Introduction

Nowadays there is an ever increasing demand for optically pure \( \beta \)-amino acids mainly by the pharmaceutical industry and peptide syntheses. In particular, the biological characteristics of the \( \beta \)-amino acids, along with their use as precursors of various heterocycles and as chiral auxiliaries in enantioselective syntheses have aroused lively interest in their chemistry. This induced rapid development of synthetic procedures for the preparation of enantiopure \( \beta \)-amino acids and their congeners. Until recently most of the biocatalytic approaches for them relied on kinetic resolution with hydrolytic enzymes, such as lipases, acylases and hydantoinases.

An attractive method for the synthesis of enantiopure, non-natural \( \beta \)-amino acids is based on the use of phenylalanine 2,3-aminomutase (PAM). According to their stereochemical preference there are two kinds of phenylalanine 2,3-aminomutases (PAMs), the one of plant origin (EC 5.4.3.10) is producing (\( R \))-\( \beta \)-phenylalanine, while the other one is of bacterial origin (EC 5.4.3.11) and is leading to (\( S \))-\( \beta \)-phenylalanine. Both are members of the class I lyase-like family also including tyrosine 2,3-aminomutase (Tam, EC 5.4.3.6), phenylalanine ammonia-lyase (PAL, EC 4.3.1.24 and EC 4.3.1.25). Tyrosine ammonia-lyase (TAL, EC 4.3.1.23 and EC 4.3.1.25), histidine ammonia-lyase (HAL, EC 4.3.1.3). All of them utilize the same protein-derived prosthetic group, 3,5-dihydro-5-methylidene-4(3H)-imidazol-4-one (MIO) (Fig. 1A), formed autocatalytically in the active site from an XSG motif which is typically Ala–Ser–Gly. Less frequently MIO could be formed from a Thr–Ser–Gly (in PaPAM and in SmPAM), a Ser...
Ser–Gly (in HAL from *Fusobacterium nucleatum*) or a Cys–Ser–Gly (in HAL from *Streptomyces griseus*) motif as well. \(^\text{12}\)

The broad range of aromatic and heteroaromatic amino acids tolerated as substrates by these enzymes was also exploited for the preparation of a wide range of non-natural aryl and heteroaryl \(\alpha\)- and \(\beta\)-amino acids. \(^\text{13}\)

PAM from *Taxus canadensis* (TcPAM) forming \((R)\)-\(\beta\)-phenylalanine was used for the partial biotransformation of \((S)\)-\(\alpha\)-phenylalanine and its derivatives into their \((R)\)-\(\beta\)-isomer, \(^\text{14}\) while the closely related *Taxus chinensis* (TchPAM) as biocatalyst was employed in the enantioselective ammonia addition to \((E)\)-cinnamate producing a mixture of enantiopure \((S)\)-\(\alpha\)- and \((R)\)-\(\beta\)-phenylalanine. \(^\text{15}\)

In a later study, significant shift of the regiosomeric preference towards the \(\beta\)-isomers was achieved by site directed mutagenesis. \(^\text{16}\)

Wanninayake *et al.* have exploited some unnatural amino acids as amino group donors. \(^\text{17}\)

The amino group of these substrates was transferred by TcPAM intermolecularly to another arylacrylate skeleton to form mixtures of \(\alpha\)- and \(\beta\)-arylalanines.

Acting on \(\alpha\)-phenylalanines the \((S)\)-isomer-prefering PAM (EC 5.4.3.11) forms \((S)\)-\(\beta\)-phenylalanine transposing exclusively the amino group of the \((S)\)-\(\alpha\)-phenylalanine to give \((S)\)-\(\beta\)-phenylalanine (Fig. 1B). \(^\text{6}\)

Among the known members of the PAM family producing \((S)\)-\(\beta\)-phenylalanine are AdmH from *Pantoea agglomerans* (PaPAM) \(^\text{6}\) and EncP from *Streptomyces maritimus* (SmPAM). \(^\text{7}\)

These enzymes have important applications in the preparation of the antibiotic andrimid, \(^\text{18}\) further various chiral phenylalanine derivatives \(^\text{19}\) as well as in the synthesis of the anticancer drug Taxol. \(^\text{20}\)

The crystal structure of PaPAM complexed with phenylalanine to its active site, supported a reaction mechanism proceeding through two \(N\)-MIO containing intermediates. \(^\text{6}\)

This was also in agreement with QM/MM calculations on TAL \(^\text{21}\) and PAL \(^\text{22}\) supporting ammonia elimination via an \(N\)-MIO
intermediate and suggesting the formation of similar N-MIO complexes as a common feature of the mechanism for all MIO-enzymes. According to these propositions, the steps starting from either \(\alpha\) or \(\beta\)-phenylalanine are quite similar such as: (i) formation of a covalent enzyme–substrate complex via Michael addition of the amino group of the substrate onto MIO, (ii) ammonia elimination from the covalent N-MIO intermediate resulting in a cinnamate binding intermediate state (Fig. 1A).

