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ARTICLE

Expanding the substrate scope of phenylalanine ammonia-lyase from *Petroselinum crispum* towards styrylalanines

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This study focused on the expansion of the substrate scope of phenylalanine ammonia-lyase from *Petroselinum crispum* (PcPAL) towards the L-enantiomers of racemic styrylalanines *rac*-**1a-d** – which are less studied and synthetically challenging unnatural amino acids – by reshaping the aromatic binding pocket of the active site of PcPAL by point mutations. Ammonia elimination from L-styrylalanine (L-**1a**) catalyzed by non-mutated PcPAL (wt-PcPAL) happened with 777-fold lower k_{cat}/K_M value as the deamination of the natural substrate, L-Phe. Computer modeling of the reactions catalyzed by wt-PcPAL indicated an unproductive and two major catalytically active conformations and detrimental interactions between the aromatic moiety of L-styrylalanine, L-**1a**, and the phenyl ring of F137 residue in the aromatic binding region of the active site. Replacing the residue F137 by smaller hydrophobic residues, resulted a small mutant library (F137X-PcPAL, X being V, A, and G), from which F137V-PcPAL could transform L-styrylalanine with comparable activity that of the wt-PcPAL with L-Phe. Furthermore, F137V-PcPAL showed superior catalytic efficiency in the ammonia elimination reaction of several racemic styrylalanine derivatives (*rac*-**1a-d**) providing access to D-**1a-d** by kinetic resolution, even though the D-enantiomers proved to be reversible inhibitors. The enhanced catalytic efficiency of F137V-PcPAL towards racemic styrylalanines *rac*-**1a-d** could be rationalized by molecular modeling indicating the more relaxed enzyme-substrate complexes and the promotion of conformations with higher catalytic activities as the main reasons. Unfortunately, ammonia addition onto the corresponding styrylacrylates **2a-d** failed both with wt-PcPAL and F137V-PcPAL. The low equilibrium constant of the ammonia addition, the poor ligand binding affinities of **2a-d**, and non-productive binding states of the unsaturated ligands **2a-d** within the active sites of either wt-PcPAL or F137V-PcPAL – as indicated by molecular modeling – might be responsible for inactivity of the PcPAL variants in the reverse reaction. Modeling predicted the F137V mutation beneficial for the KR of 4-fluoro-, 4-cyano- and 4-bromostyrylalanines, but non-effective for the KR process of 4-trifluoromethylstyrylalanine.

Introduction

The synthesis of unnatural aromatic amino acids in enantiopure form is still a challenge of synthetic chemistry, highlighted by the significant interest towards these building blocks in the development of therapeutic peptides and proteins.¹⁻³

Since phenylalanine – as a pharmacophore element – has an important role in biologically active peptides and proteins, development of synthetic methods leading to sterically more demanding analogues of this amino acid has gained increased

attention.⁴⁻⁷ An attractive enzymatic route for the synthesis of non-natural phenylalanine analogues is provided by aromatic ammonia-lyases and aminomutases which show high structural and sequence similarities, and have a common electrophilic prosthetic group, the autocatalytically forming 3,5-dihydro-5-methylidene-4H-imidazol-4-one (MIO).⁸⁻¹²

Recent advances in improvement of the functional properties of the MIO-enzymes initiated their application as biocatalysts for the synthesis of non-natural amino acids, well exemplified by the phenylalanine ammonia-lyase (PAL)-based process, developed by DSM (Netherlands), for the multiton scale production of (S)-2,3-dihydro-1H-indole-2-carboxylic acid.¹³

Due to its broad substrate tolerance, phenylalanine ammonia-lyase from *Petroselinum crispum* (PcPAL) was one of the most frequently used aromatic amino acid ammonia-lyase applied as biocatalyst.^{14,15,16} Immobilization of isolated PcPAL led to increased operational stability and enabled the employment of PcPAL for the synthesis of non-natural substrates in continuous-flow packed-bed reactors,¹⁷ in magnetic nanoparticle-filled microfluidic systems,^{18,19} and provided new

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ARTICLE

Organic & Biomolecular Chemistry

perspectives in the development of stable and efficient catalysts for industrial applications.

Numerous evidences suggest that interactions between the amino group of the substrate L-Phe and the electrophilic MIO of PAL facilitate the ammonia-lyase reaction.²⁰ The crystal structures of phenylalanine 2,3-aminomutases from *Pantoea agglomerans* (PaPAM) complexed with L-phenylalanine²¹ and from *Taxus chinensis* (TchPAM) complexed with L- α,α -difluoro- β -phenylalanine,²² the tyrosine 2,3-aminomutase from *Streptomyces globisporus* (SgTAM) co-crystallized with L- α,α -difluoro- β -tyrosine,²³ all with a covalent bond between the amino group of the substrate/inhibitor and the exocyclic methylene carbon of the MIO supported a reaction mechanism proceeding through the so called *N*-MIO intermediate. This was also in agreement with QM/MM calculations on TAL²⁴ and PAL¹⁹ supporting ammonia elimination via an *N*-MIO intermediate and suggesting the formation of similar *N*-MIO complexes as a common feature of the mechanisms for all MIO-enzymes.²⁰

Proceeding from the *N*-MIO intermediate of PAL by the aid of the catalytically essential Tyr110,²⁵ the elimination is either stepwise i) through a benzylic carbanion (E1cB), ii) or through a carbocation (E1), or iii) it is a concerted process (E2). A thorough kinetic analysis of the PAL reaction,²⁶ based on kinetic isotope effects of [¹⁵N]- and/or [3-²H₂]-phenylalanine favored the mechanism via a carbanionic intermediate (E1cB). In a recent study,²⁷ a previously unrevealed, competing MIO-independent reaction pathway was reported with PALs from *Anabaena variabilis* (AvPAL), *Rhodotorula glutinis* (RgPAL), and parsley (PcPAL) which proceeds in a non-stereoselective manner and results in the generation of both L- and D-phenylalanine derivatives. Isotope-labeling studies and mutagenesis of key active site residues were consistent with amino acid deamination occurring by a stepwise E1cB elimination mechanism in this case as well.

It was already shown that phenylalanine 2,3-aminomutase from *Taxus canadensis* (TcaPAM) could transform (S)- α -styrylalanine (L-1a) into (2E,4E)-5-phenylpenta-2,4-dienoic

acid (2a) as the major product.²⁸ Considering the high structural and sequence similarities between TcaPAM and PcPAL, it seemed commendable to explore the potential of PcPAL in the kinetic resolution of α -styrylalanines and stereoselective amination of (2E,4E)-styrylacrylates in the reverse reaction (Scheme 1).

The present study – with the aim of extending the substrate scope of PcPAL towards sterically demanding phenylalanine analogues – created mutant variants of recombinant PcPAL with enlarged aromatic binding pocket enabling smooth fit of styrylalanines with increased distance between the aromatic moiety and the α -stereogenic carbon into the active site.

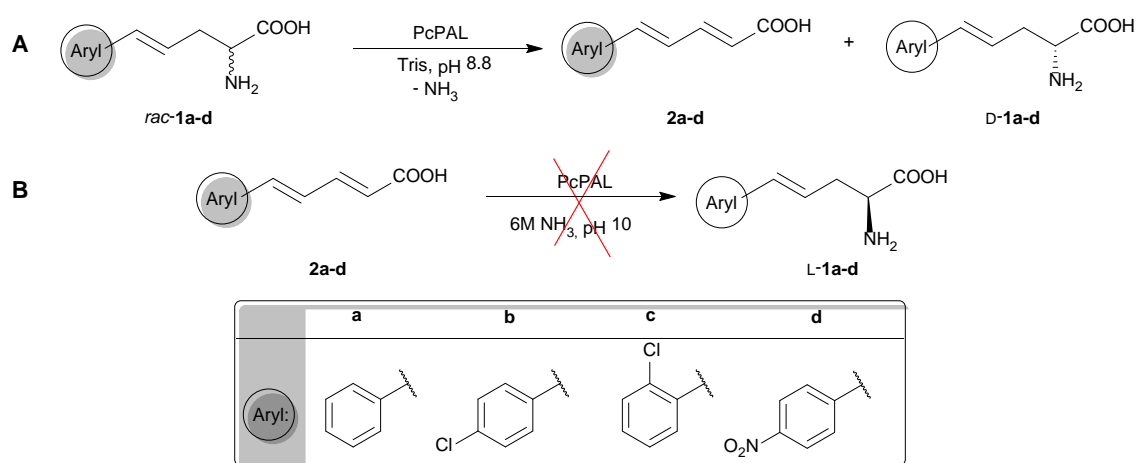
The results of this study were rationalized by molecular modeling in the framework of a straightforward physics-based approach based on ligand/*N*-MIO intermediate poses/conformation sets and several molecular descriptors to approximate changes on the potential energy surface and thus characterize certain phenomena of the catalysis.

