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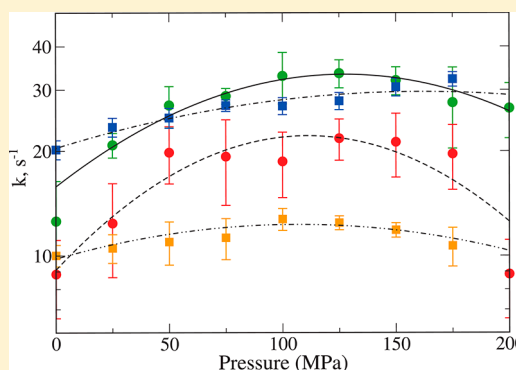
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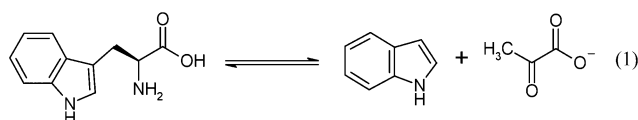
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ABSTRACT: The effects of pH and hydrostatic pressure on the reaction of H463F tryptophan indole-lyase (TIL) have been evaluated. The mutant TIL shows very low activity for elimination of indole but is still competent to form a quinonoid intermediate from L-tryptophan [Phillips, R. S., Johnson, N., and Kamath, A. V. (2002) *Biochemistry* 41, 4012–4019]. Stopped-flow measurements show that the formation of the quinonoid intermediate at 505 nm is affected by pH, with a bell-shaped dependence for the forward rate constant, k_f , and dependence on a single basic group for the reverse rate constant, k_r , with the following values: $pK_{a1} = 8.14 \pm 0.15$, $pK_{a2} = 7.54 \pm 0.15$, $k_{f,min} = 18.1 \pm 1.3 \text{ s}^{-1}$, $k_{f,max} = 179 \pm 46.3 \text{ s}^{-1}$, $k_{r,min} = 11.4 \pm 1.2 \text{ s}^{-1}$, and $k_{r,max} = 33 \pm 1.6 \text{ s}^{-1}$. The pH effects may be due to ionization of Tyr74 as the base and Cys298 as the acid influencing the rate constant for deprotonation. High-pressure stopped-flow measurements were performed at pH 8, which is the optimum for the forward reaction.

The rate constants show an increase with pressure up to 100 MPa and a subsequent decrease above 100 MPa. Fitting the pressure data gives the following values: $k_{f,0} = 15.4 \pm 0.8 \text{ s}^{-1}$, $\Delta V^\ddagger = -29.4 \pm 2.9 \text{ cm}^3 \text{ mol}^{-1}$, and $\Delta\beta^\ddagger = -0.23 \pm 0.03 \text{ cm}^3 \text{ mol}^{-1} \text{ MPa}^{-1}$ for the forward reaction, and $k_{r,0} = 20.7 \pm 0.8 \text{ s}^{-1}$, $\Delta V^\ddagger = -9.6 \pm 2.3 \text{ cm}^3 \text{ mol}^{-1}$, and $\Delta\beta^\ddagger = -0.05 \pm 0.02 \text{ cm}^3 \text{ mol}^{-1} \text{ MPa}^{-1}$ for the reverse reaction. The primary kinetic isotope effect on quinonoid intermediate formation at pH 8 is small (~ 2) and is not significantly pressure-dependent, suggesting that the effect of pressure on k_f may be due to perturbation of an active site preorganization step. The negative activation volume is also consistent with preorganization of the ES complex prior to quinonoid intermediate formation, and the negative compressibility may be due to the effect of pressure on the enzyme conformation. These results support the conclusion that the preorganization of the H463F TIL Trp complex, which is probably dominated by motion of the L-Trp indole moiety of the aldimine complex, contributes to quinonoid intermediate formation.



Tryptophan indole-lyase (TIL, tryptophanase, EC 4.1.99.1) is a pyridoxal 5'-phosphate (PLP) enzyme that catalyzes the reversible hydrolytic β -elimination of L-tryptophan to give indole and ammonium pyruvate (eq 1). This enzyme is widely



