Asymmetric Synthesis of Vabicaserin via Oxidative Multicomponent Annulation and Asymmetric Hydrogenation of a 3,4-Substituted Quinolinium Salt

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ABSTRACT

An efficient, asymmetric synthesis of the 5-HT_{2C} agonist vabicaserin in four chemical steps and 54% overall yield from commercially available benzodiazepine was achieved. The synthesis was highlighted by a novel oxidative, multicomponent reaction to affect the quinolinium ring assembly in one step followed by an unprecedented asymmetric hydrogenation of a 3,4-substituted quinolinium salt.

Vabicaserin (SCA-136, 1) is a potent and selective 5-HT_{2C} receptor full agonist (K_i = 3 nM; EC_{50} = 8 nM), demonstrating efficacy in several animal models predictive of antipsychotic activity.1 Vabicaserin advanced through phase 2 clinical trials as a potential treatment for schizophrenia, exhibiting positive trends across several PANSS measures.2 An efficient and scalable synthesis of 1, containing an unusual cyclopentadiazepinoquinoline ring system with two syn-oriented chiral centers, posed several challenges. In this paper, we describe a four-step, enantioselective synthesis of 1 that was enabled by the discovery of an oxidative, multicomponent reaction yielding two carbon—carbon and two carbon—nitrogen bonds of the pentacyclic core structure, and an unprecedented asymmetric hydrogenation of a 1,3,4,8-tetrasubstituted quinolinium salt.

Early pharmaceutical development supplies were provided by the previously disclosed racemic synthetic route to SCA-136 (Scheme 1).3 A Povarov reaction between benzodiazepine (2), cyclopentene, and formaldehyde provided racemic 1, which was resolved by classical diastereoselective salt resolution. While the synthesis was short, the overall yield for the route was only 23%. Toward a more efficient route to 1, several asymmetric approaches were pursued. One of the more promising routes consisted of reaction of chiral allylsilane 3 with the in situ generated iminium ion 1, which gave a highly diastereoselective Povarov product. However, subsequent cleavage of the carbon—silicon bond proved to be difficult.4 Installation of chirality via asymmetric hydrogenation of a fully assembled pentacyclic core was an attractive route to 1, and several permutations of this approach were evaluated. The unsaturated amide 4 and the unfunctionalized alkene 5 were found to be either unreactive or nonselective under a variety of reduction conditions. The asymmetric

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(2) PANSS: Positive and Negative Syndrome Scale used to measure symptom severity of patients with schizophrenia.

hydrogenation of tetrasubstituted quinolinium salts of type II was unprecedented. However, the potential of assembling the quinolinium salt in a single step coupled to an enantioselective hydrogenation made this route a very attractive target for study.

Assembly of a quinolinium salt II directly from a derivative of readily available benzodiazepine \(6\) was initially envisioned to occur through an economical multicomponent reaction (MCR) with components at the targeted oxidation state. However, all attempts to produce structures of type II with benzodiazepine derivatives in the absence of oxidants were unsuccessful. We then considered a more complex, oxidative MCR sequence, which required annulation, elimination, and oxidation all occurring in the same pot.

The substrate selected for the oxidative MCR study, \(N\)-tosylbenzodiazepine \(7\), was obtained in 95% yield from \(6\) by treatment with \(p\)-TsCl and NaHCO\(_3\) in EtOAc (Scheme 2). We were encouraged when the oxidative MCR conditions developed by Kozlov were examined.

