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Binding Energy and Catalysis by D-Xylose Isomerase: Kinetic, Product and X-Ray Crystallographic Analysis of Enzyme-Catalyzed Isomerization of (*R*)-Glyceraldehyde[‡], ¶

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Abstract

D-Xylose isomerase (XI) and triosephosphate isomerase (TIM) catalyze the aldose-ketose isomerization reactions of D-xylose and D-glyceraldehyde 3-phosphate (DGAP), respectively. D-Glyceraldehyde (DGA) is the triose fragment common to the substrates for XI and TIM. The XI-catalyzed isomerization of DGA to give dihydroxyacetone (DHA) in D₂O was monitored by ¹H NMR spectroscopy and $k_{\text{cat}}/K_{\text{m}} = 0.034 \text{ M}^{-1} \text{ s}^{-1}$ was determined for this isomerization at pD 7.0. This is similar to $k_{\text{cat}}/K_{\text{m}} = 0.017 \text{ M}^{-1} \text{ s}^{-1}$ for the TIM-catalyzed carbon deprotonation reaction of DGA in D₂O at pD 7.0 [Amyes, T. L.; O'Donoghue, A. C. and Richard J. P. (2001) *J. Am. Chem. Soc.* 123, 11325–11326]. The much larger activation barrier for XI-catalyzed isomerization of D-xylose ($k_{\text{cat}}/K_{\text{m}} = 490 \text{ M}^{-1} \text{ s}^{-1}$) than for the TIM-catalyzed isomerization of DGAP ($k_{\text{cat}}/K_{\text{m}} = 9.6 \times 10^6 \text{ M}^{-1} \text{ s}^{-1}$) is due to: (i) The larger barrier to conversion of cyclic D-xylose to the reactive linear sugar (5.4 kcal/mol) than for conversion of DGAP hydrate to the free aldehyde (1.7 kcal/mol). (ii) The smaller intrinsic binding energy [Jencks, W. P. (1975) *Adv. Enzymol. Relat. Areas Mol. Biol.* 43, 219–410] of the terminal ethylene glycol fragment of D-xylose (9.3 kcal/mol) than of the phosphodianion group of DGAP (ca. 12 kcal/mol). The XI-catalyzed isomerization of DGA in D₂O at pD 7.0 gives a 90% yield of [1-¹H]-DHA and a 10% yield of [1-²H]-DHA, the product of isomerization with deuterium incorporation from solvent D₂O. By comparison, the transfer of 3H from labeled hexose substrate to solvent is observed only once in every 10⁹ turnovers for the XI-catalyzed isomerization of [2-³H]-glucose in H₂O [Allen, K. N., Lavie, A., Farber, G. K., Glasfeld, A., Petsko, G. A., and Ringe, D. (1994), *Biochemistry* 33, 1481–1487]. We propose that truncation of the terminal ethylene glycol fragment of D-xylose to give DGA results in a large decrease in the rate of XI-catalyzed isomerization with hydride transfer compared with that for proton transfer. An ultra-high resolution (0.97 Å) X-ray crystal structure was determined for the complex obtained by soaking crystals of XI with 50 mM DGA. The triose binds to XI as the unreactive hydrate, but ligand binding induces metal cofactor movement and conformational changes in active site residues similar to those observed for XI•sugar complexes.

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[¶]The coordinates of the refined structure have been deposited with the Protein Data Bank as entry #3U3H.

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INTRODUCTION

Two mechanisms have evolved for the enzyme-catalyzed isomerization of α -hydroxyaldehydes (aldoses) to give α -hydroxyketones (ketoses). Isomerization of sugar phosphates, as exemplified by conversion of D-glyceraldehyde 3-phosphate (DGAP)¹ to give dihydroxyacetone phosphate catalyzed by triosephosphate isomerase (TIM, Scheme 1A) (1–8), proceeds by a proton transfer mechanism through an enediolate intermediate (8, 9), while isomerization of sugars, as exemplified by reactions catalyzed by D-xylose isomerase (XI, Scheme 1B) (10–16), proceeds by intramolecular hydride transfer. By comparison, the general-base-catalyzed isomerization of triosephosphates in water proceeds by a proton transfer reaction mechanism (17, 18), while competing proton and hydride transfer reaction pathways are observed for the nonenzymatic isomerization of glyceraldehyde to give dihydroxyacetone (DHA) in alkaline aqueous solution (19, 20).