The mechanism proceeds further with (iii) ammonia re-addition and (iv) the release of the product. Occasionally, a cinamic acid can appear as a by-product (Fig. 1A), supposedly due to intermittent opening of the Tyr78-containing loop resulting in a leak from the main cycle at the cinnamate binding intermediate state.

SmPAM, described earlier as a lyase, has been shown lately to be closely related to PaPAM (63% overall sequence identity and 76% sequence similarity). More recently, Weise et al. investigated SmPAM in the context of ammonia addition to several aryl-substituted \(E\)-cinnamic acid analogues. They found that SmPAM converted a range of arylacrylates to a mixture of \(S\)-\(\alpha\)- and \(S\)-\(\beta\)-arylalanyls. The enzyme exhibited variable regioselectivity, much affected by ring substituents, but introduction of certain active site mutations could shift regioselectivity in either direction. However, in SmPAM-catalysed isomerization of \(S\)-\(\alpha\-) and \(S\)-\(\beta\-)arylalanyls the enantioselectivity was incomplete in many cases.

Substrate specificity of PaPAM was tested with a wide range of aromatic and heteroaromatic \(S\)-\(\alpha\-)arylalanyls. Electronic and steric effects of substituents at the aromatic ring significantly influenced both catalytic efficiency and the formation of arylacrylates as by-products. It was observed that 3-substituted \(S\)-\(\alpha\-)arylalanyls were transformed faster than the 2- or 4-substituted isomers. To explain these observations, computational analysis of substrate–PaPAM structural interactions was performed by substrate docking studies. Recently it was shown that recombinant whole cell \(E\). coli expressing PaPAM could also produce enantiopure \(S\)-\(\beta\-)arylalanyls from \(S\)-\(\alpha\-)arylalanyls.

Worth noticing, that PaPAM failed to catalyse the transformation of several 2-substituted \((S\)-\(\alpha\))-phenylalanines studied.

### Results and discussion

The time course of the reaction (ESI material) showed that after a relatively short time period (2–6 hours) an equilibrium state is reached in case of both enzymatic reactions, using \(\text{rac}\)-\(\beta\)-phenylalanines (Fig. 1C) or \(\text{rac}\)-\(\alpha\)-phenylalanines (Fig. 1B) as substrates. Therefore the results after 20 h show the final product distribution near to the equilibrium state, after longer reaction times further product formation was not observed.

#### Transformation of \((\pm)-\alpha\)-arylalanyls (\(\text{rac}\)-1a–l) with PaPAM

According to previous results, PaPAM failed to catalyse transformation of 2-substituted \((S\)-\(\alpha\))-phenylalanines bearing large electron withdrawing substituents. Our study starting from \((\pm)-\alpha\)-arylalanyls \(\text{rac}\)-1a–l (Fig. 2A) confirmed the validity of this observation also for the racemic mixtures (Table 1).

In the presence of PaPAM, no product could be detected with \((\pm)-\alpha\)-(2-chlorophenyl)alanine \(\text{rac}\)-1g) or \((\pm)-\alpha\)-(2-nitrophenyl) alanine \(\text{rac}\)-1j), while with \((\pm)-\alpha\)-(2-fluorophenyl)alanine \(\text{rac}\)-1d being substituted with the smallest halogen atom moderate mutase-activity was observed resulting in the formation of enantiopure \((S\)-\(\beta\))-(2-fluorophenyl)alanine [(\(S\)-2d).

Conversion of the \((\pm)-3\)-substituted-\(\alpha\)-phenylalanines \(\text{rac}\)-1c, h, k with PaPAM within 20 h was moderately faster than that of the 2-substituted substrate \(\text{rac}\)-1d, but still slower than with \((\pm)-\alpha\)-phenylalanine \(\text{rac}\)-1a).

