Biocatalytic One-Carbon Ring Expansion of Aziridines to Azetidines via a Highly Enantioselective [1,2]-Stevens Rearrangement

David C. Miller,† Ravi G. Lal,‡ Luca A. Marchetti,†§ and Frances H. Arnold*†

† Division of Chemistry and Chemical Engineering, California Institute of Technology, Pasadena, California 91125
‡ Present Address: Department of Biosystems Science and Engineering, ETH Zürich, 4058 Basel, Switzerland

ABSTRACT: We report enantioselective one-carbon ring expansion of aziridines to make azetidines as a new, abiological activity of engineered ‘carbene transferase’ enzymes. A laboratory-evolved variant of cytochrome P450OM3, P411-AzetS, not only overrides the inherent reactivity of aziridinium ylides to undergo chelotropic extrusion of ethylene, it also exerts unparalleled stereocontrol (99:1 er) over a [1,2]-Stevens rearrangement, a notoriously challenging reaction class for asymmetric catalysis. These unprecedented selectivities enable an entirely new strategy for the synthesis of chiral azetidine products from readily available synthetic precursors. The utility of this reaction is highlighted by the synthesis of an enantiopure azetidine on gram scale. The exquisite selectivity of the enzyme enables new-to-nature ring-expansion chemistry that overcomes a longstanding synthetic problem.

Ring-size manipulation has emerged as a powerful strategy to convert readily available cyclic structures into ring-expanded or ring-contracted compounds that are more difficult to synthesize using conventional means.1 In particular, “cut and sew” strategies relying on transition-metal catalyzed oxidative addition to form C-C bonds have emerged as powerful tools for insertion of carbon monoxide or two-carbon fragments such as olefins and alkynes into existing rings to effect one- or two-carbon ring expansions, respectively.2 For nitrogen-containing heterocycles, one possible strategy is to induce a [1,2]-Stevens rearrangement to enact a one-carbon ring expansion.3 Pioneering work by Hata, West, and Couty has demonstrated the power of this approach for 4- to 5-membered ring expansions, wherein treatment of an azetidine with a diazo compound in the presence of a copper catalyst provides facile access to the corresponding pyrrolidine.4 Conceptually, carbene transfer followed by an intramolecular [1,2]-Stevens rearrangement complements “cut and sew” reactions for non-carbonylative, one-carbon homologation of nitrogen-containing compounds. Given the prevalence of nitrogen heterocycles across numerous sectors of chemical industry, especially pharmaceuticals,5 general strategies to extend this approach to other nitrogen-containing compounds would represent a powerful new approach for the synthesis of important pharmacophores.