TIM satisfies at least two criteria for perfection in enzymatic catalysis (3), the most important being that the second-order rate constant $k_{\text{cat}}/K_{\text{m}} = 2 \times 10^8 \text{ M}^{-1} \text{ s}^{-1}$ for isomerization of the *free carbonyl* form of the substrate is close to the maximum possible for a diffusion-limited reaction (21).² By comparison, $k_{\text{cat}}/K_{\text{m}} \approx 10^3 \text{ M}^{-1} \text{ s}^{-1}$ for the XI-catalyzed isomerization of D-xylose (22–24) falls far short of the limiting rate constant of ca. $10^8 \text{ M}^{-1} \text{ s}^{-1}$ for a diffusion-limited reaction catalyzed by a perfect enzyme (3). We are interested in understanding how this difference in catalytic efficiency is related to the difference in the structure of the substrates of the TIM- and XI-catalyzed isomerization reactions (15, 25).

Truncation of the phosphoryl dianion from DGAP or the terminal $-\text{CH}(\text{OH})\text{CH}_2\text{OH}$ fragment from D-xylose gives D-glyceraldehyde (DGA), the fragment common to the substrates for the two enzymes (Scheme 2). TIM catalyzes the slow isomerization of the *free carbonyl* form of DGA to give DHA in D_2O with $k_{\text{cat}}/K_{\text{m}} = 0.34 \text{ M}^{-1} \text{ s}^{-1}$ that is much smaller than $k_{\text{cat}}/K_{\text{m}} = 2 \times 10^8 \text{ M}^{-1} \text{ s}^{-1}$ for isomerization of DGAP, and only marginally larger than $k_{\text{B}} = 6.5 \times 10^{-3} \text{ M}^{-1} \text{ s}^{-1}$ for deprotonation of DGA by the small Brønsted base 3-quinuclidinone (26).² We report here that the observed kinetic parameter $k_{\text{cat}}/K_{\text{m}}$ for the XI-catalyzed isomerization of DGA to give DHA in D_2O is similar to that for the TIM-catalyzed isomerization under the same reaction conditions (26). The greater efficiency of TIM compared to XI for catalysis of the reaction of their whole substrates is therefore related to a difference in the activation of the respective enzymes by interactions with the nonreacting phosphodianion of DGAP and the terminal ethylene glycol fragment of D-xylose, respectively.

We have complemented these kinetic studies with the report of an ultra-high resolution (0.97 Å) X-ray crystal structure of the XI•triose complex obtained by soaking crystals of XI with DGA. This X-ray crystal structure shows that the more abundant (95% of total DGA) hydrate of DGA (DGAH) was bound to XI. By contrast, TIM shows selectivity for binding and catalysis of the reactive free carbonyl form of the substrate DGAP (27). The formation of the complex of XI with the unreactive hydrated triose induces movement of the metal cofactor and associated shifts in the position of active site residues that are remarkably

¹Abbreviations: DGA, D-glyceraldehyde; *h*-DGA, [1(*R*)-¹H]-glyceraldehyde; *d*-DGA, [2(*R*)-²H]-glyceraldehyde; DGAH, hydrate of D-glyceraldehyde; DGAP, (*R*)-glyceraldehyde 3-phosphate; DHA, dihydroxyacetone; *h*-DHA, [1-¹H]-dihydroxyacetone; *d*-DHA, [1-²H]-dihydroxyacetone; NADH, nicotinamide adenine dinucleotide, reduced form; NMR, nuclear magnetic resonance; PGA, 2-phosphoglycolate; TIM, triosephosphate isomerase; XI, D-xylose isomerase.

²The rate constants for DGA and DGAP reported in the Introduction are calculated for the reaction of the minor (5%) *free carbonyl* form of the substrates. They are 20-fold larger than the corresponding observed rate constants that refer to the reaction of total DGA or DGAP given in Tables 1 and 3.

similar to the conformational changes observed for complexes of XI with pentose and hexose sugar substrates.

