Asymmetric synthesis of chiral cycloalkenone derivatives via palladium catalysis†

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The palladium-catalyzed oxidative desymmetrization of meso-dibenoates yields γ-benzoyloxy cycloalkenones in good yields and with excellent levels of enantioselectivity. These compounds serve as precursors to a broad range of substituted cycloalkenones, including well-established synthetic building blocks and elaborated cycloalkanone derivatives. The ability to prepare both enantiomers of the oxidative desymmetrization products enables a unified strategy toward stereochemically diverse epoxyquinoid natural products.

Introduction

Chiral cycloalkenones are prominent structural motifs in organic synthesis.1 Chiral γ-substituted cycloalkenones in particular are important building blocks in both natural product and pharmaceutical synthesis. Heteroatom-substituted cycloalkenones such as 1–5 (Fig. 1) have been used extensively in these endeavors.2 Chemoselective elaboration of these building blocks may also provide densely substituted products in which every carbon is differentially functionalized. This is exemplified by cyclohexenone derivative 6, a general strategy that encompasses a number of natural products.

A number of syntheses of such chiral cycloalkenones have been reported, many of which involve enzymatic processes or multi-step derivations of chiral pool materials.3 These well-established approaches can provide products with very high levels of enantioenrichment, but they may not permit access to both product enantiomers without requiring lengthier or otherwise less practical synthetic sequences. This consideration is of particular importance given that both enantiomers of certain building blocks (e.g., 1 and ent-1) have been employed in synthesis.4

Asymmetric catalysis may present an opportunity to prepare both enantiomers of a target with equal facility through catalyst control. These preparations may also benefit from high atom and step economy.5,6 Several innovative, catalytic asymmetric syntheses of γ-substituted cycloalkenones have been described.7 These methods often target or perform best for a specific cycloalkenone ring size or type of γ-substituent. A general method for the synthesis of chiral cycloalkenones of various ring sizes and with different γ-substituents would complement these approaches.

Herein we report such a strategy using asymmetric palladium catalysis. This process affords enantioenriched cyclopentenones, cyclohexenones, and cycloheptenones bearing various heteroatom substituents at the γ-position. This protocol also enables the efficient, asymmetric synthesis of more densely functionalized cyclohexenone-derived natural products.

Our group has demonstrated that palladium-catalyzed allylic alkylation presents a unique method for the oxidation of allylic esters (Fig. 2).8 In this process, ionization of allylic ester 7 generates π-allylpalladium intermediate 8, which undergoes selective O-alkylation with nitronate 9 to yield 10. Fragmentation then provides α,β-unsaturated product 11 and oxime 12, which can be recycled to 9.9

Meso-1,4-allylic dibenoates are excellent substrates for this process. When compounds 13–15 (Fig. 3) are subjected to the reaction conditions in the presence of our chiral ligand L1, an oxidative desymmetrization proceeds. This delivers

Fig. 1 Diverse chiral cycloalkenones.

Fig. 2 Pd-catalyzed allylic oxidation.
γ-benzoyloxy cycloalkenones 16–18 in good yields and with excellent enantioselectivity. Importantly, either enantiomer of the desired product can be obtained from the same meso precursor simply by selecting either (R,R)- or (S,S)-L1. With the aforementioned building blocks and natural product motifs serving as motivation, we sought to expand upon this entry into chiral γ-substituted cycloalkenones. We envisioned pairing this oxidative desymmetrization with the palladium-catalyzed allylic alkylation of heteroatom nucleophiles. This would enable access to a diverse set of γ-substituted cycloalkenones (Fig. 4). Complementing this, ester hydrolysis would provide γ-hydroxy cycloalkenones.

We also anticipated that the γ-substituents of the cycloalkenones so produced would direct the diastereoselectivity of subsequent intra- and intermolecular reactions involving the enone moiety. This would enable the rapid assembly of molecular complexity with excellent stereocontrol. Further manipulations would install additional functionality at the remaining positions, leading to fully elaborated structures of the type 6.

