Manganese-Catalyzed Cross-Coupling of Aryl Halides and Grignard Reagents by a Radical Mechanism

Antonacci, Giuseppe; Ahlburg, Andreas; Fristrup, Peter; Norrby, Per-Ola; Madsen, Robert

Published in:
European Journal of Organic Chemistry

Link to article, DOI:
10.1002/ejoc.201700981

Publication date:
2017

Document Version
Peer reviewed version

Link back to DTU Orbit

Citation (APA):
Manganese-Catalyzed Cross Coupling of Aryl Halides and Grignard Reagents by a Radical Mechanism

Giuseppe Antonacci,[a] Andreas Ahlburg,[a] Peter Fristrup,[a] Per-Ola Norrby,[b,c] and Robert Madsen*[a]

Abstract: The substrate scope and the mechanism have been investigated for the MnCl2-catalyzed cross coupling reaction between aryl halides and Grignard reagents. The transformation proceeds rapidly and in good yield when the aryl halide is a chloride containing a cyano or an ester group in the para position or a cyano group in the ortho position. A range of other substituents gave no conversion of the aryl halide or led to the formation of side products. A broader scope was observed for the Grignard reagents where a variety of alkyl- and arylmagnesium chlorides participated in the coupling. Two radical clock experiments were performed which in both cases succeeded in trapping an intermediate aryl radical. The cross coupling is therefore believed to proceed by a $S_{\text{RN}}$ mechanism, where a triorganomanganate complex serves as the most likely nucleophile and single electron donor. Other mechanistic scenarios were excluded based on the substrate scope of the aryl halide.

Introduction

The palladium-catalyzed cross coupling reaction has been one of the most important discoveries in organic chemistry over the past 50 years. The reaction has had a tremendous impact on the pharmaceutical industry where it accounts for about 10% of all reactions used in the synthesis of drug candidates. The reaction, however, suffers from one major drawback which is the use of the metal palladium. This metal does not occur naturally in the human body and all palladium compounds are considered toxic. Furthermore, palladium is a precious metal with a low annual production. This has prompted a thorough search for alternative catalysts where nickel complexes have been extensively investigated, but are more toxic than the palladium counterparts. Recently, copper,[7] iron[8] and cobalt[9] complexes have gained much attention, but often high catalyst loadings are required. As a result, there is still a demand for effective, cheap and non-toxic catalysts for the cross coupling reaction.

This has inspired research into manganese catalysts since manganese is one of the cheapest metals and is also present in all living organisms. Although, the general application of manganese in homogeneous catalysis is rapidly increasing, the metal has still only found limited applications for the cross coupling reaction. To date, only four publications describe the manganese-catalyzed coupling between aryl/alkenyl halides and Grignard reagents where MnCl2 is used as the catalyst in all cases.[10-11] This includes the coupling of activated aryl halides,[9] reative heterocyclic chlorides[11] and alkenyl halides[11] with both alkyl- and arylmagnesium halides. No information is provided about the mechanism of these manganese-catalyzed reactions.

We envisaged that the scope of the MnCl2-catalyzed coupling between aryl halides and Grignard reagents could be expanded, possibly by gaining an understanding of the reaction mechanism. Some of us have previously studied the reactivity of Grignard reagents[12] and investigated the mechanism of the iron-catalyzed cross coupling[13] and the Barbier allylation.[14] We decided to use the MnCl2-catalyzed cross coupling between activated aryl halides and aryl/alkyl Grignard reagents as a starting point for our investigation.[9] In this transformation, $o$-chlorobenzonitrile undergoes a successful reaction with the organomagnesium halides in THF solution with 10% of the catalyst.[9] In addition, both $o$- and $p$-chlorobenzaldehyde $N$-butylimine can be coupled with the Grignard reagents under the same conditions.[9] However, this is a very narrow range of substrates and it would be interesting to exploit the transformation with a broader array of aryl halides. Herein, we describe the substrate scope and limitations for the manganese-catalyzed cross coupling of aryl halides with Grignard reagents and elucidate part of the reaction mechanism.