MATERIALS AND METHODS

Materials

Reagent grade organic chemicals and inorganic salts from commercial sources were used without further purification. Water was distilled and then passed through a Milli-Q water purification system. Deuterium oxide (99.9% atom D), deuterium chloride (35% w/w, 99.5% atom D) and CD₃COOD (99.5% atom D) were purchased from Cambridge Isotope Laboratories. D-Glyceraldehyde (90 wt % in water) was purchased from Aldrich. NADH (disodium salt), D-xylose, D-fructose, and sorbitol dehydrogenase from sheep liver (lyophilized powder) were purchased from Sigma. A crystalline suspension of D-xylose isomerase from *Streptomyces rubiginosus* was purchased from Hampton Research. D-Threono-hydroxamic acid was prepared by a published procedure (28).

General methods

Imidazole and NaCl were dried in a vacuum oven overnight before preparation of buffered solutions in D₂O. Values of pD were determined by adding 0.40 to the observed reading of the pH meter at 25 °C (29). Stock solutions (2 M) of D-xylose and D-fructose in D₂O at *I* = 0.1 (NaCl) were kept at room temperature for one day to ensure formation of an equilibrium mixture of sugar anomers. Commercial D-glyceraldehyde (90 wt % in water) exists mainly as the dimer. This was dissolved in D₂O, the D₂O was removed *in vacuo*, and the compound was then dissolved in 0.1 M NaCl in D₂O to give a ca. 50 mM solution at *I* = 0.1 (NaCl). This solution was kept at room temperature for ≥ 1 week, until ≥ 95% of monomeric glyceraldehyde was observed by ¹H NMR analysis. ¹H NMR showed the presence of only ca. 0.04% of DHA. Solutions of authentic DHA (50 mM at *I* = 0.1, NaCl) were prepared in D₂O and were used on the day that they were prepared.

Enzymes were dialyzed against the appropriate buffer at 5 °C. Crystalline XI used in kinetic studies was freed of ammonium sulfate by extensive dialysis against 30 mM imidazole buffer in H₂O (pH 7.0 or 8.0) that contained 12.5 mM MgCl₂ (*I* = 0.1, NaCl). The protein was then dialyzed exhaustively against 30 mM imidazole buffer in D₂O (pD 7.0 or 8.0) that contained 12.5 mM MgCl₂ at *I* = 0.1 (NaCl). Typically, a 3 – 4-fold volume excess of dialysis buffer over sample was used, and the buffer was changed 7 times. The concentration of XI in stock solutions used for kinetic and product studies was determined from the absorbance at 280 nm using $\epsilon = 0.96 \text{ mg}^{-1} \text{ ml cm}^{-1}$ (30) and a molecular weight of 43,000 Dalton/monomer (23). The concentration of XI in stock solutions used for crystallization studies was determined using the Bio-Rad protein assay (Bio-Rad, Hercules, CA). Sorbitol dehydrogenase was dialyzed against 30 mM imidazole buffer in D₂O (pD 7.0 or 8.0) that contained 12.5 mM MgCl₂ (*I* = 0.1, NaCl). The enzyme was stable (< 5% loss of activity) at 5 °C for more than 6 weeks. One unit of enzyme activity is defined as the amount of enzyme that will catalyze the transformation of 1 μmole of substrate per minute under our assay conditions at 25 °C.

Enzyme assays

Enzyme assays were performed at 25 °C. The activity of sorbitol dehydrogenase in D₂O was determined by monitoring the enzyme-catalyzed oxidation of NADH by D-fructose at 340 nm. The standard assay solution (1.0 mL) contained 100 mM D-fructose, 0.15 mM NADH, 10 mM MgCl₂ and 0.06 units of sorbitol dehydrogenase in 24 mM imidazole buffer at pD 7.0 or 8.0 (*I* = 0.1, NaCl). XI was assayed in D₂O by coupling the isomerization of D-xylose to the reduction of the product D-xylulose by NADH using sorbitol dehydrogenase. The

standard assay solution (1.0 mL) contained 50 mM D-xylose, 0.23 mM NADH, 10 mM MgCl_2 , 0.012 units of XI and 0.6 units of sorbitol dehydrogenase in 24 mM imidazole buffer at pD 7.0 or pD 8.0 ($I = 0.1$, NaCl).