Comparison of the conversions of the \((\pm)-4\)-substituted-\(\alpha\)-phenylalanines \(\text{rac}\)-1f, i, l with PaPAM for 20 h indicated that with \((\pm)-4\)-fluoro- \(\text{rac}\)-1f and \((\pm)-4\)-bromo- \(\alpha\)-phenylalanine \(\text{rac}\)-1c about 1.2 times higher conversions were attained than with \((\pm)-4\)-chloro- \(\text{rac}\)-1i or \((\pm)-4\)-nitro- \(\alpha\)-phenylalanine \(\text{rac}\)-1l).

Reactions of \((\pm)-\alpha\)-arylalanine bearing thiophen-2-yl group as a small heteroaromatic moiety \(\text{rac}\)-1b with PaPAM were also investigated. It was found that the conversion of \((\pm)-\alpha\)-

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Fig. 2  PaPAM-catalysed transformation of (A) \((\pm)-\alpha\)-arylalanyls \(\text{rac}\)-1a–l and (B) \((\pm)-\beta\)-arylalanyls \(\text{rac}\)-2a–l.
(thiophen-2-yl)alanine (rac-1b) was similar to (±)-α-phenylalanine (rac-1a).

In an earlier study with (S)-1a, negligible ammonia-lyase activity was predicted for PaPAM at low temperatures (under 30 °C). In another study and also in our present one, however, formation of significant amount of (E)-arylcyclates was observed in many cases (3a, b, d, e, f, h, l in Table 1) indicating substantial lyase-activity of PaPAM. Note that with (±)-4-bromo- (rac-1c), (±)-4-chloro (rac-1i) and (±)-3-nitro-α-phenylalanine (rac-1k) as substrates no such activity was observed. Relatively high amounts of arylcyclates were formed in reactions with substrates bearing the smallest aromatic moieties i.e. with (±)-α-phenylalanine (rac-1a), (±)-α-(thiophen-2-yl)alanine (rac-1b) and (±)-2-fluroo-α-phenylalanine (rac-1d).

Results with the (±)-α-arylcyclates (rac-1a-l) demonstrated that mutase and/or lyase activity of PaPAM was much affected by the nature of the aromatic moiety of the substrates. Importantly, high enantiopurity of the products [>98% ee] for (S)-2a-f, h, i, k, l indicated high enantiopurity of the PaPAM-catalysed isomerization in the α → β-direction. The observed high stereoselectivity of the PaPAM-catalysed isomerization of the (S)-α- and (S)-β-arylcyclates was a major advantage compared to the incomplete stereoselectivity of SmPAM-catalysed isomerizations.

### Transformation of (±)-β-arylcyclates (rac-2a-l) with PaPAM

In order to explore the potential of kinetic resolutions of (±)-α- and β-arylcyclates for the preparation of antipodal products, we extended our study to the reactions of (±)-β-arylcyclates rac-2a-l (Fig. 2B and Table 2). The similar time course profiles of the product formation in PaPAM catalysed reactions from (S)-β-phenylalanine and rac-β-phenylalanine (ESI† material) supported that the unreactive (R)-β-phenylalanine did not act as a significant inhibitor. In contrast to the (±)-2-substituted α-phenylalanines with large substituents (rac-1g, j), which were apparently not accepted as substrates by PaPAM, all the (±)-2-substituted β-phenylalanines in the present study (rac-1d, g, j) were smoothly transformed. On the other hand, sluggish or no conversion was observed with PaPAM using as substrates (±)-3- and (±)-4-substituted β-phenylalanines bearing bulky electron withdrawing substituents (rac-2c, h, i, k, l).

The conversions of (±)-β-(thiophen-2-yl)alanine (rac-2b) and (±)-3-fluroo-β-phenylalanine (rac-2e) with PaPAM were higher than that of (±)-β-phenylalanine (rac-2a). The (±)-4-fluroo-β-phenylalanine (rac-2f) was converted similarly as (±)-β-phenylalanine (rac-2a), while (±)-4-bromo-β-phenylalanine (rac-2c) was transformed to the α-isomer (S)-1c and a small amount of 4-bromocinnamate (3e) at significantly lower conversion than rac-2a. In all cases, except for (±)-2-nitro-β-phenylalanine (rac-2j), reactions catalysed by PaPAM proceeded with the formation of the enantiopure (S)-α-isomer [(S)-3a-g] and some arylaclylate (3a-g). Transformation of (±)-2-nitro-β-phenylalanine (rac-2c) was an exception owing to the absence of the by-product (3j) and incomplete stereoselectivity (ee(S)-3j = 92%).