Results and Discussion

The studies began by investigating the reaction between cyclohexylmagnesium chloride and various para-substituted halobenzenes (Table 1). The coupling afforded a 94% yield with $p$-chlorobenzonitrile (entry 1) while methyl $p$-chlorobenzoate gave 65% yield (entry 2). The transformation was performed in THF since the coupling with $p$-chlorobenzonitrile gave a higher yield in this solvent than in diethyl ether, dioxane, DME or toluene. In addition, the best results with this substrate were obtained with MnCl2 as the catalyst while a lower yield was achieved with MnBr2 and no coupling occurred with MnF2, MnI2 or in the absence of a manganese salt. The use of additives such as LiCl and MgBr2 also led to lower yields. MnCl2 is not soluble in THF, but dissolves upon addition of the Grignard reagents to afford a brown solution. Chloride appears to be the preferred leaving group since only a 43% yield was obtained.

[a] G. Antonacci, A. Ahlburg, Dr. P. Fristrup, Prof. Dr. R. Madsen
Department of Chemistry
Technical University of Denmark
2800 Kgs. Lyngby (Denmark)
E-mail: rm@kemi.dtu.dk
[b] Prof. Dr. Per-Ola Norrby
Department of Chemistry and Molecular Biology
University of Gothenburg
Kungelgården 4, 412 96, Göteborg (Sweden)
[c] Prof. Dr. Per-Ola Norrby
Pharmaceutical Sciences
AstraZeneca
Pepparedsleden 1, 431 83 Mölndal (Sweden)
Supporting information for this article is given via a link at the end of the document.
with p-bromobenzonitrile (entry 3) while p-iodobenzonitrile underwent complete dehalogenation (entry 4).

Table 1. Coupling with cyclohexylmagnesium bromide.

<table>
<thead>
<tr>
<th>Entry</th>
<th>X</th>
<th>Y</th>
<th>Yield [%]</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Cl</td>
<td>CN</td>
<td>94</td>
</tr>
<tr>
<td>2</td>
<td>Cl</td>
<td>COOMe</td>
<td>65</td>
</tr>
<tr>
<td>3</td>
<td>Br</td>
<td>CN</td>
<td>43</td>
</tr>
<tr>
<td>4</td>
<td>I</td>
<td>CN</td>
<td>0</td>
</tr>
<tr>
<td>5</td>
<td>F</td>
<td>CN</td>
<td>0</td>
</tr>
<tr>
<td>6</td>
<td>Cl</td>
<td>CF₃</td>
<td>0</td>
</tr>
<tr>
<td>7</td>
<td>Cl</td>
<td>NO₂</td>
<td>0</td>
</tr>
<tr>
<td>8</td>
<td>Br</td>
<td>CONMe₂</td>
<td>0</td>
</tr>
</tbody>
</table>

[a] Isolated yield.

Attempts to extend the coupling to a variety of other para-substituted halobenzenes were not successful. No reaction was observed when p-fluorobenzonitrile and p-chlorobenzotrifluoride were mixed with the Grignard reagent under the optimized conditions (entries 5 and 6) which are important observations for understanding the mechanism of the coupling. The trifluoromethyl and the cyano group are both electron-withdrawing groups with Hammett constants around 0.6, and the vast difference in reactivity between these groups indicates that an oxidative addition to the aryl chloride is not part of the reaction pathway. The fact that the chloro-substrate reacts well with the Grignard reagent while the fluoro compound is unreactive shows that the transformation does not proceed by a SₐAr mechanism through an intermediate Meisenheimer adduct with the addition as the rate-determining step.

A number of other para-substituted halobenzenes were also unreactive or led to side reactions. p-Chloronitrobenzene reacted with the Grignard reagent at the nitro group (entry 7) which is a known transformation for organomagnesium halides[16] whereas no reaction was observed with N,N-dimethyl p-bromobenzamide (entry 8). p-Chlorobenzaldehyde and -acetophenone underwent addition to the carbonyl group while chlorobenzenes with a methyl, phenyl, bromo, methoxy or methythio substituent in the para position did not react with cyclohexylmagnesium chloride (results not shown). The meta-substituted substrate, m-chlorobenzonitrile, did not react either under the optimized conditions.