Results and discussion

In pursuit of these goals, it was necessary to perform the synthesis of cyclohexenone 17 from meso-diester 14 on larger (multi-gram) scale. It was soon discovered that the isolated yields on such a scale were highly variable and generally lower than yields on smaller scale. This was the case even when the consumption of 14 was high (TLC). Contemporaneously, other operations involving 17 revealed its sensitivity toward aqueous or alcoholic base. Suspecting that such a base (e.g., KOH, from the KH used to generate nitronate 9) may have been the cause of the lowered yields on larger scale, we instituted several operational modifications to improve the robustness of the oxidative desymmetrization reaction. Specifically, in the preparation of 9, the commercial KH dispersion in mineral oil was washed with THF, rather than the more typical hexanes, to remove not only the oil but also any KOH present in the material. In addition, the reactions were limited in duration while monitoring for the consumption of the meso-dibenzoate (TLC). Once the dibenzoate was fully or largely consumed, the reaction mixture was thoroughly quenched with pH 7 buffer prior to extractive workup.

Validation of these modifications came in the form of increased and more consistent reaction yields. For example, the oxidation of 14 → 17 proceeded in a reproducible ca. 67% yield on multi-gram scale (9 mmol, a 35-fold increase in scale from our original report) while maintaining excellent enantioselectivity (Fig. 5). Applying this modified protocol to 13 and 15 was similarly successful.

The palladium-catalyzed allylic alkylation (Pd-AA) of cycloalkenones 16–18 with heteroatom nucleophiles proved highly effective.9 Chiral γ-substituted cyclopentenone, cyclohexenone, and cycloheptenone products were obtained in good yields and with very good to excellent levels of enantioselectivity (Table 1). Alkylation with carboxylic, phenolic, and alcoholic nucleophiles successfully delivered oxygenated building blocks ent-2 and 4 as well as products 19–21. This synthesis of ent-2 is particularly significant, as alkyl alcohols are typically poor nucleophiles in palladium-catalyzed allylic alkylation.10 The preparation of acetoxynone 19 is also noteworthy, as our attempts to engage the diacete analogue of 14 in the oxidative desymmetrization were unsuccessful. Thus, this strategy of Pd-catalyzed oxidative desymmetrization and Pd-catalyzed transesterification represents a new and useful solution to this limitation. Alkylation with nitrogen nucleophiles was also successful, with potassium phthalimide as well as alkyl amines reacting smoothly to provide products 22–25.

Notably, the products of conjugate addition were not observed in any of these cases. Moreover, despite the sensitivity of 17 toward hydroxide or alkoxide base, the basic nature of the reactions did not hinder us from obtaining good yields of the desired products.
The transformations proceeded with high levels of chirality transfer. This occurred despite the potential for racemization via \( \pi-\sigma-\pi \) interconversion involving an \( O \)-bound palladium enolate, a means by which the enantiotopic faces of the \( \pi \)-allylpalladium species might be equilibrated as in the case of butenolide electrophiles. The substitution appears to proceed primarily through overall double inversion, with the absolute stereochemistry of the starting material being conserved in the product.

Further elaboration of these Pd-AA products to more complex cycloalkanone derivatives could be achieved through reactions that engaged the enone moiety with the \( \gamma \)-substituent. For example, exposure of ethanolamine derivative 24 to buffered TBAF effected desilylation and oxa-Michael addition, providing bicyclic morpholine derivative 26 (Fig. 6). The rapid, stereocontrolled synthesis of this compound further highlights the utility of this tandem oxidative desymmetrization/allylic substitution process: the former reaction sets high stereochemical purity at the \( \gamma \)-position that is relayed in the diasteroselective conjugate addition, and the latter enables the introduction of a useful bis-nucleophile unit for a convergent synthesis of 26.

Compounds 16–18 were also converted to the corresponding enantioenriched \( \gamma \)-hydroxycycloalkenones 1, 5, and 27 (Fig. 7). In evaluating methods to hydrolyze these base-sensitive derivatives, Table 1 presents the scope of Pd-catalyzed allylic alkylation (Pd-AA) of 16–18.