Results and discussion

Initial studies and F137X-PcPAL mutant set construction

First, to assess the applicability of wt-PcPAL to the styrylalanine compound family, the commercially available L-styrylalanine (L-1a) was tested as substrate in the ammonia elimination reaction (Scheme 1), monitored by HPLC and UV-based assays. The ¹H-NMR spectra of the reaction mixtures confirmed the formation of (2E,4E)-5-phenylpenta-2,4-dienoic acid (2a) as product. Kinetic studies of the ammonia elimination catalyzed by wt-PcPAL showed 14.7-fold lower *k*_{cat} and 777-fold lower *k*_{cat}/*K*_M values for the deamination of L-1a in comparison with the deamination of the natural substrate, L-Phe (Table 1).

The low enzyme activity, observed with L-styrylalanine (L-1a), was rationalized by molecular modeling. Evaluation of energetics indicated as the most disadvantageous interaction



Scheme 1 The (A) ammonia elimination from racemic styrylalanines 1a-d and the (B) attempted ammonia addition onto styrylacrylates 2a-d catalyzed

Table 1 Kinetic parameters for wt-PcPAL and for F137X-PcPAL mutants in ammonia elimination from L-phenylalanine and from L-styrylalanine (L-**1a**)

PcPAL	L-Phenylalanine			L-Styrylalanine		
	K_M (μM)	k_{cat} (s^{-1})	k_{cat}/K_M ($\text{M}^{-1} \text{s}^{-1}$)	K_M (μM)	k_{cat} (s^{-1})	k_{cat}/K_M ($\text{M}^{-1} \text{s}^{-1}$)
wt	83 ± 5	0.694 ± 0.020	8361 ± 291	4384 ± 158	0.0471 ± 0.0003	10.7 ± 0.4
F137V	86 ± 10	0.173 ± 0.001	2011 ± 131	186 ± 6	0.4220 ± 0.0280	2269 ± 168
F137A	1732 ± 15	0.283 ± 0.001	163 ± 2	1173 ± 70	0.1320 ± 0.0026	112.5 ± 7.2
F137G	4969 ± 153	0.052 ± 0.003	10.4 ± 0.9	4120 ± 270	0.0345 ± 0.0038	8.3 ± 0.2

Table 2 Kinetic parameters for wt- and F137V-PcPALs in the ammonia elimination of racemic styrylalanines *rac*-**1a-d**

Substrate	wt-PcPAL			F137V-PcPAL		
	K_M (μM)	k_{cat} (s^{-1})	k_{cat}/K_M ($\text{M}^{-1} \text{s}^{-1}$)	K_M (μM)	k_{cat} (s^{-1})	k_{cat}/K_M ($\text{M}^{-1} \text{s}^{-1}$)
<i>rac</i> - 1a	395 ± 6	0.00620 ± 0.00070	15.6 ± 0.9	201 ± 12	0.2760 ± 0.0130	1373 ± 97
<i>rac</i> - 1b	154 ± 7	0.00024 ± 0.00001	1.5 ± 0.03	78.3 ± 2	0.0786 ± 0.0044	1004 ± 86
<i>rac</i> - 1c	28 ± 1	0.00034 ± 0.00002	12.1 ± 0.6	94.7 ± 2	0.1570 ± 0.0020	1657 ± 10
<i>rac</i> - 1d	287 ± 3	0.00320 ± 0.00030	11.1 ± 0.5	326 ± 10	0.0099 ± 0.0001	30.2 ± 1.9

the one between the phenyl ring of the *N*-MIO intermediate from L-**1a** and the phenyl ring of residue F137 in the aromatic binding pocket of wt-PcPAL (Figure 1). Moreover, enumeration of the possible conformations indicated three major species. In panel **A** of Figure 1, two apparently catalytically active conformations, resembling the active arrangement of phenylalanine in PcPAL,¹⁹ are depicted. These two *N*-MIO intermediate conformations can result in two different arrangements of the diene product (**2a**), the *s-cis* [referred as *pro-s-cis* (*psc*)] or the *s-trans* [referred as *pro-s-trans* (*pst*)] conformations. Panel **B** of Figure 1 depicts a third conformation adopting a somewhat different arrangement with significantly less stress exerted by F137. However, this specimen would lead to (2*Z*,4*E*)-5-phenylpenta-2,4-dienoic acid which was not detected by NMR in the reaction mixture. Thus, conformation L-**1a_u** could be considered as an unproductive state. These modeling results implied that the

reaction rate might be improved by reducing the clash between the aromatic moieties of L-**1a** in its *N*-MIO state and F137 within the active site of wt-PcPAL, and simultaneously by relocating the population of the unproductive conformation. Notable, that earlier, in case of 4-substituted cinnamates as substrates in the reverse amination reaction with PcPAL, mutations of the F137 site were efficacious.²⁹ Therefore, to provide more space for the phenyl ring of L-**1a**, residue Phe137 in the hydrophobic pocket of wt-PcPAL was changed to smaller hydrophobic residues Val, Ala, and Gly in mutants F137V-, F137A-, and F137G-PcPAL, respectively. The kinetic parameters of the wild-type and the mutant PcPALs were determined in the ammonia elimination reaction for the natural substrate L-Phe and for L-**1a** (Table 1).

The ratio of k_{cat} and K_M – often referred as the “catalytic efficiency” – is a useful index for comparing the efficiency of an enzyme acting on alternative substrates.³⁰ Not surprisingly, all

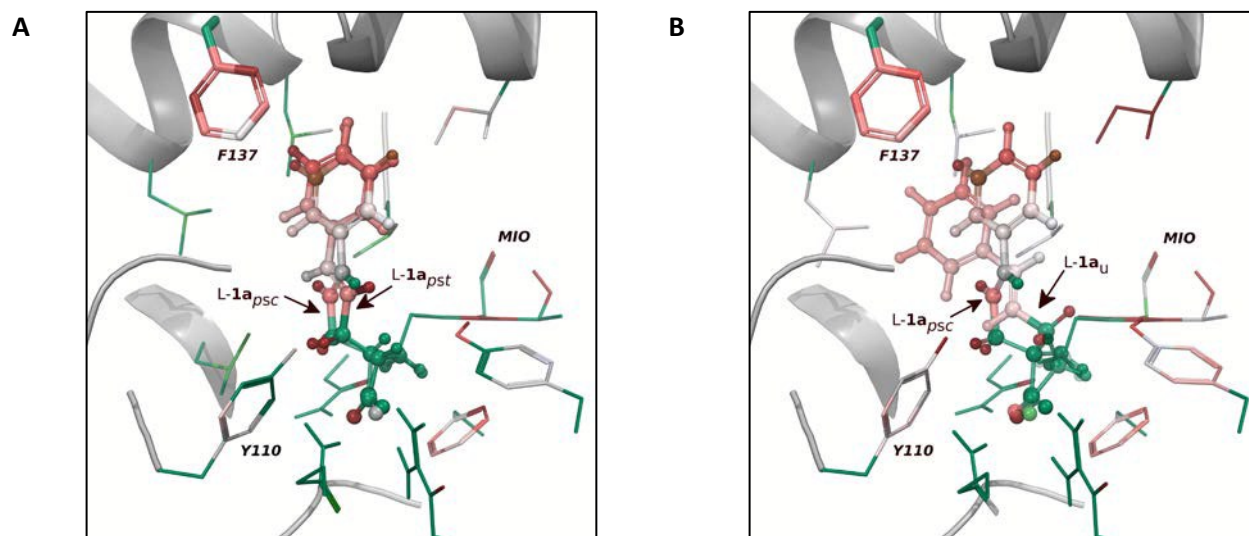


Figure 1 Active site models of the three major conformations of the *N*-MIO complex of L-**1a** with wt-PcPAL [colors refer to the atomic contributions to the total energy ranging from green (beneficial, ≤ -5 kcal mol⁻¹) to red (detrimental, ≥ 2 kcal mol⁻¹)]. (A) The two allegedly catalytically active conformations L-**1a_{psc}** and L-**1a_{pst}** leading to the product **2a** in *s-cis* (synperiplanar) or *s-trans* (antiperiplanar) arrangements, respectively. (B) the apparently catalytically inactive conformation L-**1a_u** compared to the apparently catalytically active L-**1a_{psc}** conformation.