The coupling could be extended to other Grignard reagents as shown in the reaction with p-chlorobenzonitrile (Table 2, entries 1 – 7). The transformation gave moderate to good yields with a variety of different aryl- and alkylmagnesium halides. The corresponding α-chlorobenzonitrile underwent a similar coupling with the Grignard reagents and the yields were close to the results obtained for the meta substrate (Table 2, entries 8 – 12). Both substrates were also reacted with allylmagnesium chloride, but the results were difficult to reproduce although the substitution product was obtained in moderate yields in some cases. In addition, the different Grignard reagents were reacted with p-chlorobenzotrifluoride, p-chloroanisole and m-chlorobenzonitrile, but no conversion of these chlorobenzenes was observed which is in line with the results in Table 1. The reaction between methyl p-chlorobenzoate and phenylmagnesium chloride gave substitution at the ester group and no reaction occurred with the halide. The same substitution to produce the ketone was observed when p-chlorophenylmagnesium bromide and p-methoxyphenylmagnesium bromide and allylmagnesium chloride were reacted with methyl p-chlorobenzoate.

Table 2. Coupling with p- and m-chlorobenzonitrile.

<table>
<thead>
<tr>
<th>Entry</th>
<th>R’</th>
<th>R”</th>
<th>R</th>
<th>X</th>
<th>Yield [%]</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>H</td>
<td>CN</td>
<td>ClH₂</td>
<td>Br</td>
<td>93</td>
</tr>
<tr>
<td>2</td>
<td>H</td>
<td>CN</td>
<td>p-MeOC₆H₄</td>
<td>Br</td>
<td>83</td>
</tr>
<tr>
<td>3</td>
<td>H</td>
<td>CN</td>
<td>p-ClC₆H₄</td>
<td>Br</td>
<td>79</td>
</tr>
<tr>
<td>4</td>
<td>H</td>
<td>CN</td>
<td>p-MeC₆H₄</td>
<td>Br</td>
<td>77</td>
</tr>
<tr>
<td>5</td>
<td>H</td>
<td>CN</td>
<td>(C₆H₄)₂Cl</td>
<td>Cl</td>
<td>68</td>
</tr>
<tr>
<td>6</td>
<td>H</td>
<td>CN</td>
<td>(C₆H₄)₂CH</td>
<td>Cl</td>
<td>63³⁺</td>
</tr>
<tr>
<td>7</td>
<td>H</td>
<td>CN</td>
<td>(C₆H₄)₂CH</td>
<td>Br</td>
<td>58</td>
</tr>
<tr>
<td>8</td>
<td>CN</td>
<td>H</td>
<td>Cyclohexyl</td>
<td>Cl</td>
<td>91</td>
</tr>
<tr>
<td>9</td>
<td>CN</td>
<td>H</td>
<td>C₆H₄</td>
<td>Br</td>
<td>90</td>
</tr>
<tr>
<td>10</td>
<td>CN</td>
<td>H</td>
<td>p-MeOC₆H₄</td>
<td>Br</td>
<td>80</td>
</tr>
<tr>
<td>11</td>
<td>CN</td>
<td>H</td>
<td>p-ClC₆H₄</td>
<td>Br</td>
<td>79</td>
</tr>
<tr>
<td>12</td>
<td>CN</td>
<td>H</td>
<td>p-MeC₆H₄</td>
<td>Br</td>
<td>78</td>
</tr>
</tbody>
</table>

[a] Isolated yield. [b] Yield based on NMR since the isolated product could not be obtained completely pure.

The influence of the temperature and the reaction time was investigated with p-chlorobenzonitrile and phenylmagnesium chloride. No reaction occurred at -12 °C while at 0 °C about 5% of the product was formed after 2 hours. At 6 °C almost 80% of the chloronitrile was consumed after only 1 minute followed by very little further consumption of the starting material over the next 30 min. At room temperature the coupling essentially went
to completion within 1 minute after which time the solvent was
refluxing due to the exothermic nature of the reaction.

