Formadide as a superior nitrogen nucleophile in palladium(II)-mediated synthesis of imidazolidines.

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Formamide as a Superior Nitrogen Nucleophile in Palladium(II) Mediated Synthesis of Imidazolidines

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Abstract: Formamides emerge as superior nitrogen nucleophiles in palladium(II) catalyzed oxidative 5-exo cyclizations of formyl aldehydes amines derived from N-Boc protected allylic amines. The product imidazolidines are readily transformed into protected vicinal diamines.

The successful concept of the "detachable connection" approach employing a tethered oxygen nucleophile in the synthesis of vicinal aminoaiknols from allylic amines using palladium(II) catalysis was recently demonstrated by us. The key oxidative 5-exo cyclization of N,O-hemiacetals to oxazolidines ensured high regio- and stereoselectivities. With Pd(OAc)$_2$ as the catalyst, molecular oxygen was used as a clean stoichiometric oxidant without the need for a co-oxidant. Dimethyl sulfoxide, used as the solvent, bears a unique character in this remarkable process by inducing the formation of "giant" palladium clusters which are most likely the catalytically active species.

Our present efforts involve the application of similar methodology to the synthesis of vicinal diamines, as depicted in Scheme 1. In this communication we report several Pd(II) catalyzed oxidative cyclizations in which different tethered nitrogen nucleophiles are incorporated into imidazolidines. As suitable precursors for these cyclizations aminals of type 2 were prepared from N-Boc protected allylic amines, formaldehyde and a protected nitrogen source. Their reactivities towards the oxidative cyclization conditions are reported. Furthermore, the convenient conversion of one imidazolidine into the corresponding diamine is described.

Scheme 1. "Detachable connection" approach to allylic diamines

Organometal-based techniques such as intramolecular aminopalladation, amidopalladation and palladium mediated amidocarbonylation of monoolefins dienes and allenes for the synthesis of five and six membered nitrogen containing heterocycles have been widely adopted. Acetamides and, especially, p-toluenesulfonylamides were the first nitrogen nucleophiles reported to be successful in 5-exo cyclizations. Since then, carbamates and urea derivatives have also been applied and have often been found superior to p-
Toluenesulfonamides but sometimes inferior. The optimum balance between nitrogen nucleophilicity and acidity apparently varies from case to case.

Two parent allylic carbamates were selected for testing different nitrogen nucleophiles. N-Boc-2-cyclopentenylamine 5 was chosen because its hemiacetals were found to be the most reactive towards Pd(II) mediated 5-endo cyclization. More flexible aminals derived from the open chain N-Boc-3-pent-2-(E)-enylamine 10 were expected to be less reactive. In addition, steric induction by the methyl group could be observed in the formation of the corresponding imidazolidine 14. These substrates would enable us to discriminate between the different nitrogen nucleophiles in terms of both reactivity and stereoselectivity.

Scheme 2 shows the preparation of the aminals. Reaction of the parent carbamates 5 and 10 with paraformaldehyde and cesium carbonate in dioxane (60 °C, 2 h) gave N-hydroxymethylcarbamates 6 and 11, respectively, in good yields. These N,O-hemiacetals were stable at room temperature and were purified by flash chromatography (silica gel, ethyl acetate/hexanes). Treatment of 6 and 11 with a catalytic amount of p-toluenesulfonic acid in formamide directly gave formamide aminals 8b and 13b, respectively, via the N-acetyliminium ions in excellent yields. Acetoxyethyl carbamates 7 and 12 obtained from the hemiacetals (Ac₂O, DMAP, pyridine, 0 °C) were similarly converted into aminals by reaction with acetamide (8c), methyl carbamate (8e, 13e), benzyl carbamate (8f, 13f) or p-toluenesulfonamide (8g, 13g) in dichloromethane (reflux, 20 h) in the presence of a catalytic amount of PPTS, or with urea (8e) in acetic acid (50 °C, 2 h).