mutants showed decreased – declining with side chain volume – catalytic efficiencies in the ammonia elimination reaction of the natural substrate, L-Phe, in comparison with wt-PcPAL. However, it can be clearly seen that F137V has superior catalytic efficiency with L-1a as substrate, with the nearly 240-fold increased k_{cat}/K_M value compared to wt-PcPAL. The F137A mutation also provided an increased k_{cat}/K_M value with L-1a (10.5-fold enhancement), while the F137G mutation showed no beneficial effect on the transformation of L-1a. Interestingly, k_{cat} values showed also substantial variances. The trends and maxima were almost the same as in the case of the k_{cat}/K_M values. Regarding L-Phe, k_{cat} declined with decreasing side chain volume, except for a surprising spike with F137A-PcPAL. Similarly, F137V-PcPAL with L-1a showed a tremendous, almost 9-fold increase; F137A showed only a moderate 2.8-fold increase; and F137G showed no improvement in k_{cat} compared to wt-PcPAL.

A comparison of the covalent *pro-s-cis* intermediates from L-1a within wt-PcPAL and F137V-PcPAL is shown in panel A of Figure 2. The aromatic moiety of the N-MIO intermediate from L-1a is forced towards I460 within the active site of wt-PcPAL, unlike in the case of the more relaxed complex within F137V-PcPAL, meanwhile the alanyl substructure of the substrate – which is directly involved in bond rearrangement – is kept relatively unchanged in both cases. It is worth noting that upon binding of L-1a to wt-PcPAL the active site supposedly undergoes conformational change at the I460-side chain which reshapes the active site and alters the energetics of distinct substrate conformations.

Synthetic applications in kinetic resolution

Next, we wanted to explore the potential of F137V-PcPAL in the synthetically valuable ammonia elimination and addition reactions of various styrylalanines and their acrylic counterparts. Therefore, racemic styrylalanines *rac*-1a-d as well as the corresponding styrylacrylates 2a-d were

Supplementary Information, ESI) and tested with F137V-PcPAL and wt-PcPAL as biocatalysts (Scheme 1) (DOI: 10.1039/C7OB00562H). In the ammonia elimination reaction all styrylalanines (*rac*-1a-d) were accepted as substrates by wt-PcPAL as well as by F137V-PcPAL. However, the F137V mutant showed significantly improved catalytic activities (k_{cat} and k_{cat}/K_M) in comparison with the wild-type enzyme in all cases (Table 2). The synthetic usefulness of F137V-PcPAL towards styrylalanines in the ammonia elimination reactions was further tested by kinetic resolutions (KRs) with 0.1 mmol *rac*-1a-d at 5 mM concentration using 0.5–1.0 mg of pure PcPALs. The conversion of the PcPAL-catalyzed reactions from racemic styrylalanines *rac*-1a-d (Figure 3), and the enantiomeric excess of the residual substrates D-1a-d were both monitored (Table 3). In case of longer reaction times (>48 h), a fresh enzyme batch was added to the reaction mixture at the end of each 48 h period to overcome enzyme inactivation during long incubation times.

Time course analysis of the conversion of the KRs of *rac*-1a-d (Figure 3) also indicated the beneficial catalytic properties of F137V-PcPAL compared to that of wt-PcPAL. The theoretically possible conversion in a kinetic resolution (KR) with exclusive selectivity (50%) could be almost perfectly approached ($c > 49\%$) with the F137V mutant in reactions of *rac*-1a,c after moderate reaction times (24 h and 134 h, respectively) and in the cases *rac*-1b,d after longer incubation times (274 h and 300 h, respectively). The unusually long time required for the compounds *rac*-1b,d can be significantly shortened by using higher enzyme to substrate ratio. This might be easily performed by applying immobilized forms of PcPAL in microreactors.¹⁷

On the other hand, in reactions of racemic styrylalanines *rac*-1a-d with wt-PcPAL, almost complete conversion of the L-enantiomer could be obtained only with *rac*-1a ($c > 49\%$ after 274 h), while the conversions of *rac*-1b-d could not approach the theoretically possible value within 500–600 h.

KRs of *rac*-1a-c with wt-PcPAL showed that enantiomeric

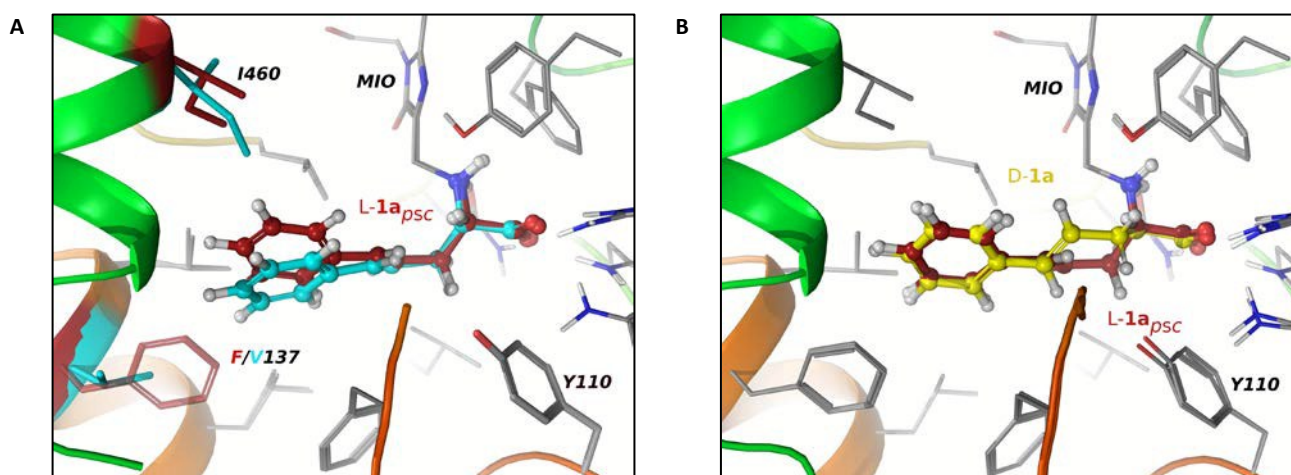


Figure 2 Molecular models of various N-MIO intermediates in PcPALs. (A) Arrangement of covalent intermediate L-1a_{psc} in wt-PcPAL (relevant residues are in red) and in F137V-PcPAL (relevant residues are in cyan). Note the different side chain conformations of I460. (B) Arrangement of the unproductive covalent complex from D-1a (yellow) compared to the productive intermediate L-1a_{psc} (red) in wt-PcPAL.

synthesized (see Schemes S1 and S2 in Electronic excess of residual D-1a-c fractions were in close agreement

Table 3 Kinetic resolution of styrylalanines *rac*-**1a-d** and enantiomeric excess of the residual *D*-**1a-d** in the ammonia elimination reactions catalyzed by PcPAL variants.

