Alkali-Metal-Mediated Manganation(II) of Functionalized Arenes and Applications of ortho-Manganated Products in Pd-Catalyzed Cross-Coupling Reactions with Iodobenzene

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Abstract: Extending the recently introduced concept of “alkali-metal-mediated manganation” to functionalised arenes, the heteroleptic sodium manganate reagent [(tmeda)Na(tmp)(R)Mn-(tmp)] (1; TMEDA = N,N,N',N'-tetramethylethylenediamine, TMP = 2,2,6,6-tetramethylpiperidide, R = CH₂SiMe₃) has been treated with anisole or N,N-dimethylbenzamide in a 1:1 stoichiometry in hexane. These reactions afforded the crystalline products [(tmeda)Na(tmp)(o-C₆H₄OMe)Mn-(tmp)] (2) and [(tmeda)Na(tmp)(o-(C(O)N(iPr)₂C₆H₄)Mn(CH₂SiMe₃)) (3), respectively, as determined by X-ray crystallographic studies. On the basis of these products, it can be surmised that reagent 1 has acted, at least partially and ultimately, as an alkyl base in the first reaction liberating the silane Me₃Si, but as an amido base in the second reaction liberating the amine TMPH. Both of these paramagnetic products 2 and 3 have contacted ion-pair structures, the key features of which are six-atom, five-element (NaN₄CCO) and seven-atom, five-element (NaNMnCCO) rings, respectively. Manganates 2 and 3 were successfully cross-coupled with iodobenzene under [PdCl₂(dppf)] (dppf = 1,1'-bis(diphenylphosphino)ferrocene) catalysis to generate unsymmetrical biaryl compounds in yields of 98.0 and 66.2 %, respectively. Emphasizing the importance of alkali-metal mediation in these manganation reactions, the bisalkyl Mn reagent on its own fails to metatalate the said benzamide, but instead produces the monomeric, donor-acceptor complex [Mn(R)₂{(iPr)₂NC(Ph)(=O)}₂] (5), which has also been crystallographically characterised. During one attempt to repeat the synthesis of 2, the butoxide-contaminated complex {[(tmeda)Na(R)(OBu)(o-C₆H₄OMe)Mn]} (6) was obtained. In contrast to 2 and 3, due to reduced sterical constraints, this complex adopts a dimeric arrangement in the crystal, the centrepiece of which is a twelve atom (NaOCCMnO)₆ ring.

Keywords: aromatic compounds · cross-coupling · manganese · manganation · sodium

Introduction

Recently we added manganese(II) to the growing list of unlikely metals that can perform direct metalation of aromatic compounds, when driven by alkali-metal mediation. Such metalations (metal–hydrogen exchange reactions) are usually the domain of highly reactive, highly polar organometallics, most typically alkylithium or lithium amide compounds.[1] Previously, to bind MnII to a carbon atom of an aromatic compound would normally require a prelithiation step, followed by a salt metathesis (commonly with a MnII halide).[3] However, despite its widespread applicability, this two-step approach is not all-conquering as lithiation suffers from limited functional group tolerance and low kinetic stability, while the ionicity of metal halide salts can lead to solubility problems in common organic solvents. On their own, organomanganese(II) complexes, as comparatively low-polarity organometallics, are generally too weakly basic to effect C–H deprotonation at a reasonable rate, but when paired with an alkali metal in a heterobimetallic ate, they can transform into super-manganating reagents. This alkali-

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metal-mediated manganation (AMMMn) and alkali-metal-mediated manganation (magnesiation, zinication, aluminiation)\(^9\) in general can potentially circumvent the obstacles stated above, but they are profoundly more than direct alternative methods to indirect metathetical manganations (magnesiations etc.), being synergic metalations capable of delivering unprecedented products and structures inaccessible by means of metathesis. The initial examples of AMMMn demonstrated this emphatically, producing in the case of benzene, the tetrasmuth dimanganese amido inverse crown with a benzene diide core \([\text{Na}_4\text{Mn}_{2}(\text{tmp})_6\text{C}_6\text{H}_4\text{]}}\),\(^7\) (TMP = 2,2,6,6-tetramethylpiperidide) and with ferrocene, the dilithium dimanganese trinuclear ferrocenophane \([\text{tmeda})_2\text{Li}_2\text{Mn}_2\{\text{Fe}(\text{C}_5\text{H}_4)_3\}]\) (TMEDA = \(N\_N\_N\_N\_\text{-tetramethylethylenediamine}\)).\(^7\) Describing here the first study in which AMMMn has been applied to functionalised arenes, we report the successful cross-coupled with iodobenzene under palladium, the first structures of their type, and show how they can be cross-coupled with iodobenzene under palladium catalysis to generate unsymmetrical biaryl.

