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Complete List of Authors:	Richter, Nina; Austrian Centre of Industrial Biotechnology (ACIB), Farnberger, Judith; Austrian Centre of Industrial Biotechnology (ACIB), Pressnitz, Desiree; University of Graz, Lechner, Horst; University of Graz, Department of Chemistry Zepeck, Ferdinand; Sandoz, Kroutil, Wolfgang; Karl-Franzens University Graz, Department of Chemistry, Organic and Bioorganic Chemistry

ARTICLE

A system for ω -transaminase mediated (*R*)-amination using L-alanine as amine donor

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N. Richter^a, J. E. Farnberger^b, D. Pressnitz^b, H. Lechner^b, F. Zepeck^c and W. Kroutil^{b*}Received 00th January 2012,
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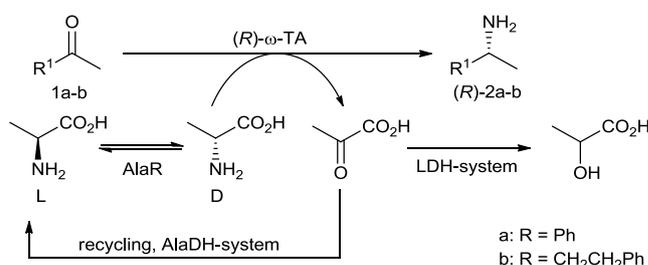
Chiral amines are important building blocks for fine chemicals and pharmaceuticals. Consequently, various biocatalytic routes in particular using ω -transaminases (ω -TAs) have been developed recently. Although catalysts for the synthesis of both enantiomers are available, the application of alanine dependent (*R*)-selective ω -TAs is less favourable due to the requirement of the more expensive D-alanine as amine donor. Here we describe an efficient method for (*R*)-amination using ω -TAs in combination with an alanine racemase (AlaR). In this case, the readily available L-alanine can be used as amine donor leading to improved atom efficiency and significantly reduced costs.

Introduction

ω -Transaminases (ω -TAs) have been proven to be powerful catalysts frequently used in the synthesis of chiral amines.¹⁻⁸ As an alternative to classical chemical strategies,⁹⁻¹² they offer a green and sustainable synthetic route and can provide access to both enantiomers of the amine. This is enabled by the recent identification of numerous ω -TAs with (*S*)- as well as (*R*)-stereoselectivity.¹³⁻¹⁹ However, due to the unfavoured reaction equilibrium of the reductive amination of ketones employing alanine as amine donor as well as co-product inhibition, the reaction requires *in-situ* techniques to remove the co-product pyruvate to set-up efficient processes. As a consequence, many studies addressed these limitations of ω -TAs over the last decade, and a variety of methods have been developed to efficiently shift the equilibrium of the desired reductive amination reaction.^{2,3,6} This progress enabled the development of various interesting applications employing ω -TAs²⁰⁻³⁰ and their exploitation on an industrial scale.¹⁵ Nevertheless, the selection of the amine donor is a crucial parameter: while a donor such as 2-propylamine seems to be a good choice with respect to economic efficiency, its use is restricted to certain ω -TAs.^{3,15,17,18,29,31-32} Therefore, a natural donor like alanine might be preferred; However, while L-alanine is readily available to be employed with (*S*)-selective enzymes, (*R*)-selective enzymes require the far more expensive D-alanine. This limitation makes their application less efficient, compared to their (*S*)-selective counterparts, and alternative strategies are required, in particular when amine donors such as 2-propylamine are not an option.

Results and discussion

To overcome this limitation for applications with (*R*)-selective ω -TAs we report here the combination of an alanine racemase (AlaR) with the amination system. Thus, by using L-alanine as starting material the required more expensive D-alanine is provided *in-situ* by racemisation (Scheme 1). This racemisation concept has been reported for the synthesis of optically pure α -amino acids using α -transaminases; however, the strategy has not been reported for the asymmetric amination of ketones employing ω -TAs.³³ In particular, the combination of the AlaR with the frequently used alanine dehydrogenase (AlaDH) to remove the co-product of the amination, pyruvate, by reductive amination to L-alanine (AlaDH-system)³⁴ enables the recycling of the amine donor also in the case of (*R*)-selective ω -TAs.