![Scheme 2. Preparation and Oxidative Cyclization of Aminals](image)

The aminals 8 and 13 were converted to their corresponding imidazolidines 9 and 14 under standard oxidative cyclization conditions using Pd(OAc)$_2$ (0.05 equiv) and one atmosphere of molecular oxygen. The
results of these reactions are summarized in Table 1. With the rigid cyclopentenyl derivatives ureas (8d) surprisingly turned out to be the least reactive nucleophile. Acetamide (8c) and carbamates (8e,f) gave satisfactory results but were found to be less reactive than formamide (8b) and p-toluenesulfonamide (8a). With the more flexible pentenyl derivatives, however, formamide (13d) emerged as the most successful nucleophile. Stereoselectivities were found to be equally moderate with formamide and p-toluenesulfonamide (13a) and poor with methyl carbamate (13e). An additional advantage of the use of formamide as the nitrogen source is the facile mono deprotection of cyclization products 9b and 14b to the N-Boc imidazolidines 9g and 14g, respectively, by mild hydrolysis (KOH, MeOH, 94%).

<table>
<thead>
<tr>
<th>Aminal</th>
<th>mp (°C)¹¹</th>
<th>Reaction Time (h)²</th>
<th>Yield (%)³</th>
<th>Imidazolidine</th>
<th>mp (°C)¹¹</th>
<th>Isomer Ratio⁵</th>
</tr>
</thead>
<tbody>
<tr>
<td>8a</td>
<td>105-107</td>
<td>1</td>
<td>71</td>
<td>9a</td>
<td>143-144</td>
<td></td>
</tr>
<tr>
<td>8b</td>
<td>67-69</td>
<td>2</td>
<td>86</td>
<td>9b</td>
<td></td>
<td></td>
</tr>
<tr>
<td>8c</td>
<td>70-72</td>
<td>4</td>
<td>70 (95)</td>
<td>9c</td>
<td>104-106</td>
<td></td>
</tr>
<tr>
<td>8d</td>
<td>119-120</td>
<td>4*</td>
<td></td>
<td>9d</td>
<td></td>
<td></td>
</tr>
<tr>
<td>8e</td>
<td>62-63</td>
<td>4</td>
<td>83</td>
<td>9e</td>
<td></td>
<td></td>
</tr>
<tr>
<td>8f</td>
<td>71-73</td>
<td>4</td>
<td>84</td>
<td>9f</td>
<td></td>
<td></td>
</tr>
<tr>
<td>13a</td>
<td>76-78</td>
<td>4*</td>
<td>98 (66)</td>
<td>14a</td>
<td></td>
<td>(35/65)</td>
</tr>
<tr>
<td>13b</td>
<td>-</td>
<td>2</td>
<td>90</td>
<td>14b</td>
<td></td>
<td>(35/65)</td>
</tr>
<tr>
<td>13e</td>
<td>45-47</td>
<td>4*</td>
<td>98 (50)</td>
<td>14e</td>
<td></td>
<td>(47/53)</td>
</tr>
</tbody>
</table>

⁵ Cis/trans: stereochemistry was established by means of ¹H NMR NOE experiments.

The N-Boc imidazolidines 9e and 9f were also mono deprotected (Scheme 3) with the purpose of facilitating removal of the methylene tether. However, we were unable to convert imidazolidines 14g, 17, or 18 into the corresponding vicinal diamines by acid or base mediated hydrolysis. We therefore turned to electrochemistry once again for removal of the tether. Anodic oxidation in methanol mediated by a catalytic amount of NaCl₁³ surprisingly yielded amidine 15 quantitatively. Attempts to prepare the corresponding amidines from imidazolidines 17 and 18 in a similar way failed.

Scheme 3. Synthesis of mono protected imidazolidines and diamine
Conversion of amidine 15 to protected diamine 16 was finally achieved by reaction with acetic anhydride in acetic acid/water (1:1).

In conclusion, we have demonstrated the advantages of formamide as a nitrogen nucleophile in palladium catalyzed synthesis of imidazolidines because of (1) its high reactivity and (2) the ease of its deprotection and thereby the feasibility of stereocontrolled synthesis of vicinal diamines. This class of compounds have found widespread use as chiral ligands in asymmetric reactions and chiral derivatizing reagents for determination of enantiomeric composition. We are currently applying the methodology presented here to the synthesis of a wider variety of vicinal diamines.

Acknowledgement

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References and Notes

11. All new compounds were characterized by means of 1H and 13C NMR (spectra were recorded at elevated temperatures because of strongly hindered rotation in biannulated) and IR spectroscopy. Solids showed correct elemental analysis data. Melting points are uncorrected.
13. Anodic oxidation in MeOH (0.07 M) using 1 mol% NaCl was performed with a potentiostat/galvanostat operating at 50mA. Carbon electrodes were used in an undivided cell as described in: Shono, T.; Matsunuma, Y.; Tanaka, K. Org. Synth. 1985, 65, 206-213.

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