Entry	PcPAL	Substrate	Time (h)	<i>c</i> (%)	<i>Ee</i> _{theor} (%) ^a	<i>Ee</i> _{obs} (%)
1	F137V	<i>rac</i> - 1a	24	50	100	>99
2		<i>rac</i> - 1b	274	50	100	>99
3		<i>rac</i> - 1c	134	50	100	>99
4		<i>rac</i> - 1d	300	50	100	>99
5	wild-type	<i>rac</i> - 1a	274	50	100	>98
6		<i>rac</i> - 1b	504	29	41	41
7		<i>rac</i> - 1c	600	36	56	55
8		<i>rac</i> - 1d	600	31	45	41

^a $Ee_{theor} = c/(1-c)$ for a fully selective kinetic resolution

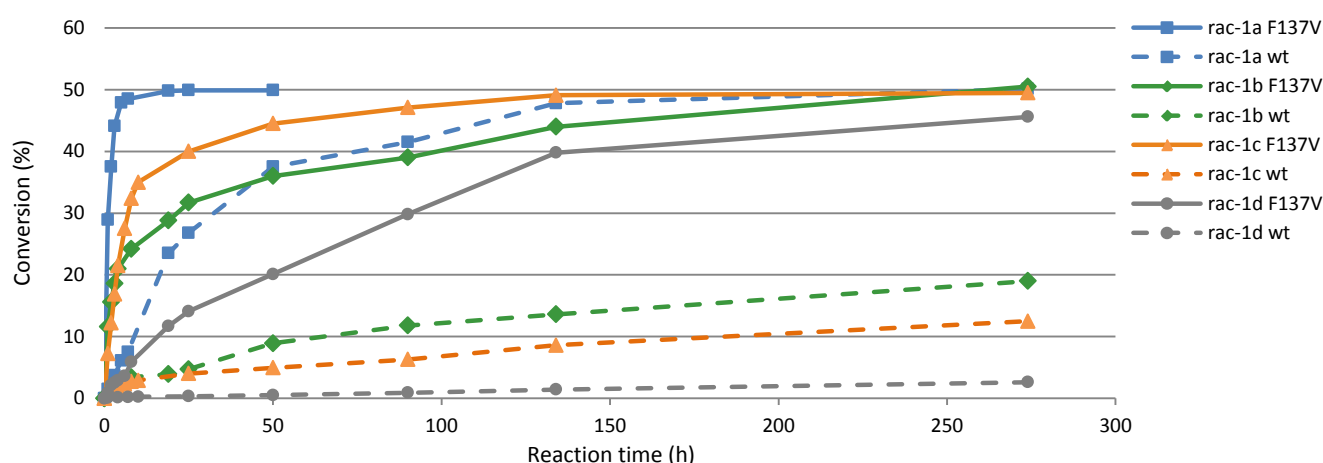


Figure 3 Comparison of the time profiles of the conversions in the ammonia elimination reactions from *rac*-**1a-d** catalyzed by F137V-PcPAL (continuous lines) and wt-PcPAL (dashed lines).

with the theoretical values of a fully selective KR indicating excellent enantiomer selectivity and preference of the process to digest the L-enantiomers, L-**1a-c** (Table 3). Although the enantiomeric excess of the residual *D*-**1d** fractions were somewhat smaller as expected for a fully selective KR, it was not possible to determine whether this happened due to experimental error of the HPLC process or due to a non-perfect enantiomer selectivity of the KR from *rac*-**1d**.

KRs of *rac*-**1a-d** with F137V-PcPAL, however, showed that the residual *D*-**1a-d** fractions had enantiomeric excess values over 99% indicating unquestionably exclusive enantiomer selectivity in all cases. This was also obvious from the time course curves of conversions approaching asymptotically the 50% value when F137V-PcPAL was applied as biocatalyst in the KRs (Figure 3). Time course profiles of the reactions and the comparison of the kinetic constants with L-**1a** (Table 1) and with *rac*-**1a** (Table 2) suggests that *D*-styrylalanine inhibits the ammonia elimination of L-**1a** both with wt-PcPAL and F137V-PcPAL. If *D*-**1a** had no interaction with wt-PcPAL than only a 2-fold higher apparent K_M of the racemate *rac*-**1a** would be observed as compared to the pure L-enantiomer L-**1a** without affecting the k_{cat} value. However, when *rac*-**1a** was applied as substrate instead of the pure L-enantiomer L-**1a**, the V_{max} (thus implicitly the apparent k_{cat} value) decreased 7.6-fold, while the K_M value decreased 11-fold. With F137V-PcPAL, the differences became less significant, k_{cat} decreased only 1.5-fold, and K_M

showed only a minor variation, within the range of experimental errors. Because the residual fractions from the KRs of *rac*-**1a-d** had quite high enantiomeric excess at 50% conversion (*ee* >99%, see Table 3), the *D*-enantiomers *D*-**1a-d** were apparently not converted. This implied *D*-**1a** being a reversible inhibitor with considerable affinity forming an unproductive enzyme-*D*-substrate complex. The covalent, unproductive *N*-MIO complex from *D*-**1a** showed astonishing overlap with the productive intermediate L-**1a_{pst}** within the active site of wt-PcPAL (panel B in Figure 2).

Rationalization of the reactions of styrylalanines *rac*-**1a-d** catalyzed by the PcPAL variants

To get a more detailed view of the catalytic process, a quantitative model was created and validated. Because *N*-MIO complexes were assumed as a common feature of the mechanism for all MIO-enzymes,²⁰ the enzyme-substrate complexes within the active site of the PcPAL variants were constructed as *N*-MIO intermediates of L- and *D*-**1a-d** applying an induced-fit covalent docking protocol which was previously used to rationalize the reactions of various substrates within another MIO enzyme, the phenylalanine 2,3-aminomutase from *Pantonea agglomerans*.³¹

The non-covalent substrate-binding states of L- and *D*-**1a-d** – directly preceding their *N*-MIO intermediate states – were also modeled. Because the calculations did not reveal any

ARTICLE

Organic & Biomolecular Chemistry

unproductive states for these non-covalent complexes, they will not be discussed hereafter.

The occurrence of the non-stereoselective, MIO-independent reaction pathway²⁷ was ruled out experimentally. The attempted ammonia elimination from *rac*-**1a-d** by wt-PcPAL with inactivated MIO group (obtained by treatment with sodium borohydride)³² showed no product formation. The conversion of 4-nitrophenylalanine with the sodium borohydride treated wt-PcPAL served as positive control for integrity of the residual parts of the active site.

To investigate the catalytic activities quantitatively with all substrate and enzyme mutant combinations, the apparent k_{cat} values – as determined from the maximum reaction rates according to the left-hand side of Eq. 1 – were decomposed into their constituents.

At substrate saturation, when the concentration of the free enzyme is negligible, four species of enzyme–substrate intermediate complexes ($E_{\text{L-psc}}$, $E_{\text{L-pst}}$, $E_{\text{L-u}}$ and E_{D}), can be present in thermal equilibrium (allosteric inhibition was not considered in this study). As discussed previously, from the L-enantiomer two catalytically active enzyme–substrate intermediate complexes (the *s-cis* and *s-trans* L-**1a-d**–N-MIO states, abbreviated as $E_{\text{L-psc}}$, $E_{\text{L-pst}}$) and an unproductive one (the unproductive L-**1a-d**–N-MIO states, abbreviated as $E_{\text{L-u}}$) could be formed, while another unproductive complex could be formed from the D-enantiomers (the unproductive D-**1a-d**–N-MIO states, abbreviated as E_{D}). Thus, according to Eq. 1, the catalytic activity is the sum of the activities of $E_{\text{L-psc}}$ and $E_{\text{L-pst}}$.

$$V_{\text{max}} = k_{\text{cat}}[E^*] = k_{\text{cat,L-psc}}[E_{\text{psc}}] + k_{\text{cat,L-pst}}[E_{\text{pst}}] \quad (1)$$

Expressing the substrate concentrations as molar fractions and substituting the Eyring equation yields Eq. 2

$$k_{\text{cat}} = \frac{k_{\text{B}}T}{h} e^{\Delta G_{\text{L-psc}}^{\ddagger}/RT} x_{\text{psc}} + \frac{k_{\text{B}}T}{h} e^{\Delta G_{\text{L-pst}}^{\ddagger}/RT} x_{\text{pst}} \quad (2)$$

where T is the temperature, k_{B} is the Boltzmann constant, h is the Planck constant, ΔG^{\ddagger} is the corresponding Gibbs energy of activation, and x is the corresponding molar fraction. To approximate the Gibbs energy of activation, two descriptors

were used, one was associated with the acidity of the β -proton (${}_{\text{CB}}^R\Delta E$, Table 4), and the other one was associated with the steric strain (${}_{\text{ster}}^R\Delta E$, Table 4). Applying these descriptors, Eq. 3 is derived,

$$k_{\text{cat}} = a(e^{b \cdot \Delta E_{\text{L-psc}}^{\ddagger}/RT + \ln x_{\text{L-psc}}} + e^{b \cdot \Delta E_{\text{L-pst}}^{\ddagger}/RT + \ln x_{\text{L-pst}}}) \quad (3)$$

where a and b are constants, and ΔE^{\ddagger} is the sum of ${}_{\text{strain}}^R\Delta E$ and ${}_{\text{CB}}^R\Delta E$.

