Aminophosphonate Inhibitors of Dialkylglycine Decarboxylase: Structural Basis for Slow Binding Inhibition†‡

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ABSTRACT: The kinetics of inhibition of dialkylglycine decarboxylase by five aminophosphonate inhibitors are presented. Two of these [(R)-1-amino-1-methylpropanephosphonate and (S)-1-aminoethanephosphonate] are slow binding inhibitors. The inhibitors follow a mechanism in which a weak complex is rapidly formed, followed by slow isomerization to the tight complex. Here, the tight complexes are bound 10-fold more tightly than the weak, initial complexes. The slow onset inhibition occurs with half-lives of 1.3 and 0.55 min at saturating inhibitor concentrations for the AMPP and S-AEP inhibitors, respectively, while dissociation of these inhibitor complexes occurs with half-lives of 13 and 4.6 min, respectively. The X-ray structures of four of the inhibitors in complex with dialkylglycine decarboxylase have been determined to resolutions ranging from 2.6 to 2.0 Å, and refined to R-factors of 14.5—19.5%. These structures show variation in the active site structure with inhibitor side chain size and slow binding character. It is proposed that the slow binding behavior originates in an isomerization from an initial complex in which the PLP pyridine nitrogen—D243 OD2 distance is ~2.9 Å to one in which it is ~2.7 Å. The angles that the C—P bonds make with the p orbitals of the aldimine π system are correlated with the reactivities of the analogous amino acid substrates, suggesting a role for stereoelectronic effects in Schiff base reactivity.

Pyridoxal phosphate (PLP)† dependent enzymes are ubiquitous in the nitrogen metabolism of all organisms, catalyzing a wide variety of reactions at the α-, β-, and γ-carbons of amine and amino acid substrates (I—3). One group of these catalyzes α-decarboxylation of amino acids to yield amine products formed by replacement of the substrate carboxylate with a proton. Another class catalyzes the transfer of amino groups between amines or amino acids and α-keto acids, in a reaction known as transamination. Dialkylglycine decarboxylase is an unusual enzyme that combines both decarboxylation and transamination half-reactions in its normal catalytic cycle (Scheme 1). This unusual combination requires that the decarboxylation reaction proceeds with transfer of the substrate amino group to the coenzyme to form the PMP enzyme as an intermediate in the catalytic cycle. This is effectively an oxidative decarboxylation with respect to the amino acid substrate, and is accomplished by

![Scheme 1: Half-Reactions Catalyzed by Dialkylglycine Decarboxylase](image)

specific protonation of coenzyme C4′ instead of replacing the substrate carboxylate with a proton. Zhou et al. (4) demonstrated that the C—C bond breaking at Cα and the C—H bond making at C4′ occur simultaneously via a concerted transition state, thus achieving extraordinarily high specificity for oxidative decarboxylation.

In 1966, Dunathan (5) proposed that PLP dependent enzymes can control reaction specificity through stereoelectronic effects. This would occur by aligning the scissile bond such that it is parallel to the p orbitals of the conjugated π system, thereby giving maximal transition state orbital overlap and maximal resonance stabilization. The unusual reaction specificity of DGD allowed Sun et al. (6) to use alternate substrates to demonstrate qualitatively the role of stereoelectronic effects in control of reaction specificity. This analysis employed a functional model of the active site with three binding subsites whose locations and specificities have been described (6, 7). These are subsite A, near Q52 and K272, which is the locus of bond making and breaking; subsite B, near R406 and S215; and subsite C, near M141. Two carboxylate-containing inhibitor structures were previously reported (8), showing the carboxylate groups bound
in subsite B. The phosphonate groups of the aminophosphonate inhibitors employed here were expected to bind in subsite A, where decarboxylation of α-amino acid substrates occurs. This would have provided a clear picture of the interactions made by substrate carboxylates in their catalytically productive conformation. It is found that the phosphonate groups are bound in a location between what are considered the A and B subsites.

The kinetics of inhibition by the aminophosphonate inhibitors are also presented. Two of the five that were analyzed are slow binding inhibitors. These act via a weak enzyme— inhibitor complex, followed by slow isomerization to the final, tight complex. The dissociation constants for the initial weak complexes of the tight binding and for the complexes of the normally behaving inhibitors are close to the values of the corresponding amino acid substrates. The slow binding behavior is proposed to arise from an isomerization of the PLP— inhibitor aldimine geometry from an initial state in which the PLP pyridine nitrogen—D243 OD2 distance is ~2.9 Å to one in which it is shortened to ~2.7 Å.