Scheme 1 Amination employing (*R*)-selective ω -TAs and L-alanine as donor. D-alanine is provided by an alanine racemase (AlaR) by racemisation of L-alanine. The co-product pyruvate is recycled/removed using either an alanine dehydrogenase (AlaDH) or lactate dehydrogenase (LDH).

Cloning and recombinant expression of the SeAlaR and PpAlaR

For the racemisation of L-alanine the PLP-dependent AlaR from *Streptomyces coelicolor* (SeAlaR)³⁵ was selected as well

as an AlaR from *Pseudomonas putida* (PpAlaR). Both enzymes were cloned in suitable expression vectors and heterologously expressed in *E. coli*.

While the PpAlaR mainly accumulated as insoluble inclusion bodies, good expression was achieved for SeAlaR. However, initial activity studies showed that both enzymes were active towards the racemisation of L-alanine. Based on the better heterologous expression, lyophilised cell free extracts of SeAlaR were used for all further experiments.

Characterisation of the SeAlaR

The SeAlaR was characterised to identify optimal conditions for its application in combination with the other required enzymes in the desired amination reaction (Scheme 1).

First the racemisation of 250 mM L-alanine was studied over a period of 30 min, using different amounts of enzyme (mg/mL). The results indicated that an efficient racemisation within that time was achieved using 5 mg as well as 2.5 mg of the lyophilised cell free extract (Fig. 1a). In addition to the enzyme loading the influence of DMSO, a co-solvent commonly used with ω -TAs, was studied. For this purpose, activities were determined in the presence of 10 vol% DMSO. As depicted in Fig. 1a the racemisation rate in the absence of DMSO was slightly higher compared to the reaction in the presence of DMSO. Nevertheless, the results indicate that DMSO can be employed in a reaction with SeAlaR, if DMSO is required to be added to improve the solubility of lipophilic substrates. As a further crucial parameter the stability of the SeAlaR under ω -TA reaction conditions was investigated. The enzyme was incubated under process conditions (30 °C, in the presence of pyruvate, acetophenone, 1-phenylethylamine and PLP) and the activity in the racemisation of L-alanine after different incubation times was monitored (Fig. 1b/c). The study showed that AlaR is a rather stable catalyst and even after an incubation time of 24 h the racemisation still proceeds equally well as at the start of the reaction (Fig 1b). The same is true for the racemisation in the presence of DMSO (10 vol%). Here the racemisation still proceeded well although slightly slower compared to the reaction without co-solvent (Fig. 1c).

Combination of racemisation and transamination reaction

For the amination employing (*R*)-selective ω -TAs and D-alanine as amine donor two systems were used to remove the co-product pyruvate in order to shift the equilibrium: an AlaDH to transform pyruvate to L-alanine (AlaDH-system, Scheme 1) or a LDH to reduce pyruvate to lactate (LDH-system, Scheme 1).²⁻³ A comparative study was performed using either only D-alanine or L-alanine in combination with the SeAlaR.

SeAlaR and the AlaDH-system

The amination of acetophenone (**1a**) employing the ω -TA from *Aspergillus terreus* (AT ω -TA)^{13,14,17-19} and five equivalents of D-alanine (Fig 2a, black bars) was compared to the reaction with L-alanine and SeAlaR. For the biotransformations using L-alanine 10 equivalents were used to provide five equivalents of the desired D-enantiomer after racemisation by the SeAlaR (Fig. 2a, grey bar). In addition, the same reaction was performed using *rac*-alanine (10 eq., white bar) and L-alanine (5 eq., light grey bar) without addition of SeAlaR. The highest