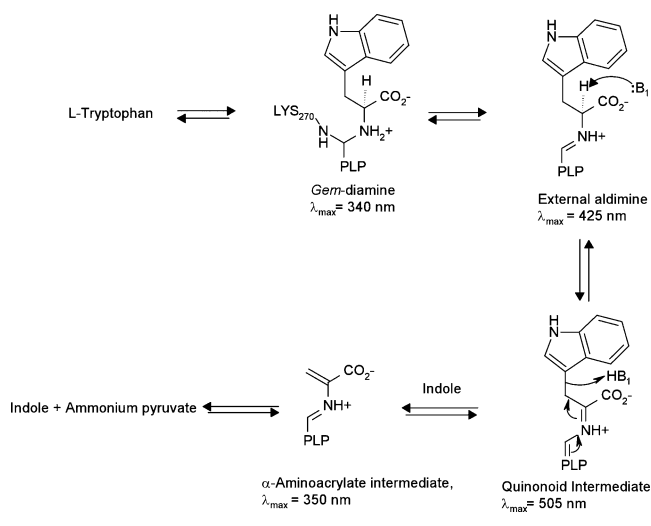
distributed in enterobacteria, particularly in *Escherichia coli*.¹ Although indole was long considered to be simply a waste product of bacterial metabolism of L-Trp, recently it has been discovered that indole is a bacterial signaling molecule that regulates gene expression,² biofilm formation,^{3–5} plasmid stability,⁶ pathogenicity,⁷ and antibiotic resistance.⁸ Thus, because TIL is not found in eukaryotes, the enzyme is a potential target for novel selective antibacterial drugs.

The mechanism of indole elimination catalyzed by TIL is intriguing, because indole is a poor leaving group for a conventional β -elimination. The proposed mechanism involves the initial formation of an external aldimine with the PLP, followed by α -deprotonation to give a quinonoid intermediate (Scheme 1).^{9–11} Subsequently, the transfer of a proton to the indole C₃ occurs, followed by or concerted with, elimination with C _{β} –C₃ bond cleavage.¹¹ In previous studies, we examined the H463F mutant TIL, which was designed to evaluate whether His463 formed a hydrogen bond to the indole NH of the Trp substrate. This mutant enzyme has very low activity for indole elimination (step 4 in Scheme 1) but still can efficiently form a quinonoid intermediate,¹² which becomes kinetically trapped. Therefore, H463F TIL is useful in the study of the TIL mechanism up to formation of the quinonoid complex (steps

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Scheme 1. Mechanism of TIL



1–3 in Scheme 1). In subsequent pressure-jump (*p*-jump) experiments, we found that the rate constant of quinonoid intermediate formation from L-Trp, but not L-Met, with H463F TIL increased with pressure up to 40 MPa (400 bar; 1 bar = 0.987 atm),¹³ the upper pressure limit of the pressure-jump instrument. As this differential pressure response may be of mechanistic origin, we have now used stopped-flow spectrophotometry, which has a much larger experimental pressure range (200 cf. 40 MPa), to further examine the effects of pH and pressure on formation of the H463F-L-Trp quinonoid intermediate from L-Trp and α -[²H]-L-Trp. The results of these experiments are presented herein.

EXPERIMENTAL PROCEDURES

Materials. L-Tryptophan was a product of US Biochemicals. Triethanolamine and other laboratory reagents were obtained from Fisher Scientific. All enzyme solutions were prepared in deionized distilled water.

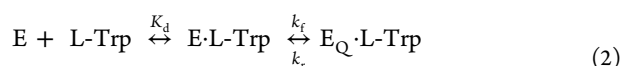
Enzyme. H463F mutant TIL was prepared as described previously.¹² The protein concentration was determined from the absorbance at 280 nm [$A_{280}(1\%) = 9.19$]¹⁴ or using the Bio-Rad Bradford protein dye reagent¹⁵ with wild-type TIL as a standard. The enzyme activity was measured with S-(*o*-nitrophenyl)-L-Cys,¹⁶ which is less sensitive than L-Trp to the effects of mutation in the active site.

α -[²H]-L-Tryptophan. L-Tryptophan (0.162 g) was dissolved in 20 mL of ²H₂O, and 0.4 mL of H463F TIL (16 mg/mL) in aqueous phosphate buffer was added. The reaction mixture was left at room temperature for 7 days, at which time ¹H NMR showed the α -proton was >95% exchanged. The reaction mixture was applied to a column (1.5 cm \times 10 cm) of Dowex-50 (H⁺); the column was washed with water until the washings were neutral, and the amino acid was eluted with 1 M NH₃. The eluate (30 mL) was evaporated in vacuo, and the solid residue was suspended in 95% ethanol and filtered, giving 0.12 g of cream-colored powder.

Instruments. Atmospheric-pressure stopped-flow experiments were performed at ambient temperature (~ 22 – 23 °C) with an Applied Photophysics SX.18MV-R stopped-flow spectrophotometer in single-wavelength mode at 505 nm. The pressure-dependent stopped-flow experiments were performed at 20 °C in a HiTech HP-SF Scientific stopped-flow spectrophotometer (TgK Scientific, Bradford on Avon,

U.K.). The protein was exchanged into 0.1 M triethanolamine hydrochloride (pH 8.0) and 0.1 M KCl, with a PD-10 gel filtration column before the experiments. This buffer was used because triethanolamine has a low ΔpK_a value with pressure ($\Delta V^\ddagger = 4.5 \pm 0.3$ l¹⁷), minimizing any possible change in pH with pressure. For pH dependence, the buffer consisted of 0.1 M triethanolamine phosphate and 0.1 M KCl, adjusted to the desired nominal pH from 6.5 to 9.0 in 0.5 pH unit steps. The enzyme was exchanged with a PD-10 gel filtration column equilibrated with the various buffers immediately prior to the experiments. The actual pH of the buffer after dilution was measured to plot the results in Figure 3.