**EXPERIMENTAL PROCEDURES**

**Materials.** Succinic acid and TEA were purchased from Fisher. PLP was obtained from Aldrich. NADPH was from Roche. AIB, pyruvic acid, 2°ADH, and R- and S-AEP were purchased from Aldrich. PEG4000 and sodium pyruvate were from Fluka. MES was from Research Organics. AMPP, ACPP, and AEP (Figure 1) were synthesized according to a literature procedure (9). The racemic mixtures of AMPP and AEP were used without resolution, and the reported \( K_i \) values are the observed values divided by 2, since no evidence for binding of both isomers was observed.

**Steady State Kinetic Assay of DGD.** AIB decarboxylation was followed by coupling acetone produced from AIB to the 2°-ADH reaction as described previously (6). A Kontron UVIKON 9420 spectrometer was employed to monitor the decrease in NADPH absorbance at 340 nm that occurs during reduction of acetone to 2-propanol. Pyruvate was used as the α-keto acid substrate for transamination in all assays.

**Determiner of \( K_i \).** AMPP and S-AEP are slow binding inhibitors of DGD. The overall dissociation constant, \( K_i \) (10), for these inhibitors was measured by preincubation of DGD in a 1 mM solution containing 100 mM TEA—succinate (pH 7.8), 100 mM dipotassium succinate, 2 mM sodium pyruvate, 1 unit/mL 2°-ADH, 50 mM PLP, 300 μM NADPH, and various inhibitor concentrations for 30 min. This preincubated solution was then mixed with 20 μL of 1 M

**Inhibition Kinetics.** Initial rates for six concentrations of 2-aminoisobutyrate at each of four concentrations of AMPP were measured to characterize the inhibition mechanism. AMPP is a slow binding inhibitor, as described below. Characterization of the inhibition mechanism employed the initial rates measured before the onset of slow binding inhibition. The Lineweaver—Burke plot for these data is presented in Figure 2, and clearly indicates competitive inhibition. The data were fitted to eq 1, which describes competitive inhibition, to obtain the \( K_i \) value.

\[
v_i = \frac{V_{\text{max}}[\text{AIB}]}{K_M + \frac{[\text{I}]}{K_i} + [\text{AIB}]} \tag{1}
\]

Here, \( v_i \) is the initial rate, \( V_{\text{max}} \) the maximal rate, \( K_M \) the Michaelis constant, \( I \) the AMPP inhibitor, and \( K_i \) the inhibition constant.
AIB, and the decrease in NADPH absorbance at 340 nm was monitored. This preincubation time was determined to be sufficient for full equilibration of inhibitor binding; thus, the measured rates reflect the slow binding inhibition of DGD. The value of $K_i^*$ was determined by fitting initial rates to eq 2:

$$v_i = \frac{V_{max}[AIB]}{K_M(1 + [I]/K_i) + [AIB]} \left(1 + \alpha [I]/K_i^*\right)$$  \hspace{1cm} (2)

where $K_i^*$ is the slow binding dissociation constant and $\alpha$ is the ratio of the final reaction volume to the volume of the preincubation mixture (i.e., 0.98).

**Measurement of Slow Binding Isomerization Rate Constants.** Scheme 2 shows the kinetic mechanism for competitive inhibition with slow binding of the inhibitor. To measure $k_2$, DGD was incubated with 50 mM TEA-succinate (pH 7.8), 50 mM dipotassium succinate, 20 μM PLP, and 8 mM AMPP or 2 mM S-AEP for 1 h. Then, 20 μL of this preincubated DGD solution was mixed with 1 mL of a cocktail containing 100 mM TEA-succinate (pH 7.8), 100 mM dipotassium succinate, 2 mM sodium pyruvate, 1 unit/mL 2°ADH, 50 μM PLP, 300 μM NADPH, and 25, 50, or 100 mM AIB, and the reactions were monitored at 340 nm. The ratio of the final inhibitor concentration to that of AIB was very small, thus enabling simple measurement of $k_2$ in Scheme 2 by enforcing irreversible conditions on the dissociation of the inhibitor. The reaction curves were fitted to eq 3

$$A_{340} = A_{340}^0 - \left[v_f + \left(\frac{v_i - v_f}{k}\right)(1 - e^{-kt})\right]$$  \hspace{1cm} (3)

where $A_{340}$ is the NADPH absorbance at 340 nm, $v_i$ the initial reaction rate, $v_f$ the final reaction rate, and $k$ the apparent isomerization rate constant. The values of $k_1$ were determined using eq 4, and the values of $k_2$, $K_i$, and $K_i^*$ as determined above.

