Generality-Oriented Optimization of Enantioselective Aminoxyl Radical Catalysis


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Abstract
Catalytic enantioselective methods that are general over a broad range of substrates facilitate application in synthetic discovery and development settings; however, truly general catalysts for asymmetric synthesis are rare. Herein, we report a strategy for the oxidative desymmetrization of meso-1,4 diols predicated on a non-traditional optimization protocol utilizing a panel of screening substrates rather than a singular model substrate. Critical to this approach was rational modulation of a peptide sequence incorporating a novel, aminoxyl-based catalytic residue. A general catalyst emerged, providing high selectivity in delivery of enantioenriched lactones across a broad range of diols.

Main Text
Substrate generality is a longstanding aim in the field of enantioselective catalysis.1–3 Even so, specificity-oriented optimization has been broadly adopted in the field of small-molecule enantioselective catalysis, resulting in many catalysts that are tailored for one “model” substrate, but often exhibit diminished selectivity upon surveying a diverse substrate scope (Fig. 1A).4 To address the limited transferability frequently encountered in asymmetric catalysis, we envisioned an optimization strategy with generality as the primary target in catalyst development. This strategy requires parallel screening of a large, diverse catalyst library against a judiciously selected panel of model substrates that represent the chemical space of the target substrate class, rather than a singular substrate.5 Such an approach is reminiscent of the “one-pot-multi-substrate” method to increase screening throughput.6–9 However, this pooled-substrate method introduces challenges in chemical compatibility and product analysis, and thus has seen limited implementation in catalyst development. Our interest focused on advancing the notion of generality in the context of previously elusive transformations using a new catalytic platform. Toward this objective, we envisioned that small synthetic, catalytic peptides would be well-suited for such a study, given the modularity of their constituent amino acids and synthetic accessibility, which provides a strong foundation for a generality-driven optimization campaign.10 Peptide optimization can also bear a resemblance to the venerable directed evolution of enzymes through survey of nth-dimensional catalyst space.11,12 The current study combines the vast amount of catalyst space surveyed with an expansive substrate space, yielding information rich data that can be utilized for subsequent catalyst optimization. We investigated this strategy in an underexplored area in asymmetric synthesis — enantioselective aminoxyl radical catalysis.13
Aminoxyl radicals are a class of persistent open-shell molecules that have found broad applications in organic synthesis. They have been shown as excellent catalysts for the oxidation of various functional groups such as alcohols, amines, and alkenes via intermediate oxoammonium ions. In contrast to the vast number of reactions known to be catalyzed by aminoxyl radicals, enantioselective variants remain rare. Foundational work by Bobbitt and recent advances by Iwabuchi provided salient precedents in asymmetric alcohol oxidation. Nevertheless, these catalysts require lengthy synthetic preparation and lack the structural modularity needed for generality-oriented optimization. Toniolo and co-workers demonstrated that...
oxoammonium ions are compatible within peptidic frameworks and observed enantioselectivity in the kinetic resolution of a singular substrate with modest selectivities ($k_{rel} < 3$). In light of these precedents, we hypothesized that incorporation of an achiral aminoxyl moiety into diverse peptide-based scaffolds could provide a catalyst platform that would enable exploration of generality within the reaction space accessible to aminoxyl catalysis. Accordingly, we designed and synthesized an aminoxyl monomer (Azc-OMe, Fig. 1C) that could be readily incorporated into peptides enabling the preparation of a library of >70 chiral catalysts. Herein, we report the development of the highly enantioselective desymmetrization of meso-diols using aminoxyl-based oligopeptides, for which a generality-oriented optimization campaign led to a unique and versatile catalyst that delivers high selectivity for structurally diverse chiral lactones.