Accordingly, treating 7 with paraformaldehyde, cyclopentanone, and triflic acid in \(n\)-butanol at 100 °C yielded the desired quinolinium triflate \(8\). However, an equal amount of the \(N\)-methyl derivative \(9\) was also formed. We propose the formation of \(9\) occurs through hydride transfer from dihydroquinoline intermediate IV to iminium ion III via an NADH-type redox process, producing equal quantities of \(8\) and \(9\). Several oxidants were tested for their potential to compete with III, but most gave only modest improvements in overall conversion. A previous report indicated catalytic iodine alone could effect this transformation, with air as a stoichiometric oxidant. Applying this protocol to 7 led predominantly to iodination of the aromatic ring, with no product formation. To redirect oxidation, we considered deactivation of the aromatic ring by protonation with acid prior to the addition of iodine. To that effect, the addition of HI prior to iodine not only led to inhibition of ring iodination, observing only trace amounts of 9, the modified conditions also affected conversion of 7 to 8b at substantially lower temperature (30 °C). Consequently, the quinolinium salt 8b was accessed in 89% isolated yield using this new, one-pot annulation/elimination/oxidation MCR sequence.

The enantioselective reduction of highly substituted heterocycles, such as quinolines or pyridines, remains an elusive target in an active field of research. While there are a variety of methods described for highly selective asymmetric reduction of 2-substituted quinolines, there are only two reports describing substrates with substitution at the 3-position, one report with substitution at the 4-position, and no reports with substitution at the 8-position. Further, while many of the prescribed protocols invoke N-activation, there are no examples of \(N\)-alkylquinolinium salts as substrates for asymmetric hydrogenation. Consequently, we were not surprised to find that applying the current state of the art reaction conditions to highly substituted 8, or model substrates thereof, resulted in little or no enantioselectivity. The most promising lead came from the use of the phosphoramidite-iridium catalyst system described by de Vries and Feringa, in which we observed low reactivity and modest enantioselectivity (Figure 1, Table 1, entries 2–4) Analagous to the findings


(7) Combinations of cyclopentanone with formic derivatives, such as the Vilsmeier reagent or DFM dimethyl acetal, did not yield product. Cyclopentanone-2-carboxaldehyde also did not yield the desired product under a variety of conditions examined.


(10) When the reaction was run using paraformaldehyde-\(d_2\), 9 was formed with incorporation of three deuterium atoms (94% of theoretical) and 8a with one deuterium atom, consistent with single deuterium transfer from the intermediate hydroquinoline IV to the iminium ion III. For a related example of an imine acting as an oxidant, see ref 8b.

(11) Iodination was not noted in ref 8c, although the authors indicate the reaction was limited in scope to a single substrate, 2-aminophenylacene.

(12) In this unoptimized process, an excess of cyclopentanone and paraformaldehyde was used to drive the reaction to completion.


(15) When the reaction was run using paraformaldehyde-\(d_2\), 9 was formed with incorporation of three deuterium atoms (94% of theoretical) and 8a with one deuterium atom, consistent with single deuterium transfer from the intermediate hydroquinoline IV to the iminium ion III. For a related example of an imine acting as an oxidant, see ref 8b.
Optimization of the achiral phosphine loading yielded a key observation; increasing the concentration of the achiral phosphine led to an increase in reactivity (Table 2, entries 1–4). We postulated that the role of the excess phosphine may be to act as a base in addition to serving as a ligand for iridium. Apparent confirmation of this conclusion is evidenced by the analogous effect of added base at low phosphine charge (Table 2, entries 5–7), yielding high conversion without significant loss in enantioselectivity. With the addition of 2,6-di-tert-butylpyridine (2,6-DtBP), catalyst loading could be reduced to ≤0.5 mol %, making this the first practical method for the enantioselective hydrogenation of highly substituted N-alkylquinolinium salts.14

The initial asymmetric hydrogenation was performed using the triflate salt 8a. The iodide 8b acquired by the oxidative MCR conditions provided similar results. However, we found that added chloride increased enantioselectivity (Table 3). This is consistent with recent reports implicating the involvement of mixed halide catalysts.15 This procedure now provided conditions to achieve up to 94% of the desired enantiomer in the asymmetric