Azetidines are valuable isosteres of pyrrolidines and piperidines, as they often have enhanced metabolic stability and potency compared to their 4- and 6-membered ring congeners.6 Azetidines are underexplored in medicinal chemistry due to the lack of robust synthetic strategies to access them,7 particularly in enantioregulated form.8 State-of-the-art methodologies such as [2+2] cycloadditions of imines and olefins,9 have been developed to address the lack of robust methods for the synthesis of azetidines.10 However, direct synthesis of enantioenriched azetidines using asymmetric catalysis has remained elusive. Application of a ring-expansion strategy for the asymmetric, one-carbon homologation of aziridines via carbene insertion would be a powerful new entry for the synthesis of chiral azetidines. However, this approach comes with two major selectivity challenges that, to date, have not been addressed in a general fashion. The first is the innate reactivity of the intermediate aziridinium ylides, which undergo highly favorable chelotropic extrusion of ethylene in many contexts.11 While these intermediates can be harnessed in [2,3]-Stevens rearrangements and other ring-opening reactions,12 no examples of [1,2]-Stevens rearrangements from aziridinium ylides have been reported. Secondly, the diradical mechanism of the [1,2]-Stevens rearrangement13 has made it a challenging reaction for asymmetric synthesis. Few asymmetric variations of this reaction have been reported.14 Enantiopure quaternary ammonium salts can undergo [1,2]-Stevens rearrangements with α-toc chirality transfer;15 however, escape of the radical pair from the solvent cage is often competitive with radical recombination16 and erosion of enantiopurity is often observed in these reactions. The joint selectivity challenges presented by the one-carbon ring expansion of aziridines into azetidines not only require a potential catalyst to select the [1,2]-Stevens rearrangement in preference to chelotropic extrusion of ethylene but also to exert enantiocontrol over a potential diradical intermediate. We are unaware of any successful examples of a one-carbon ring expansion of aziridines through a [1,2]-Stevens rearrangement strategy.
Nature utilizes ring size manipulation in the biosynthesis of natural products, with common strategies for biocatalytic one-carbon ring expansion including oxidative ring expansions and carboxylation. We hypothesized that engineered carbone transferases could potentially extend enzymatic ring expansions through a [1,2]-Stevens-type mechanism (Figure 2). Over the past decade, enzymes derived from cytochrome P450 BM3, such as cytochromes P411, and other hemoproteins have emerged as powerful catalysts for carbene transfer reactions, and formation of strained rings such as cyclopropanes and cycloprenes with excellent stereoselectivities has been reported. The most common reactions of the iron-carbenoid intermediate are additions across π-systems or X–H bond insertions: C–N bond insertion through Stevens rearrangements of any kind have yet to be reported. The enzyme may be able to control the stereoselectivity of this reaction. Hemoproteins demonstrate high stereoselectivity in radical reactions, both in their native activity and in abiological reactions cultivated through protein engineering. In addition, we envisioned that the active site of an enzyme could mimic solvent cage effects which are known to exert control over radical recombination in [1,2]-Stevens rearrangements. Provided an enzyme could overcome both of these selectivity challenges, this approach could be a facile and powerful strategy to access enantioenriched azetidines.

We initiated our studies by screening a panel of hemoproteins for the model reaction of benzyl aziridine carbonate with ethyl diazoacetate (EDA) as a carbene precursor to provide enantioenriched azetidine (Table 1) in suspensions of Escherichia coli (E. coli) whole cells. The highest activity for this reaction was observed with a variant of P411BM3-CIS with the additional mutations P248T, I263G, and L437F (“Parent F2”), providing the product with 3.7% yield, 73 total turnover number (TTN), and 90:10 er for the (S)-enantiomer (Entry 1). Parent F2 is derived from hemoproteins originally engineered for the cyclopropanation of heteroatom-substituted olefins and is 17 mutations away from its wild-type progenitor, cytochrome P450BM3 from...
Bacillus megaterium which natively catalyzes the oxidation of long-chain fatty acids. Control experiments revealed that hemin is unable to catalyze this reaction (see SI for details). Encouraged by this promising initial activity and high enantioselectivity, we chose Parent F2 as a starting point for directed evolution to improve enzyme performance using iterative site-saturation mutagenesis (SSM) of residues located in the heme domain (Entries 2–11) and screening for improved azetidine yield by gas chromatography.

Table 1: Lineage and Reaction Optimization

<table>
<thead>
<tr>
<th>Entry</th>
<th>Variant</th>
<th>Mutations Relative to Prior Generation</th>
<th>TTN</th>
<th>Yield (%)</th>
<th>e.r.</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Parent F2</td>
<td>None</td>
<td>73</td>
<td>3.6</td>
<td>90:10</td>
</tr>
<tr>
<td>2</td>
<td>F2.1</td>
<td>G263Y</td>
<td>70</td>
<td>3.5</td>
<td>75:25</td>
</tr>
<tr>
<td>3</td>
<td>F2.2</td>
<td>T327V</td>
<td>126</td>
<td>6.3</td>
<td>56:44</td>
</tr>
<tr>
<td>4</td>
<td>F2.3</td>
<td>A330T</td>
<td>193</td>
<td>9.6</td>
<td>59:41</td>
</tr>
<tr>
<td>5</td>
<td>F2.4</td>
<td>H266P</td>
<td>394</td>
<td>19.7</td>
<td>62:38</td>
</tr>
<tr>
<td>6</td>
<td>F2.5</td>
<td>M177Q</td>
<td>699</td>
<td>34.9</td>
<td>94:6</td>
</tr>
<tr>
<td>7</td>
<td>F2.6</td>
<td>T436G</td>
<td>945</td>
<td>47.3</td>
<td>93:7</td>
</tr>
<tr>
<td>8</td>
<td>F2.7</td>
<td>L233F</td>
<td>997</td>
<td>48.8</td>
<td>94:6</td>
</tr>
<tr>
<td>9</td>
<td>F2.8</td>
<td>T149M</td>
<td>1041</td>
<td>52.0</td>
<td>99:1</td>
</tr>
<tr>
<td>10</td>
<td>F2.9</td>
<td>R474Q</td>
<td>1123</td>
<td>59.7</td>
<td>99:1</td>
</tr>
<tr>
<td>11</td>
<td>P411-AzetS</td>
<td>M118K</td>
<td>1198</td>
<td>59.9</td>
<td>99:1</td>
</tr>
</tbody>
</table>