Data Analysis. The concentration dependence of the stopped-flow data was fit to eq 3, assuming the kinetic model shown in eq 2



$$k_{\text{obs}} = k_f[L\text{-Trp}]/(K_d + [L\text{-Trp}]) + k_r \quad (3)$$

where k_{obs} is the rate constant at any L-Trp concentration, k_f is the rate constant for the forward reaction (α -deprotonation), K_d is the dissociation constant for L-Trp, and k_r is the rate constant for the reverse reaction (α -protonation). The pH dependence data were fit to eq 4 for a bell-shaped pH profile with a limiting minimum value, k_{min} .

$$k_{\text{obs}} = k_{\text{max}}/(1 + 10^{-\text{pH}}/10^{-\text{p}K_{a1}} + 10^{-\text{p}K_{a2}}/10^{-\text{pH}}) + k_{\text{min}} \quad (4)$$

Global fitting was performed with a shared $\text{p}K_{a1}$ value using Origin 8.1 (OriginLab). The pressure dependence data were fit to eq 5

$$k_{\text{obs}} = k_o \exp(-\Delta V^\ddagger p/RT) \exp(\Delta\beta^\ddagger p^2/2RT) \quad (5)$$

where ΔV^\ddagger is the activation volume, $\Delta\beta^\ddagger$ is the transition state compressibility, and R is the ideal gas constant (8.314 cm³ MPa mol⁻¹ K⁻¹).

Modeling. Density function theory (DFT) calculations of the external aldimine and quinonoid intermediates were performed using B3LYP/6-31G(d) with Gaussian 03 revision D.¹⁸ The two models containing the relevant PLP, truncated by removing the PO₄²⁻ moiety, and the Trp moieties were geometry optimized in the gas phase. All models were aligned over the six PLP ring atoms.

RESULTS AND DISCUSSION

pH Dependence of L-Trp Quinonoid Complex Formation. While the rate of turnover of the H463F mutant TIL is much lower than the wild-type rate,¹² this enzyme readily reacts with L-Trp at pH 8 to form a quinonoid complex, with a λ_{max} at 505 nm, that is similar to wild-type TIL (Figure 1A). This quinonoid complex is kinetically trapped because of the relatively rapid α -deprotonation reaction, together with the very slow elimination (steps 1–3 and 4, respectively, in Scheme 1). Fitting the stopped-flow data for the first 200 ms of the reaction to a single exponential gives a reasonable fit (Figure 1B). The apparent rate constant for quinonoid intermediate formation, k_{obs} , exhibits a hyperbolic dependence on L-Trp concentration (Figure 2), indicating that there is a rapid binding equilibrium, followed by a slower step in which the quinonoid species is formed (eq 2). Fitting the concentration dependence data in Figure 2 to eq 3 provides the apparent rate constants at pH 8 for deprotonation ($k_f = 49.4 \pm 2.4 \text{ s}^{-1}$) and reprotonation ($k_r =$

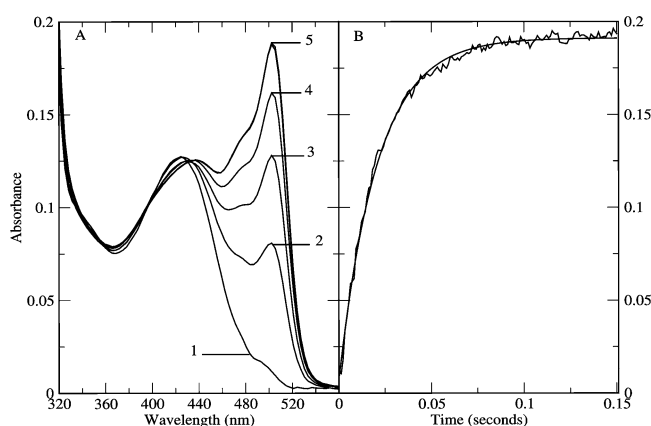


Figure 1. Spectroscopic changes during the reaction of H463F TIL with L-Trp. (A) Reaction of H463F TIL (1 mg/mL) in 0.05 M potassium phosphate (pH 8.0) with 20 mM L-Trp. Scans are shown at 0.3, 10.3, 20.3, 40.3, and 80.3 ms (1–5, respectively). (B) The time course at 504 nm fit to a single-exponential function.

