COMMUNICATION
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Stereoselective synthesis of γ-hydroxynorvaline through combination of organo- and biocatalysis
Stereoselective synthesis of γ-hydroxynorvaline through combination of organo- and biocatalysis†

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An efficient route for the synthesis of all four diastereomers of PMP-protected α-amino-γ-butyrolacton to access γ-hydroxynorvaline was established. The asymmetric key steps comprise an organocatalytic Mannich reaction and an enzymatic ketone reduction. Three reaction steps could be integrated in a one-pot process, using 2-PrOH both as solvent and as reducing agent. The sequential construction of stereogenic centres gave access to each of the four stereoisomers in high yield and with excellent stereocontrol.

Progressive legislative regulations with respect to sustainability, safety and quality improvements have changed the focus of contemporary chemical research to designing more economic processes. In this context, (asymmetric) catalysis has become the method of choice to replace stoichiometric amounts of reagents and auxiliaries, thereby facilitating an efficient and energy saving process. While each of the catalysis sub-fields (metal-, organo- and biocatalysis) has reached an impressive level of sophistication, efficiency can be further increased by combining methods from these different sub-fields. This allows exploiting and combining specific features of each type of catalyst in a highly cooperative fashion. For example, numerous reports have been published based on the combination of transition metals and organocatalysts;2 furthermore, biocatalytic transformations have been reported based on the combination of bio- and organocatalysts;2 hence, access to only one or two out of four stereoisomers is provided. However, as the biological activity of bioactive compounds is related to each stereoisomer,3 a convenient access to all four diastereomers, e.g. for SAR-studies, is desired.

In this communication, a novel combination of organo- and biocatalysis is reported, which provides access to optically pure α-amino-γ-butyrolactones either in a step-wise fashion or in a cascade reaction.

In our study we focused initially on the bioreduction of racemic α-methyl-ketoester rac-1, which is readily available by a proline catalysed Mannich reaction.13,14 A set of homo- and heterologously expressed alcohol dehydrogenases (ADHs)15 were assayed on analytical scale (5.6 mg, 20 mM) employing 2-propanol for NAD(P)+ regeneration. Asymmetric reduction of the carbonyl group of amino-ketoester 1 afforded the corresponding diastereomers of alcohol 2, which cyclised spontaneously to the desired 2-amino-lactones 3 (Scheme 1).

† Electronic supplementary information (ESI) available: Detailed description of each experiment, full characterisation of all compounds and copy of HPLC traces and NMR data. See DOI: 10.1039/c4cc06230b

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of 83% and 86%, respectively. In all cases, the enantiomeric excess (ee) of all diastereomers of lactones 3 was > 99%.

Among the enzymes tested, ADH-LB and evo-1.1.270 gave unexpected results as in both cases mainly one diastereomer (3S,5R)-3 was detected (entries 6 and 7). Since the conversions remained below 50%, we assumed a selective kinetic resolution with respect to the existing stereogenic centre in 5-position to the carbonyl group. In order to gain a better understanding of such a rare case of distant stereorecognition by ADHs, further investigations were performed: racemic and enantiopure Mannich-bases rac-1 ([S]-1) and rac-1 ([R]-1) were incubated with ADH-LB and the reaction progress was monitored over time (Fig. 2). Thereby it was found that rac-1 gave the (3S,5R)-3 diastereomer as sole product with 40% conversion after eight hours, while [R]-1 was not transformed. The pure enantiomer [S]-1 reacted fastest, reaching 70% conversion within the same time.

The experimental observations were additionally supported by in silico docking studies: only substrate [S]-1 was found to fit in the active site enabling a productive binding mode; the substrate occupies therein a pro-R conformation in which the carbonyl group can be activated by two hydrogen bonds through amino acids Tyr152 and Ser142 (Fig. 3). Attempts to dock the opposite enantiomer [R]-1 into the active site failed to provide a reasonable productive binding mode (see ESI†). Hence, the structure of ADH-LB allows a chiral recognition of a remote stereocentre enabling a kinetic resolution. Attempts to couple the diastereoselective reduction of 1 with a racemisation of 1 in a dynamic kinetic resolution failed so far due spontaneous degradation of 1 via elimination of anisidine over time.