Non-linear estimation of the constants in Eq. 3 gave statistically significant regression ($R^2=0.8$, $p=0.002$) and parameters ($a=90.63$, $p=0.023$; $b=0.0791$, $p=0.014$). The complete derivation of Eq. 3 is described in the ESI.

Evaluation of the F137V mutation based on computed data (Table 4) clearly demonstrated that the affinities of L-**1a-d** in the *pro-s-cis* conformation (${}_{\text{D}}^R\Delta E_{\text{L-psc}}$) increased substantially, as it was assumed. Moreover, the affinities of L-**1a-d** in the *pro-s-trans* conformation increased even more, as their population increased relative to the *pro-s-cis* conformation ($x_{\text{L-pst}}$ vs. $x_{\text{L-psc}}$), with the exception of L-**1d** with F137V-PcPAL. The unproductive complexes of **1a-d** (x_{D} and $x_{\text{L-u}}$), however, showed a more complex situation. By the F137V-mutation, in some cases unproductive states were displaced (e.g. **1c**), and in some other cases they were introduced (e.g. **1d**). These results indicated that the simple increase of active site volume is not sufficient from the standpoint of enzyme design, and a more sophisticated active site shape altering scheme – taking the relative spatial requirements of the intermediate states and unproductive states into account – should be employed. Inspection of the descriptors for the acidity of the *pro-S* β -hydrogen (${}_{\text{CB}}^R\Delta E$) – which was derived from single point DFT calculations on a truncated part of the intermediate model – revealed that in almost all cases acidity of the *pro-s-trans* conformations compared to the *pro-s-cis* conformations (${}_{\text{CB}}^R\Delta E_{\text{L-pst}}$ vs. ${}_{\text{CB}}^R\Delta E_{\text{L-psc}}$) was comparable or higher, except for **1b**/wt-PcPAL and **1d**/F137V-PcPAL. Thus, the higher proportion of the *pro-s-trans* conformations ($x_{\text{L-pst}}$), enabling more facilitated reaction due to the more acidic *pro-S* β -hydrogens, could contribute to the increased activity of F137V-PcPAL. Moreover, the acidity descriptors ${}_{\text{CB}}^R\Delta E_{\text{L-psc}}$ and ${}_{\text{CB}}^R\Delta E_{\text{L-pst}}$ varied remarkably in wt- and F137V-PcPAL, mostly due to the changes in dihedral angle

Table 4 Calculated relative binding energies (${}_{\text{D}}^R\Delta E$), rounded values of molar fractions of the various covalent N-MIO complexes (x), energy-based descriptors of the acidity of their *pro-S* β -hydrogen (${}_{\text{CB}}^R\Delta E$) and steric strain of activation (${}_{\text{strain}}^R\Delta E$) for intermediates of compound **1a-d** in PcPAL variants. In the subscript L and D correspond to the actual enantiomer; *psc*, *pst*, and *u* correspond to the *s-cis*, *s-trans* product forming, and the unproductive **1a-d**–N-MIO states.

PcPAL	Substrate	${}_{\text{D}}^R\Delta E_{\text{L-psc}}^{\text{b}}$ (kcal mol ⁻¹)	$x_{\text{L-psc}}^{\text{c}}$ (%)	$x_{\text{L-pst}}^{\text{c}}$ (%)	x_{D}^{c} (%)	$x_{\text{L-u}}^{\text{c}}$ (%)	${}_{\text{strain}}^R\Delta E_{\text{L-psc}}^{\text{b}}$ (kcal mol ⁻¹)	${}_{\text{strain}}^R\Delta E_{\text{L-pst}}^{\text{b}}$ (kcal mol ⁻¹)	${}_{\text{CB}}^R\Delta E_{\text{L-psc}}^{\text{b}}$ (kcal mol ⁻¹)	${}_{\text{CB}}^R\Delta E_{\text{L-pst}}^{\text{b}}$ (kcal mol ⁻¹)	${}_{\text{CB}}^R\Delta E_{\text{L-u}}^{\text{b}}$ (kcal mol ⁻¹)
wild-type	1a	13.0	99.7	0.2	0.1	0.0	11.0	16.2	-2.2	3.8	6.1
	1b	22.5	2.1	0.0	0.0	97.9	22.5	28.5	-1.5	-1.8	-0.3
	1c ^a	13.6	41.5	0.0	0.7	57.8	12.9	20.4	11.9	8.2	-3.6
	1d	33.7	93.4	0.0	0.0	6.6	33.9	42.4	-11.4	-11.5	0.1
	1a	-3.3	95.2	4.8	0.0	0.0	-0.7	1.1	-4.6	-10.7	-6.2
F137V	1b	-3.2	99.9	0.1	0.0	0.0	1.4	5.7	-5.4	-7.7	-2.4
	1c ^a	1.4	60.9	5.0	0.8	0.0	5.3	6.8	11.3	0.5	-10.8
	1d ^a	22.7	0.3	18.1	0.9	80.7	25.3	22.9	-31.6	-20.4	11.3

^a In these cases multiple *psc* and *pst* conformations were obtained, but only the lowest energy representatives were taken into account. ^b Quantities are related to the same property of L-Phe with wt-PcPAL, in the form of ${}^R\Delta E = \Delta E - \Delta E_{\text{L-Phe,wt-PcPAL}}$. For the detailed description of the calculation method, see the ESI. For the reasoning of the necessary relativization, see ref. 30. ^c Molar fractions were calculated based on Boltzmann probabilities incorporating all substrate conformations.

between the phenyl ring and the conjugated double bond of the styryl moiety. The more planar the skeleton was; the better dispersion of the negative charge could happen due to the higher degree of conjugation. This implied that in an efficient enzyme design protocol the substrate affinity and electronic effects should be monitored simultaneously.

This exploratory model validates the very concept of our computations and it can be used as a tool to analyze various factors of importance in catalysis and to provide valuable input for further studies. Although this model was not expected to predict the apparent k_{cat} with high accuracy, it enabled us to predict the order of magnitude of the catalytic activity for several further congeneric substrates.

With our model, we examined the behavior of further substituted styrylalanines bearing bromo, cyano, fluoro, and trifluoromethyl substituents at position 4 (Table 5). Calculations with wt-PcPAL predicted *rac*-4-fluorostyrylalanine and *rac*-4-cyanostyrylalanine to be poor substrates behaving similarly to *rac*-styrylalanine, *rac*-1a. Modeling forecast *rac*-4-

bromo- and *rac*-4-trifluorostyrylalanines to be extremely poor substrates or reversible inhibitors ($k_{\text{cat-pred}} \leq 0.0001 \text{ s}^{-1}$). The F137V mutation calculated no significant improvement only for *rac*-4-trifluorostyrylalanine ($k_{\text{cat-pred}} \leq 0.0001 \text{ s}^{-1}$). Modeling within this mutant indicated comparably high activity for *rac*-4-fluorostyrylalanine ($k_{\text{cat-pred}} = 0.18 \text{ s}^{-1}$) as for *rac*-styrylalanine (*rac*-1a: $k_{\text{cat}} = 0.28 \text{ s}^{-1}$). Somewhat lower catalytic activities were predicted for *rac*-4-bromo- and *rac*-4-cyanostyrylalanines ($k_{\text{cat-pred}} = 0.120 \text{ s}^{-1}$ and 0.098 s^{-1} , respectively) being in the range of the activity of *rac*-4-chlorostyrylalanine (*rac*-1c: $k_{\text{cat}} = 0.16 \text{ s}^{-1}$).