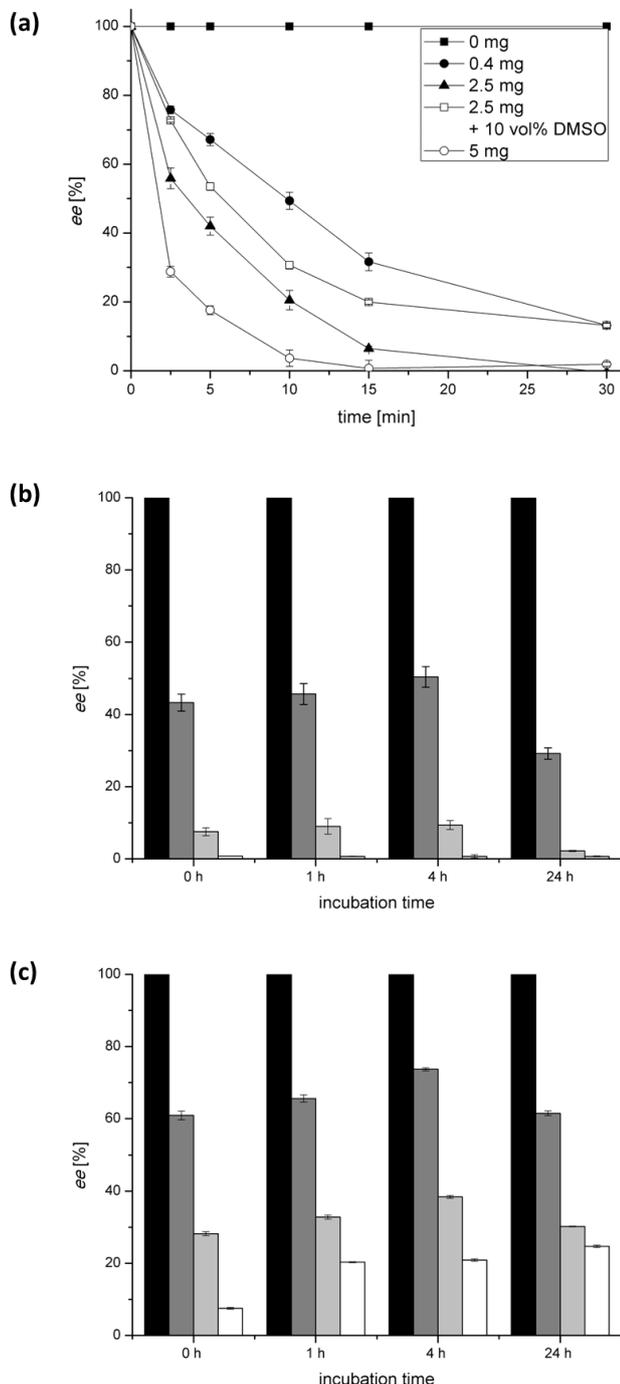
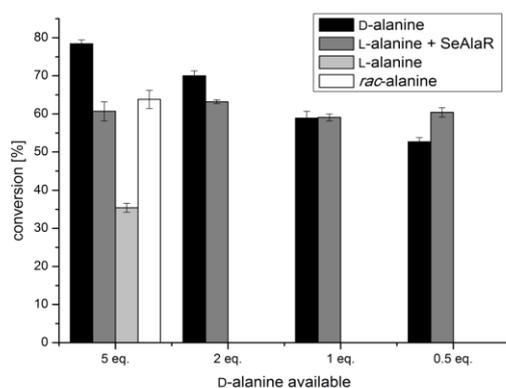


Fig. 1 Activity and stability of the SeAlaR. **(a)** Racemisation profile of L-alanine catalysed by the SeAlaR. The racemisation of 250 mM L-alanine was monitored over a period of 30 min using different amounts of AlaR (mg/mL) in the presence and absence of DMSO as co-solvent. **(b/c)** Racemisation profile of SeAlaR after an incubation in the absence **(b)** and in the presence of DMSO **(c)** under process conditions (30 °C, in the presence of pyruvate, acetophenone, 1-phenylethylamine and PLP). Samples were taken after 0 min (black bars); 5 min (dark grey bars); 15 min (light grey bars) and 30 min (white bars).

conversion was achieved using five equivalents of D-alanine in the absence of SeAlaR (78%). The addition of L-alanine to produce five equivalents of D-alanine *in-situ* led to a conversion of 61%. Since *rac*-alanine in the control reaction led to a conversion in a similar range (64%) it seemed that inhibition occurs to a certain extent due to the presence of the L-enantiomer. However, when providing only one equivalent of D-alanine both systems led to the same conversion and at 0.5 equivalents even a higher conversion was achieved starting with L-alanine and SeAlaR. This clearly demonstrated that by employing the SeAlaR-catalysed racemisation of L-alanine the efficiency of the desired amination reaction was improved.