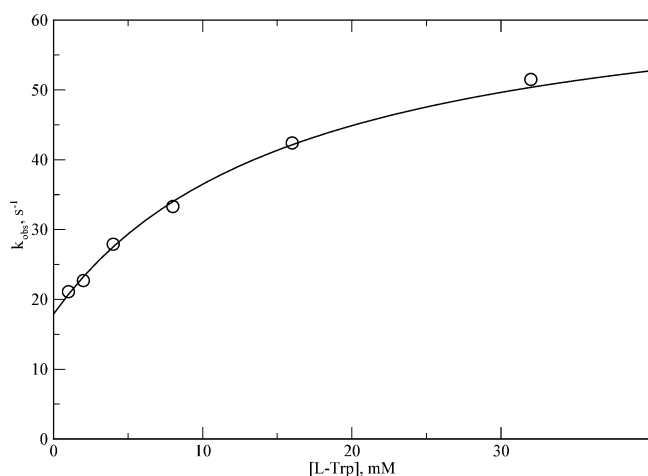


Figure 2. Concentration dependence of quinonoid intermediate formation. The reactions were performed with H463F TIL (1 mg/mL) in 0.1 M triethanolamine phosphate (pH 8.0) and 0.1 M KCl. The line is the result of fitting the data to eq 3.

$17.9 \pm 1.0 \text{ s}^{-1}$). These constants are ~ 1 order of magnitude slower than what is seen for quinonoid intermediate formation from L-Trp with wild-type TIL.^{9–11} The concentration-dependent stopped-flow measurements like that shown in Figure 2 were also performed at pH values ranging from 6.5 to 9.5, and the rate constants found after fitting are pH-dependent, with the results shown in Figure 3. Fitting the data in Figure 3 for both k_f and k_r simultaneously to eq 4 gives the following values: $\text{pK}_{a1} = 8.14 \pm 0.15$, $\text{pK}_{a2} = 7.54 \pm 0.15$, $k_{f,\text{min}} = 18.1 \pm 1.3 \text{ s}^{-1}$, $k_{f,\text{max}} = 179 \pm 46.3 \text{ s}^{-1}$, $k_{r,\text{min}} = 11.4 \pm 1.2 \text{ s}^{-1}$, and $k_{r,\text{max}} = 33 \pm 1.6 \text{ s}^{-1}$. The lines in Figure 3 are the fitted curves using these parameters. Because the rate constant does not decrease logarithmically at high and low pH but rather reaches a minimum, the acid with a pK_a of 8.14 and the base with a pK_a of 7.54 are not the groups directly involved in the deprotonation but rather must be auxiliaries whose ionization states modulate the enzyme reactivity, perhaps by affecting the enzyme conformation. The deprotonation is influenced by both the acidic and basic ionizations, but the reprotonation is influenced by only the base. It is interesting in this regard that the PLP spectrum of wild-type TIL exhibits pH-dependent

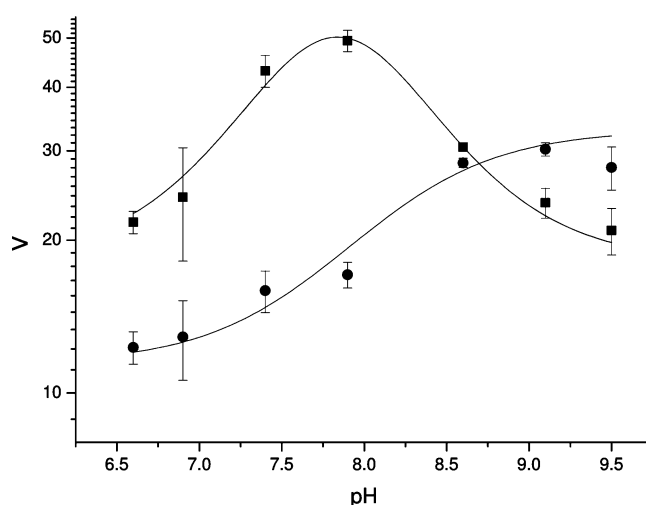


Figure 3. pH dependence of quinonoid intermediate formation: k_f (●) and k_r (■). The reactions were performed as shown in Figure 2, in 0.1 M triethanolamine phosphate and 0.1 M KCl, at pH values ranging from 6.5 to 9.5. The lines are the curves from fitting the data to eq 4.