We initiated our study with the desymmetrization of meso-diols (Fig. 2A). In the presence of a chiral catalyst, diol substrate S underwent selective oxidation at one of the enantiotopic hydroxyl groups, with the other hydroxyl group cyclized onto the incipient aldehyde to furnish an intermediate lactol. Subsequent second oxidation delivered chiral lactone L, a class of compounds displaying biological activity and utility as building blocks in organic and materials synthesis (vide infra). We initially utilized a traditional approach by screening a collection of aminoxyl-embedded peptide catalysts in the oxidation of 1,4-diol S1, employing trichloroisocyanuric acid (TCCA) as terminal oxidant and sodium bicarbonate as base. This effort yielded peptide P1 that provided lactone L1 with substantial enantioselectivity (ee = 75%). However, P1 showed diminished selectivity when tested against an array of additional 1,4-diols displaying a variety of steric and electronic profiles, giving a median ee of only 16% (Fig. 2E, column P1). This result underscores the challenge of identifying highly selective catalysts that tolerate a broad scope when focusing on a singular substrate during catalyst optimization.

Aiming for a more general method, we assessed a diverse library of peptide scaffolds with high-throughput experimentation (HTE) using 24-well plate parallel screening and GC analysis. Employing this workflow, the data obtained with a given catalyst is discussed below in terms of the median ee ($\text{ee}_{\text{med}}$), which served as the primary optimization target; excellent conversion for each substrate allowed us to remain agnostic to yield throughout optimization. Substrate selection was guided by mapping the chemical space of commercially available 1,4-meso-diols to ensure diversity of the screening set (Fig. 2B). This analysis revealed two clusters of diols, with the larger cluster representing di-substituted meso-diols and the smaller cluster consisting of a unique group of 2,2,3,3-tetrasubstituted 1,4-diols. From this collection of compounds, 15 structurally diverse substrates with a variety of functionality were included in the model set that effectively sampled the chemical space, consisting of eight monocyclic (S1, S3, S5, S6, S9, S13–15) and five polycyclic (S2, S4, S7, S8, S12) substrates with various ring sizes, as well as one acyclic diol (S10) (Fig. 2C). Additionally, a 1,5-diol (S11) was included to further increase diversity in the optimization set.

With this set of model substrates, catalysts were then evaluated as a function of their constituent residues (Fig. 2D). Beginning with P1 ($\text{ee}_{\text{med}}$ = 16%), two single point changes exhibited increases in $\text{ee}_{\text{med}}$: inversion of stereochemistry at $i+3$ (P2) provided $\text{ee}_{\text{med}}$ of 31% and replacement of the $N\text{Me}_2$ group at the C-terminus with $N\text{HMe}$ (P3) provided an $\text{ee}_{\text{med}}$ of 15% but displayed considerable improvement for three substrates (S1, S6, S9). Auspiciously, P3b, an analog of P3, was crystalline; allowing for the secondary structure of this catalyst to be determined by X-ray crystallography (Fig. 2F) and confirmed by ROESY NMR to adopt a 310-helical motif in solution. Notably, the confluence of changes producing P2 and P3 from P1 was found to be beneficial; P4, possessing both the optimized relative stereochemistry of P2 and the $N\text{HMe}$ C-terminus of P3, provided a significant enhancement in selectivity with an $\text{ee}_{\text{med}}$ of 69% (P4). Further modifications to P4 showed that two additional point changes furnished substantial enhancement in $\text{ee}_{\text{med}}$: replacement of the 5-membered proline (Pro) with the 6-membered homologue pipericolic acid (Pip) at $i+1$ (P5) led to an increase of $\text{ee}_{\text{med}}$ to 74%; concurrently, the inclusion of an $N\text{HCH}_2\text{CF}_3$ substituent at the C-terminal position resulted in peptide P6 that greatly
Figure 2. Generality-oriented optimization through combinatorial screening of catalysts with a substrate library and iterative modification of catalyst structure. (A) Chiral lactone formation via meso-diol desymmetrization. (B) Chemical space of 1,4-diols (UMAP) with molecular fingerprints of commercial meso-1,4-diols from a Reaxys® search. (C) Meso-diol substrates: optimization set. (D) Catalyst structure optimization towards a general catalyst for meso-diol desymmetrization. (E) Optimization data. (F) Crystal structures of peptides P3b and P7.
outperforms catalysts with other C-termini, presenting an ee$_{\text{med}}$ of 83%. Catalyst P7, combining the Pip residue found in P5 with an elongated perfluorinated C-terminus (NHCH$_2$C$_2$F$_3$) and a change to biphenylalanine (Bip) at i+3 gave a further boost of ee$_{\text{med}}$ to 93% with 10 of 15 substrates showing >90% ee, and an additional four substrates displaying ee >60%. X-ray analysis and ROESY NMR were also performed on P7, revealing that both the aminoxyl radical and the active oxoammonium forms of the catalyst share secondary structural features. Notably, the solid-state structure of P7 shows a departure from the 310-helix observed in P3b, displaying instead two successive β-turns (Fig. 2F). Provided the success of the parallel optimization campaign, particular challenges for certain substrates persisted. For example, throughout the optimization, one diol (S15) was recalcitrant towards significant selectivity enhancement, delivering only 26% ee with P7. Parenthetically, a structurally distinct catalyst P8 was identified that provided 71% ee for the corresponding lactone L15.