Sites were selected for mutagenesis based on success in previous directed evolution campaigns of P450BM3 as well as prior knowledge of residues responsible for substrate binding and catalysis in the heme domain of this protein scaffold. Ten beneficial mutations were identified during this campaign, resulting in a more efficient ‘azetidine synthase’ (P411-AzetS) with a net improvement of 16-fold in TTN and 99:1 e.r. With P411-AzetS in hand, we next examined the impact of varying the reaction conditions on the product yield (Entries 12–17). Notably, increasing the scale from 4 μmol to 100 μmol resulted in an increase in the reaction yield. When the concentrations of 1 and EDA were doubled to 20 mM and 30 mM, respectively, a decrease in reaction yield was observed (although TTN increased). The ring expansion reaction also proceeded in clarified cell lysate, albeit with decreased yields when compared to analogous reactions performed with whole-cell suspensions. Lastly, decreasing the reaction temperature from 22 to 4 °C did not have a meaningful impact on the reaction yields when run in whole-cell suspensions.

Next, we sought to examine the substrate scope of this reaction. When this reaction was run at 0.5-mmol scale, azetidine 2 could be formed in 75% yield, 1490 TTN, 67% yield, and 99:1 e.r. Other aromatic groups could be used in lieu of a phenyl group with uniformly high enantioselection observed in all cases.

Scheme 1: Substrate Scope

Reactions were performed on the designated scale and run for 16 h with 10 mM of 1, 15 mM of EDA, and 5 μM of protein. TTN and yields were determined via GC analysis of crude reaction mixtures relative to internal standard and represent the average of three experiments. The enantiomeric ratio (e.r.) was determined by chiral GC.

Sites were selected for mutagenesis based on success in previous directed evolution campaigns of P450BM3 as well as prior knowledge of residues responsible for substrate binding and catalysis in the heme domain of this protein scaffold. Ten beneficial mutations were identified during this campaign, resulting in a more efficient ‘azetidine synthase’ (P411-AzetS) with a net improvement of 16-fold in TTN and 99:1 e.r. With P411-AzetS in hand, we next examined the impact of varying the reaction conditions on the product yield (Entries 12–17). Notably, increasing the scale from 4 μmol to 100 μmol resulted in an increase in the reaction yield. When the concentrations of 1 and EDA were doubled to 20 mM and 30 mM, respectively, a decrease in reaction yield was observed (although TTN increased). The ring expansion reaction also proceeded in clarified cell lysate, albeit with decreased yields when compared to analogous reactions performed with whole-cell suspensions. Lastly, decreasing the reaction temperature from 22 to 4 °C did not have a meaningful impact on the reaction yields when run in whole-cell suspensions.

Next, we sought to examine the substrate scope of this reaction. When this reaction was run at 0.5-mmol scale, azetidine 2 could be formed in 75% yield, 1490 TTN, 67% yield, and 99:1 e.r. Other aromatic groups could be used in lieu of a phenyl group with uniformly high enantioselection observed in all cases.