interconversion of ketoenamine and enolimine tautomers, absorbing at 423 and 338 nm, respectively, showing a pK_a of 7.5–7.8.^{14,19,20} Thus, the acidic group observed in these experiments with a pK_a of 8.14 is possibly Cys298, because mutation of this Cys residue in the active site to Ser affects the PLP spectrum and the pH dependence of the reaction¹⁴ (Scheme 2). The basic group with a pK_a of 7.54 may be the tyrosinate form of Tyr-74, which is likely to be the proton donor to the leaving group for the elimination (Scheme 2), because Y74F mutant TIL exhibits no detectable elimination activity with L-Trp but retains activity with other substrates with good leaving groups.¹² Furthermore, the crystal structures of the paralogous tyrosine phenol-lyase (TPL) F448H mutant with 3-F-Tyr bound show that Tyr71, homologous with Tyr74 in TIL, is located near the γ -carbon of the substrate and forms a hydrogen bond to the active site Lys257 in the quinonoid complex. Although the pK_a of the phenol of free tyrosine in solution is 10.1, an active site tyrosine with a pK_a of 7.5 is not unprecedented. As an example, it has been shown in alanine racemase that a tyrosinate is a catalytic base, with a pK_a value of 7.1–7.4.²¹

Pressure Dependence of L-Trp Quinonoid Complex Formation. Previous p -jump kinetic measurements of the reaction of H463F TIL with L-Trp at pressures of up to 40 MPa found that the reaction showed a significant increase in rate constant under pressure.¹³ However, we were unable to perform p -jumps at higher pressures because of instrumental limitations. Hence, we were interested in using high-pressure stopped-flow spectrophotometry to obtain kinetic data at higher pressures. Because the rate constant and amplitude at 505 nm of quinonoid intermediate formation are optimal at pH 8 (Figure 3), the subsequent pressure-dependent stopped-flow kinetics experiments were performed at pH 8 and 293 K. The pressure dependence was performed at 1.75 and 19 mM L-Trp, as at these concentrations, the observed rate constant, k_{obs} , is approximately equal to k_r and $k_f + k_r$, respectively (Figure 2). Thus, k_f can be approximated by $k_{\text{obs}}(19 \text{ mM}) - k_{\text{obs}}(1.75 \text{ mM})$. The observed rate constants were found to increase with pressure, up to a maximum at ~ 100 MPa, and then decrease (Figure 4A). Thus, we fit the pressure-dependent data with a

Scheme 2. Role of Tyr71 and Cys298 in Quinonoid Intermediate Formation

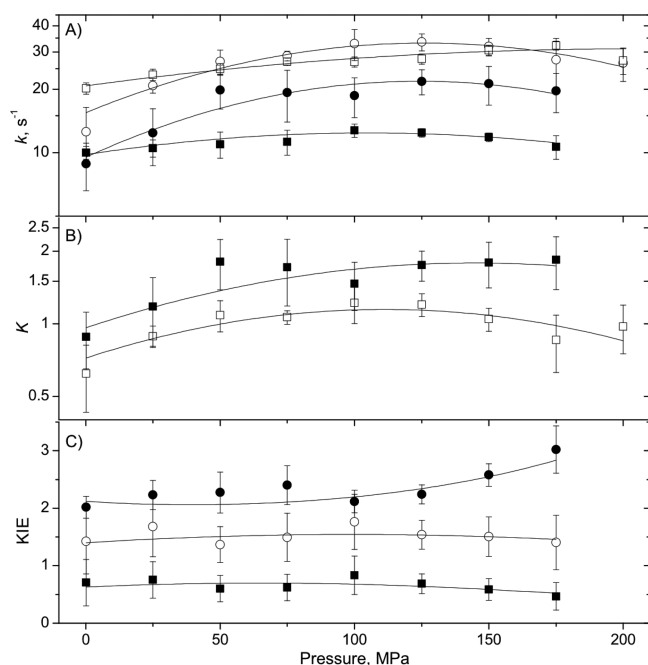
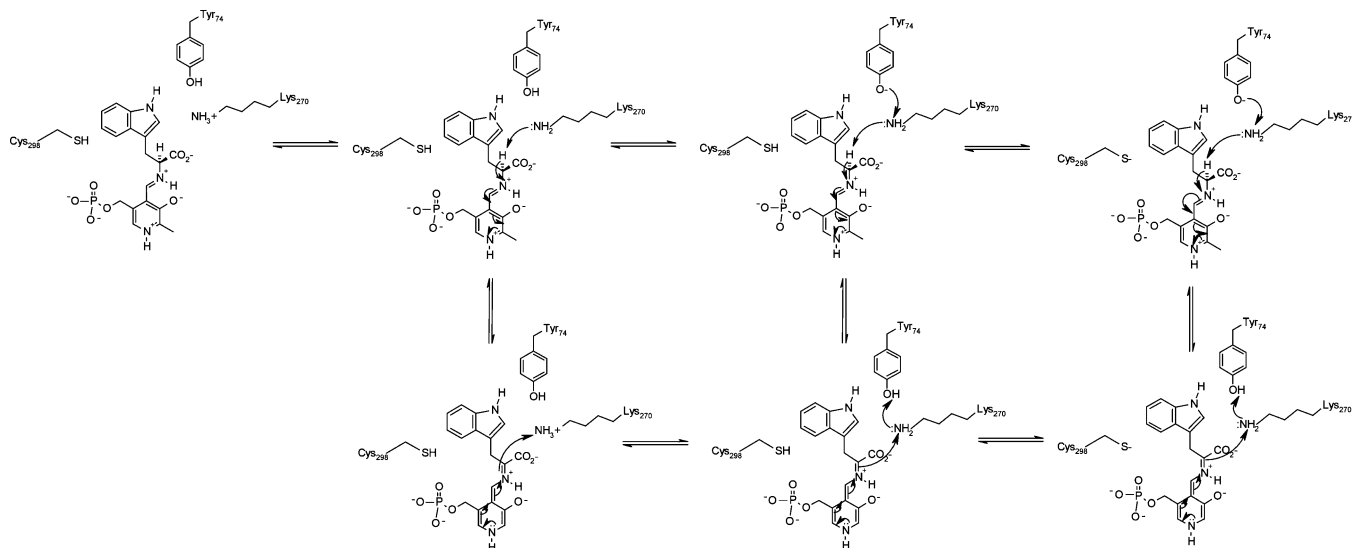