Study for ammonia addition onto styrylacrylates 2a-d with PcPALs

Regrettably, no 1a-d could be detected by HPLC in the attempted ammonia addition reactions of styrylacrylates 2a-d with either wt-, F137V-, F137A-, or F137G-PcPAL even after 20 days (Scheme 1, for details see Section 5.6 of ESI). The possible reasons of these results were analyzed by computer modeling in the cases of wt- and F137V-PcPAL. It was not surprising, that

Table 5 Calculated relative binding energies ($^R\Delta E$), rounded values of molar fractions of the various covalent *N*-MIO complexes (x), energy-based descriptors of the acidity of their *pro*-S β -hydrogen ($^R\Delta E$) and steric strain of activation ($^R\Delta E$) for intermediates of further 4-substituted styrylalanines in PcPAL variants, and predicted ($k_{\text{cat-pred}}$) values for the corresponding reactions. In the subscript l and d correspond to the actual enantiomer; *p*sc, *p*st, and *u* correspond to the *s*-*cis*, *s*-*trans* product forming, and the unproductive 4-substituted styrylalanine-*N*-MIO states.

PcPAL	4-Substituent ^a	$^R\Delta E_{\text{c-psc}}^b$ (kcal mol ⁻¹)	$x_{\text{l-psc}}^c$ (%)	$x_{\text{l-pst}}^c$ (%)	x_{d}^c (%)	$x_{\text{l-u}}^c$ (%)	$^R\Delta E_{\text{c-psc}}^b$ (kcal mol ⁻¹)	$^R\Delta E_{\text{c-pst}}^b$ (kcal mol ⁻¹)	$^R\Delta E_{\text{c-psc}}^b$ (kcal mol ⁻¹)	$^R\Delta E_{\text{c-pst}}^b$ (kcal mol ⁻¹)	$k_{\text{cat-pred}}$ (s ⁻¹)
wild-type	F-	13.2	88.0	0.0	0.0	12.0	13.5	20.3	-0.3	-1.5	0.014
	CF ₃ -	38.7	0.2	0.0	0.0	99.8	40.1	48.0	-3.7	-3.8	<0.0001
	Br-	23.3	2.1	0.0	0.0	97.9	24.2	32.3	-1.7	-2.4	0.0001
	NC-	33.7	93.4	0.0	0.0	6.6	33.9	42.4	-11.4	-11.5	0.004
F137V	F-	-4.3	99.3	0.7	0.0	0.0	0.6	3.6	-5.7	-6.9	0.180
	CF ₃ -	15.5	0.0	0.0	0.0	100.0	21.5	15.5	-6.8	-8.2	<0.0001
	Br-	-2.6	95.9	0.0	0.0	4.1	2.9	8.2	-5.4	-1.6	0.120
	NC-	8.3	100.0	0.0	0.0	0.0	13.1	17.8	-13.7	-15.0	0.098

^a Substituent of the further 4-substituted styrylalanines. ^b Quantities are related to the same property of L-Phe with wt-PcPAL, in the form of $^R\Delta E = \Delta E - \Delta E_{\text{L-Phe;wt-PcPAL}}$. For the detailed description of the calculation method, see the ESI. For the reasoning of the necessary relativization, see ref. 30. ^c Molar fractions were calculated based on Boltzmann probabilities incorporating all substrate conformations.

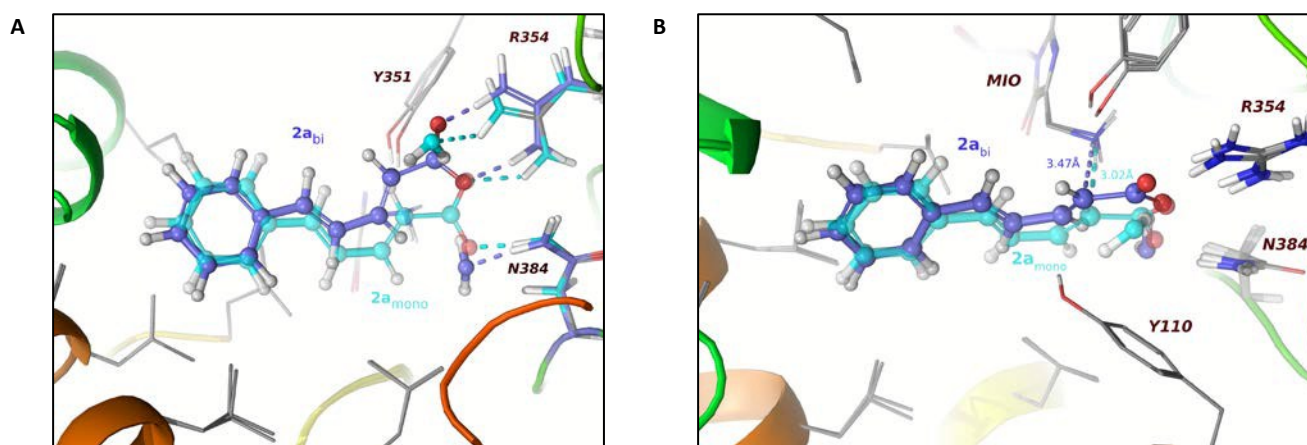


Figure 4 Models of possible styrylacrylate 2a-arrangements within amino-enzyme (NH_2 -*N*-MIO) forms of PcPAL variants. (A) Hydrogen bonds forming among the carboxylate moiety, a water molecule and R354, N384 of the bidentate and monodentate salt bridge forming poses of 2a within F137V-PcPAL. (B) The distances of the N atom of NH_2 of the NH_2 -*N*-MIO from the C_2 atom of the two lowest energy mono- ($2a_{\text{mono}}$) or bidentate ($2a_{\text{bi}}$) carboxylate binding poses of 2a within F137V-PcPAL. In both panels, the model depicting $2a_{\text{mono}}$ (in cyan) corresponds to a product from the productive *N*-MIO intermediate (L-1a_{psc}), the model depicting $2a_{\text{bi}}$ (in blue) shows a low energy inactive state.

Table 6 Calculated relative ligand binding energies ($^R\Delta E$) and rounded values of molar fractions of mono- and bidentate carboxylate binding modes (x) for **2a-d** [mono and bi in the subscript relate to the lowest energy ligand pose that forms a monodentate (putatively active pose) or bidentate (putatively inactive pose) salt bridge].

Compound	PcPAL	wt-PcPAL			F137V-PcPAL		
		$^R\Delta E_{\text{mono}}^a$ kcal mol ⁻¹	x_{mono}^b (%)	x_{bi}^b (%)	$^R\Delta E_{\text{mono}}^a$ kcal mol ⁻¹	x_{mono}^b (%)	x_{bi}^b (%)
2a		17.2	1.0	97.5	2.2	15.8	71.8
2b		25.8	18.0	81.9	-0.2	77.9	14.4
2c		20.4	0.1	75.8	2.0	0.6	91.3
2d		41.8	0.0	92.6	10.4	6.3	93.3

^a Quantities are related to the same property of (*E*)-cinnamate with wt-PcPAL, in the form of $^R\Delta E = \Delta E - \Delta E_{(E)\text{-cin;wt-PcPAL}}$. For the detailed description of the calculation method, see the ESI. For the reasoning of the necessary relativization, see ref. 30. ^b Molar fractions were calculated based on Boltzmann probabilities incorporating all ligand poses.