### Effects of pH and ammonia concentration on the PaPAM-catalysed isomerization of (±)-α- and β-(thiophen-2-yl)alanine (rac-1b and rac-2b)

Prompted by the fact that conversions from (±)-α- and (±)-β-(thiophen-2-yl)alanine (rac-1b and rac-2b) with PaPAM were similar to those from the natural substrates i.e. (±)-α- and β-phenylalanine (rac-1a and rac-2a) but more of the by-product [(E)-3-(thiophen-2-yl)acylate, 3b] was formed (Tables 1 and 2), transformations of these substrates were studied in more detail by varying pH or ammonia concentration. An alteration of pH of the buffer solution in the range of 7–9 [at 100 mM (NH₄)₂CO₃] was indifferent to conversion (data not shown), unlike changing the (NH₄)₂CO₃ concentration in the range of 50–1000 mM (at pH 8) which significantly influenced product compositions (Table 3).

Analysis of the PaPAM-catalysed reactions of (±)-α- (thiophen-2-yl)alanine (rac-1b) at various (NH₄)₂CO₃ concentrations with 20 h incubation showed that increasing (NH₄)₂CO₃ concentration...
concentration above 100 mM resulted in decreasing conversion to both (S)-β-(thiophen-2-yl)alanine and the elimination product. At 1000 mM (NH₄)₂CO₃ concentration both (S)-β-(thiophen-2-yl)alanine and arylacrylate formation dropped to 12%. Interestingly, when starting from (±)-β-(thiophen-2-yl)alanine (rac-2b), the best conversion to (S)-α-(thiophen-2-yl)alanine (S)-1b (27%) was achieved at 100 mM buffer concentration. At higher (NH₄)₂CO₃ concentrations smaller conversions were observed.

Study of the dependence of the PaPAM-catalysed reactions of (±)-α- and β-(thiophen-2-yl)alanine (rac-1b and rac-2b, respectively) on (NH₄)₂CO₃ concentration indicated that elevated ammonia concentrations lowered both the rate of isomerization and ammonia elimination as side reaction in both directions of the reaction.

**Computational modelling of PaPAM-catalysed isomerizations**

To rationalize the effect of substituents of aromatic and heteroaromatic (±)-α- and β-arylalanes (rac-1a-1 and rac-2a-1) on PaPAM-catalysed isomerizations computational and statistical analysis was performed based on modelling and comparison of the N-MIO states (S)-1a-1, k, l, i, MIO and (S)-2a-1, k, l, i, MIO formed from the corresponding (S)-α- and (S)-β-arylalanes. Because of the experimentally observed stereospecificity of isomerizations, in the computations only the (S)-enantiomers were considered. Due to the mixed mechanism indicated by the incomplete enantioslectivity in the PaPAM-catalysed reaction of rac-2j, data calculated for the reactions of (S)-α- and (S)-β-2-nitrophenylalanine ([S]-1j and [S]-2j) were omitted from the comparison of reactivities.

In molecular mechanics the bonded terms measure the strain compared to a hypothetical zero-point energy. Thus potential energies derived from molecular mechanics calculations cannot be compared directly if the molecular structures to be related do not have exactly the same atom connectivity. In this modelling study it was assumed that for all investigated compounds the common alanine part of α- and β-arylalanes underwent in the corresponding N-MIO intermediates similar structural changes[21] (Fig. 1). Therefore, it was possible to compare the difference of the energies calculated for the N-MIO intermediates from the (S)-α- and (S)-β-arylalanines ([S]-1b-i, k, l, MIO and [S]-2b-i, k, l, MIO) relative to the corresponding values obtained for (S)-α- and (S)-β-phenylalanine ([S]-1a-N,MIO and [S]-2a-N,MIO), explicitly: (E(S,S) - E(S,1)) - (E(S,2)-E(S,1)), thus cancelling the hypothetical terms representing the difference in heat of formation for the α- and β-alanine part (Table 4).