### Results and Discussion

The sodium monoalkyl-bisamidomanganate \([\text{tmeda})\text{Na}-(\text{tmp})(\text{R})\text{Mn}(\text{tmp})]\) (1; \(\text{R} = \text{CH}_2\text{SiMe}_3\)) was used as the AMMMn reagent in the new reactions. Only one previous reaction of 1 has been reported,\(^8\) namely with benzene, in which it acted as an alkyl (R) base to produce the phenyl derivative \([\text{tmeda})\text{Na}(\text{tmp})(\text{Ph})\text{Mn}(\text{tmp})]\) and tetrakisilane. Here, prepared in situ in hexane, 1 was treated with a molar equivalent of the appropriate functionalised arene (anisole or \(N\_N\_\text{-disopropylbenzamide};\) Scheme 1) selected on the basis of their respective contrasting weak and strong DoM ability.\(^9\) Both reactions afforded a crystalline product in yellowish-green \([\text{tmeda})\text{Na}(\text{tmp})(\text{o-C}_6\text{H}_4\text{OMe})\text{Mn}(\text{tmp})]\) (2) and orange \([\text{tmeda})\text{Na}(\text{tmp})(\text{o-C}_6\text{H}_4\text{N}-(\text{iPr}))\text{Mn}(\text{tmp})]\) (3), respectively. Isolated yields were 66 and 31 %, respectively.

In both 2 and 3, the arene molecules have been selectively ortho-deprotonated, with a Mn\(^{1+}\) atom occupying the vacated hydrogen site (see structures below), thus, to the best of our knowledge these syntheses represent the first examples of directed ortho-manganations of substituted arenes.\(^7\) However, there is a major distinction between 2 and 3. Following the benzene precedent, 1 operates ultimately as an alkyl base towards anisole, generating tetrakisilane and 2, which consequently has a monoaryl-bisamido composition. In contrast, 1 exhibits amido basicity towards the tertiary amide, leading to the monoalkyl-monoaryl-monoamido 3 with concomitant elimination of TMPh. This unexpected result establishes that 1 is a versatile manganating reagent with a basicity switch (potentially tuneable to alkyl or amido) seemingly dependent on the nature of the aromatic substrate. Clearly distinct mechanisms must therefore be available to 1 in its reactions with anisole and \(N\_N\_\text{-disopropylbenzamide}.\) Of course, at this stage all we can definitely conclude from structurally characterizing the products from these reactions is that manganating reagent 1 functions ultimately as an alkyl base towards anisole and as an amido base towards benzamide. Thermodynamically the loss of volatile Me\(_4\)Si from such manganations is likely to be greatly preferred. However, as recently discussed in the context of alkyl–Zn–zincates,\(^8\) there may be an additional intermediate step in which TMP acts firstly as the base, but then re-enters the coordination sphere of the alkali metal as TMP(H) ligand, before reacting with the alkyl ligand to eliminate alkane and to restore the TMP anion to the heterobimetalllic structure. Thus the distinction between 2 and 3 may be attributed to kinetic factors. Intriguingly, subjecting the same tertiary amide to the related zincate base \([\text{tmeda})\text{Na}(\text{tmp})(\text{Bu})\text{Zn}(\text{Bu})]\), induces ortho-zincation ultimately not through amido (TMP) basicity, but through alkyl (Bu) basicity.\(^9\) Thus the basicity switch in these synergic metalations is also dependent on the identity of the o-bonding divalent metal (and coligand set) within the base, as changing these parameters will also change the kinetic profile of the metalation reaction.

Due to the paramagnetic nature of 2 and 3, NMR spectroscopy could not be used for their characterisation. We therefore turned to X-ray crystallography. The molecular structures of 2 and 3 (Figure 1 and Table 1) share common

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**Scheme 1. Ambibasic behaviour of reagent 1 towards anisole and \(N\_N\_\text{-disopropylbenzamide affording the ortho-manganated products 2 and 3, respectively.**
and only one of two crystallographically independent molecules is shown.