Figure 4. Effect of hydrostatic pressure on the reaction of 1 mg/mL *E. coli* H463F mutant TIL with L-Trp and α -[^2H]-L-Trp at 293 K and pH 8.0. (A) $k_f(\text{H})$ (○), $k_r(\text{H})$ (□), $k_f(\text{D})$ (●), and $k_r(\text{D})$ (■). (B) Equilibrium constant $K = k_f/k_r$ on H (□) and D (■) transfer. (C) KIEs on k_f (○), k_r (●), and K (■). The data are fit to eq 5, and fitted parameters are listed in Table 1.

nonlinear equation, including both the apparent activation volume (ΔV^\ddagger) and compressibility ($\Delta\beta^\ddagger$, eq 5). The results of the fits are shown in Figure 4A, with the parameters listed in Table 1. The initial increase in k_f with pressure is due to a ΔV^\ddagger of $-29.4 \pm 2.9 \text{ cm}^3 \text{ mol}^{-1}$, in good agreement with the previous results of p -jump experiments in the pressure range of 10–40 MPa, where we obtained a ΔV^\ddagger of $-26.5 \text{ cm}^3 \text{ mol}^{-1}$.¹³ It is interesting that the ΔV^\ddagger for k_r of $-11.5 \text{ cm}^3 \text{ mol}^{-1}$ is much smaller than for k_f . Although we have analyzed the data with eq 5 assuming that the increase in the rate constants is due to ΔV^\ddagger , it is also possible that the pK_a values are pressure-dependent. At pressures of >100 MPa, the decrease in k_f is the

result of the $\Delta\beta^\ddagger$ of $-0.23 \text{ cm}^3 \text{ mol}^{-1} \text{ MPa}^{-1}$. As was found with ΔV^\ddagger , $\Delta\beta^\ddagger$ is smaller for k_r (only $-0.05 \text{ cm}^3 \text{ mol}^{-1} \text{ MPa}^{-1}$). We found previously that the binding of Trp may be affected by pressure,¹³ especially at subsaturating concentrations. However, in this work we see a large $\Delta\beta^\ddagger$ even in the 19 mM experiments, which is not likely to be due to effects on binding.

The equilibrium constant ($K = k_f/k_r$) is 0.72 at 1 bar and increases with pressure, with a ΔV of $-20.4 \text{ cm}^3 \text{ mol}^{-1}$ and a $\Delta\beta$ of $-0.18 \pm 0.05 \text{ cm}^3 \text{ mol}^{-1} \text{ MPa}^{-1}$ (Figure 4B and Table 1). In this case, the ΔV describes the reduction in the reaction volume between the L-Trp external aldimine and the quinonoid intermediate, while the negative compressibility indicates that the quinonoid intermediate is also less compressible than the L-Trp external aldimine. The smaller volume and reduced compressibility of the quinonoid intermediate are likely due to the formation of a closed (active) conformation of the enzyme, as was recently observed in crystal structures of quinonoid complexes of the paralogous enzyme, TPL, while the aldimines are in more open conformations.^{22,23}