Our molecular modelling studies based on an exhaustive search for possible conformations of the covalently bound N-MIO intermediates of the (S)-α- and (S)-β-arylalanines ([S]-1b-i, k, l, MIO and [S]-2b-i, k, l, MIO), indicated that unlike the reactions of the majority of the regioisomers ([S]-1, 2a-c, e, f, h, i, l, see Fig. 3A[20] for [S]-1h-N,MIO and [S]-2h-N,MIO) in certain pairs of isomers ([S]-1, 2d, [S]-1, 2g and [S]-1, 2k) the arrangements of the aromatic ring in the lowest energy conformations of the enzyme-bound substrate complexes were dissimilar and differing by a 180° flip [see Fig. 3B for (S)-1g, S,MIO and (S)-2g,S,MIO]. Assuming that in the N-MIO complexes in the tightly packed active site of PaPAM there is no room for a flip neither of the complete arylacrylate molecule nor of the aromatic ring only, it can be predicted that in such cases one of the two N-MIO intermediate states is not in its lowest energy state. Thus the calculated lowest energy differences in such cases should be higher, corresponding to the allowed arrangements without involving a sterically hindered ring flip (Table 4).

Statistical analysis of the experimental and computational data revealed that compounds [S]-1, 2a-i, k, l could be classified

### Table 3

| Substrate | (NH₄)₂CO₃ (mM) | x₁b | x₂b | x₃b |
|-----------|---------------|-----|-----|-----|
| rac-1b    | 50            | 0.68| 0.17| 0.15|
| rac-1b    | 100           | 0.68| 0.17| 0.15|
| rac-1b    | 200           | 0.72| 0.16| 0.12|
| rac-1b    | 1000          | 0.76| 0.12| 0.12|
| rac-2b    | 50            | 0.19| 0.76| 0.05|
| rac-2b    | 100           | 0.27| 0.69| 0.04|
| rac-2b    | 200           | 0.23| 0.75| 0.02|
| rac-2b    | 1000          | 0.13| 0.84| 0.03|

a x₁b, x₂b, and x₃b represent the relative molar fractions of the reaction components as determined by 1H-NMR measurements.

### Table 4

| Substrates (Ar) | Difference of conversionsᵃ | Categoryᵇ | Energy differenceᶜ (kcal mol⁻¹) |
|----------------|-----------------------------|-----------|-------------------------------|
| (S)-1, 2a (phenyl) | −0.05 | E | 0.0 |
| (S)-1, 2b (thiophen-2-yl) | −0.01 | E | 6.1 |
| (S)-1, 2c (4-bromophenyl) | −0.12 | A | −3.6 |
| (S)-1, 2d (2-fluorophenyl) | 0.19 | B | 1.8 (1.0) |
| (S)-1, 2e (3-fluorophenyl) | 0.09 | E | −4.1 |
| (S)-1, 2f (4-fluorophenyl) | 0.01 | E | 0.2 |
| (S)-1, 2g (2-chlorophenyl) | 0.46 | B | 3.1 (0.6) |
| (S)-1, 2h (3-chlorophenyl) | −0.16 | A | −3.0 |
| (S)-1, 2i (4-chlorophenyl) | −0.10 | A | −2.8 |
| (S)-1, 2j (2-nitrophenyl) | −0.26 | A | −7.2 (2.0) |
| (S)-1, 2k (3-nitrophenyl) | −0.11 | A | −6.7 |

ᵃ Difference of conversions is defined as: (1 - x₂) - (1 - x₁), where x₁ and x₂ are the corresponding molar fraction values listed in Tables 1 and 2. ᵇ Categories based on regioisomeric preference are defined as follows: A: (1 - x₂) - (1 - x₁) < −0.09; B: −0.09 ≤ (1 - x₂) - (1 - x₁) < 0.09; E: (1 - x₂) - (1 - x₁) > 0.09. ᵈ Energy difference of N-MIO states (S)-1b-N,MIO and (S)-2b-N,MIO is defined and normalized for (S)-1a-N,MIO and (S)-2a-N,MIO as: (E(S,S) - E(S,1)) - (E(S,2)-E(S,1)). For compounds with different ring orientations in their lowest energy conformations of the α- and β-N-MIO intermediate states, energy values are selected for structures consistent with the reaction of lower or no conversion. Energy differences pertinent to the other direction are presented in parentheses. ᵇ Due to the mixed mechanism indicated by the incomplete stereoselectivity in the reaction of rac-2j with PaPAM, data for (S)-1g,N,MIO and (S)-2g,N,MIO were not included.
into three categories (A, E and B, in Table 4) based on the difference of conversions [defined as: \(\text{Statistical power than parametric methods but:} \]
\[\text{Conversion} \times (1 - \text{Conversion})\], where \(x_1, x_2\) are the corresponding molar fraction values listed in
\[\text{Tables 1 and 2}\]. The three categories: A: \(1 - x_2 > 1 - x_1 \geq -0.09\); E: \(-0.09 \leq (1 - x_2) - (1 - x_1) \leq 0.09\); B: \(1 - x_2 - (1 - x_1) > 0.09\) could be correlated with the regioisomeric preferences (Table 4, Fig. 4). One-way ANOVA test – treating the categories (Table 4) as the independent and the conversion differences (Tables 1 and 2) as the dependent variables – indicated significant difference between categories A and B at the
\[\alpha = 0.050\] level. In addition, further nonparametric tests – having lower statistical power than parametric methods but without the requirement of normal distribution of data – were
also performed. The Kruskal–Wallis ANOVA and Mann–Whitney U tests resulted only somewhat higher p values (0.100 and 0.053, respectively) than the threshold. Moreover, the median test showed also significant differences between the categories. Overall, the statistical tests supported the finding that categories A and B are truly separate.