Scheme 1: Substrate Scope

Reactions were performed on 0.5-mmol scale unless otherwise specified. Analytical yields and TTN were determined by GC-FID. Yields for isolated and purified material are designated in parentheses. The e.r. was determined by Chiral GC. For 0.5-mmol scale reactions, all numbers reported represent the average of two trials. For 10-mmol scale reaction, numbers reported represent one run.
Notably, a thiophene-bearing aziridine could undergo cyclopropanation expansion to azetidine 3 with no observed cyclopropanation byproducts. This selectivity is notable not only because thiophenes are known to react with EDA-derived metal carbenoids under mild conditions, but also because Parent F2 was originally engineered to perform cyclopropanation of heteroatom-substituted olefins. Fluorine substituents were also tolerated on the arenne ring at the para, meta, and ortho positions to furnish fluorinated products 4-6. In addition to EDA, other diazoacetate compounds could participate in one-carbon ring expansion with at least 99:1 er (7-8). When methyl diazoacetate was used as the carbene precursor to yield 9, a notable decrease in er (81:19) was observed. One hypothesis for this decrease in enantioselectivity is that the smaller aliphatic chain allows for greater conformational freedom of the iron porphyrin carbene intermediate or the putative diradical intermediate. This explanation is consistent with prior work on enzyme-mediated carbene transfer reactions using perfluoroalkyl-stabilized diazo compounds as carbene precursors, where the substrate chain length has a profound influence on the absolute stereochemistry of the reaction. The reaction could also be scaled up from 0.5-mmol scale to 10-mmol scale to furnish 2 in 1220 TTN, 61% yield, and 99:1 er with an isolated yield of 1.44 g (55% yield), demonstrating that gram-scale production of enantioenriched azetidines is feasible using this platform. Taken together, the unprecedented chemoselectivity observed demonstrates the tremendous synthetic potential of this approach for the synthesis of enantioenriched azetidines. Currently, P411-AzetS and its lineage perform poorly with other classes of nitro-groups enable activation of unstrained C-C bonds and selectivity to the synthesis of other classes of azetidines are ongoing.

In summary, we have demonstrated unprecedented hemoprotein-catalyzed [1,2]-Stevens rearrangement in the context of a one-carbon ring expansion of aziridines to azetidines. This system not only represents a rare example of a highly enantioselective [1,2]-Stevens rearrangement of ammonium ylides, it also exhibits unprecedented chemoselectivity for rearrangement of aziridinium ylides over chelotropic extrusion of ethylene. We anticipate that this platform for azetidine synthesis can be extended to other classes for synthesis of other enantiomer-rich azetidine cores, providing facile access to valuable chiral building blocks. We are optimistic that the reactivities and selectivities observed can be applied to other [1,2]-Stevens rearrangements as well, providing the grounds for future work in this area towards the synthesis of chiral amines.

ASSOCIATED CONTENT
The Supporting Information is available free of charge at: Supporting Information Placeholder

AUTHOR INFORMATION

Corresponding Author
Frances H. Arnold – Division of Chemistry and Chemical Engineering, California Institute of Technology, Pasadena, California 91125, United States; orcid.org/0000-0002-4027-364X; Email: frances@cheme.caltech.edu

Authors
David C. Miller – Division of Chemistry and Chemical Engineering, California Institute of Technology, Pasadena, California 91125, United States; orcid.org/0000-0002-4560-8824

Ravi G. Lal – Division of Chemistry and Chemical Engineering, California Institute of Technology, Pasadena, California 91125, United States; orcid.org/0000-0001-6943-4147.

Luca A. Marchetti – Present address: Department of Biosystems Science and Engineering, ETH Zürich, 4058 Basel, Switzerland; orcid.org/0000-0002-6100-4465

Notes
The authors declare no competing financial interest.

ACKNOWLEDGMENT
Research was sponsored by the U.S. Army Research Office and accomplished under contracts W911NF-19-D-0001 and W911NF-19-2-0026 for the Institute for Collaborative Biotechnologies. D.C.M. was supported by a Ruth Kirschstein NIH Postdoctoral Fellowship (F32GM128247). The authors acknowledge Dr. Sabine Brinkmann-Chen, Nathaniel Goldberg, and Dr. Nicholas Porter for assistance preparing the manuscript.

REFERENCES


Highly chemo- and stereoselective rearrangement of aziridinium ylides