the relative binding energies of **2a-d** indicated poor binding to wt-PcPAL which was remarkably improved with F137V-PcPAL (Table 6, $^R\Delta E_{\text{mono}}$). Modeling revealed allegedly unproductive ligand poses for **2a-d** with both PcPAL variants in almost all cases with higher binding affinity as compared to the possibly reactive arrangements of **2a-d**. The case of **2a** within F137V-PcPAL is shown in Figure 4. Panel **A** depicts the lower energy arrangement of the putatively inactive conformation of **2a** with a bidentate salt bridge between R354 and the carboxylate group (in blue) compared to the putatively catalytically active conformation forming a monodentate salt bridge (in cyan). Panel **B** of Figure 4 depicts models of the possible arrangements of **2a** within F137V-PcPAL with an elongated distance between the N atom of NH₂ of the NH₂-N-MIO and the C₂ atom in the model for putatively inactive conformation of **2a** (in blue) as compared to the corresponding distance in the model including **2a** in reactive conformation (in cyan). Grand canonical Monte Carlo simulations indicated a water molecule for both binding poses of **2a** forming a hydrogen bond with the carboxylate group of the ligand, albeit in remarkably different positions. Irrespectively to the monodentate or bidentate nature of the carboxylate moiety regarding to the arginine R354, the three oxygen atoms of the carboxylate group and the water always formed hydrogen bonds with residues R354 and N384 (Figure 4, Panel A). These findings agreed with experimental results found in X-ray

Table 7. Primers used for site-directed mutagenesis

Primer	Sequence (5'-3')	T_m^{pp} (°C)*	T_m^{no} (°C)*
F137V-for	GATTAGAGTCTTGAACGCCGGAATTTTCGGAACGGATC	56	64
F137V-rev	GGCGTTCAAGACTCTAATCAATTCCTTCTGCAAGGCTCC	56	64
F137A-for	GATTAGAGCCTTGAACGCCGGAATTTTCGGAACGGA	58	67
F137A-rev	GCGTTCAAGGCTCTAATCAATTCCTTCTGCAAGGCTCCTC	58	67
F137G-for	GATTAGAGGCTTGAACGCCGGAATTTTCGGAACGGA	58	67
F137G-rev	GCGTTCAAGCCTCTAATCAATTCCTTCTGCAAGGCTCCTC	58	67
PcPAL-for1	CAGAACATTACTCCTGCTTG	-	-
PcPAL-for2	TTGACGTATCCAGAAACAAGG	-	-

* T_m^{pp} and T_m^{no} represents the melting temperature for the primer-primer overlapping sequence and for the primer sequence matched to the template

structure of a TchPAM mutant (PDB ID: 4CQ5)²². In the structure of this enzyme homologous to PcPAL, a continuous electron density between R354 and N384 and the carboxylate of the ligand was observed in the active site complexed with (*E*)-cinnamic acid and a water which was consistent with multiple ligand poses in similar arrangements to that was found in our models for **2a** in PcPAL.

Modeling results suggest that low binding affinities of **2a-d** (especially to wt-PcPAL) and substrate inhibition by the inactive substrate poses of **2a-d** in both PcPAL variants could contribute to the failure of the ammonia addition. Modeling, however, could not represent results which could completely exclude the possibility of successful ammonia addition, especially in the cases of **2a,b** with F137V-PcPAL. Therefore, further reasons of the failure of ammonia addition can be proposed: i) the thermodynamic equilibrium of ammonia addition can be too low to result in detectable amounts of **1-1a-d**, ii) the low water solubility and rather poor affinity of **2a-d** result in non-detectable reaction rates / product concentrations, and iii) probably the enzyme adopts significantly different conformation in 6M ammonia that makes the computational results invalid under the conditions of the ammonia addition.

Experimental

The detailed description of the materials, analytical methods, chemical synthesis of substrates *rac*-**1a-d**, **2a-d**, curves from kinetic measurements, HPLC chromatograms used for the determination of conversion and enantiomeric excess values, as well as the details of the methods used for molecular modeling are attached as Electronic Supplementary Information (ESI).

Site-directed mutagenesis

The site-directed mutagenesis was performed following the protocol described by Naismith and Liu.³³ The PCR reaction contained 10 ng of template DNA (recombinant plasmid encoding PcPAL³⁴), 1 μM solution of primer pair (Table 7), 200 μM dNTPs and 3 units of Pfu DNA polymerase, filled up to 50 μL with water. The PCR cycles were initiated at 95 °C for 5 min, followed by 15 amplification cycles.

Each amplification cycle consisted of 95 °C for 1 min, at temperature of T_m^{no} -5 °C for 1 min and 72 °C for 15 min. The PCR cycles were finished with an annealing step at T_m^{pp} -5 °C for 1 min and the final extension step at 72 °C for 30 min. The PCR products were treated with 5 units of *DpnI* at 37 °C for 2 h and then 10 µL of each PCR reactions was analyzed by agarose gel electrophoresis. An aliquot of 3 µL from the above PCR product was transformed into 100 µL suspension of *E. coli* XL1-Blue competent cells (OD_{600} 2.2) by heat shock. The transformed cells were spread on a Luria-Bertani (LB) plate containing carbenicillin (34 µg/mL) and chloramphenicol (50 µg/mL) and incubated at 37 °C for 16 h. Two colonies from each plate were grown and the plasmid DNA was isolated. To verify the mutations, 400 ng of plasmid DNA was mixed with 50 pmol of T7-for, T7-rev and PcPAL-for1, PcPAL-for2 sequencing primers (Table 6) in a final volume of 15 µL. DNA sequencing was carried out by Genomed (Debrecen, Hungary).

Protein expression

Expression and purification of the wild-type PcPAL and the F137V, F137A, F137G mutants were performed using the protocol described earlier.³⁴ The protein purity and homogeneity was determined using SDS-PAGE and size-exclusion chromatography (Fig. S37, Fig. S38).

Enzyme activity measurements

Enzyme assays were carried out in triplicate at 30 °C in 1 mL quartz cuvette using 0.1 M Tris and 0.12 M NaCl (pH 8.8) as buffer at constant concentration of wt-PcPAL or F137V-PcPAL (for details, see the ESI), varying the substrate concentration from 0.025 and 7.8 mM, until substrate saturation occurred. The kinetic measurements were based on UV-spectroscopy by monitoring the production of the acrylic derivatives **2a-d** at wavelengths where the corresponding amino acids *rac*-**1a-d** showed no absorption (Table S1). Kinetic constants (K_M , v_{max} , and k_{cat}) were obtained from Michaelis-Menten curves by non-linear fitting obtained by MATLAB (see the ESI).

Preparative scale ammonia elimination reactions from *rac*-**1a-d**

The enzymatic reactions were performed in 20 mL of Tris (0.1 M, 0.12 M NaCl, pH 8.8), containing styrylalanines (*rac*-**1a-d**, 5 mM) and purified PcPAL (0.5 mg of F137V-PcPAL or 1 mg of wt-PcPAL). Conversion and the enantiomeric excess of the residual *D*-**1a-d** were monitored using reverse phase high performance liquid chromatography (HPLC) analysis (see the ESI). Samples (30 µL) were taken from the reaction mixture after different time intervals, quenched by adding an equal volume of MeOH, vortexed and centrifuged (13400 rpm, 12100 g, 2 min). The supernatant was used directly for HPLC analysis after transferring through a 0.22 µm nylon membrane filter. In case of enzyme reactions over longer reaction time, another batch of enzyme was added after each 48 h period.

Conclusions

In this study, the substrate scope of PcPAL was extended, by a small library of mutants, towards styrylalanine derivatives, a useful class of synthetically challenging phenylalanine analogues.

Although L-styrylalanine, **L-1a**, was accepted as substrate by wt-PcPAL in the ammonia elimination reaction, low reaction rates were observed with the wild-type enzyme. Molecular modeling suggested that the most serious clash between the aromatic moiety of L-styrylalanine, **L-1a**, and the side chain of residue F137 in the hydrophobic binding pocket can be reduced by introducing residues smaller than F137 by point mutations.

Single point mutations of F137 resulted a small mutant library (F137X-PcPAL, X being V, A, and G), from which F137V-PcPAL showed significantly increased catalytic efficiency and could transform L-styrylalanine, **L-1a**, with comparable activity that of the wt-PcPAL with L-Phe. Furthermore, F137V-PcPAL could be applied as an effective biocatalyst in the synthetically valuable kinetic resolutions by ammonia elimination from racemic styrylalanine analogues *rac*-**1a-d** providing access to *D*-styrylalanines **D-1a-d**, with highly improved catalytic efficiency as compared to the wt-PcPAL in the same reactions.