The computational study extended with statistical analysis revealed that the cases when one of the regioisomers was converted much faster (or is the only one to be converted) the energetics of the regioisomeric N-MIO-type enzyme–substrate complexes was one of the most important factors governing the outcome of the reaction. In the case of the intermediate group (category E) involving small aromatic moieties contribution of the steric effects are much less pronounced.

The computational results listed in Table 4 suggest that the reactions of 2-substituted \(\alpha\)-phenylalanines [(S)-1d, g and (S)-2d, g: category B in Table 4 and Fig. 4] are strongly disfavoured because the energy calculated for the \(\alpha\)-N-MIO intermediates [(S)-1d, gN-MIO] is much lower than that calculated for the corresponding \(\beta\)-N-MIO intermediates [(S)-2d, gN-MIO]. This was in full agreement with experimental observations indicating slow or no reaction from the \((\pm)\)-2-substituted \(\alpha\)-phenylalanines (rac-1d, g, j) with PaPAM, in contrast to high conversions from 2-substituted \((\pm)\)-\(\beta\)-phenylalanines (rac-2d, g, j) under the same conditions. The situation for the 3- and 4-substituted phenylalanines with bulky substituents [(S)-1c, h, i, k, l and (S)-2c, h, i, k, l: category A in Table 4 and Fig. 4] was the opposite. In good correlation with the experimental regioisomeric preferences, much lower energies were calculated for the \(\beta\)-N-MIO intermediates [(S)-2c, h, i, k, lN-MIO] than for the corresponding \(\alpha\)-N-MIO intermediates [(S)-1c, h, i, k, lN-MIO]. This could explain sluggish or no conversion of \(\beta\)-arylalanines containing bulky substituents at positions 3 or 4 (rac-2c, h, i, k, l) with PaPAM and much higher conversions of the \(\alpha\) compounds (rac-1c, h, i, k, l) under the same conditions. In
addition, the energy differences corresponding to the direction of reaction with higher conversions and assuming a disallowed flip of the aromatic ring (Table 4, values in parentheses) showed remarkable differences compared to the sterically non-restricted values and the hindered flip of the aromatic ring further amplified the difference in reactivity mentioned before. Because of the hindered flip, in case of certain regioisomeric amino acid pairs the aromatic ring in a given N-MIO intermediate adopts different conformation depending on whether the actual N-MIO intermediate is at the substrate side [step (ii) in our postulated mechanism, Fig. 1A] or at the product side [step (iii) in our postulated mechanism, Fig. 1A].

**Conclusion**

According to our results, regioselectivity and activity of PaPAM towards α- and β-arylalanines are mainly influenced by the nature of the aromatic moiety of the substrates. PaPAM catalyses the synthesis of the corresponding (S)-β-enantiomer from racemic 3- and 4-substituted α-phenylalanines more smoothly than that starting from racemic 2-substituted α-phenylalanines, the latter being poor or no substrates of the enzyme. Contrarily, racemic 2-substituted β-phenylalanines were good substrates and provided the (S)-α-enantiomers smoothly while the racemic 3- or 4-substituted β-phenylalanines were almost no substrates. Importantly, in all but one case (for racemic 2-nitro-β-phenylalanine), the isomerizations were stereospecific giving mixture of unreacted (R)-α- or (R)-β-arylanilines with an enantiomeric excess depending on conversion, and enantiopure (S)-α- or β-arylanilines as products, along with various amounts of arylacrylate as by-product. Computational and statistical analysis revealed significant correlation between the energetics of the N-MIO intermediate states forming from the (S)-α- or β-arylanilines and the regioisomeric preferences of PaPAM in case of substrates with bulky electron withdrawing substituents on the aromatic ring. In several cases “hysteresis” was postulated: the conformation (thus energy) of a given regioisomeric N-MIO intermediate formed from the given regioisomeric substrate differed from that conformation which resulted in the given regioisomer as product.