Figure 1. Molecular structures of 2 (top) and 3 (bottom) with 30% probability displacement ellipsoids. H atoms have been omitted for clarity. In addition for 3, minor disorder components have been omitted for clarity, and only one of two crystallographically independent molecules is shown.

general features. Classifiable as contacted ion pairs, both structures display a trigonal planar Mn coordination (N,N,C in 2; C,N,C in 3) comprising one terminal (TMP in 2, R in 3) and two bridging ligands (TMP in both; o-deprotonated anisole and benzamide in 2 and 3, respectively). Defining the metalations as manganations, the deprotonated ortho-C atoms form o bonds with the Mn atoms, but do not interact with the Na atoms. Instead the aryl ligands bridge to Na through their O-heteroatoms to close six-membered (NaNMnCCO) and seven-membered (NaNMnCCCO) rings in 2 and 3, respectively, with bidentate TMEDA ligands completing the distorted tetrahedral (N,N,O) environments of Na. A search of the Cambridge Structural Database\textsuperscript{[19]} revealed 2 and 3 to be unique with no hits found for ether- or amide-substituted aryls with ortho-Mn\textsuperscript{II} atoms\textsuperscript{[11]} or indeed in the latter case with any ortho-transition-metal atoms. Furthermore alkali-metal-aryl-manganated structures of any type are surprisingly rare with no sodium examples at all (besides those previously made by AMMMn\textsuperscript{[1,5]}). Power has reported\textsuperscript{[12]} three lithium examples, significantly all synthesised from metathesis reactions, and only one, [Li(thf)]\textsubscript{2}[MnMes\textsubscript{3}], has trigonal planar Mn, but in a very different (cf. 2 and 3) solvent-separated ion-pair arrangement. Unfortunately, disorder within 3 limits the precision of its metrical parameters, though the connectivities are definite, and the main features of its two independent molecules are similar to those of 2. The best comparison with 2 is provided by the unsubstituted phenyl analogue [(tmp)(Ph)Mn]\textsubscript{4} (4). Lack of aryl substitution gives 4 a smaller four-membered (NaNMnC) ring with a modestly longer Mn–C bond (2.207(4) \textgreek{Å} to ipso-C cf. 2.189(2) \textgreek{Å} to ortho-C in 2) made by the Mn atom lying almost coplanar with the aryl ring plane (deviation 0.339 \textgreek{Å} cf. 0.558 \textgreek{Å} in 2; 0.221 and 0.220 \textgreek{Å} for the two independent molecules of 3). With respect to the MnN\textsubscript{2}C planes, the aryl rings have dihedral angles of exactly 90\textdegree{} by symmetry in 4 and 71.4\textdegree{} in 2. A long Na–ipso-C bond in 4 (2.731(4) \textgreek{Å}) is replaced by a significantly shorter Na–O(Me) bond in 2 (2.5357(16) \textgreek{Å}) reflecting the greater bond strength of donor–acceptor dative interactions versus cation–π interactions;\textsuperscript{[13]} the Na–O bonds in 3 are even shorter, at 2.313(6) and 2.309(5) \textgreek{Å}.

It is worth emphasising that the organomanganese reagent Mn(CH\textsubscript{3}SiMe\textsubscript{3})\textsubscript{3} on its own failed to metalate either anisole or N,N-diisopropylbenzamide and did not act as a nucleophile towards the latter. Thus, no reaction was observed at all when the bisalkyl reagent was mixed with anisole, whereas the reaction with N,N-diisopropylbenzamide afforded the Lewis acid/Lewis base association complex [Mn-
The molecular structure of 5 (Figure 2 and Table 1) has a Mn atom in a distorted tetrahedral environment made up of two alkyl groups and two neutral benzamide ligands, which coordinate to the metal through their oxygen atoms. To the best of our knowledge the only precedent for a crystallographically characterised homometallic[14] compound containing a neutral tertiary aromatic amide coordinated to a metal is that of the related manganese bis(diisopropylbenzamide).[15] The Mn–C bond lengths in 5 (2.159(2), 2.165(2) Å) are nearly identical to those found in the related monomeric complex [Mn(CH₂SiMe₃)₂][(-)-sparteine][16] (2.158(2), 2.165(17) Å). In addition, the Mn–O bond length (2.172(4), 2.1415(15) Å) are similar to that in the THF adduct of the related manganese bis(alkyl) species [Mn(CH₂SiMe₃)₂](thf)][17] (2.19(2) Å), although in the latter the Mn centre is tricoordinate due to the heavier silylation of the ligand, whereas in 5 there are four ligands around the metal. The structure of the unsolvated manganese compound is known to be a polymer,[18] thus the formation of 5 illustrates the superior Lewis basicity of N,N-disopropylbenzamide relative to that of anisole, which cannot accomplish the same cleavage of polymeric [Mn(CH₂SiMe₃)₂]₆. Cocomplexation of the bis(alkyl)manganese reagent, or fragment thereof, with an alkali metal amide leads to a discrete molecular structure as seen with 1, hence the problem of having to cleave a polymeric structure is not an issue. This is one advantageous factor of the mixed-metal synergy inherent in reagents such as 1.