X-Ray crystallographic analysis

XI from *S. rubiginosus* was prepared for crystallization by overnight dialysis of the commercial crystalline suspension (Hampton Research, Inc.) against 4 L of distilled H_2O followed by concentration to 20 mg/mL using an Amicon centrifugal concentrator (Millipore, Inc.) with a 10,000 MWCO membrane. XI was crystallized by the vapor diffusion method with hanging drop geometry over wells containing a solution of 0.2 – 0.3 M $\text{Mg}\cdot\text{formate}$ at pH 7.0. Large crystals (ca. 0.4 x 0.4 x 0.2 mm) were obtained after two days of growth at room temperature (ca. 22 °C). The structure of the complex of XI with D-glyceraldehyde was obtained by soaking crystals of unliganded XI for 9 h in a solution containing 0.4 M $\text{Mg}\cdot\text{formate}$ and 50 mM DGA. Crystals were transferred sequentially to solutions containing 0.4 M $\text{Mg}\cdot\text{formate}$, 50 mM DGA, and 5 %, 15 %, or 30 % 2-methyl-2,4-pentanediol (MPD) and flash-cooled in a gaseous N_2 stream at 100K. Diffraction data were collected at the National Synchrotron Light Source (NSLS), beamline X12B, using an ADSC Quantum 4 CCD detector. The data were integrated and scaled using DENZO and SCALEPACK (31).

Model refinement

The initial phases were derived from the 1.5 Å resolution structure of the unliganded *S. rubiginosus* enzyme (PDB entry 1OAD) (32) in space group $\text{P}2_12_12_1$. The coordinates of the unliganded enzyme were used as the search model in determining the orientation of the molecule in the I222 unit cell with EPMR (33). Once the initial phases were obtained, the structure was refined to 1.5 Å resolution using CNS (34). Solvent molecules were added that showed reasonable hydrogen bond distances and geometry to donor and acceptor atoms. Care was taken to avoid modeling the density for the bound ligands as solvent. After the water structure was deemed to be satisfactory and the R_{cryst} reached approximately 19% (~21% R_{free}), further refinement was done using SHELX-97 (35). Following isotropic refinement at full resolution, the model was refined anisotropically. Refinement of the individual anisotropic displacement parameters resulted in a 5.3% decrease in R_{free} . Comparable drops in R_{cryst} accompanied the changes in R_{free} . Iterative cycles of fitting in COOT (36) followed by refinement in SHELXL continued until the R factors converged. At this point, all protein hydrogen atoms except for amino and hydroxyl hydrogens were added using the riding model in SHELXL. This led to a ~1% decrease in the R_{free} value.

^1H NMR analyses

^1H NMR spectra at 500 MHz were acquired at 25 °C using a Varian Unity Inova spectrometer with a 90 degree pulse angle, a 6000 Hz sweep width, a 4 – 6 s acquisition time, and a relaxation delay between pulses of 120 s, as described in our earlier work (20, 26, 37, 38). Normalized areas of the peaks corresponding to a *single* proton of DGA (A_{DGA}), *h*-DHA ($A_{h\text{-DHA}}$) and *d*-DHA ($A_{d\text{-DHA}}$) in mixtures of these compounds were determined from the integrated areas of the peaks due to the C-1 proton of DGA hydrate ($f_{\text{hyd}} = 0.95$) and the protons due to the free keto forms of *h*-DHA and of *d*-DHA ($f_{\text{hyd}} = 0.16$), respectively, as described previously (20). The concentration of DGA was determined relative to the known concentration of the imidazole buffer from the ratio of the peak areas for the C-4 and C-5 protons of imidazole and the C-1 proton of DGA hydrate using $f_{\text{hyd}} = 0.95$ for the fraction of DGA present as the hydrate in D_2O (20, 26). A similar method was used to determine the concentration of D-threono-hydroxamic acid using the peak area for its C-2 proton.

