.1 (CH, CH=CH Im), 122.4 (CH, CH=CH Im), 127.9 (CH, SPh-CH), 129.2 (CH, SPh-CH), 131.7 (CH, SPh-CH), 174.2 (C, C=O), 188.9 (C, NCN). *Anal. Calc.* for $\text{C}_{43}\text{H}_{44}\text{N}_4\text{O}_2\text{PdS}_2$: C, 63.03; H, 5.41; N, 6.84. Found: C, 63.01; H, 5.48; N, 6.79%.

4.28. Equilibrium reactions between silver(I)- and palladium(0)-NHC derivatives

To a 1.5×10^{-2} M solution in CD_2Cl_2 of palladium(0)NHC complexes **6a** or **6b**, equimolecular amounts of the silver derivatives **4a** or **4b** were added at RT and the $^1\text{H NMR}$ spectra of the ensuing reaction mixtures were recorded.

4.29. Reaction between palladium(0)NHC derivatives and dmbd

To a 1.5×10^{-2} M CD_2Cl_2 solution of palladium(0)NHC of complexes **5a**, **5b**, **6a** or **6b** a stoichiometric amount of dmbd ([complex]: [dmbd] \approx 1:2) was added at 253 K and the $^1\text{H NMR}$ spectra of the ensuing reaction mixtures were recorded. In the RT ^1H and ^{13}C NMR spectra in CDCl_3 of the reaction mixture between **6a** and dmbd the formation of species **12** was observed. $^1\text{H NMR}$ (CDCl_3 , $T = 298$ K, ppm) δ : 3.40 (s, 3H, OCH₃), 3.42 (s, 3H, OCH₃), 3.68 (s, 3H, OCH₃), 3.72 (s, 3H, NCH₃), 3.73 (s, 3H, OCH₃), 5.07 (bd, 1H, NCH₂), 5.83 (bd, 1H, NCH₂), 6.78 (d, $J = 1.5$ Hz, 1H, CH=CH Im), 7.08 (d, $J = 1.5$ Hz, 1H, CH=CH Im), 7.33 (ddd, $J = 7.7$, 5.4, 1.0 Hz, 1H, 5-Pyr), 7.56 (d, $J = 7.7$ Hz, 1H, 3-Pyr), 7.84 (td, $J = 7.7$, 1.7 Hz, 1H, 4-Pyr), 8.66 (d, $J = 5.4$ Hz, 1H, 6-Pyr). $^{13}\text{C}\{^1\text{H}\}$ NMR (CDCl_3 , $T = 298$ K, ppm) δ : 37.4 (CH₃, NCH₃), 50.3 (CH₃, OCH₃), 50.9 (CH₃, OCH₃), 51.2 (CH₃, OCH₃), 51.2 (CH₃, OCH₃), 56.5 (CH₂, NCH₂), 121.3 (CH, CH=CH Im), 121.4 (CH, CH=CH Im), 124.1 (CH, 5-Pyr), 124.7 (CH, 3-Pyr), 138.5 (CH, 4-Pyr), 141.1 (C, C=C), 148.4 (C, C=C), 152.7 (C, 6-Pyr), 154.1 (CH, 2-Pyr), 164.3 (C, C=O), 166.1 (C, C=O), 175.1 (C, C=O), 175.7 (C, C=O), 177.5 (C, C=C), 180.5 (C, NCN). *Anal. Calc.* for $\text{C}_{22}\text{H}_{23}\text{N}_3\text{O}_8\text{Pd}$: C, 46.86; H, 4.11; N, 7.45. Found: C, 46.53; H, 4.05; N, 7.38%.

Appendix A. Supplementary material

Supplementary data associated with this article can be found, in the online version, at <http://dx.doi.org/10.1016/j.ica.2012.04.018>.

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