(a) AlaDH-system



(b) LDH-system

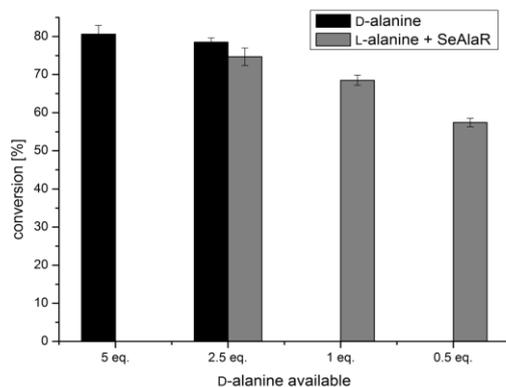


Fig. 2 Conversions of the ω -TA-catalysed amination of **1a** using AT ω -TA with and without SeAlaR. (a) conversions for the AlaDH-system after 24 h in the presence (grey bars) and in the absence (black bars) of SeAlaR at different equivalents of D-alanine. (b) Conversions for the LDH-system in the presence (grey bars) and in the absence (black bars) of SeAlaR at varied amounts of alanine. With the exception of the control reaction where L-alanine was used (a, light grey bar), results are always correlated to the equivalents of D-alanine available in the reaction mixture.

Interestingly, the control reaction with L-alanine (5 eq.) in the absence of SeAlaR also led to a moderate conversion. Additional experiments using purified ω -TA for the transformation could prove that this conversion is due to the presence of an AlaR in the *E. coli* host.^{36,37} Thus, the ω -TA employed was highly specific for D-alanine not accepting L-alanine and the *E. coli* AlaR slowly racemised L-alanine

consequently leading to the observed conversion (for results see SI section 2). However, the studies also showed that the racemisation by the *E. coli* AlaR was insufficient, in particular with respect to reaction time and conversion. This becomes obvious when comparing the reaction rates of the transformations with D-alanine and L-alanine within the first ten hours: while the reaction was almost completed using D-alanine (72%) as amine donor, a conversion of only 16% was reached in the case of L-alanine (SI Fig. S3).

SeAlaR and the LDH-system

Similar to the AlaDH-system, the LDH-system was studied in the presence and absence of SeAlaR. Again the influence of different alanine concentrations was investigated. A maximal conversion of 81% for the amination of **1a** was reached using five equivalents of D-alanine without racemisation by the SeAlaR. The best conversion (75%) was obtained for the racemisation system using 5 equivalents of L-alanine leading to 2.5 equivalent of d-alanine after racemisation by the SeAlaR (Fig. 2b). Actually, when comparing both systems (AlaDH and LDH) comparable conversions (78% and 81%) were obtained using five equivalents of D-alanine in the absence of the SeAlaR (Fig. 2a and b, black bar). However, in the presence of SeAlaR-catalysed racemisation slightly different results were obtained. A combination of SeAlaR with the LDH-system led to a conversion of 75%, whereas the AlaDH/SeAlaR-system led to a conversion of 63%. This variation might be caused by the fact that D-alanine is consumed when using the LDH-system and not recycled as for the AlaDH-system, hence the concentration of both alanine enantiomers is reduced over time. The decreasing amount of L-alanine present might have a positive effect on the reaction assuming an inhibition by this enantiomer.

Since a reduction of alanine equivalents had no significant effect in the case of the AlaDH-system (Fig. 2a) the influence of lower alanine concentrations was investigated also with the LDH-system. However, in this case the conversion decreased with lower amounts of alanine in the presence of SeAlaR (Fig. 2b). This decrease might be due to the constantly decreasing alanine concentration in the LDH system, while the concentration in the AlaDH-system stayed constant.