To further probe the influence of the Grignard reagent, a
competition experiment was set up in which p-chlorobenzonitrile
was allowed to react with a mixture of phenyl- and
cyclohexylmagnesium chloride (i.e., a contest between the
reactions in Table 1, entry 1 and Table 2, entry 1). This resulted
in immediate formation of p-cyclohexylbenzonitrile and very little
of p-phenylbenzonitrile which shows that the most nucleophilic
Grignard reagent is also the most reactive. An additional
competition experiment was set up in which
cyclohexylmagnesium chloride was allowed to react with a mixture of
p-chlorobenzonitrile and methyl p-chlorobenzoate (i.e.,
a contest between the reactions in entry 1 and 2 in Table 1). In
this case, the two substitution products were formed in equal
amounts and the p-cyano and the p-methyl ester substituents
therefore display a similar influence on the reactivity of the aryl
halide.

A Hammett study was also considered because it may
provide information about the nature of the intermediate species
in the coupling. Since the reaction gives the best results with
o- and p-chlorobenzonitrile, differently substituted analogs of
these were investigated as possible substrates for the kinetic
study (Figure 1 and Scheme 1). Unfortunately, analogs 1–4 all
led to mixtures of several products when reacted with
cyclohexylmagnesium chloride. Only with methyl substituted
analog 5–7 was it possible to obtain one coupling product 8–10
upon reaction with the cyclohexyl Grignard reagent and MnCl₂
(Scheme 1). The yields ranged from 88% and 81% with 5
and 7 to 62% with 6. It is noteworthy that compound 6 can be
coupled at all since the halide and the cyano group are positioned meta
to each other.

These results show that the substrate scope of the cross
coupling is limited under the present conditions. However, the
transformation is still very fast with a narrow range of ortho- and
para-substituted aryl halides and the mechanism must therefore
involve a pathway where these substituents are essential. As
mentioned above, the reaction is not operating by a classical
S₄Ar route or through an oxidative addition pathway as known
for the corresponding palladium- and nickel-catalyzed reactions.
This raises the question whether a radical pathway is involved,
i.e., a S₅Ar mechanism. Alkali metal enolates and a few other
carbanions have previously been reacted with halobenzenes
through a S₅Ar pathway, but whether Grignard reagents are
able to react with aryl radicals is still a matter of debate.

Several radical experiments were therefore conducted in order to
trap an intermediate aryl radical. First, the reaction between
p-chlorobenzonitrile and cyclohexylmagnesium chloride was
repeated in the presence of cyclohexa-1,4-diene in an attempt to
dehalogenate the aryl chloride. However, the coupling still
proceeded smoothly under these conditions and gave p-
cyclohexylbenzonitrile as the only product. Then, a radical clock
experiment was designed in which allyl ether 13 and the
corresponding but-3-enyl compound 14 were reacted with
cyclohexyl Grignard and MnCl₂ (Scheme 2). The two olefinic
chlorobenzonitriles were prepared by allylation from the
respective phenol 11 and benzyl bromide 12. Compound 14
could not be obtained completely pure, but contained about 30%
of a byproduct where the olefin had migrated. The reaction with
cyclohexylmagnesium chloride gave in both cases a mixture of
several compounds, but the main products arose from
cyclization with the olefin and addition to the nitrile. The
cyclization products 15 and 16 were isolated in 9% and 7%
yield, respectively. Only very small amounts (1–2%) were
observed by GCMS from the direct cross coupling between the
aryl halide and the Grignard reagent, but the products could not
be isolated or further quantified.