The next step was the synthesis of all four stereoisomers of amino-lactone 3; for this purpose, the biocatalysed reduction of ketoester 1 was optimised using alcohol dehydrogenases ADH-A and evo-1.1.200. Investigating 2-ProH as a cosolvent revealed that a substrate concentration of 30 mM (7.95 mg mL⁻¹, 10 vol% 2-ProH) led to completion of the reaction in the case of the [S]-selective ADH-A within 24 h (for details, see ESI†). The [R]-selective enzyme evo-1.1.200 transformed even 50 mM (13.3 mg mL⁻¹, 30 vol% 2-ProH) to completion within 24 h. The optimum pH was determined to be 6.3 in order to keep the partial racemisation of the optically pure Mannich adduct 1 at a maximum (see ESI†). In the subsequent semi-preparative scale transformations [56 mg [S]-1 > 99% conversion was achieved; however, ring closure was incomplete, as mainly the optically pure amino-alcohol 2 rather than the lactone 3 was isolated (ca. 50-70%; see ESI†). Assuming that the corresponding methyl ester of 1 would undergo faster lactonisation than the ethyl analogue, a transesterification step (HCl–MeOH) was integrated into the reaction sequence. The optimised conditions finally allowed conversion of the pure amino-ester ([R]-1) into the corresponding Mannich reaction of acetone and imine 4 into the corresponding lactones 3. The syn- and anti-diastereomers were isolated in excellent yields (78-82%) and with remarkable stereocontrol (ee > 98:2 and ee > 99% in all cases; see Scheme 2).

As an alternative to the conventional step-by-step synthesis (Scheme 2) two sequential approaches were established. In a first approach, aldimine 4 was formed in 2-ProH, which also served as a solvent for the subsequent Mannich reaction and as

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**Table 1** Selected results of the ADH catalysed reduction of ketoester rac-1 yielding lactone 3 (analytical scale, double determination)†

<table>
<thead>
<tr>
<th>Entry</th>
<th>Enzyme</th>
<th>Conv. [%]</th>
<th>d.r.</th>
<th>ee&lt;sub&gt;syn&lt;/sub&gt; [%]</th>
<th>ee&lt;sub&gt;anti&lt;/sub&gt; [%]</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>ADH-A</td>
<td>99</td>
<td>52/48</td>
<td>&gt;99 (3R,5S)</td>
<td>&gt;99 (3S,5S)</td>
</tr>
<tr>
<td>2</td>
<td>evo-1.1.030</td>
<td>99</td>
<td>51/49</td>
<td>&gt;99 (3R,5S)</td>
<td>&gt;99 (3S,5S)</td>
</tr>
<tr>
<td>3</td>
<td>ADH-T</td>
<td>83</td>
<td>12/88</td>
<td>&gt;99 (3R,5S)</td>
<td>&gt;99 (3S,5S)</td>
</tr>
<tr>
<td>4</td>
<td>ADH-LS</td>
<td>86</td>
<td>71/29</td>
<td>&gt;99 (3S,5R)</td>
<td>&gt;99 (3R,5R)</td>
</tr>
<tr>
<td>5</td>
<td>evo-1.1.200</td>
<td>99</td>
<td>52/48</td>
<td>&gt;99 (3S,5R)</td>
<td>&gt;99 (3R,5R)</td>
</tr>
<tr>
<td>6</td>
<td>ADH-LB</td>
<td>49</td>
<td>99/1</td>
<td>&gt;99 (3S,5R)</td>
<td>—</td>
</tr>
<tr>
<td>7</td>
<td>evo-1.1.270</td>
<td>49</td>
<td>99/1</td>
<td>&gt;99 (3S,5R)</td>
<td>—</td>
</tr>
</tbody>
</table>