As a further example, the amination of 4-phenyl-2-butanone (**1b**) was studied initially using the AT ω -TA. Fig. 3 shows that the amination of **1b** led to conversions close to 100% using both the LDH- and the AlaDH-system. Even when employing only one equivalent of D-alanine a conversion of 95% was obtained. It is important to note, that the additional racemisation catalyst (SeAlaR) did not have any influence on the optical purity of the product which was still very high (*ee* >99%). This was also true for the amination of **1a** where exclusively (*R*)-1-phenylethylamine (**2a**) was detected (*ee* >99%).

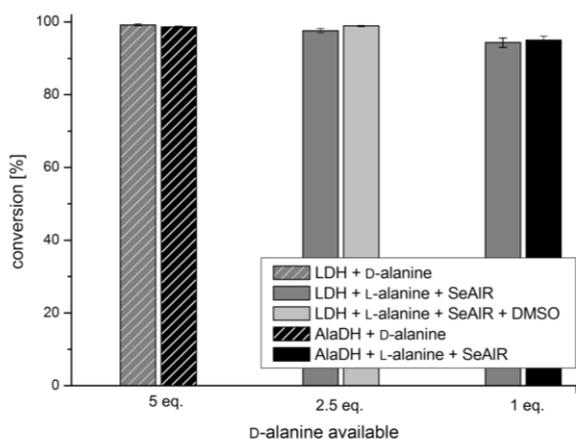


Fig. 3 Amination of **1b** applying the best conditions (see text) of both the AlaDH- and LDH-system with and without the SeAlaR using the AT ω -TA. Conversions after 24 h for the AlaDH-system (black bars) and LDH-system (grey bars). Control reactions using D-alanine as amine donor are indicated by the hatched bars

Finally, the system was tested with a second ω -TA originating from *Arthrobacter* sp. (ArR)^{17,38} and 4-phenyl-2-butanone **1b** as substrate. Transaminations with five equivalents of D-alanine were compared to reactions with SeAlaR using the identified optimal conditions for both pyruvate removal systems [AlaDH/FDH (1 eq.) and LDH/GDH (2.5 eq.)]. Again, it was clearly shown that L-alanine can be used successfully as amine donor in the presence of the SeAlaR. Moreover, comparable conversions were obtained with respect to the use of D-alanine without the SeAIR (Fig. 4).

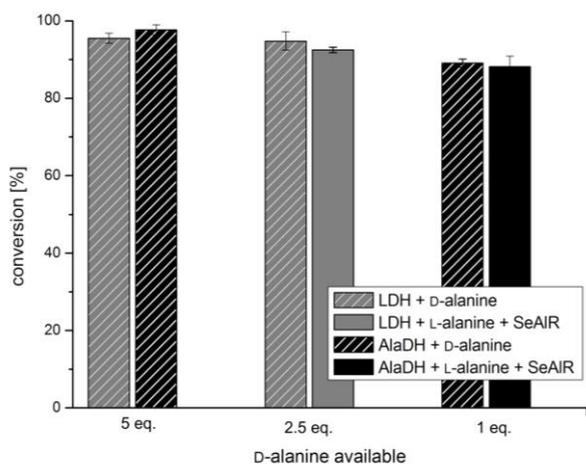


Fig. 4 Results of the amination of **1b** employing the ArR ω -TA comparing both systems with and without addition of SeAlaR. Conversions are given for the LDH- (grey bars) and AlaDH-system (black bars) after 24 h. Control reactions using D-alanine as amine donor are indicated by the hatched bars

Comparison of efficiency

In summary, it was clearly demonstrated that L-alanine can be applied for (*R*)-selective ω -TAs when used in combination with an AlaR. The most prominent advantage of this method is that the cost of a process using (*R*)-selective ω -TAs can be significantly reduced by using much cheaper L-alanine as amine

source as well as being able to reduce the equivalents of amine donor employed. Table 1 outlines the relative costs for the production of (*R*)-4-phenyl-2-butylamine [(*R*)-**2b**] for the systems applied. By just employing 10 equivalents of L-alanine instead of 5 equivalents of D-alanine the costs of the reaction can be reduced 6.8-fold. Moreover, since an efficient production was also possible using less equivalents of L-alanine the price could be even reduced by 97%, with respect to the amine source, compared to the previously applied reaction conditions.