$$K_i^* = K_i \left(\frac{k_2}{k_1 + k_2}\right)$$  \hspace{1cm} (4)

**Spectral Kinetic Analysis of Isomerization of DGD–AMPP Complexes.** The spectral changes in the coenzyme on reaction of DGD-PLP with AMPP were used to determine independently the values of $k_1$ and $k_2$ for AMPP. DGD-PLP, 4 mg/mL in 100 mM TEA-succinate, 100 mM dipotassium succinate, and 5 μM PLP, was mixed with various concentra-

**Crystallization, X-ray Diffraction Data Collection, and Structural Refinements.** Crystals of DGD were obtained by
the hanging drop vapor diffusion method using a slight modification of the protocol described in ref 11. Drops of 1 

μL containing 30 mg/mL protein, 5 mM potassium phosphate (pH 7.5), 20 μM PLP, and 0.2% (w/v) sodium azide were mixed with 1 μL of reservoir solution containing 15% PEG4000 (w/v), 15 mM MES-KOH (pH 6.4), 75–200 mM sodium pyruvate, and 20 μM PLP, and equilibrated against 1 mL of this reservoir solution. Crystals generally appeared overnight at room temperature, and grew to a maximum size of 0.3 mm × 0.3 mm × 0.5 mm after 1–2 weeks. During the crystallization, microseeding was added. Inhibitors were diffused into DGD crystals by soaking crystals in a stabilizing solution containing inhibitor. For AMPP, the stabilizing solution contained 30% PEG4000, 15 mM MES-KOH (pH 6.4), 20 μM PLP, and 50 mM AMPP. The crystals were soaked in this stabilizing solution for 2 days at room temperature. Since MES is also a ligand of DGD (7, 12), it competes with aminophosphonate binding to DGD. Thus, ACPP or AEP was diffused into crystals soaked in a stabilizing solution without MES. These solutions contained 30% PEG4000, 40 mM ACPP (pH 6.4) or 40 mM AEP (pH 6.4), 50 mM KCl, and 20 μM PLP. Here, ACPP and AEP were used as buffers since their pKa values are appropriate. Both isomers of AEP were diffused into DGD crystals by using a stabilizing solution containing 30% PEG4000, 15 mM Bis-Tris-propane-HCl (pH 6.4), 30 mM KCl, 20 μM PLP, and 20 mM S-AEP or 20 mM R-AEP.

Diffraction data for the DGD–AMPP complex were collected at room temperature, while data for the DGD–ACPP and DGD–AEP complexes were collected at 100 K on an RAXIS IV image plate detector using Cu Kα radiation. DGD–R-AEP data were collected on a Bruker CCD detector at 100 K. High-resolution data for the DGD–S-AEP complex were collected at 100 K using synchrotron beamline 9-1 at Stanford Synchrotron Radiation Laboratory at a wavelength of 0.98 Å. Crystals used at 100 K were harvested from the stabilizing solution described above additionally containing 26% glycerol and flash-frozen. Reflection intensities were integrated with DENZO and merged using SCALEPACK. The merged data were reduced to structure factors using the combination of SCALEPACK2MTZ, CAD, and TRUNCATE programs from the CCP4 program suite. The structures of DGD inhibitor complexes were determined simply by using the DGD-K structure (PLP, MES ligand, and water molecules removed) as an initial model for simulated annealing using CNS, since the space group and unit cell parameters were unchanged. During refinement, the program O was used for model building and electron density interpretation. The final refined structures from CNS were further refined using the TNT refinement package. Only water molecules with B-factors of <85 Å² were retained during the course of the refinement. Data collection and refinement statistics are presented in Table 2.