![Figure 1. Substrates investigated for the Hammett study.](image)

![Scheme 1. Coupling of chloromethylbenzonitriles.](image)

![Scheme 2. Radical clock experiments.](image)
The mechanistic proposal in Scheme 3 should be compared with the recently published cross coupling reaction between aryl iodides/bromides and aryl Grignard reagents in the absence of a catalyst.[25] This reaction was performed in toluene at 110 °C for 24 h and allowed for coupling of ether and alkyl substituted aryl moieties.[26] The mechanism was subsequently investigated and a radical clock experiment failed to produce the cyclization product from an aryl radical.[26] DFT calculations suggested a pathway where the starting aryl halide Ar–X is converted by SET into [Ar–X]− which reacts with Ar′–MgBr to furnish a magnesium ion-radical cage [Ar′–ArMgBrX]−.[19] The latter is transformed into a ArMgAr′ radical anion from which [Ar–Ar′]− is formed followed by SET to Ar–X.[19]

Conclusions

In summary, we have managed to exclude several commonly proposed catalytic cycles for the manganese-assisted coupling of Grignard reagents with aryl chlorides, and by inference, limited the mechanistic possibilities to one plausible reaction mechanism, $S_{RN1}$. In line with this mechanism, a narrow aryl halide scope is observed, where only substituents allowing a single electron reduction followed by a facile halide dissociation give coupling. The proposed radical intermediate can be trapped by an internal radical clock substituent, but will prefer coupling with the Grignard reagent over base-stable intermolecular radical traps like cyclohexadiene. Substrates that will react directly with Grignard reagents, such as nitro-aromatics, ketones and aryl iodides, are not competent coupling partners. On the Grignard side, the scope is wider and allows for coupling of a variety of alkyl and arylmagnesium halides.

Experimental Section

General Information: All solvents were of HPLC grade and were not further purified. Gas chromatography was performed on a Shimadzu GCMS-QP2010S instrument fitted with an Equity 5, 30 m × 0.25 mm × 0.25 μm column. Flash column chromatography separations were performed on silica gel 60 (40 – 63 μm). NMR spectra were recorded on a Bruker Ascend 400 spectrometer. Chemical shifts were measured relative to the signals of residual CHCl$_3$ (δ = 7.26 ppm) and CDCl$_3$ (δ = 77.16 ppm). HRMS measurements were made using ESI with TOF detection. All Grignard reagents were obtained from commercial suppliers and titrated with a 0.06 M solution of I$_2$ in Et$_2$O to determine the concentration: cyclohexylmagnesium chloride (1.6 M in Et$_2$O), phenylmagnesium bromide (0.9 M in THF), p-methoxyphenylmagnesium bromide (0.3 M in THF), p-chlorophenylmagnesium bromide (0.5 M in Et$_2$O), p-tolylmagnesium bromide (0.9 M in THF), n-butylmagnesium chloride (1.6 M in THF), isobutylmagnesium chloride (1.8 M in THF) and isopropylmagnesium bromide (0.8 M in THF).

General Procedure for Cross Coupling: A dry three-neck Schlenk tube was equipped with a stir bar and a nitrogen inlet. The flask was flushed with nitrogen and charged with MnCl$_2$ (25 mg, 0.2 mmol) and dry THF (6 mL). The mixture was stirred for about 10 min to completely dissolve MnCl$_2$ followed by addition of the aryl halide (2 mmol) and cooling to 0 °C in an ice bath. A solution of the Grignard reagent (4 mmol) was added dropwise over 5 min and the ice bath was removed. The mixture was stirred for 1 h at ambient temperature. Decane (0.4 mL, 2 mmol) was

![Scheme 3. Proposed mechanism for manganese-catalyzed cross coupling.](image-url)
injected as an internal standard for determining the yield by GC and the reaction was quenched with saturated ammonium chloride solution (10 mL). The mixture was extracted with EtOAc (4 x 10 mL) and the combined organic layers were concentrated and the residue purified by flash column chromatography (70/30 pentane/CHCl3).