Table 1: Comparing the systems in the absence and presence of the SeAlaR with respect to the relative costs of the amine donor as well as E-factor for the production of (*R*)-**2b**.

process	equivalents D-ala ^a	alanine	relative costs [%] ^b	E-factor
AlaDH/FDH system ^c	5	D	100	13.75
LDH/FDH-system	5	D	100	13.78
LDH/GDH/SeAlaR system	2.5	L + AlaR	5.2	14.57
AlaDH/FDH/SeAlaR system	1	L + AlaR	2.1	13.21

^a equivalents of D-alanine, when starting from L-alanine + AlaR twice the equivalents of L-alanine were used to provide comparable equivalents of D-alanine

^b prices for the calculation were taken from Sigma Aldrich (D-alanine, # A7377) and Carl Roth (L-alanine, # 3076) (26.10.2014)

In a recently published multi-step synthesis of a dual orexin receptor antagonist a ω -TA catalysed amination was performed on preparative scale (22.3 mol) using D-alanine as amine source.³⁹ The desired intermediate (piperidone) was produced with 74% yield and excellent stereoselectivity. However, due to its solubility limit only 3.6 equivalents of D-alanine were used, prolonging the reaction time to 31 h. Still the 7.2 kg of D-alanine used on this scale are rather expensive. In theory, by applying the described concept the costs for the amine donor could be reduced by 85.5% when using equal equivalents of D-alanine (7.2 eq. of L-alanine). Moreover, when using the optimised AlaDH-system and consequently reducing the equivalents a reduction of even 96% of the costs is possible, demonstrating the impact for amination reactions on industrial scale.

In addition to the significant reduction in costs, also the E-factor was calculated to address the environmental impact of the different systems (Table 1, for calculations see SI). The process without the additional racemisation step using the AlaDH-system can be considered as a rather green process (E-factor 13.75). Nevertheless, by introducing the racemisation step a further reduction of the E-factor by 5.5% was possible when employing the AlaDH/SeAlaR-system (E-factor 12.99). The additional fermentation, which is required to produce the SeAlaR, has to be taken into account, too. However, it has to be considered that the possibility to use L-alanine instead of D-alanine is also a significant improvement towards a greener process which is unfortunately not considered in the calculation of the E-factor, since using an amine source from the readily available chiral pool of nature makes the whole transformation more environmental friendly using an amine source from the readily available chiral pool of nature makes the whole

transformation more environmentally friendly. In particular, since a bio-based production of L-alanine using organisms such as *Corynebacterium glutamicum* as well as *E. coli* at the expense of glucose is possible.^{40,41}

In addition, the introduced racemisation step improves the efficiency by enabling the recycling of the amine donor when using the AlaDH-system. Based on the L-selectivity of the employed AlaDH, L-alanine is produced which can then be racemised to provide the amine donor for the (R)-selective ω -TAs. Theoretically, only one equivalent of L-alanine should be sufficient to aminate the starting ketone with complete conversion and indeed, studies starting with 1 eq. of L-alanine (providing 0.5 eq. of D-alanine after racemisation) led to similar results compared to those observed with 10 times more amine donor (Fig. 2). The benefit of recycling the amine donor in the case of the AlaDH-system becomes even more obvious when comparing the atom efficiency of the AlaDH process, whereby atom efficiency is increased from 50% without to 71% with addition of AlaR (for calculations see SI).

Conclusion

The application of (R)-selective ω -TAs requiring D-alanine as amine donor is limited by the 13.8-times more expensive D-enantiomer compared to L-alanine. The here introduced racemisation step catalysed by an AlaR from *Streptomyces coelicolor* (SeAlaR) led to an economically improved and greener (R)-amination reaction system using L-alanine as amine donor. The experiments showed that comparable conversions were achieved using either D-alanine as amine donor or L-alanine with the SeAlaR. In addition, it could be shown that the equivalents of amine donor can be significantly reduced using the described system. In particular, when the AlaR is used in combination with an AlaDH for the removal of the co-product pyruvate, a significantly improved process was established due to the recycling of alanine. As a consequence, the costs for the amine donor alanine can be reduced by up to 97 % while simultaneously decreasing the E-factor of the process as well as increasing the atom efficiency.