RESULTS AND DISCUSSION

The aminophosphonates used in this study (Figure 1) are structural analogues of the natural substrates of DGD (2,2-dialkylglycines), in which the reactive carboxylate is substituted with a phosphonate group that is chemically stable in the active site. This allows the determination of the structures of these substrate analogues as Schiff bases with PLP, providing approximations to the structures of the

Table 1: Inhibition Constants for Aminophosphonate Inhibitors

<table>
<thead>
<tr>
<th>inhibitor</th>
<th>homologous substrate</th>
<th>k_diss (s⁻¹)</th>
<th>K_a (mM)</th>
<th>K_i (mM)</th>
<th>K_i⁺ (mM)</th>
<th>k_1 (×10³ s⁻¹)</th>
<th>k_2 (×10³ s⁻¹)</th>
</tr>
</thead>
<tbody>
<tr>
<td>AEP</td>
<td>L-phenylglycine</td>
<td>0.20⁺(0.03)</td>
<td>8.3⁺(2.2)</td>
<td>13⁺(3)</td>
<td></td>
<td></td>
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<tr>
<td>ACPP</td>
<td>aminocyclopentanolate</td>
<td>0.98⁺(0.03)</td>
<td>4.5⁺(0.4)</td>
<td>6.0⁺(0.8)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>AMPP</td>
<td>l-isoleucine</td>
<td>1.0⁺(0.04)</td>
<td>4.9⁺(0.8)</td>
<td>0.5⁺(0.06)</td>
<td>7.9⁺(0.9)</td>
<td>6.0⁺(0.6)</td>
<td>1.3⁺(0.3)</td>
</tr>
<tr>
<td>R-AEP</td>
<td>l-alanine</td>
<td>0.03⁺⁺(0.003)</td>
<td>7⁺(1)</td>
<td>19⁺(1)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>S-AEP</td>
<td>d-alanine</td>
<td>0.20⁺⁺(0.02)</td>
<td>7.1⁺⁺(1.5)</td>
<td>1.1⁺⁺(0.1)</td>
<td>0.13⁺⁺(0.01)</td>
<td>19⁺⁺(1.4)</td>
<td>2.5⁺⁺(0.2)</td>
</tr>
</tbody>
</table>

a k_diss and K_a of homologous substrates are from ref 6 and are for the decarboxylation of the specified substrates. b k_diss and K_a of l-isoleucine were calculated from data in refs 6 and 25. c Conditions: 100 mM TEA-succinate, 100 mM dipotassium succinate, 50 μM PLP, 2 mM AIB, 2 mM sodium pyruvate, 300 μM NADPH, 1 unit/mL 2⁺⁺ADH, 0.1 mM DGD, and 0–20 mM inhibitors. d Conditions: same as footnote c except that inhibitors were preincubated with DGD for 30 min. e k_i is the rate constant observed for the restoration of activity from the DGD–AMPP and DGD–S-AEP complexes. Reaction conditions: 100 mM TEA-succinate, 100 mM dipotassium succinate, 50 μM PLP, 2 mM sodium pyruvate, 300 μM NADPH, 1 unit/mL 2⁺⁺ADH, 50, 100, or 200 mM AIB, 0.1 mM DGD–AMPP or DGD–S-AEP complex, and 0.16 mM AMPP or 0.04 mM S-AEP. Slow binding complexes were prepared by preincubating with AMPP or S-AEP for 30 min. Preincubation conditions: 5 μM DGD, 50 mM TEA-succinate, 50 mM dipotassium succinate, 8 mM AMPP or 2 mM S-AEP, and 50 mM PLP. k_1 was calculated from the equation K_i⁺ = (KK_i)(k_1 + k_2). f k_1 and k_2 were obtained from spectral kinetic data for the reaction between DGD-PLP and AMPP. Reaction conditions: 2 mM DGD-PLP, 50 mM PLP, and 0–35 mM AMPP. g Structurally, this is the closest α-aminocarboxylic acid substrate for which kinetic data are available.