**Experimental**

**Reagents and analytical methods**

The starting materials were purchased from Sigma-Aldrich (St. Louis, MO, USA) or Carl Roth (Karlsruhe, Germany) and used without purification. Solvents were purified and dried by standard methods. The racemic amino acids rac-1a-l rac-2a-l and the α,β-unsaturated acids 3a-l were synthesized using published methods.26

The 1H- and 19F-NMR spectra were recorded at 21 °C on Bruker spectrometers operating at 400 MHz, 101 MHz and 600 MHz, 151 MHz, respectively. Enantiomer separations were obtained on Agilent 1260 HPLC instrument using either Chiralpak® ZWIX+(+) [4 mm × 250 mm] column and a mixture of methanol (containing 100 mmol L⁻¹ formic acid and 50 mmol L⁻¹ diethylamine), acetonitrile and water in proportion of 49 : 49 : 2 (v/v/v) as eluent at a flow rate of 1 mL min⁻¹ or Crownpak® CR-I(+) column (150 × 3.0 mm × 5 µm) and a mixture of HClO₄ solution (3.6 g L⁻¹, pH 1.5): acetonitrile as mobile phase at a flow rate of 0.4 mL min⁻¹. NMR spectra and HPLC data are presented as ESI† material.

**Expression and purification of PaPAM**

The gene of PaPAM from *Pantoea agglomerans* (encoding 541 AAs – Uniprot code: Q84FL5) was optimized to the codon usage of *E. coli*. The 1626 bps long synthetic gene was produced and cloned into pET-19b vector using the *Xhol* and *Bpu1102I* cloning sites. The recombinant PaPAM carrying an N-terminal (His)₅₆α-tag was produced in *E. coli* BL21(DE3)pLysS cells. For the expression step a colony of the transformed plasmid was grown overnight at 37 °C in 5 mL of LB medium containing carbencillin (50 µg mL⁻¹) and chloramphenicol (30 µg mL⁻¹). The overnight culture was added to LB medium (0.5 L) in an Erlenmeyer flask and grown at 37 °C until OD₆₀₀ reached 0.7–0.8. Then the temperature was lowered to 25 °C and the cells were induced by the addition of IPTG (1 mM). The culture was shaken at 220 rpm at 25 °C for 19 h longer. All of the subsequent procedures were carried out in an ice-bath. The cells were harvested by centrifugation (25 min, 5000 × g) and re-suspended in 50 mL of lysis buffer (150 mM NaCl, 50 mM TRIS, pH 7.5) supplemented with DNase, RNase, lysozyme, PMSF (2 mM) and an EDTA-free protease-inhibitor cocktail. The cells were then lysed by sonication and the cell debris was removed by centrifugation (10 000 × g, 30 min).

The proteins were purified on a column filled with nickel-nitrilotriacetic acid agarose gel (Ni-NTA) following the manufacturer’s protocol.27 The expressed protein was eluted from the column with imidazole (500 mM in low salt buffer, pH 7.5). The purity of the protein in the eluted fractions was verified by SDS-PAGE analysis. The protein fractions were dialyzed against phosphate-buffered saline (50 mM) at 4 °C followed by concentration by centrifugal ultrafiltration (using vertically-oriented ultrafiltration membrane VIVASPIN 10 000 MWCO, 5000 × g, 4 °C, to final concentration of 3–5 mg mL⁻¹). The concentration of PaPAM in the final solutions was determined by Bradford’s method.28

**PaPAM-catalysed biotransformations of (±)-α- and β-arylalanines**

Into the solution of the substrate (rac-1a-f, h, i, k, l and rac-2a-g, j, 4 mg) in (NH₄)₂CO₃ buffer (100 mM, pH 8.0, 2 mL), PaPAM (1.6 mg) was added and the reaction mixture was stirred at room temperature for 20 h. For HPLC measurements reaction samples (30 µL) were taken and the enzyme was precipitated with 30 µL MeOH. After filtration, the samples were diluted with the corresponding mobile phase and injected on HPLC.