Experimental

Chemicals

Commercial grade chemicals were purchased from different suppliers and used without further purification. DNA-modifying enzymes were obtained from Thermo Scientific or New England Biolabs. L-lactate dehydrogenase (LDH, evo-1.1.150), formate dehydrogenase (FDH, evo-1.1.230) and glucose dehydrogenase (GDH, evo-1.1.060) (all evocatal, Germany) are commercially available. The L-alanine dehydrogenase (AlaDH) was used as purified enzyme which was prepared as described previously.²³ Both used ω -transaminases were employed as lyophilised whole cell preparation prepared as previously reported.¹⁷

Cloning and recombinant protein expression of SeAlaR and PpAlaR

The previously reported AlaR from *Streptomyces coelicolor* (SeAlaR, pEG238)³⁵ and the novel AlaR from *Pseudomonas putida* (PpAlaR, pEG236) were designed as codon-optimised genes (both sequences can be found in SI) (GeneArt, Germany). Both genes were subsequently cloned into the expression vector pET21a (Invitrogen, USA) using the restriction sites *NdeI/NotI* to construct a variant containing a C-terminal His-Tag.

The AlaRs were expressed in *E. coli* BL21(DE3)pLysS in lysogeny broth medium containing ampicillin (100 mg mL⁻¹). Cultures were grown at 37 °C until an OD₆₀₀ of 0.5-0.7 was reached. Enzyme expression was induced by addition of isopropyl- β -D-thiogalactopyranoside (0.5 mM), and the temperature was reduced to 25 °C. After 16-20 h cells were harvested by centrifugation and stored at -20 °C. To prepare cell free crude extract, cells (10% v/w) were suspended in sodium phosphate buffer (10 mM, pH 8), and disrupted by ultra sonication (2 \times 1 min, 0.1 sec pulse on and 0.4 sec pulse off, 40% output). The crude extract was cleared by centrifugation (20 min, 14,000 rpm), freeze dried and stored at -20 °C.

Determination of racemisation rate

To test the activity of both AlaRs the racemisation of L-alanine was followed over time. Different amounts of AlaR (0.4-5 mg mL⁻¹) were incubated with L-alanine (250 mM) in sodium phosphate buffer (50 mM, pH 8) containing pyridoxal-5'-phosphate (PLP; 1 mM) at 30°C and 500 rpm. For the determination of the co-solvent tolerance of the AlaR, DMSO (10 vol%) was added to the reaction. After 2.5; 5; 10 and 30 min samples were withdrawn (10 μ L), quenched with MeOH (10 μ L) and subsequently diluted 500-fold with HPLC-MS eluent (MeOH:H₂O = 65:35 with 0.1% of formic acid). Denatured protein was removed by centrifugation (3 min, 13,000 rpm) and the supernatant was analysed by HPLC-MS (1260 Infinity equipped with a quadrupole 6120, both Agilent, USA) using a chiral Chirobiotic-T column (150 \times 4.5 mm, Astec, Sigma-Aldrich, Switzerland) and the following conditions: MeOH / H₂O / HCO₂H (65:35:0.1) at a flow rate of 1 mL min⁻¹; elution times; L- (2.3min), D-alanine (2.7 min).

Stability of the SeAlaR

For the determination of the SeAlaR stability, the enzyme was incubated under process conditions (30 °C, in the presence of pyruvate, acetophenone, 1-phenylethylamine and PLP) and the activity was determined after different incubation times. For this study the SeAlaR (2 mg mL⁻¹) was incubated in sodium phosphate buffer (50 mM, pH 8) containing sodium pyruvate (12.5 mM), acetophenone (12.5 mM), 1-phenylethylamine (12.5 mM) and PLP (1 mM) at 30 °C and 600 rpm. After 0 h, 1 h, 4h and 24 h L-alanine (250 mM) was added to start the reaction and samples were taken over 30 min and analysed as described before.