We then probed the scope and limitations associated with catalyst P7 through desymmetrization on a 0.2-mmol scale (Fig. 3A). Chiral lactones are valuable and versatile building blocks in organic and polymer synthesis. While highly selective methods are available for synthesizing specific types of enantioenriched lactones, a general approach that is agnostic of substrate structure and substitution remains elusive and desirable. We found that ketal protected erythronic acid γ-lactones L1 and L18 were formed in good yields and excellent enantioenrichment. Polycyclic diols were efficiently desymmetrized (L2, L4, L7, L8, L19), providing valuable precursors for pharmaceuticals and chiral ligands. Cyclopentane lactone L3 and fungistatic cyclohexane lactone L9 were formed with 96% ee and 94% ee, respectively. Further evaluation of cyclopropyl diols revealed a requirement for substitution on the ring for high selectivity. While unsubstituted L15 was accessed in only 26% ee, dichloro- and dimethylcyclopropane lactones L5 and L22 were produced with excellent enantioselectivity, the latter of which is a key intermediate in the preparation of numerous natural and unnatural products including COVID drug Nirmatrelvir (Paxlovid) (Fig. 3B). Pyrrolidine-bearing S6 was desymmetrized with excellent yield (92%) and selectivity (95% ee). Linear 1,4-diols were also efficient substrates in this reaction, as lactone L10 was obtained in 80% yield and 91% ee. In a dramatic test of our approach, the desymmetrization of a prochiral linear 1,5-diol bearing a mere methyl group at the central carbon was transformed to δ-lactone L11 with 76% ee, providing access to a monomer for a biodegradable isotactic polymer. Notably, the catalyst is also competent in desymmetrizing a prochiral-at-phosphorous 1,5-diol, affording lactone L20 in 82% ee. An additional 1,5-diol exhibited lower selectivity, with L17 being enriched to only 37% ee. Tetrasubstituted diols are also tolerated with bicyclic substrate S12 yielding a [4.3.1] propellane in 80% yield and 79% ee; tetrasubstituted lactone L16, however, was obtained in only 33% ee. Diphenylcyclohexyl diol S14, which throughout optimization gave poor selectivities, was desymmetrized up to 56% ee. Urea L13, a valuable chiral intermediate in the chemical synthesis of biotin, was accessed in 60% ee. Markedly, a kinetic resolution of intermediate H13 was observed, which preferentially oxidized its minor enantiomer to L13 and allowed for the isolation of H13 in 52% yield and 97% ee when the reaction was stopped at an early stage.