Table 2: Data Collection and Refinement Statistics

<table>
<thead>
<tr>
<th></th>
<th>AMPP</th>
<th>ACPP</th>
<th>AEP</th>
<th>S-AEP</th>
</tr>
</thead>
<tbody>
<tr>
<td>unit cell dimensions (Å) (a, b, c)</td>
<td>152.8, 86.4</td>
<td>151.2, 84.8</td>
<td>150.8, 84.6</td>
<td>150.8, 85.1</td>
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<tr>
<td>resolution (Å)</td>
<td>20.0–2.40</td>
<td>20.0–2.20</td>
<td>20.0–2.60</td>
<td>20.0–2.00</td>
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<td>Rmerge (%)</td>
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<td>0.077</td>
<td>0.082</td>
<td>0.059</td>
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<td>no. of reflections</td>
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<td>28332</td>
<td>16888</td>
<td>38642</td>
</tr>
<tr>
<td>completeness (%)</td>
<td>100.0</td>
<td>96.1</td>
<td>93.8</td>
<td>99.2</td>
</tr>
<tr>
<td>mean</td>
<td>14.4</td>
<td>19.2</td>
<td>17.8</td>
<td>19.4</td>
</tr>
<tr>
<td>no. of protein atoms</td>
<td>3252</td>
<td>3252</td>
<td>3252</td>
<td>3252</td>
</tr>
<tr>
<td>no. of solvent atoms</td>
<td>168</td>
<td>210</td>
<td>138</td>
<td>235</td>
</tr>
<tr>
<td>no. of heteroatoms</td>
<td>26</td>
<td>27</td>
<td>30</td>
<td>24</td>
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<tr>
<td>mean B-factor (Å²)</td>
<td>38.2</td>
<td>41.6</td>
<td>43.3</td>
<td>35.4</td>
</tr>
<tr>
<td>entire molecule</td>
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<td>38.1</td>
<td>39.5</td>
<td>31.6</td>
</tr>
<tr>
<td>main chain</td>
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<td>45.6</td>
<td>47.6</td>
<td>39.7</td>
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<tr>
<td>side chains</td>
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<td>48.9</td>
<td>46.5</td>
<td>45.2</td>
</tr>
<tr>
<td>solvent</td>
<td>38.5</td>
<td>41.6</td>
<td>43.3</td>
<td>35.4</td>
</tr>
<tr>
<td>msd from identity</td>
<td>0.025</td>
<td>0.027</td>
<td>0.030</td>
<td>0.029</td>
</tr>
<tr>
<td>bond distances (Å)</td>
<td>1.78</td>
<td>1.94</td>
<td>2.07</td>
<td>1.98</td>
</tr>
<tr>
<td>planar groups (Å)</td>
<td>0.016</td>
<td>0.012</td>
<td>0.015</td>
<td>0.016</td>
</tr>
</tbody>
</table>

a R_mer = Σ|I(hkl) − (hkl)|/Σ|I(hkl)|, where I(hkl) is the measured intensity of reflection j and (hkl) is the mean intensity over j reflections. b R_cryst = Σ|F_obs(hkl) − |F_calc(hkl)|/Σ|F_obs(hkl)|, where F_obs and F_calc are the observed and calculated structure factors, respectively. No σ cutoff was applied. c The R_cryst and R_mer values after CNS refinement were as follows: 17.0 and 20.5% for AMPP, 21.1 and 24.0% for ACPP, 17.7 and 22.8% for AMPP, and 20.1 and 22.1% for S-AEP, respectively.
reactive external aldimine intermediates with the corresponding amino acid substrates.

Inhibition Kinetics. The inhibition of DGD was studied in detail for AMPP. Figure 2 presents the double-reciprocal plot for AMPP inhibition using the initial linear portions of the progress curves, before the onset of significant slow binding inhibition (see below). The fitted curves show the results of least-squares analysis employing eq 1. The data strongly indicate that AMPP is, in its initial interaction with DGD, a competitive inhibitor with respect to the AIB substrate. This was expected given the close structural correspondence between the aminophosphonate inhibitor and amino acid substrate. The other aminophosphonate inhibitors were therefore assumed to act by competitive inhibition with respect to AIB. In support of this conclusion, no evidence for amino acid binding to the PMP form of DGD has been found (13), which would open the possibility of noncompetitive or uncompetitive inhibition.

Progress curves for the AIB—pyruvate reaction (Scheme 1) catalyzed by DGD are, in general, not highly linear. This was demonstrated to be due to a slow isomerization of DGD-PLP from a more to a less active form (14). The same progress curves in the presence of AMPP were much more nonlinear (Figure 3), and the rate of change of the initial velocity was dependent on inhibitor concentration (data not shown). This suggested that AMPP is a slow binding inhibitor of DGD. Slow binding inhibition has been reviewed in the literature by Morrison and Walsh (10).

Slow binding inhibition by AMPP was confirmed by two other observations. Preincubation of DGD with AMPP for 30 min yields an apparent $K_i$ value (i.e., $K_i^*$) that is 10-fold smaller than that observed without preincubation. A slow regain of activity after the large dilution of the preformed DGD–AMPP complex into a reaction mixture lacking AMPP (Figure 3) and with saturating AIB is also observed.