ω -TA-catalysed amination of acetophenone and 4-phenyl-2-butanone

AlaDH-system Transamination reactions were performed on 1 mL scale as follows: lyophilised *E. coli* cells containing the ω -TA (20 mg) were rehydrated in sodium phosphate buffer (200 μ L, 50 mM, pH 8) containing PLP (5 mM, 1 mM final concentration in the reaction) at 30°C and 120 rpm for 15 min. To provide racemic alanine for the reaction the crude AlaR (2 mg) was incubated in sodium phosphate buffer (200-400 μ L, 50 mM, pH 8) containing L-alanine (1-10 eq., 25-250 mM final concentration) at 30°C and 120 rpm for 15 min. For control reactions either D- or L-alanine was incubated without adding AlaR. After this initial incubation both mixtures were combined and the volume was adjusted to 1 mL by adding sodium phosphate buffer (50 mM, pH 8) containing NAD⁺ (1 mM final concentration), ammonium formate (150 mM final concentration), FDH (20 U) and AlaDH (10 U). The reaction was started by the addition of substrate (3.7 mg, 25 μ mol, 25 mM final conc.) and incubated at 30 °C and 800 rpm. Samples were stopped at different time points by addition of an aqueous NaHCO₃ solution (10 vol%) and extracted with ethyl acetate (2 \times 400 mL). The organic phase was dried (Na₂SO₄) and analysed by gas chromatography (Agilent 7890 A system with a FID detector) equipped with an Agilent DB-1701 column (30 m, 0.25 mm, 0.25 μ m) using the following temperature programmes: for acetophenone: 60 °C, hold for 5 min, 10 °C min⁻¹ to 160 °C, 60 °C min⁻¹ to 280 °C, hold for 2 min, retention time: 1-phenylethylamine 10.2 min, acetophenone 11.2 min; and for 4-phenyl-2-butanone: 120 °C, 10 °C min⁻¹ to 180 °C, 60 °C min⁻¹ to 280 °C, hold for 2 min, retention times: 1-methyl-3-phenylpropylamine 3.3 min, 4-phenyl-2-butanone 3.7 min.

For the determination of enantiomeric excess samples were derivatised to the corresponding acetamides by adding pyridine (2 μ L) and acetic anhydride (6 μ L) to the extracted organic phase. Samples were incubated at 40 °C and 800 rpm for 2 h and quenched by the addition of sat. NaHCO₃ (25 μ L). Then the organic phase was analysed *via* gas chromatography (Agilent 7890 A system with a FID detector) equipped with a chiral Varian Dex CB column (25 m, 0.32 mm, 0.25 μ m). The following method was used: 120 °C, 5 °C min⁻¹ to 180, hold 2 min, retention times: 1-methyl-3-phenylpropylamine: (*S*) 9.7 min, (*R*) 9.9 min, 1-phenylethylamine: (*S*) 6.4 min, (*R*) 6.7 min.

LDH-system Transamination reactions were performed on 1 mL scale as follows: lyophilised *E. coli* cells containing the ω -TA (20 mg) and the AlaR (2 mg) were preincubated as described above. After this initial incubation both mixtures were combined and the volume was adjusted to 1 mL by adding sodium phosphate buffer (50 mM, pH 8) containing NAD⁺ (1 mM final concentration), glucose (150 mM final concentration), LDH (140 U) and GDH (35 U). The reaction was started by the addition of substrate (3.7 mg, 25 μ mol, 25 mM final concentration) and incubated at 30°C and 800 rpm. Samples were withdrawn at different time points, and treated and analysed as described for the AlaDH-system.

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^a ACIB GmbH, c/o Heinrichstraße 28, 8010 Graz, Austria
^b Department of Chemistry, Organic and Bioorganic Chemistry, NAWI Graz, University Graz, Heinrichstraße 28, 8010 Graz, Austria. E-mail: wolfgang.Kroutil@uni-graz.at; Tel.: +43-(0)-316-380-5350.
^c Sandoz GmbH, Biocatalysis Lab, Biochemiestraße 10, 6250 Kundl/Tirol, Austria

† Footnotes should appear here. These might include comments relevant to but not central to the matter under discussion, limited experimental and spectral data, and crystallographic data.

Electronic Supplementary Information (ESI) available: [details of any supplementary information available should be included here]. See DOI: 10.1039/b000000x/

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Introducing an alanine racemase allowed to use L-alanine as initial amine source for the biocatalytic amination leading to (*R*)-amines.