Catalyst P7 exhibits high chemoselectivity for oxidation of diols over lactols (see Fig. 4b), which enabled direct access to these valuable synthons by reducing the oxidant stoichiometry. We showcased this method in the concise formal synthesis of two pharmaceutical candidates, improving upon previous routes in step count. Thus, tricyclic T21, a key intermediate in the synthesis of an HIV protease inhibitor, was obtained through oxidation of acetonide S21 followed by one-pot acid-mediated transacetalization in 64% overall yield with 77% ee (Fig. 3B). In a second example, lactol H8, an intermediate in the synthesis of thromboxane receptor antagonist ifetoban, currently in clinical trials, was obtained on 2 mmol scale in 96% ee and 86% yield via an expedient two-step process from commercial meso-acid anhydride A8. In contrast, traditional syntheses required enantioselective methanolation of A8 to form mono-ester M8 followed by a sequence of redundant functional group interconversions and redox manipulations.
(A) Substrate scope using catalyst P7

<table>
<thead>
<tr>
<th>Substrate</th>
<th>Yield</th>
<th>ee</th>
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<tbody>
<tr>
<td>L1</td>
<td>77%</td>
<td>96% ee</td>
</tr>
<tr>
<td>L2</td>
<td>74%</td>
<td>99% ee</td>
</tr>
<tr>
<td>L3</td>
<td>75%</td>
<td>96% ee</td>
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<tr>
<td>L4</td>
<td>70%</td>
<td>97% ee</td>
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<td>L5</td>
<td>2 mol%</td>
<td>98% yield, 95% ee</td>
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<tr>
<td>L6</td>
<td>92%</td>
<td>95% ee</td>
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<td>L7</td>
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<td>L10</td>
<td>80%</td>
<td>91% ee</td>
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<tr>
<td>L11</td>
<td>68%</td>
<td>76% ee</td>
</tr>
<tr>
<td>L12</td>
<td>80%</td>
<td>79% ee</td>
</tr>
<tr>
<td>L13</td>
<td>67%</td>
<td>60% ee</td>
</tr>
<tr>
<td>L14</td>
<td>70%</td>
<td>56% ee</td>
</tr>
<tr>
<td>L15</td>
<td>69%</td>
<td>26% ee (94% ee, see SI)</td>
</tr>
<tr>
<td>L16</td>
<td>91%</td>
<td>33% ee</td>
</tr>
<tr>
<td>L17</td>
<td>68%</td>
<td>37% ee</td>
</tr>
<tr>
<td>L18</td>
<td>89%</td>
<td>94% ee</td>
</tr>
<tr>
<td>L19</td>
<td>49%</td>
<td>92% ee</td>
</tr>
<tr>
<td>L20</td>
<td>29%</td>
<td>82% ee</td>
</tr>
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</table>

(B) Utility in the synthesis of drug molecules

1. P7, TCCA (0.34 equiv) (oxidation to lactol)
2. TsOH

<table>
<thead>
<tr>
<th>Key intermediate</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>HIV protease inhibitor synthesis</td>
<td>Ref 30</td>
</tr>
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</table>

P22a, Intermediate in Paxlovid

(C) Expanding functional group compatibility using electrochemistry

<table>
<thead>
<tr>
<th>Optimal electrochemical conditions:</th>
</tr>
</thead>
<tbody>
<tr>
<td>P7 (5 mol%), DIAD (2.2 equiv)</td>
</tr>
<tr>
<td>base (A or B), TBAPF6 (0.25 M), CH3CN</td>
</tr>
<tr>
<td>NaHCO3 (5 equiv), CO2 (1 atm; buffer), -50 °C</td>
</tr>
<tr>
<td>B = 2.6-difluorobutyly (4 equiv), r.t.</td>
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Incompatible with chemical oxidation

Figure 3. Substrate scope. Absolute stereochemistry was assigned by analogy to L1, L4, and L6.10 0.001 mol% P7, 96 h, -50 °C, then 2 mol% ACT, 0.3 equiv TCCA, r.t., 6 h.1 Isolated as the methyl ester after SiO2 catalyzed methanalysis.10 10 mol% P7. (a) Substrate scope using P7. (b) Utility of products in drug molecules. (c) Expansion of functional group compatibility using electrochemistry.