<table>
<thead>
<tr>
<th>Nucleophile</th>
<th>Electrophile</th>
<th>Product</th>
<th>Yield [%]</th>
<th>ee [%]</th>
</tr>
</thead>
<tbody>
<tr>
<td>NaOAc</td>
<td>16</td>
<td>4</td>
<td>64</td>
<td>93</td>
</tr>
<tr>
<td>NaOAc</td>
<td>17</td>
<td>19</td>
<td>75</td>
<td>96</td>
</tr>
<tr>
<td>OH</td>
<td>17</td>
<td>19</td>
<td>62</td>
<td>91</td>
</tr>
<tr>
<td>HO</td>
<td>17</td>
<td>20</td>
<td>80</td>
<td>98</td>
</tr>
<tr>
<td>Br</td>
<td>17</td>
<td>21</td>
<td>66</td>
<td>92</td>
</tr>
<tr>
<td>N(H(nPr)₂)</td>
<td>17</td>
<td>23</td>
<td>69</td>
<td>99</td>
</tr>
<tr>
<td>TBSO-NH₂</td>
<td>17</td>
<td>24</td>
<td>85</td>
<td>87</td>
</tr>
<tr>
<td>Kn</td>
<td>18</td>
<td>25</td>
<td>63</td>
<td>94</td>
</tr>
<tr>
<td>PMB</td>
<td>18</td>
<td>25</td>
<td>69</td>
<td>99</td>
</tr>
</tbody>
</table>

\( a \) Reactions performed with 0.1 or 0.2 mmol 16–18 (99% ee), 2.5 mol% \((n°C_{6}H_{7})_{3}PdCl_{2}\), and 15 mol% PPh\(_{3}\) at 0.2 M and 23 °C for 0.5–20 h. For complete details, see the ESL. \( b \) Determined by chiral HPLC. \( c \) Performed at 0 °C with 2.5 mol% Pd\(_{2}db\(_{3}\))CHCl\(_{3}\) and 7.5 mol% \((S,S)\)-L\(_{1}\). 10% of 16 (79% ee) was recovered. \( d \) Cs\(_{2}CO\(_{3}\) added. \( e \) 7.5 mol% dpf used in place of PPh\(_{3}\). PMB = para-methoxybenzyl, TBS = tert-butylidimethylsilyl.

Fig. 6 Elaboration of 24 to bicyclic ketone 26. Reagents and conditions: TBAF · 3H\(_{2}\)O, AcOH, THF, 0 → 23 °C, 2 h, 77%.
compounds, Me$_3$SnOH—which is most typically used for the hydrolysis of methyl esters to their acids$^{14}$—emerged as the optimal reagent for this transformation. To the best of our knowledge, this represents its first use for the saponification of an acyloxy group at a secondary stereocenter, to yield a chiral alcohol (Fig. 6), while others display diastereoselectivity.

Embodied by the chemo-, enantio-, and diastereoselective preparation of the products illustrated in Table 1, Fig. 6 and 7, we next examined the synthesis of more densely functionalized cyclohexenone-derived natural products. We targeted several members of the epoxyquinoid class of natural products, compounds 28–35 (Fig. 8).$^{16}$ These biologically active natural products bear an intriguing stereochemical feature: although structurally similar to each other, certain members exhibit the opposite sense of absolute stereochemistry ($e.g.$, 28 vs. 29–33), while others display different relative stereochemistry between the epoxide and the alcohol ($e.g.$, 31 and 32 vs. 34 and 35). Moreover, for at least one member of this family (harveynone, 29), both enantiomers are naturally occurring yet are derived from dissimilar sources and exhibit different biological activities.$^{16a}$