This latter experiment allowed the determination of $k_2$ (Scheme 2), the rate constant for conversion of the final, tight complex to the initial, weak complex, since formation of the initial, weak complex is apparently a rapid equilibrium process and inhibitor dissociation under these conditions is essentially irreversible. The value of $k_1$, the rate constant for conversion of the initial, weak complex to the final, tight complex, was calculated from the measured value of $k_2$ and the measured $K_i$ value, using eq 4. The latter was obtained from the AMPP concentration dependence of the reaction velocities obtained after preincubation for 30 min, using eq 2. The values of these rate constants are reported in Table 1.

The values of $k_1$ and $k_2$ were measured independently by monitoring directly the reaction of DGD-PLP with AMPP. The PLP in the active site provides a spectroscopic handle on reactions at the active site (4, 15). In the case of AMPP, small but readily observable changes in the absorbance spectrum of enzyme-bound PLP were observed after mixing the enzyme and inhibitor (Figure 4). Spectral data were collected as a function of time and analyzed globally using the program SPECFIT. A single-exponential process accounted well for the data, and the apparent rate constants are plotted against total AMPP concentration (i.e., sum of $R$ and $S$ isomers) in the inset of Figure 4. The curve shows the least-squares fit to eq 5, with $K_i$ being held at its value measured above. The $y$-intercept yields the value of $k_2$, while the maximal value is the sum of $k_1$ and $k_2$. The values of $k_1$ and $k_2$ obtained from this experiment are reported in Table 1. They agree quite well with the values obtained from steady state kinetic experiments.

The behavior of $S$-AEP was found to be similar to that of AMPP, indicating that it is also a slow binding inhibitor. Similar experiments were thus performed to obtain the values of $K_i$, $K_i^*$, and $k_1$ and $k_2$. The results of these experiments are reported in Table 1. The three other inhibitors employed here (APEP, ACPP, and $R$-AEP) showed no signs of slow binding behavior. Their $K_i$ values are reported in Table 1.

Other examples of slow binding inhibitors of PLP dependent enzymes exist in the literature. Atherton et al. (16) first synthesized 1-aminophosphonophosphate as an alanine racemase inhibitor. Walsh’s group later demonstrated that it is a slow binding inhibitor of Gram-positive but not Gram-negative bacterial alanine racemases (17, 18). It was further demonstrated by NMR and crystallography that it forms a Schiff base with PLP in the active site (19, 20), but the structural origin of the slow binding behavior was not clearly elucidated. 1-Aminocyclopropaneposphonate was also shown to be a slow binding inhibitor of alanine racemase, as well as of 1-aminocyclopropanecarboxylate deaminase (21). Other examples of slow binding inhibitors of PLP enzymes that follow a pre-equilibrium association mechanism, i.e., mechanism B of Morrison and Walsh (10), like the aminophosphonates is L-hydrazinosuccinate and L-aminoxyxsuccinate in their interaction with aspartate aminotransferase (22, 23).

Structures of the Aminophosphonate Inhibitor Complexes. Several structures of DGD exist, including those of the MES-ligated internal aldimine forms of DGD-Li, DGD-Na, DGD-K, and DGD-Rb (7, 12, 24), as well as the structures of DGD-K bound to aminocyclopropane-1-carboxylate, phosphopyridoxyl-AIB, and D- and L-cycloserine (8). The four structures of aminophosphonate complexes presented here are also for the DGD-K form. The $R$-AEP complex with DGD dirfected only to low resolution and is thus not reported here. The kinetic data are included for comparison since it is not slow binding but is very similar in structure to $S$-AEP.

Electron density for the four inhibitors bound in the active site is presented in Figure 5. In each case, the inhibitor conformation is unambiguously determined from the electron density, even in the medium-resolution structures (AMPP and AEP).

The superimposed backbone traces of the four inhibitor structures are presented in Figure 6A. Although not included in the figure, the MES-ligated DGD-K backbone trace is indistinguishable from the four structures that are shown. Clearly, as previously concluded on the basis of the other inhibitor structures (8), there is no large-scale conformational change that accompanies substrate analogue binding, as is observed with the prototypical PLP dependent enzyme aspartate aminotransferase.