† Reaction conditions: ketoester rac-1 (5.6 mg, 20 mM), KPi buffer (100 mM, pH 6.8), NADP⁺ (1.0 mM), 10 vol% 2-ProH, ADH (for units see ESI), 30 °C, 700 rpm on a thermostaker (horizontal position), 24 h. Determined via GC on an achiral stationary phase. Determined via HPLC on a chiral stationary phase.
the reducing agent for the enzyme-mediated reduction (after dilution with buffer solution). All three steps were accomplished in one pot, after which the transesterification-lactonisation cascade was performed in a separate vessel. The whole sequence afforded the syn-configured lactone (3S,5R)-3 in a remarkable 47% yield with a d.r. of 86:14 in favour of the syn-diastereomer (Scheme 3a: 5 steps in total, 2 pots, 90% average yield per step). The diminished d.r. in comparison to the previous results can be rationalised by the usage of 2-ProH for the proline-catalysed reaction, which under these conditions afforded (S)-1 with 72% ee.

In a second approach, the Mannich reaction was carried out with the preformed aldimine 4 in acetone as solvent, and amino-ketoester (R)-1 was obtained at prolonged reaction time (16 h) with >99% conversion and in optically pure form (ee > 99%). After evaporation of the solvent, the crude product was subjected to reduction by ADH-A in buffer–2-ProH. The final transesterification-lactonisation cascade furnished the diastereomically pure amino lactone (3S,5S)-3 in 51% yield with an excellent d.r. of 99:1 (Scheme 3b: 4 steps, 2 pots, 85% average yield per step).

Final PMP-deprotection of the diastereomerically pure α-amino-lactone 3 was achieved employing TCCA (trichloroisocyanuric acid) in MeOH,16b providing the hydrochloride salt of 6. Unexpectedly, other oxidative deprotection methods [Pd(OAc)2, CAN, HIO4, laccase, etc.]18 were unsuccessful. Intensive optimisation of the purification of 6 with respect to the solvent, acid, reaction time and work-up procedure finally provided access to γ-hydroxynorvaline: the natural product was isolated diastereomerically pure after a base promoted ring opening (58% yield over two steps) as demonstrated for the (2R,4S)-diastereomer (Scheme 4).

In conclusion, a straightforward synthesis of all four diastereomers of PMP-protected α-amino-γ-butyrolactone 3 through combination of organo- and biocatalysis. Reagents and conditions: (a) o-1-proline (0.25 eq.), DMSO, 3 h, rt; (b) (R)-/S)-1 (30 mM), ADH-A, NAD+, buffer (100 mM, pH 6.3), 2-ProH (10 vol%), 30 °C, 24 h; (c) (R)-/S)-1 (50 mM), evol-1.2.200, NAD+, buffer (100 mM, pH 6.3), 2-ProH (30 vol%), 30 °C, 24 h; (d) HCl-2-PrOH (2.0 eq.), MeOH, 16 h, rt. PMP = para-methoxyphenyl.

including the organo- as well as the biocatalysed transformations were performed in one pot, integrating 2-propanol first as solvent and then as reducing agent.

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Notes and references
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15 ADHs employed during the studies: ADH-A originates from Rhodococcus ruber; see: (a) K. Edegger, C. C. Gruber, T. M. Poesl, S. R. Wallner, I. Lavandera, K. Faber, F. Niehaus, J. Eck, R. Oehrlein, A. Hafrer and W. Kroutil, Chem. Commun., 2006, 2402; (b) Eozymes were obtained from Evocatal (origin not stated); (c) ADH-T was obtained from Codecis Inc. (origin not stated); (d) ADH-LS originates from Leifsonia sp.; see: K. Inoue, Y. Makino and N. Itoh, Appl. Environ. Microbiol., 2005, 3633; (e) ADH-LB originates from Lactobacillus brevis; see: W. Hummel and B. Riebel, Biotecnol. Lett., 2003, 25, 51.


17 The ring-closure was found to be promoted by the GC injector pretended full conversion to the lactone 3. In an additional control experiment the purified alcohol 2 was subjected to GC and GC-MS analysis, showing only the signal for the lactone 3.