We proposed a unified strategy toward the structural cores of 28–35 from chiral cyclohexenones 17 and ent-17 (Fig. 9). This approach would exploit the ability of an appropriately $\gamma$-substituted cyclohexenone to direct the diastereoselectivity of an intermolecular nucleophilic epoxidation. For instance, $\gamma$-hydroxycyclohexenone 1 is known to undergo hydroxyl-directed syn epoxidation.$^{17}$ Thus, the preparation of 1 from 17 establishes access to cis product 36, the core structure of 34 and 35. In contrast, 17 could be expected to undergo anti-selective epoxidation. Hydrolysis would then deliver trans product 37, featuring the absolute and relative stereochemistry of 28. Performing the latter procedure with ent-17 would provide ent-37, a precursor to 29–33. Through this strategy, the core structures of all of the stereochromically diverse products 28–35 could be accessed from a common precursor, meso-diester 14.

Focusing on the synthesis of (−)-tricholomenyn A (28), we investigated the epoxidation of 17 (Fig. 10). Pleasingly, upon exposure to $\text{H}_2\text{O}_2$ and catalytic benzyltrimethylammonium hydroxide (Triton B),$^{18}$ cleanly underwent epoxidation to give 38 as a single diastereomer ($^1\text{H NMR}$) and in 99% ee (chiral HPLC). Thus, the oxygenation and the absolute and relative stereochemistry of the trans epoxypiquinoids were set, with excellent enantio- and diastereoselectivity, in three steps from commercial 1,3-cyclohexadiene.$^{19}$ The fact that the oxidative desymmetrization delivered 17 with a $\gamma$-benzoyloxy substituent proved to be a significant synthetic advantage, as the epoxidation of the $\gamma$-acetoxy analogue led to poorer diastereoselectivity and to the formation of byproducts. These results are consistent with the observations of Bayón, Figueredo, and co-workers.$^{20}$

Hydrolysis and acetylation proceeded smoothly, affording 39. Completion of the synthesis then required oxidation of the ketone to an enone, $\alpha$-iodination, and side chain installation. A number of oxidation conditions were investigated on both 38 and 39 ($e.g.$, selenylation/elimination, Saegusa-Ito oxidation, bromination/elimination, IBX oxidation of a silyl enol ether,$^{21}$ and Pd(TFA)$_2$-catalyzed aerobic oxidation$^{22}$), all of which led to no or trace product and/or decomposition. The oxidative desymmetrization delivered 17 with a $\gamma$-benzoyloxy substituent proved to be a significant synthetic advantage, as the epoxidation of the $\gamma$-acetoxy analogue led to poorer diastereoselectivity and to the formation of byproducts. These results are consistent with the observations of Bayón, Figueredo, and co-workers.$^{20}$

As oxidations of this sort have been performed on $\gamma$-silyloxy, $\alpha$,$\beta$-epoxyketones,$^{18}$ we suspected that the
incompatibility was due to the ester moiety, the electrophilic nature of which promoted side reactions. However, rather than lengthening the synthesis with a protection/deprotection sequence involving alcohol 37, we investigated procedures in which the ketone enolate of 39 could be both formed and reacted at low temperature. We turned to the method of Mukaiyama,24 which involved selenylation of the lithium enolate of 39 with 40 and subsequent elimination, both of which proceeded efficiently at −78 °C to deliver 41.

The conversion of 41 to iodoenone 42 initially proved challenging, as typical conditions (I2 in CH2Cl2/pyridine25) led to decomposition. This may have been the result of interactions between the acetoxy group of 41 and these co-solvent amounts of pyridine. Success was realized by applying the conditions of Benhida26 as (applied by Hayashi26b), which employ only stoichiometric pyridine and a more activated iodinating reagent.