Figure 6B shows an overlay of the active site structures of the four inhibitor complexes, as well as the MES-ligated internal aldimine form as a reference. Formation of the external aldimine between the inhibitor and PLP requires a rotation of the PLP ring plane forward toward the solvent to prevent steric clashes with K272. The tilt angle observed here for the aminophosphonate inhibitors is 18–23°, which is larger than the value of 13° observed previously for the
aminocyclopropane carboxylate structure (8). Of the four inhibitor structures, only APEP has the PLP–inhibitor Schiff base not planar with the pyridine ring. This is similar to what was found in the aminocyclopropane carboxylate structure. The phenyl ring of APEP stacks tightly with the indole ring of W138 at a distance of \(3\,\text{Å}\). The twisting of the Schiff base out of the plane of the pyridine ring (thereby breaking the hydrogen bond between the Schiff base nitrogen and PLP O3) is apparently required to maintain this minimal van der Waals interaction distance.

The indole ring of W138 changes its angle to accommodate the variations in the size of the inhibitor side chain, with the larger side chains (AMPP, APEP, and ACPP) pushing the benzyl ring of the indole heterocycle toward M141. This causes the M141 side chain to rotate about the C=S bond, moving the terminal methyl group away from W138 to avoid steric clashes.

The edge of the cyclopentyl ring of ACPP is directed toward and in van der Waals contact with Y301*. Accommodation of the large cyclopentyl ring as it is bound requires breaking of the hydrogen bond normally found between Y301* and Q52 (Figure 5). Surprisingly, this hydrogen bond loss does not dramatically lower the affinity of DGD for ACPP, compared to the affinities for the other inhibitors in which it is maintained.

Yet another variation in structure between the inhibitor complexes occurs near the active site. R251 is located above and behind Q246, which hydrogen bonds to PLP O3 in the MES-ligated structure. This arginine plays no obvious role in the mechanism of either the decarboxylation or transamination reactions. The guanidino group shows two conformations, with concomitant water molecule reorganization, in the four structures presented here (Figure 7A). Neither of these appears to be energetically highly favorable since no counterion interactions are made and hydrogen bonding is minimal. The variation in conformation does not account for slow binding behavior since S-AEP and ACPP show one conformation and AMPP and APEP another.

One difference between the inhibitor structures that is correlated with the slow binding behavior of AMPP and S-AEP is an extra water molecule located below Met141 in the AMPP and S-AEP structures. This water is shown in Figures 6B and 7B as a large sphere. Figure 7B shows the details of the water binding site. In the AMPP structure, this water apparently forms only one hydrogen bond, to a water just below it in this figure. The S-AEP structure shows this water forming two hydrogen bonds, one with the same water as in the AMPP structure and another with R120. The difficulty with assigning this structural difference to the source of slow binding behavior is that there is no clear
structural explanation for the absence of this water in the ACPP and APEP structures. In the AMPP and $S$-AEP structures, no steric clashes are removed and no extra hydrogen bonds are enabled.

One other structural difference was found to correlate with the slow binding behavior of AMPP and $S$-AEP. The PLP pyridine nitrogen–D243 OD2 distance is $\sim 2.7$ Å for AMPP and $S$-AEP, while it is $\sim 2.9$ Å for ACPP and APEP. OD2 of D243 makes two other hydrogen bonding interactions: (1) to the ND1 atom of H139 and (2) to the ND2 atom of N115. These are also found to vary with the pyridine nitrogen–D243 OD2 distance.

Figure 8 presents the distance correlations between the three hydrogen bonding interactions that the OD2 atom of D243 makes for the four phosphonate structures presented here as well as DGD-K, DGD-Na, and the aminocyclopropane-carboxylate inhibitor structures. A monotonic decrease in the H139–D243 interaction distance is observed as the pyridine nitrogen–D243 interaction distance increases. The N115–D243 interaction distance begins to decrease only after the pyridine nitrogen–D243 interaction distance increases beyond 2.8 Å. Clearly, there are compensatory changes in hydrogen bonding distances to accommodate the presumed negative charge on D243. The slope of a line fitted to the H139 data in Figure 8 has a slope of $-0.86 \pm 0.08$ when all data are included and $-1.0 \pm 0.1$ when the DGD-Na point is excluded.

The two slow binding phosphonate inhibitors (AMPP and $S$-AEP), with their shorter $\sim 2.7$ Å interaction distance between the pyridine nitrogen and D243 OD2, are more
similar to the DGD-K and aminocyclopropanecarboxylate structures, which presumably have relaxed pyridine nitrogen–D243 interactions. The ACPP and APEP structures suggest that shorter pyridine nitrogen–D243 interaction distances are not possible for these due to steric interactions between, mainly, W138 and the larger inhibitor side chains.