To the best of our knowledge alkali metal arylmanganates have not been previously studied in the context of transition-metal-catalysed cross-coupling reactions with aryl halides. Precedents exist for organomanganese(II) halides of general formula ArMnX in excellent work reported by Cahiez.[19] Since 2 and 3 have covalent Mn–C (aryl) bonds within anionic ate-activated structures, we reasoned they could make excellent coupling agents as playing only a secondary, Lewis acidic role in the structures, sodium should not be detrimental even though ionic arylsodium reagents are not generally suitable for coupling applications. Test reactions with iodobenzene in hexane in the presence of 2% [PdCl₂(dppf)] (Scheme 3; dppf = 1,1’-bis(diphenylphosphino)ferrocene) confirmed this analysis. In the case of 2, the coupled product 2-methoxybiphenyl was obtained in an impressive isolated yield of 98.0%. Coupling was also successful with 3, though the yield of isolated N,N-disopropyl-2-phenylbenzamide was lower at 66.2%. Interestingly when the experiment with 2 was repeated in the absence of the Pd catalyst direct coupling was still observed albeit in a reduced yield (32.0%). The same catalyst-free experiment with 3 was even more successful with a less reduced yield of 47%.

High purity in the reagents employed and complete exclusion of oxygen in the system are crucially important for the success of these metalation applications. Thus, when the reaction that afforded 2 was repeated with a sample of n-butyllithium contaminated by n-butoxide, compound [(tmeda)Na(R)(OBu)(o-C₆H₄OMe)Mn₂][20] (6) was serendipitously obtained (Scheme 4). This new heterotrianionic com-

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**Scheme 2. Contrasting reactivity of [Mn(CH₂SiMe₃)₂]₆ towards anisole and N,N-disopropylbenzamide.**

![Scheme 2](image)

**Figure 2. Molecular structure of 5 with 30% probability displacement ellipsoids. H atoms and minor disordering have been omitted for clarity.**

**Figure 2.**

**Scheme 3. Palladium-catalysed cross-coupling reactions of 2 and 3 with iodobenzene.**

![Scheme 3](image)
pound bears a resemblance to 2, as both contain a molecule of anisole selectively ortho-manganated and a \([\text{Na(tmeda)}]^+\) ion. However they differ in the rest of the anionic ligands supporting the Mn atom. Thus, for 2 two TMP ligands are found, whereas in 6 there is an alkyl and a butoxide anion. A plausible pathway that could account for the synthesis of 6 would involve the initial formation of a heteroleptic manganate species \([(\text{tmeda})\text{Na}\text{Mn}O\text{Bu}]\) that would react with anisole as an amido base liberating TMPH as coproduct of the reaction. However we must mention that all the attempts to prepare this potentially new mixed-metal reagent have been unsuccessful to date.

The centrosymmetric molecular structure of 6 (Figure 3 and Table 1) was successfully determined by X-ray crystallographic studies. Unlike 2 and 3 which are monomeric structures, 6 comprises two identical bridging cationic \([(\text{tmeda})\text{Na}\text{Mn}O\text{Bu}](\text{o-HOMe})(\text{CH}_2\text{SiMe}_3))\) units dimerised through alkoxide and alkyl bridges. This structure can be envisaged as a cationic twelve-membered \([(\text{NaOCCMnC})_2]^+\) ring hosting in its interior two butoxide ligands (Figure 4). This structural motif is reminiscent of that previously found for the series of inverse crown complexes \([(\text{M}\text{I}\text{M}\text{II})(\text{NiPr}_2)_2X_2](\text{M}\text{I}=\text{Li or Na}; \text{M}\text{II}=\text{Mg}; \text{X}=\text{OR, H})]\) (Figure 4), and for several other homometallic and heterometallic systems. These compounds exhibit a cationic eight-membered ring hosting two anionic ligands. Due to the am-bidentate nature of the ortho-metalated anisole ligand, C-bonding to Mn and O-bonding to Na, an expansion in ring size from eight atoms in 7 to twelve atoms in 6 is observed.