The reverse ammonia addition reactions onto (*E*)-styrylacrylates **2a-d** – which would be useful for the synthesis of L-styrylalanines **L-1a-d** – failed with either the wild-type or the mutant enzyme variants (F137V, F137A, F137G), presumably due to the formation of energetically favorable unproductive binding states of the styrylacrylates **2a-d**.

Molecular modeling – used as an aid to explain the results of this study – revealed unusually high affinities of the *D*-enantiomers **D-1a-d** towards wt-PcPAL, but proved to be unproductive. Moreover, an unproductive and two productive conformations were revealed for the L-enantiomers **L-1a-d**. Thus, the proportion of the unproductive complexes acting as potent reversible inhibitors and compared to the active arrangements of the L-enantiomers **L-1a-d** turned out to be important.

By the mutation F137V in PcPAL, the less strained arrangement of substrates **L-1a-d** within the active site along with the more pronounced populations of conformations with higher catalytic activities simultaneously led to the beneficial catalytic efficiency of this mutant. Modeling the behavior of further 4-substituted styrylalanines with PcPAL variants predicted the F137V mutation beneficial for the KRs of 4-fluoro-, 4-cyano- and 4-bromostyrylalanines, but non-effective for the KR process of 4-trifluoromethylstyrylalanine.

The results of this study opens new perspectives for substrate scope expansion of PcPAL towards synthetically challenging phenylalanine analogues with increased length of the carbon skeleton between the aromatic moiety and the α -stereogenic carbon.

ARTICLE

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Notes and references

- J. A. Robinson, S. J. DeMarco, F. Gombert, K. Moehle, D. Obrecht, *Drug Discov. Today*, 2008, **13**, 944–951.
- N. Srinivas, P. Jetter, B. J. Ueberbacher, M. Werneburg, K. Zerbe, J. Steinmann, B. Van der Meijden, F. Bernardini, A. Lederer, R. L. A. Dias, P. E. Misson, H. Henze, J. Zumburn, F. O. Gombert, D. Obrecht, P. Hunziker, S. Schauer, U. Ziegler, A. Kach, L. Eberl, K. Riedel, S. J. DeMarco, J. A. Robinson, *Science*, 2010, **327**, 1010–1013.
- D. J. Craik, D. P. Fairlie, S. Liras, D. Price, *Chem. Biol. Drug Des.*, 2013, **81**, 136–147.
- S. E. Gibson, N. Guillo, M. J. Tozer, *Tetrahedron*, 1999, **55**, 585–615.
- I. Berezowska, N. N. Chung, C. Lemieux, B. C. Wilkes, P. W. Schiller, *J. Med. Chem.*, 2009, **52**, 6941–6945.
- X. Yu, P. Talukder, C. Bhattacharya, N. E. Fahmi, J. A. Lines, L. M. Dedkova, J. LaBaer, S. M. Hecht, C. Shengxi, *Bioorg. Med. Chem. Lett.*, 2014, **24**, 5699–5703.
- C. Solanas, B. G. de la Torre, M. Fernandez-Reyes, C. M. Santiveri, M. A. Jimenez, L. Rivas, A. I. Jimenez, D. Andreu, C. Cativiela, *J. Med. Chem.*, 2009, **52**, 664–674.
- N. Turner, *Curr. Opin. Chem. Biol.* 2011, **15**, 234–240.
- M. M. Heberling, B. Wu, S. Bartsch, D. B. Janssen, *Curr. Opin. Chem. Biol.*, 2013, **17**, 250–260.
- L. Poppe, J. Rétey, *Curr. Org. Chem.*, 2003, **7**, 1297–1315.
- L. Poppe, J. Rétey, *Angew. Chem. Int. Ed.*, 2005, **44**, 3668–3688.
- (a) L. Poppe, *Curr. Opin. Chem. Biol.*, 2001, **5**, 512–524; (b) J. Rétey, *Biochim. Biophys. Acta*, 2003, **1647**, 179–184.
- B. de Lange, D. J. Hyett, P. J. D. Maas, D. Mink, F. B. J. van Assema, *ChemCatChem*, 2011, **3**, 289–292.
- A. Gloge, J. Zon, A. Kóvári, L. Poppe, J. Rétey, *Chem. Eur. J.*, 2000, **6**, 3386–3390.
- C. Paizs, A. Katona, J. Rétey, *Chem. Eur. J.*, 2006, **12**, 2739–2744.
- C. Paizs, M. I. Toşa, L. C. Bencze, J. Brem, F. D. Irímie, *Heterocycles*, 2011, **82**, 1217–1228.
- (a) J. H. Bartha-Vári, M. I. Toşa, F. D. Irímie, D. Weiser, Z. Boros, B. G. Vértessy, C. Paizs, L. Poppe, *ChemCatChem*, 2015, **7**, 1122–1128; (b) J. H. Bartha-Vári, L. C. Bencze, E. Bell, L. Poppe, G. Katona, F. D. Irímie, C. Paizs, M. I. Toşa, *Per. Polytechn. Chem. Eng.* 2017, **61(1)**, 59–66.
- F. Ender, D. Weiser, B. Nagy, L. C. Bencze, C. Paizs, P. Pálovics, L. Poppe, *J. Flow Chem.*, 2016, **6**, 43–52.
- D. Weiser, L. C. Bencze, G. Bánóczy, F. Ender, R. Kiss, E. Kókai, A. Szilágyi, B. G. Vértessy, Ö. Farkas, C. Paizs, L. Poppe, *ChemBioChem*, 2015, **16**, 2283–2288.
- H. A. Cooke, C. V. Christianson, S. D. Bruner, *Curr. Opin. Chem. Biol.* 2009, **13**, 460–468.
- S. Strom, U. Wanninayake, N. D. Ratnayake, K. D. Walker, J. H. Geiger, *Angew. Chem. Int. Ed.*, 2012, **51**, 2898–2902.
- G. Wybenga, W. Szymanski, B. Wu, B. L. Feringa, D. B. Janssen, B. W. Dijkstra, *Biochemistry* 2014, **53(19)**, 3187–3198.
- C. V. Christianson, T. J. Montavon, G. M. Festina, H. A. Cooke, B. Shen, S. D. Bruner, *J. Am. Chem. Soc.* 2007, **129**, 15744–15745.
- S. Pilbák, Ö. Farkas, L. Poppe, *Chem. Eur. J.*, 2012, **18**, 7793–7802.
- (a) D. Röther, L. Poppe, G. Morlock, S. Vieregutz, J. Rétey, *Eur. J. Biochem.*, 2002, **269**, 3065–3075; (b) S. Pilbák, A. Tomin, J. Rétey, L. Poppe, *FEBS J.* 2006, **273(5)**, 1004–1019.
- J. D. Hermes, C. A. Roeske, M. H. O’Leary, W. W. Cleland, *Biochemistry*, 1982, **21**, 5106–5114.
- S. L. Lovelock, R. C. Lloyd, N. J. Turner, *Angew. Chem. Int. Ed.*, 2014, **53**, 4652–4656.
- U. Wanninayake, Y. DePorre, M. Ondari, K. D. Walker, *Biochemistry*, 2011, **50**, 10082–10090.
- S. Bartsch, U. T. Bornscheuer, *Prot. Eng. Des. Sel.*, 2010, **23**, 929–933.
- R. Eienthal, M. J. Danson, D. W. Hough, *Trends Biotechnol.* 2007, **25(6)**, 247–249.
- A. Varga, G. Bánóczy, B. Nagy, L. C. Bencze, M. I. Toşa, Á. Gellért, F. D. Irímie, J. Rétey, L. Poppe, *RSC Adv.*, 2016, **6**, 56412–56420.
- B. Schuster, J. Rétey, *Proc. Natl. Acad. Sci. USA*, 1995, **92**, 8433–8437.
- H. Liu, J. H. Naismith, *BMC Biotechnology*, 2008, **8**, 91–101.
- N. A. Dima, A. Filip, A., L. C. Bencze, M. Oláh, P. Sátorhelyi, B. G. Vértessy, L. Poppe, C. Paizs, *Stud. Univ. Babes-Bolyai Ser. Chem.*, 2016, **61**, 21–34.

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