While the optimal conditions using TCCA tolerated substrates with diverse steric and electronic profiles, the scope was limited to compounds that were compatible in the presence of a strong chemical oxidant. Indeed, substrates containing alkenes were chlorinated, resulting in diminished yields (0–25%). Such products are nonetheless important synthetic intermediates; for example, unsaturated L23 was leveraged in the preparation of a thromboxane-A2 antagonist.39,40
To address this issue, an electrochemical protocol was developed. The key was to identify a sacrificial oxidant that would not decompose sensitive substrates but would allow for the desired electrochemical reaction to take place in the presence of an electric driving force. In traditional anodic oxidation reactions, protic acids or alkyl halides are used as sacrificial oxidants, but these species proved incompatible in our reaction system likely due to the liberation of Lewis basic anions during their cathodic reduction that led to detrimental catalyst binding. Rapid screening using parallel reactor HTeChem led to the identification of two classes of oxidants that provided high yields and enantioselectivities: dialkyl azodicarboxylates and disubstituted peroxides. We found that diisopropyl azodicarboxylate (DIAD) proved to be optimal, providing products \[ \text{L23–25} \] in good yields and high enantioselectivity, while itself was reduced to innocuous diisopropyl hydrazine-1,2-dicarboxylate (Fig. 3C). In this case, either an organic base 2,6-di-tert-butylpyridine or an inorganic buffer consisting of NaHCO\(_3\) and CO\(_2\) were used.

To establish the upper limit of catalytic turnover, we surveyed decreased catalyst loadings using diethyl ketal \[ \text{S18} \] and found that even 0.001 mol\% catalyst (10 ppm; 2,000-fold decrease from standard 2 mmol\% conditions) promoted the reaction without any loss in yield or enantioselectivity. Thus, \[ \text{P7} \] achieved \( \sim 1.2 \times 10^5 \) turnovers, to our knowledge the highest number recorded for an oligopeptide organocatalyst. The catalyst could also be recycled and reused multiple times as shown in the synthesis of \[ \text{H8} \], further demonstrating the potential practicality of the reported protocol.

Finally, we conducted studies to gain understanding of the mechanistic underpinnings of the observed substrate generality and high enantioselectivity (Fig. 4A). The composition of the resting state of the catalyst was interrogated using in situ diffusion-ordered spectroscopy (DOSY) NMR experiments and direct analysis in real time (DART) high-resolution mass spectrometry, showing the formation of oxoammonium-alkoxide complexes \[ \text{(C)} \] between the catalyst and the diol substrate. Further kinetic isotope effect studies support that the rate-determining step of the reaction is a Cope-elimination from the resting state adduct to provide the desymmetrized aldehyde intermediates \[ \text{(A)} \]. More enantioselective catalysts showed a higher degree of selectivity towards diol over mono-ol oxidation with up to a 26-fold rate difference. These findings depict a structure wherein the spectator alcohol participates in hydrogen bonding with a functional group on the peptide backbone (Fig. 4A), in addition to the covalent association of the reacting alcohol and the oxoammonium catalytic unit. Consistent with this, substrate features can be used to quantitatively classify when a substrate should perform well in this reaction. Using a threshold value of 90\% ee, substrates can be classified based on the difference between the buried volumes \( \Delta V_{\text{bur}} \) of each face of the lactone motif (Fig. 4C), revealing that substrates with a minimum \( \Delta V_{\text{bur}} \) of greater than 10\% provide excellent enantioselectivity.

Taken together, positing that the diol motif provides robust two-point binding with the catalyst, the substituents on the substrate backbone imparts steric differentiation between the opposing diastereometric transition states, thereby providing a basis for the observed high enantioselectivity. This mechanism of stereoinduction is effective for a wide range of diols with distinct structures and functional groups, resulting in the observed substrate generality.
Figure 4. Mechanistic analysis. (a) Mechanistic analysis. (b) Intermolecular competition studies. (c) Featurization of substrates.

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The computational portion of this work was supported by the Center for High Performance Computing (CHPC) at the University of Utah.

References:
5. During preparation of this manuscript, Jacobsen and co-workers reported a pooled-product method for enantioselectivity analysis, which enabled high-throughput evaluation of the generality of known catalysts in the context of an asymmetric Pictet-Spengler reaction. See ref. 4.