**4-Cyclohexylbenzoate**[25] Table 1. Entry 1. Isolated as a colorless oil in 94% yield (347 mg). H NMR (400 MHz, CDCl3): δ = 7.78–8.44 (m, 1H), 7.33–7.56 (m, 3H), 7.26 (d, J = 8.0 Hz, 1H), 7.10 (d, J = 7.8 Hz, 1H), 7.04 (d, J = 7.8 Hz, 1H), 2.67 (m, 4H), 2.10 (m, 2H). 13C NMR (100 MHz, CDCl3): δ = 147.3, 123.9, 122.0, 119.1, 109.5, 45.4, 30.0, 22.2 ppm (EI; m/z = 213 [M]+).

**Methyl 4-cyclohexylbenzoate**[25] Table 2. Entry 1. Isolated as a white solid (430 mg) containing about 25% of cyclohexyl p-cyclohexylphenyl ketone which could not be separated. Yield 65%. H NMR (400 MHz, CDCl3): δ = 7.78–8.70 (m, 2H), 7.30–7.35 (m, 2H), 5.36–5.50 (m, 1H), 5.2–5.3 (m, 1H), 1.90–1.93 (m, 4H), 1.82–1.76 (m, 1H), 1.44–1.36 (m, 4H), 1.33–1.17 (m, 1H) ppm. 13C NMR (100 MHz, CDCl3): δ = 153.6, 132.3, 127.8, 119.4, 107.9, 44.9, 34.1, 26.8, 26.1 ppm (EI; m/z = 185 [M]+).

2-Cyclohexylbenzotriazole[34] Table 2. Entry 8. Isolated as a colorless oil in 91% yield (337 mg). H NMR (400 MHz, CDCl3): δ = 7.53–7.36 (m, 2H), 7.25 (dd, J = 8.0, 1.1 Hz, 1H), 7.14 (dd, J = 7.6, 1.2 Hz, 1H), 2.93–2.77 (m, 1H), 1.83–1.60 (m, 5H), 1.42–1.25 (m, 4H), 1.22–1.02 (m, 1H) ppm. 13C NMR (100 MHz, CDCl3): δ = 151.3, 132.8, 132.7, 126.4, 126.2, 118.1, 111.7, 42.7, 33.5, 26.5, 25.8 ppm (MS; EI; m/z = 189 [M]+).

**[1,1′-Biphenyl]-2-carbonitri te**[25] Table 2. Entry 9. Isolated as a white solid in 80% yield (336 mg). H NMR (400 MHz, CDCl3): δ = 8.01–8.03 (m, 6H), 7.87–7.65 (m, 1H), 7.58–7.50 (m, 1H), 7.49–7.39 (m, 1H) ppm. 13C NMR (100 MHz, CDCl3): δ = 140.8, 138.6, 129.7, 128.3, 127.7, 121.3, 119.2, 119.6, 111.0, 114.3 ppm (EI; m/z = 199 [M]+).

**4-Methoxy-[1,1′-biphenyl]-2-carbonitri te**[25] Table 2. Entry 10. Prepared according to the general procedure where the Grignard reagent was added over 120 min at 0 °C to prevent a competing addition to the cyano group. Isolated as a white solid in 83% yield (296 mg). H NMR (400 MHz, CDCl3): δ = 8.04–8.00 (m, 6H), 7.65–7.62 (m, 2H), 7.57–7.50 (m, 2H), 7.04–6.97 (m, 2H), 3.87 (s, 3H) ppm. 13C NMR (100 MHz, CDCl3): δ = 166.4, 143.3, 137.7, 131.6, 128.3, 128.7, 121.7, 119.2, 114.7, 110.2, 55.5 ppm (EI; m/z = 209 [M]+).

**4-Chloro-[1,1′-biphenyl]-4-carbonitri te**[25] Table 2. Entry 3. Prepared according to the general procedure where the reaction mixture was stirred for 2 h at 60 °C in an oil bath to ensure complete conversion of p-chlorobenzonitrile. Isolated as a white solid in 79% yield (335 mg). δ H NMR (400 MHz, CDCl3): δ = 7.37 (d, J = 6.0 Hz, 1H), 7.12 (d, J = 6.0 Hz, 1H), 2.42–2.40 (m, 4H), 2.11–2.08 (m, 4H), 1.30–1.26 (m, 4H) ppm. 13C NMR (100 MHz, CDCl3): δ = 144.5, 137.7, 135.1, 132.9, 129.5, 128.6, 127.7, 118.9, 111.4 ppm (MS; EI; m/z = 219 [M]+).