To perform KIE experiments, H463F TIL was used to prepare α -[^2H]-L-Trp by exchange with L-Trp in D_2O (see Experimental Procedures). When the stopped-flow reactions were performed with α -[^2H]-L-Trp, small but significant kinetic isotope effects on the rate constants are observed [$\text{KIE}_f = 1.6 \pm 0.3$, and $\text{KIE}_r = 2.1 \pm 0.1$ (Figure 4B and Table 1)]. It is interesting that the KIE is larger for k_r than for k_f which implies a significant amount of internal return of the α -proton, despite the ease of H–D exchange during extended incubation. While there are significant KIEs on k_f and k_r (and on K), there are no significant isotope effects on either ΔV^\ddagger or $\Delta\beta^\ddagger$ (Table 1); i.e., the KIEs are not significantly pressure-dependent. Hydrogen transfer reactions involving a significant degree of nuclear quantum mechanical tunnelling (NQMT) (of the transferred H) have been shown to exhibit pressure dependencies of both the observed rate and KIE.^{24,25} Conversely, KIEs arising from semiclassical mechanisms (no significant NQMT) are not expected to be pressure-dependent as these KIEs arise from differences in the vibrational frequencies of the transferred atoms, which have been shown to be relatively insensitive to pressures of a few kilobars.²⁶ While we cannot rule out the involvement of NQMT during enzyme-catalyzed hydrogen transfer on the basis of a lack of a pressure-dependent KIE,^{26–28}

Table 1. Apparent Activation Volumes and Compressibilities for α -Deprotonation of L-Trp by *E. coli* H463F Mutant TIL^a

	k_i	k_r	$K = k_i/k_r$
k_0^H (s ⁻¹)	15.4 ± 0.8	20.7 ± 0.8	0.72 ± 0.06
k_0^D (s ⁻¹)	9.5 ± 1.0	9.8 ± 0.3	0.96 ± 0.12
KIE ₀	1.6 ± 0.3	2.1 ± 0.1	0.75 ± 0.15
$\Delta V^{\ddagger H}$ (cm ³ mol ⁻¹)	-29.4 ± 2.9	-9.6 ± 2.3	-20.4 ± 4.3
$\Delta V^{\ddagger D}$ (cm ³ mol ⁻¹)	-32.6 ± 6.4	-11.1 ± 2.0	-20.9 ± 7.6
$\Delta\Delta V^{\ddagger}$ (cm ³ mol ⁻¹)	3.2 ± 9.3	1.3 ± 4.3	0.5 ± 11.9
$\Delta\beta^{\ddagger H}$ (cm ³ mol ⁻¹ MPa ⁻¹)	-0.23 ± 0.03	-0.05 ± 0.02	-0.18 ± 0.04
$\Delta\beta^{\ddagger D}$ (cm ³ mol ⁻¹ MPa ⁻¹)	-0.26 ± 0.07	-0.11 ± 0.02	-0.14 ± 0.09
$\Delta\Delta\beta^{\ddagger}$ (cm ³ mol ⁻¹ MPa ⁻¹)	+0.03 ± 0.10	+0.06 ± 0.05	-0.04 ± 0.13

^aValues were obtained by fitting the data in Figure 4 to eq 5. KIE values were determined as follows: $\Delta\Delta V^{\ddagger} = \Delta V^{\ddagger H} - \Delta V^{\ddagger D}$, and $\Delta\Delta\beta^{\ddagger} = \Delta\beta^{\ddagger H} - \Delta\beta^{\ddagger D}$. K values were determined by fitting the data in Figure 4B to eq 5.

the pressure dependence of k_i and k_r and the pressure independence of the KIE suggest that either the reaction does not employ significant quantum mechanical tunneling or the effect of pressure on the observed rate constants is due to pressure perturbing a step that is kinetically coupled to, but independent of, the reaction, e.g., active site closure and/or active site preorganization.

Implications for the Enzyme Mechanism. The proposed mechanism of TIL is shown in Scheme 1. L-Trp reacts rapidly, within the dead time of the stopped-flow instrument, with the internal aldimine to form an external aldimine. Subsequent deprotonation of the α -CH of the external aldimine can then occur to form the resonance-stabilized quinonoid carbanion. Elimination of indole had been thought to occur from the quinonoid by initial transfer of a proton to C₃ to give an indolenine species, followed by C _{β} -C₃ bond cleavage. However, multiple-isotope effect experiments suggest that the proton transfer to C₃ and C _{β} -C₃ bond cleavage may be concerted.¹¹ H463F TIL has very low elimination activity (<0.1% of that of wild-type TIL) because His463 has been proposed to interact with the substrate indole NH as a base or by hydrogen bonding to facilitate the transfer of a proton to C₃.¹² However, on the basis of the recent structures of the paralogous enzyme, TPL, that show substrate strain,²¹ it is possible that His463 forms a hydrogen bond with the substrate NH to introduce strain that is necessary for efficient catalysis. His463 is obviously not required for quinonoid intermediate formation (Figure 1), although the rate constant is reduced in H463F by ~1 order of magnitude compared with that of wild-type TIL.¹² Thus, it is possible that preorganization of the enzyme–substrate complex contributes to quinonoid intermediate formation. The stereochemical course of the elimination of indole is most likely *anti*, as was found in the quinonoid structures of Y71F and F448H mutant TPL with 3-fluoro-L-tyrosine,²² so the indole ring must be situated with the C _{β} -C₃ bond antiperiplanar to the α -C-H bond when deprotonation occurs. In addition, the indole ring should be oriented approximately perpendicular relative to the C _{α} -C _{β} bond axis for elimination. Finally, the β -carbon will be moving into the plane of the PLP π -system during quinonoid intermediate formation, and the indole ring will be in motion during deprotonation (Figure 5A). Thus, the positioning of the indole ring into the correct geometry for the elimination to occur will lower the activation energy barrier for quinonoid intermediate formation by reducing the required level of motion. Two stable conformations of both the external aldimine and quinonoid structures were found, and both external aldimine conformers are compared in Figure 5B. Only

