Microwave-promoted synthesis of chiral pyridinium salts

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Abstract—The synthesis of several chiral pyridinium salts via Zincke’s reaction can be easily accomplished by domestic microwave oven irradiation. Yield enhancements, reduction of reaction time, and less racemization were observed under microwave heating when compared to conventional heating in similar conditions.

Pyridinium salts are a versatile class of compounds used as phase transfer catalysts, 1 initiators of cationic polymerization, 2 wide range antimicrobials, 3 enzyme inhibitors, 4 acylating agents, 5 dyes, and cationic surfactants. 6 Their chiral representatives, that is, pyridinium salts containing a chiral auxiliary group linked to the ring nitrogen, have been largely used as starting materials in asymmetric synthesis to obtain substituted chiral dihydropyridines, 7 tetrahydropyridines, 8 piperidines, 9 and nitrogenated complex compounds such as natural alkaloids 10 and synthetic candidates for drugs. 4b

Several synthesis routes for pyridinium salts are known, but the most commonly used one is the Menschutkin reaction, a S N 2 reaction of a pyridine derivative with an organic halide or sulfonate. 6 Nevertheless, this method in not suitable to prepare chiral pyridinium salts with a stereocenter directly attached to the ring nitrogen since there is a significant risk of racemization by a competing S N 1 process. There are two main methods to obtain these kinds of salts with negligible racemization. The first uses pyrilium salts 11, 12 and the second employs Zincke’s salts, 13 a highly electrophilic species formed by reaction between a pyridine derivative and 1-chloro-2,4-dinitrobenzene.

The original Zincke’s reaction 14 involves the attack of a primary amine 1 on Zincke’s salt 2 (Eq. 1) to produce salt 3. As the mechanism of reaction does not involve rupture of the C–N bond, if we use a primary chiral amine, the configuration of the stereocenter will be kept. This idea was extensively studied by Marazano 15 and Zincke’s reaction became an excellent method for the synthesis of chiral pyridinium salts.

Some recent uses of Zincke’s reaction have been described by Eda and co-workers 4a, 16 and Urban et al. 17 The former adapted this reaction for solid-phase application to achieve chiral quaternary ammonium salts analogs of vesamicol 4a and salts that could be useful to treat cystic fibrosis. 16 The latter applied Zincke’s reaction to make chiral naphthyridinium salts for the synthesis of cytotoxic alkaloid manzamine A. 17

Keywords: Asymmetric synthesis; Amino alcohols; Pyridines; Microwave irradiation; Zincke’s reaction.

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Faster and simpler methods are a permanent need in organic synthesis. In 1986, Gedye and Giguere demonstrated that many organic reactions could be rapidly conducted under microwave irradiation. Since then, microwave irradiation has been increasingly used as a laboratory method to accelerate synthesis. Microwave-assisted organic synthesis has several advantages over conventional methodology: remarkable reduction of reaction time (up to three orders of magnitude), improved isolated product yields (when thermal decomposition is associated with conventional heating reactions), and sometimes, effects on chemo-, regio-, and stereoselectivities are also achieved.

Due to the important role of chiral pyridinium salts in asymmetric synthesis and the absence of reported microwave-assisted Zincke's reaction, we report here a new and efficient method for the fast synthesis of chiral pyridinium salts from chiral primary amines.

We have used a modified domestic microwave oven, 2450-MHz General Electric J1145 AWA. The oven top was cut to accommodate a reflux condenser and a steel ring was used to avoid microwave leakage. The turnable dish was turned off.

A typical procedure involves irradiation of a mixture of 1 equiv of Zincke's salt and 1 equiv of primary amine using 10 mL of 1-butanol as solvent under microwave irradiation for 5–10 min. In the end of the reaction, the salt recovery by simple partition into water phase from ethyl acetate allowed its complete separation from 2,4-dinitroaniline (Eq. 1). Purification of the crude products by silica gel flash chromatography furnished the isolated salts with yields from 61% to 100% (Table 1). All salts were confirmed by 1H NMR and 13C NMR analyses. All reactions were performed under microwave and conventional heating for comparison. The results are presented in Table 1.

Table 1. Synthesis of pyridinium salts under microwave and conventional heating

<table>
<thead>
<tr>
<th>Entry</th>
<th>Primary amine 1</th>
<th>R</th>
<th>Y−</th>
<th>Time (min)</th>
<th>Yield (%)</th>
<th>Time (h)</th>
<th>Yield (%)</th>
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<tr>
<td>1</td>
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<td>−CH3</td>
<td>Cl−</td>
<td>5</td>
<td>91</td>
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<td>−CH3</td>
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<td>68</td>
</tr>
<tr>
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<td>NH2</td>
<td>−CH3</td>
<td>Cl−</td>
<td>10</td>
<td>57</td>
<td>15</td>
<td>29</td>
</tr>
<tr>
<td>4</td>
<td>NH2</td>
<td>−CH3</td>
<td>Cl−</td>
<td>10</td>
<td>100</td>
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<td>10</td>
<td>98</td>
<td>15</td>
<td>72</td>
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</table>
From Table 1, it can be seen that the reaction time was reduced by several times under microwave irradiation when compared with reactions performed under conventional heating. Marazano established that the ideal condition for this reaction under conventional heating was 1-butanol refluxing as solvent for 15 h. We have found similar results for conventional heating; however, for microwave heating, the reaction time is never larger than 10 min. Furthermore, the yield of microwave heating reactions after purification was greatly increased. It is very interesting to observe that under microwave irradiation, pyridinium salts were obtained with less racemization than by conventional heating methods. These results confirm the applicability of microwave heating to the improvement of classic reactions.