**Table 2. Impact of Achiral Phosphine Loading and Bases on Conversion and Enantioselectivity**

<table>
<thead>
<tr>
<th>entry</th>
<th>(t-Bu)3P (mol %)</th>
<th>additive (equiv)</th>
<th>conv (%)</th>
<th>er</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>1</td>
<td>none</td>
<td>22</td>
<td>90:10</td>
</tr>
<tr>
<td>2</td>
<td>40</td>
<td>none</td>
<td>50</td>
<td>87:13</td>
</tr>
<tr>
<td>3</td>
<td>80</td>
<td>none</td>
<td>85</td>
<td>85:15</td>
</tr>
<tr>
<td>4</td>
<td>100</td>
<td>none</td>
<td>78</td>
<td>83:17</td>
</tr>
<tr>
<td>5</td>
<td>3</td>
<td>DBU (1.3)</td>
<td>81</td>
<td>89:11</td>
</tr>
<tr>
<td>6</td>
<td>3</td>
<td>KO-t-Bu (1.3)</td>
<td>75</td>
<td>89:11</td>
</tr>
<tr>
<td>7</td>
<td>3</td>
<td>2,6-DtBP (2)b</td>
<td>99</td>
<td>89:11</td>
</tr>
<tr>
<td>8c</td>
<td>1.5</td>
<td>2,6-DtBP (4)b</td>
<td>96</td>
<td>89:11</td>
</tr>
</tbody>
</table>

*5 mol % of n-Bu4I for entries 1–4; 1.3 equiv of LiI for entries 5–8.

b 2,6-di-tert-butylpyridine; c 0.5 mol % of [Ir(COD)Cl]2, 1 mol % of Morphos.

**Table 3. Impact of LiCl on Enantioselective Hydrogenation of 8b**

<table>
<thead>
<tr>
<th>entry</th>
<th>mol % Ir</th>
<th>additive</th>
<th>equiv</th>
<th>conv (%)</th>
<th>er</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>0.5</td>
<td>none</td>
<td>N/A</td>
<td>99</td>
<td>90:10</td>
</tr>
<tr>
<td>2</td>
<td>0.5</td>
<td>LiCl</td>
<td>3</td>
<td>99</td>
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</tr>
<tr>
<td>3</td>
<td>0.75</td>
<td>LiCl</td>
<td>3</td>
<td>99</td>
<td>94:06</td>
</tr>
</tbody>
</table>

*2:3:1 ratio of (S)-Morphos/(t-Bu)3P/[Ir(COD)Cl]2, 2.5 equiv of 2,6-DtBP.
hydrogenation of a tetrasubstituted quinolinium salt. Crystalline 11 was isolated from the hydrogenation mixture in 82% yield.

The final conversion of 11 to 1 consisted of deprotection with HCl in acetic acid, yielding vabicaserin 1 in 92% yield and 92% ee after crystallization (eq 1). An additional recrystallization from EtOH/MTBE provided the target compound in 86% yield and 99.9+% ee.

\[
\begin{align*}
11 & \xrightarrow{1) \text{HCl, AcOH, 115 }^{\circ} \text{C}} 1 \\
& \xrightarrow{2) \text{95% EtOH, MTBE}} 79\% \text{ yield} \\
& \quad 99.9+\% \text{ ee}
\end{align*}
\]

In summary, we have developed an efficient, asymmetric synthesis of the 5-HT\textsubscript{2c} agonist vabicaserin in four chemical steps and 54% overall yield from commercially available benzodiazepine, a > 2 fold increase compared to the resolution approach to 1. The short synthesis was enabled by the discovery of two new reaction processes: an oxidative, multicomponent reaction to affect the quinolinium ring assembly in one step, and an unprecedented asymmetric hydrogenation of a highly substituted quinolinium salt. Notably, this work provides the first example of an enantioselective reduction of a quinoline heterocycle with substitution at the 4 position.

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Supporting Information Available. Copies of \textsuperscript{1}H NMR and \textsuperscript{13}NMR, as well as experimental procedures. This material is available free of charge via the Internet at http://pubs.acs.org

The authors declare no competing financial interest.