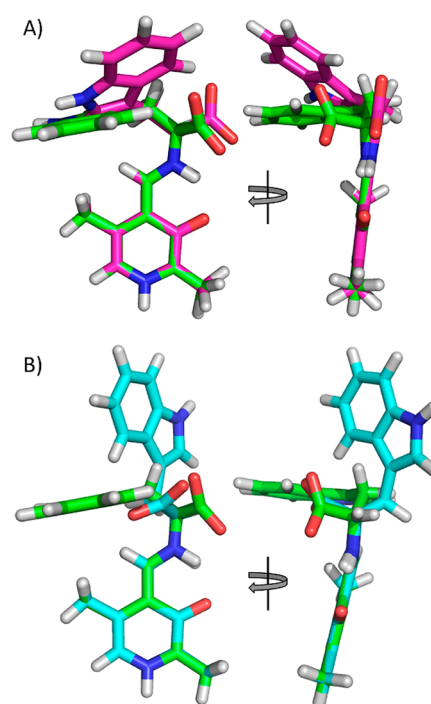


Figure 5. Active site preorganization. Orthogonal projections of models of (A) compact conformers of the aldimine intermediate (green carbons) and quinonoid intermediate (magenta carbons) and (B) comparison of the compact and extended conformers of the aldimine intermediate.

the compact quinonoid conformer is compatible with subsequent proton transfer, which has been proposed to occur to the *re*-face of the indole group.²⁹ Pressure could enhance the rate of reaction in a number of ways. By straining the external aldimine intermediate to form a more quinonoid-like geometry, e.g., by pulling the carboxylate into the plane of the PLP, or by increasing the angle between the PLP and indole rings (Figure 5A), we can enhance the reaction rate due to ground state destabilization. Alternatively, as the external aldimine may adopt multiple conformational states (Figure 5B), if pressure favors the more reactive conformer (in this case, the “compact” form), then the observed rate of reaction will increase with pressure. In this case, the rate enhancement occurs because of preorganization of the reactant state, rather than through direct perturbation of the chemical step. This preorganization of the enzyme–substrate complex results in a negative activation volume and, thus, the observed increase in rate constant with increasing pressure up to ~100 MPa. In

contrast, a competitive inhibitor, L-Met, was found in previous experiments to show no change in rate constant for quinonoid intermediate formation with pressures of up to 40 MPa with either wild-type or H463F TIL,¹³ indicating that preorganization is not significant in this case. However, as the proposed preorganization is dominated by movement of the L-Trp indole moiety (Figure 5), the lack of preorganization of L-Met is not unexpected as the S-CH₃ moiety of L-Met is much smaller than the L-Trp indole. Together, these results support the conclusion that the preorganization of the H463F TIL Trp complex, which is probably dominated by motion of the L-Trp indole moiety of the aldimine complex, contributes to quinonoid intermediate formation. Quinonoid intermediate formation may be gated by preorganization of the external aldimine, and thus, the enzyme is under a degree of dynamic control.

CONCLUSIONS

The formation of the quinonoid intermediate from L-Trp by H463F TIL is influenced by pH, with pK_a values of 7.54 and 8.14. These ionizations are not due to the active site base responsible for deprotonation but probably influence the reaction by changing the enzyme conformation. The rate constant for quinonoid intermediate formation is influenced by pressure, with an initial increase up to 100 MPa, followed by a decrease. These results are consistent with a significant preorganization of the external aldimine complex before deprotonation. The small and pressure-independent KIE seen with α -[²H]-L-Trp suggests that preorganization and/or conformational changes are rate-determining in quinonoid intermediate formation.

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Notes

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ABBREVIATIONS

TIL, tryptophan indole-lyase or tryptophanase (EC 4.1.99.1); TPL, tyrosine phenol-lyase (EC 4.1.99.2); PLP, pyridoxal 5'-phosphate.

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