**4-Methyl-[1,1′-biphenyl]-4-carbonitri te**[25] Table 2. Entry 4. Prepared according to the general procedure where the Grignard reagent was added over 120 min at 0 °C to prevent a competing addition to the cyano group. Isolated as a white solid in 77% yield (296 mg). H NMR (400 MHz, CDCl3): δ = 7.63 (d, J = 8.5 Hz, 2H), 7.58 (d, J = 8.5 Hz, 2H), 7.43–7.39 (m, 2H), 7.21 (d, J = 7.9 Hz, 2H), 2.43 (s, 3H) ppm. 13C NMR (100 MHz, CDCl3): δ = 145.7, 138.9, 136.4, 132.7, 130.0, 127.6, 127.2, 119.2, 110.7, 21.3 ppm. MS (EI; m/z = 193 [M]+).

**4-Butyl-[1,1′-biphenyl]-4-carbonitri te**[25] Table 2. Entry 5. Isolated as a colorless oil in 86% yield (217 mg). H NMR (400 MHz, CDCl3): δ = 7.41 (q, J = 7.5 Hz, 2H), 7.25 (d, J = 7.2 Hz, 2H), 2.64 (t, J = 7.8 Hz, 2H), 1.67–1.51 (p, J = 7.5 Hz, 2H), 1.33 (q, J = 7.4 Hz, 2H), 0.91 (t, J = 7.4 Hz, 3H) ppm. 13C NMR (100 MHz, CDCl3): δ = 148.6, 132.0, 129.2, 119.1, 109.4, 35.8, 33.1, 22.2, 13.8 ppm (EI; m/z = 219 [M]+).

**4-Isobutylbenzotriazole**[25] Table 2. Entry 6. Isolated as a colorless oil (290 mg) which could not be obtained completely pure. Yield 63% as estimated from the NMR spectrum. H NMR (400 MHz, CDCl3): δ = 7.42–7.38 (m, 2H), 7.12–7.08 (m, 2H), 2.38 (s, J = 7.3 Hz, 2H), 1.74 (dt, J = 13.6, 6.8 Hz, 1H), 1.82–1.73 (m, 4H) ppm. 13C NMR (100 MHz, CDCl3): δ = 147.3, 131.9, 129.8, 129.7, 118.1, 109.5, 45.4, 30.0, 22.2 ppm (EI; m/z = 193 [M]+).

**4-Isopropylbenzotriazole**[25] Table 2. Entry 7. Isolated as a yellowish oil in 98% yield (168 mg). H NMR (400 MHz, CDCl3): δ = 7.54–7.45 (m, 2H), 7.25 (d, J = 8.3 Hz, 2H), 2.88 (d, J = 6.9 Hz, 1H), 1.19 (d, J = 7.0 Hz, 6H) ppm. 13C NMR (100 MHz, CDCl3): δ = 154.4, 132.2, 127.3, 119.2, 109.6, 34.4, 23.5 ppm (MS; EI; m/z = 245 [M]+).
We thank the Danish Council for Independent Research – Technology and Production Sciences for financial support (grant 1335-00153).

**Keywords:** cross-coupling • Grignard reagent • manganese • radical reactions • reaction mechanisms

[25] For our previous studies of organometallic reagents by Hammet studies, see reference 14.
FORMED BY RADICALS: Aryl halides and Grignard reagents are coupled with MnCl₂ as catalyst. The substrate scope and the mechanism are investigated, and an aryl radical is identified as an intermediate. As a result, the cross coupling is believed to proceed through a S_n1 mechanism.

Giuseppe Antonacci, Andreas Ahlburg, Peter Fristrup, Per-Ola Norbý, Robert Madsen*

Page No. – Page No.

Manganese-Catalyzed Cross Coupling of Aryl Halides and Grignard Reagents by a Radical Mechanism