This 12-membered ring adopts a pseudo-chair conformation (Figure 5) with the sodium atoms displaced on either side of the plane defined by \(\text{C}(1)\text{Mn}(1)\text{C}(18)\)...\(\text{C}(1)\text{Mn}(1)\text{C}(18)\). The oxygen atoms of the anisole groups are also slightly out of this plane as shown by the torsion angle \(\text{O}(1)\text{-C}(2)\text{-C}(1)\text{-Mn}(1)\), 17.1°. The sodium atoms achieve pentacoordination by bonding to two anionic (C of the alkyl; O of the alkoxide) ligands, the neutral oxygen atom of the metalated anisole, and the two TMEDA nitrogen atoms. There is a slight contraction of the \(\text{Na}-\text{O}('\text{Bu})\) bond length (2.2982(18) Å) relative to that for the OMe group of the anisole ligand (2.3997(18) Å), reflecting the anionic nature of the former even though the latter oxygen atom has a lower coordination number. Surrounded by four anionic ligands, the Mn atom has a distorted tetrahedral en-
environment comprising two OnBu, one ortho-aryl C and one alkyl C atoms. The Mn–O bond lengths (2.1333(15) and 2.1318(16) Å) are modestly longer than the ones found in the mixed Li/Mn alkoxides [Li[Mn(N(SiMe)$_3$)$_2$-(OCBu)$_2$]] (2.019(4), 1.984(4) Å) and [Li$_2$[MnBr$_2$-(OCBu)$_2$](thf)$_2$)] (2.019(7) Å), which can be attributed to the dimeric nature of 6, in which each butoxide ligand is tri-coordinated, whereas the other structures have two-coordinate alkoxide ligands being monomers. The Mn–C(anisole) (2.177(2) Å) bond length is similar to the one found in 2 (2.189(2) Å).

**Conclusion**

In conclusion, the new concept of AMMMn has been successfully applied to arenes functionalised with electron-donating (MeO) or electron-withdrawing [(C)(O)N(ipr)$_2$] groups. Probing the structural changes that accompany the Mn$^{II}$–H exchange reactions revealed that the mixed-metal reagent can function ultimately as an alkyl or amido base depending on the specific arene. When the alkali metal is absent from the solution mixture, the bisalkyl Mn reagent can no longer metalate the benzamide, but instead forms a coordination complex with it. With sodium tamed within a cage of heteroatoms distant from the active carbanion–Mn centre, the bimetallic products of AMMMn can in turn be successfully cross-coupled with iodobenzene to form unsymmetrical biaryls. Overall these results serve to broaden significantly the opportunities for synthesizing new compounds and building new structures in organomanganese(II) chemistry. The sensitivity of these mixed-metal organometallic compounds to air and moisture was demonstrated through the serendipitous synthesis of an n-butoxy derivative. The adventitious presence of the nBuO anion, which has good bridging capabilities and is not sterically demanding, leads to an increase in aggregation number (monomer to dimer) of the mixed-metal complex and to coordination expansion about the manganese atom (trigonal planar to tetrahedral).

**Experimental Section**

All reactions were carried out under a protective argon atmosphere by using standard Schlenk techniques. Hexane was distilled from sodium/benzophenone. $^1$H NMR spectra were recorded on a Bruker DPX 400 MHz spectrometer; data for X-ray crystal structure determination were obtained with a Nonius Kappa CCD and a Bruker SMART 1 K CCD diffractometer using graphite monochromated Mo K$_\alpha$ radiation ($\lambda = 0.71073$ Å). Selected crystallographic data for compounds 2, 3, 5 and 6 are given in Table 2. CCDC 656944 (2), 656945 (3), 663055 (5) and 661429 (6) contain the supplementary crystallographic data for this paper. These data can be obtained free of charge from The Cambridge Crystallographic Data Centre via www.ccdc.cam.ac.uk/data_request/cif. The IR-spectra were recorded on a Nicolet Avatar 360 FT-IR spectrometer and elemental analyses were carried out on a Perkin Elmer 2400 elemental analyser. Melting/decomposition points were measured with a Büchi Melting Point B-545 apparatus. 