From 42, cross coupling with known stannane 4327 afforded (−)-tricholomenylen A (28), the analytical data for which matched literature data. The observed optical rotation ([α]D28° = −234.6° (c = 0.96, CH2Cl2)) was in line with what reported for material obtained using enzymatic resolution as the source of chirality ([α]D28° = −235.7° (c = 1.47, CH2Cl2)).27

Synthetic access to natural products 29–33 was then established through the synthesis of ent-37 from ent-17 (prepared from 14 in 98% ee using (S,S)-L1). Silylation afforded 44, from which syntheses of 31–33 are known (Fig. 11). The analytical data for 44, including optical rotation, was in line with data reported for material prepared through an enzymatic approach.18

![Fig. 11 Syntheses of epoxygenoids 29–33. Reagents and conditions: (a) TBSCl, imidazole, CH2Cl2, 23 °C, 16 h, 88%, (b) LiHMDS, THF, −78 °C, 30 min, then 40. −78 °C, 1 h, 66%.](image)

The asymmetric synthesis of 44 has itself been the subject of past studies, for both natural product synthesis and pharmaceutical research.18,28 Prior syntheses have employed enzymatic reduction,18 enzymatic resolution,29m chiral pool material,29h or asymmetric deprotonation.26e strategies that either required eight or more steps or, in the latter case, delivered more modest (77–85%) ee. The five-step procedure presented herein, which provides 98–99% ee, complements these approaches.

The oxidation of ketone 44 to enone 45 has previously been performed via silyl enol ether formation followed by either selenylation/elimination18 or stoichiometric Saegusa-Ito oxidation.26m Given our successful oxidation of ketone 39 via the Mukaiyama protocol, we considered performing this reaction on 44 to deliver 45 directly in one step. Indeed, this protocol proved very effective, providing the enone in good yield. From 45, syntheses of 29 and 30 are known.29

Conclusions

We report a strategy for the synthesis of a diverse set of chiral cycloalkenone derivatives via asymmetric palladium catalysis. Using a newly optimized reaction procedure, the oxidative desymmetrization of meso-dibenzoates 13–15 affords cycloalkenones 16–18 in good yields and with excellent enantioselectivity. These products serve as platforms from which a variety of substituted cycloalkenones can be prepared through palladium-catalyzed allylic alkylation, including established building blocks such as ent-4 and 4. The products are obtained in good yields and with high levels of enantioenrichment. This Pd-AA approach also enables the convergent introduction of useful functionality, allowing for the synthesis of more elaborate cycloalkanone derivatives. The hydrolysis of 16–18 to enantiomerically pure hydroxycycloalkenone building blocks is also demonstrated.

Cyclohexenones 17 and ent-17 provide entry points to epoxygenoid natural products 28–33 through stereoselective transformations and through reactions that elaborate the entire cyclohexenone structure. This approach to these stereocchemically diverse products proceeds with excellent levels of enantio- and diastereoselectivity, and it establishes a concise, unified strategy starting from a single meso precursor (14).

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Notes and references


4 As examples, see ref. 2e and 2f for the use of ent-1 and 1, respectively.


9 Nitronate 9 is prepared by the oxidation of oxime 12 with NaBO₃ in AcOH followed by in situ deprotonation with KHMDS, thus enabling this recycling.

10 The pH 7 quench may be the most significant of these modifications: as the workup and isolation associated with a larger scale reaction necessarily takes more time compared to a smaller scale reaction, an incompletely quenched larger scale reaction mixture may translate to a longer period of exposure to reactive impurities such as base. The results listed in Fig. 3 were obtained using this modified reaction procedure (≥1 mmol scale). For complete experimental details, see the ESI†.


13 The absolute stereochemistry of compounds 4, 19, and ent-2 was determined by comparison of their optical rotations to literature values, and the stereochemistry of the remaining Pd-AA products is assigned by analogy.


15 The absolute stereochemistry of the alcohol products was determined by comparison of their optical rotations to literature values, and the % ee was determined by chiral HPLC after their conversion to benzoates 16-18 (BzCl; pyridine; for details, see the ESI†).


19 Dibenzoate 14 is readily prepared in one step from 1,3-cyclohexadiene (J.-E. Bäckvall, K. L. Granberg, R. B. Hopkins, *Acta Chem. Scand.*, 1990, 44, 492); for a multigram synthesis, see the ESI†.