Prior to 1H- and 19F-NMR investigations the reaction was quenched with methanol, followed by filtration and evaporation of the solvent in vacuum. Deuterium sodium hydroxide (2% NaOD) solution was added and the spectrum (ESI†) was recorded at room temperature.
Biotransformation of (±)-x- or β-(thiophen-2-yl)alanine (rac-1b or rac-2b) under various conditions

pH change in the range of 7–9. Into the solution of the substrate (rac-1b or rac-2b, 4 mg) in (NH₄)₂CO₃ buffer (pH 7.0, 8.0 or 9.0; 100 mM, 2 mL), PaPAM (1.6 mg) was added and the reaction mixture was stirred at room temperature for 20 h. The reaction was stopped by heating at 90 °C for 10 min. The solvent was evaporated, the sample re-dissolved in NaOD solution (2%, 0.5 mL) and analysed by ¹H-NMR (data not shown).

Change of ammonium carbonate concentration in the range of 50–1000 mM. Into the solution of the substrate (rac-1b or rac-2b, 4 mg) in (NH₄)₂CO₃ buffer (50, 100, 200, or 1000 mM; pH 8.0; 2 mL), PaPAM (1.6 mg) was added and the reaction mixture was stirred at room temperature for 20 h. The reaction was stopped by heating at 90 °C for 10 min. Then the solvent was evaporated, the sample re-dissolved in NaOD solution (2%, 0.5 mL) and analysed by ¹H-NMR (see Table 3).

Molecular modelling of the covalent enzyme–substrate N-MIO complexes in PaPAM

The homotetrameric X-ray structure of PaPAM [PDB ID: 3UNV] was completed and adjusted using the Protein Preparation Wizard in four steps: (i) hydrogen atoms were added and bond orders were assigned, (ii) artefacts of the protein crystallization procedure were removed, except for two phosphate ions, (iii) hydrogen bond network, tautomeric states, side chain conformations of selected amino acids and ionization states were determined and optimized corresponding to the experimental assay conditions and (iv) a constrained minimization was performed. Protein pKₐ values were predicted using PROPKA. In the further modelling process for the N-MIO intermediates, in accordance with the proposed mechanism (Fig. 1), Tyr78 was set deprotonated and Tyr320 protonated. Further details on the computational methods and the model will be published in a forthcoming paper.

The refined and completed X-ray structure served as a starting point to create an overall protein model corresponding to the experimental assay conditions. The buffer solution solvated model was created by the Desmond program suite. The PaPAM model was solvated explicitly with water and additional ions were added with respect to the experimental assay conditions. The buffer solvated model was then equilibrated with a slightly modified default equilibration protocol, applying harmonic constraints to the Cartesian coordinates of protein heavy atoms. A spherical model of the active site with a radius of 27 Å and centred on the exocyclic methylene carbon of the MIO prosthetic group of chain C was cut off and capped with acetyl and N-methylamino groups.

N-MIO type covalent complexes of our substrates (S)-1a–l and (S)-2a–l were created by our induced-fit covalent docking protocol. This involved the creation of initial conformations of compounds (S)-1a–l and (S)-2a–l by docking with Glide program suite into a modified and artificially enlarged active site in which the residues Leu216, Ile219, Leu104, Val108, Met84, Leu421, Leu171, and Phe428 were exchanged to Ala residues, further the MIO prosthetic group was reduced to Thr + Gly and three water molecules in the active site were removed.

After having docked into the enlarged active site, all side chains and the MIO group were restored, a covalent bond between the nitrogen atom of the amino group and the exocyclic carbon of MIO was created, the covalently bound ligands and the residues in close proximity to them were minimized and finally, redundant conformations were eliminated with MacroModel. During restoring the mutated side-chains, conformations of several active site residues were predicted with Prime. After replacing the three active site water molecules removed earlier, a final minimization in the 6 Å proximity of the covalently bound ligand using Prime resulted in the final models and energies. OPLS2005 force field was applied in all molecular mechanics calculations and simulations.

Statistical tests (one-way ANOVA, Kruskal–Wallis ANOVA, median test and Mann–Whitney U tests) were carried out and Fig. 4 was created using Statistica. Non-significant Levene and Shapiro–Wilks tests justified the use of one-way ANOVA. The probability value of type I error (α) was chosen to be 0.05 in all the cases.

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