In summary, we have developed an operationally simple and efficient method for the synthesis of chiral pyridinium salts by microwave-promoted Zincke reaction. The conditions employed furnished good yields and less racemization than by conventional heating methods. These results confirm the applicability of microwave heating to the improvement of classic reactions.

Acknowledgements

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Table 1

<table>
<thead>
<tr>
<th>Entry</th>
<th>Primary amine 1</th>
<th>2</th>
<th>3</th>
<th>Microwave heating</th>
<th>Conventional heating</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>R</td>
<td>Y&quot;</td>
<td>Time (min)</td>
<td>Yield (%)</td>
</tr>
<tr>
<td>8</td>
<td></td>
<td>-H</td>
<td>SDS ¹</td>
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<td>Ethyl</td>
<td>Cl⁻</td>
<td>10</td>
<td>86</td>
<td>15</td>
</tr>
</tbody>
</table>

¹ SDS = dodecylsulfate.

Zincke’s salts were easily prepared by reaction of commercially available pyridine, 3-picoline, and 3-ethylpyridine (1 equiv of each) and 1 equiv of 1-chloro-2,4-dinitrobenzene under acetone reflux for 15 h. The selected primary amines were (R)-(-)-phenylglycinol, (S)(+)-, (S)(-)-phenylalaninol, (S)(+)-isoleucin, (R)(+)- and (S)(-)-α-methylbenzylamine, and the achiral amine serinol.

From Table 1, it can be seen that the reaction time was reduced by several times under microwave irradiation when compared with reactions performed under conventional heating. Marazano established that the ideal condition for this reaction under conventional heating was 1-butanol refluxing as solvent for 15 h. We have found similar results for conventional heating; however, for microwave heating, the reaction time is never larger than 10 min. Furthermore, the yield of microwave heating reactions after purification was greatly increased. It is very interesting to observe that under microwave irradiation, pyridinium salts were obtained with less racemization than by the methods described until today, possibly due to the much shorter reaction times. For example, for the compound of the table entry 9, the value of \([\alpha]_D^{20}\) reported for the solution conventional method\(^9\) was −44°. Under solid-phase\(^{4a}\) Zincke conditions, the same compound was obtained with \([\alpha]_D^{20}−50\), and by microwave irradiation, the value recorded was −53.4° (CH₃OH, c 2.95). In addition, we also report the values of −25.9° (CH₃OH, c 1.20) for the specific rotation of the compound derived from (S)(−)-α-methylbenzylamine (entry 7) for conventional heating and −32.3° (CH₃OH, c 1.05) for microwave heating. This success led to the preparation of other chiral derivatives by microwave-promoted Zincke’s reaction.

In summary, we have developed an operationally simple and efficient method for the synthesis of chiral pyridinium salts by microwave-promoted Zincke’s reaction. The conditions employed furnished good yields and less racemization than by conventional heating methods. These results confirm the applicability of microwave heating to the improvement of classic reactions.

References and notes

A representative procedure to obtain pyridinium salts is as follows: A solution of (S)-(-)-α-methylbenzylamine (0.259 g, 2.41 mmol) and Zincke’s salt (R = CH₃, Y = Cl⁻) (0.633 g, 2.14 mmol) in 1-butanol (10 mL) was irradiated with microwave for 10 min. After evaporation, the residue was solubilized in water, and to this solution was added ammonium hydroxide until pH 10. The next step was a simple partition in water phase from ethyl acetate. The aqueous phase was evaporated and the residue was chromatographed (CHCl₃/CH₃OH 100/1, 94/6, 90/10, 83/17, 50/50). Pyridinium salt (entry 7, 0.490 g, 98%) was obtained as a yellow oil: [α]D²⁰ = 32.3 (CH₃OH, c 1.05); ¹H NMR (D₂O, 200 MHz): δ 1.97 (3H, t, J = 7.1 Hz); 2.41 (3H, s); 5.99 (1H, q, J = 7.0 Hz); 7.37–7.46 (5H, m); 7.82 (1H, t, J = 7.4 Hz); 8.25 (1H, d, J = 7.9 Hz); 8.62–8.75 (2H, m); ¹³C NMR (D₂O, 50 MHz): δ 18.09; 20.12; 70.89; 127.82; 128.00; 129.82; 130.17; 137.37; 140.62; 142.84; 146.85.