Table 2. Selected crystallographic data for compounds 2, 3, 5 and 6.$^a$

| formula    | M$_r$ [g mol$^{-1}$] | crystal system | space group   | a [Å]        | b [Å]     | c [Å]   | $\alpha$ [°] | $\beta$ [°] | $\gamma$ [°] | V [Å$^3$] | $Z$ | $\rho_{calc}$ [g cm$^{-3}$] | $\rho$(Mo$_{gon}$) [mm$^{-3}$] | measured refins | independent refins | observed refins$^b$ | parameters | $R$1[$I$] (R1 all data) | wR2$^{0.2}$ (wR2 all data) | max/min peaks [e Å$^{-1}$] |
|------------|----------------------|----------------|---------------|--------------|-----------|--------|-------------|-------------|-------------|---------|-----|-----------------------------|--------------------------|----------------|------------------------|-----------------------------|----------------|----------------------|-------------------------|------------------|
| C$_8$H$_6$Mn$_2$Na$_2$O   | 581.75               | triclinic      | $P1$          | 10.059(3)    | 11.212(4) | 15.952(5) | 75.986(5)  | 74.632(5)  | 87.910(5)  | 1682.3(9) | 2   | 1.14                             | 0.433                      | 11.833          | 5325                   | 356                         | 0.039             | 0.0959                | +0.84 / –0.40          |
| C$_8$H$_6$Mn$_2$Na$_2$O$_2$S   | 625.88               | orthorhombic   | $Pca2_1$      | 21.854(4)    | 31.590(8) | 20.736(6) | 90.000(5)  | 90.000(5)  | 90.000(5)  | 7759(3)  | 8   | 1.072                            | 0.409                      | 17286           | 1085                   | 8970                        | 0.054             | 0.1299                | +0.49 / –0.29          |
| C$_8$H$_6$Mn$_2$Na$_2$O$_2$Si | 639.96               | triclinic      | $P1$          | 9.9826(3)    | 10.9135(3) | 20.2736(6) | 100.34(5)  | 95.14(5)   | 113.836(2) | 1954.91(10) | 2   | 1.087                            | 0.427                      | 39613            | 405                    | 5987                        | 0.0469            | 0.0859                | +0.609 / –0.398        |
| C$_8$H$_6$Mn$_2$Na$_2$O$_2$Si$_2$ | 923.19               | triclinic      | $P1$          | 10.949(2)    | 11.583(3) | 12.071(3)  | 72.369(3)  | 67.101(3)  | 84.063(3)  | 1343.8(5) | 1   | 0.609                            | 0.568                      | 7472             | 4469                   | 3968                        | 0.0378            | 0.1062                | +0.55 / –0.40          |

[a] All data were collected at 150 K using Mo$_{gon}$( $\lambda = 0.71073$ Å) radiation. [b] Observation criterion: $I > 2\sigma(I)$. [c] $R1 = \Sigma ||F_o|-|F_e||/\Sigma |F_o|$. [d] wR2 = $[\Sigma (wF_o^2-F_e^2)^2]/[\Sigma (wF_o^2)]^{1/2}$.
situ by reaction of BuNa (0.16 g, 2 mmol), TMP(H) (0.68 mL, 4 mmol), Mn(CH₂SiMe₃)₂ (0.46 g, 2 mmol) and TEMEDA (0.30 mL, 2 mmol). N,N-diisopropylbenzamide (0.41 g, 2 mmol) was then introduced and the mixture was stirred for 18 h affording a bright orange solution that was filtered and concentrated by removing some solvent under vacuum. Storage of this filtrate at room temperature afforded a crop of orange crystals (0.41 g, 31%) after two days; one of them was employed in an X-ray diffraction experiment. M.p. 148°C (decomp); IR (nujol): v = 1592 cm⁻¹ (C=O); elemental analysis caled (%) for C₁₄H₁₈Mn₂Na₂O₄Si₂ (625.88): C 61.41, H 10.15, N 8.95; found: C 61.40, H 10.08, N 8.45.

**Cross-coupling of 2 with isodobenzene:** Isodobenzene (0.44 mL, 4.0 mmol) was added to a crude hexane solution of 2 (0.2 mmol), followed by [PdCl₂(dppf)] (32.6 mg, 0.04 mmol). After stirring for 18 h under reflux conditions, the mixture was quenched with a saturated NH₄Cl solution (10 mL) and filtered. The yellow filtrate was concentrated by removing some solvent under vacuum. Storage of this filtrate at room temperature afforded a crop of orange crystals (0.71 g, 31%) after two days; one of them was employed in an X-ray diffraction experiment. M.p. 159°C (decomp); IR (nujol): v = 1592 cm⁻¹ (C=O); elemental analysis caled (%) for C₁₄H₁₈Mn₂Na₂O₄Si₂ (625.88): C 61.41, H 10.15, N 8.95; found: C 61.40, H 10.08, N 8.45.

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