Calculations of the energy differences incurred by elongating the pyridine nitrogen–D243 interaction distance were performed with SPARTAN at the 6-31G* level for a simple model in which the methyamine Schiff base of 3-hydroxypyridine-4-aldehyde interacts via the protonated pyridine nitrogen with formate. The geometries of the pyridine nitrogen–formate interaction were constrained to be those found for D243 in the inhibitor structures. These show that the S-AEP geometry is ~2 kcal/mol higher in energy than the AMPP geometry, while the ACPP and APEP geometries are ~6 and ~9 kcal/mol higher in energy, respectively. These calculations suffer, of course, from the simplicity of the model since it does not include the H139 and N115 interactions.

Clearly, the variation in the PLP pyridine nitrogen–D243 OD2 interaction distance has the potential to account for the observed 1.4 kcal/mol difference in binding energy between the rapidly formed DGD–AMPP or DGD–S-AEP complexes and the final, tight binding, isomerized complexes. Thus, it is proposed that the structural basis of the slow binding behavior of AMPP and S-AEP is initial formation of a Schiff base between the inhibitor and PLP that resembles...
more the ACPP and APEP structures in their PLP—inhibitor geometries and interactions, followed by slow isomerization to the observed geometries in which the PLP pyridine nitrogen—D243 OD2 interaction distance is shortened. This proposition is supported by the spectra of the initial and final AMPP complexes (obtained from global analysis of the spectral kinetic data, Figure 4) and the ACPP complex. The latter is very similar to the initial AMPP spectrum, while the final AMPP complex shows a red shift of the ~410 nm band to ~430 nm and an altered 330 nm/430 nm ratio, indicative of an altered tautomeric ratio (data not shown).

Also, the similarities of the indicative of an altered tautomeric ratio (data not shown).

The C4 analysis since its Schiff base is not coplanar with the pyridine corresponding substrates. APEP is excluded from this correlation of aldimine structure with the reactivities of their AMPP, of the substrate bound in the A subsite notwithstanding, the Q52 and K272 (to the pyridine ring plane through hydrogen bonding with the former placing the substrate carboxylate perpendicular bind between what is ideally considered the A and B subsite, near R406 and S215. The phosphonates, on the other hand, boxylate group clearly binding in the so-called “B” subsite, which there are three binding subsites. The previous car—was based on a functional model of the DGD active site in mechanism for control of reaction specificity. This analysis using alternate substrates that DGD does indeed employ this parallel to the p orbitals of the conjugated Schiff base—variation distance that is partially but not completely compensated by the covariation in the H139 and N115 interactions with D243.

**Implications for Catalysis.** Dunathan (5) proposed in 1966 that PLP dependent enzymes can control reaction specificity largely by orienting the bond to be broken such that it is parallel to the p orbitals of the conjugated Schiff base—pyridine ring π system. Sun and Toney (6) provided evidence using alternate substrates that DGD does indeed employ this mechanism for control of reaction specificity. This analysis was based on a functional model of the DGD active site in which there are three binding subsites. The previous carboxylate-containing inhibitor structures (8) show the carboxylate group clearly binding in the so-called “B” subsite, near R406 and S215. The phosphonates, on the other hand, bind between what is ideally considered the A and B subsite, the former placing the substrate carboxylate perpendicular to the pyridine ring plane through hydrogen bonding with Q52 and K272 (7). This is probably the result of the steric bulk of the three phosphate oxygens and the requirement to compensate electrostatically its double negative charge versus the single charge of a substrate carboxylate, since the “in between” site allows interaction with Q52, K272, Q246, R406, and (for S-AEP and ACPP) S215.

This suboptimal representation by the amino phosphonates of the substrate bound in the A subsite notwithstanding, the AMPP, S-AEP, and ACPP structures show an enticing correlation of aldimine structure with the reactivities of their corresponding substrates. APEP is excluded from this analysis since its Schiff base is not coplanar with the pyridine ring. The C4—N—Cα—P dihedral angle is ~139° for AMPP, ~159° for S-AEP, and ~161° for ACPP, which correspond to \( k_{\text{cat}} \) values for the analogous substrates of 14, 0.2, and 0.98 s\(^{-1}\), respectively (Table 1). Thus, the inhibitor with the C—P bond closest to being parallel to the p orbitals of the π system corresponds to the substrate with the highest \( k_{\text{cat}} \) value. This suggests that the structural features of the phosphonate inhibitors that determine binding geometry also affect the binding geometry of the corresponding carboxylate substrates, which in turn determines reactivity through stereoelectronic effects.

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