Reactions of DGA and DHA monitored by ^1H NMR spectroscopy

The XI-catalyzed reactions of DGA and DHA in D_2O at pD 7.0 or 8.0 in the presence of 24 mM imidazole buffer and 10 mM MgCl_2 at 25°C and $I = 0.1$ (NaCl) were monitored by ^1H NMR spectroscopy. The reactions (10 mL volume) were initiated by making a 5-fold dilution of a ca. 50 mM stock solution of DGA or DHA ($I = 0.1$, NaCl) into 30 mM imidazole buffer ($I = 0.1$, NaCl) containing 12.5 mM MgCl_2 to give final concentrations of 11 mM substrate, 24 mM imidazole, 10 mM MgCl_2 and 0.30 mM XI for the reaction at pD 7.0 and 0.20 mM XI for the reaction at pD 8.0. At timed intervals an aliquot (1 mL) was withdrawn and the solution was adjusted to pD ≈ 5 with 10 μL or 25 μL of 1 M CD_3COOD for reactions at pD 7.0 or 8.0, respectively. It was shown in control experiments that this drop in pD effectively quenches the slow XI-catalyzed reaction. The enzyme was removed by ultrafiltration at 5°C using a Microcon YM-10 centrifugal filtration device. The filtrate was then flash frozen over solid CO_2 and stored at -15°C . ^1H NMR analyses of the mixture were performed within one week of freezing. It has been shown that DGA is stable to nonenzymatic isomerization and deuterium exchange reactions over this period of time (20). The same procedure was used to monitor the XI-catalyzed isomerization of DGA in the presence of 0.5 mM of the competitive inhibitor D-threono-hydroxamic acid (28).

The fraction of DGA converted to DHA, $(f_{\text{DHA}})_{\text{tot}}$, during reaction of $\leq 10\%$ of the starting DGA was calculated from the normalized peak areas due to a *single* proton of the reactant, A_{DGA} , and the two products using eq 1, where $A_{h\text{-DHA}}$ and $A_{d\text{-DHA}}$ are the normalized peak areas for a *single* proton of the two products *h*-DHA and *d*-DHA, respectively, determined as described previously (see above) (20). The *apparent* first-order rate constants for the XI-catalyzed conversion of DGA to *h*-DHA and *d*-DHA, k_{obsd} (s^{-1}), were then calculated as the slopes of plots of $(f_{\text{DHA}})_{\text{tot}}$ against time according to eq 2, which were linear over the initial 10% of the reaction (see Figure 1). The ratio of the yields of *h*-DHA and *d*-DHA were obtained using eq 3, where A_{CH_2} is the observed area of the singlet at 4.300 ppm due to the $\alpha\text{-CH}_2\text{OD}$ groups of the free carbonyl form of both *h*-DHA and *d*-DHA, and A_{CHD} is the area of the 0.024 ppm upfield-shifted triplet due to the single $\alpha\text{-CHDOD}$ group of *d*-DHA (see Figure 2) (20). The fraction of the total DHA product labeled with deuterium, $f_{d\text{-DHA}}/(f_{\text{DHA}})_{\text{tot}}$, was then calculated from this ratio using eq 4.

$$(f_{\text{DHA}})_{\text{tot}} = \frac{A_{h\text{-DHA}} + A_{d\text{-DHA}}}{A_{\text{DGA}} + A_{h\text{-DHA}} + A_{d\text{-DHA}}} \quad (1)$$

$$(f_{\text{DHA}})_{\text{tot}} = k_{\text{obsd}} t \quad (2)$$

$$\frac{[d\text{-DHA}]}{[h\text{-DHA}]} = \frac{4A_{\text{CHD}}}{A_{\text{CH}_2} - 2A_{\text{CHD}}} \quad (3)$$

$$\frac{f_{d\text{-DHA}}}{(f_{\text{DHA}})_{\text{tot}}} = \frac{\{[d\text{-DHA}]/[h\text{-DHA}]\}}{1 + \{[d\text{-DHA}]/[h\text{-DHA}]\}} \quad (4)$$

The slow uncatalyzed reaction of DGA in the absence of XI in D_2O at pD 7.0 in the presence of 24 mM imidazole buffer and 10 mM MgCl_2 at 25°C and $I = 0.1$ (NaCl) was

followed directly by ^1H NMR spectroscopy (26) by monitoring the decrease in the area of the doublet due to the C-1 proton of DGA hydrate, using the signals for the C-4 and C-5 hydrogens of imidazole as an internal standard (26). The *observed* first-order rate constant for this nonenzymatic reaction of DGA, $(k_N)_{\text{app}}$, was determined as the slope of a semilogarithmic plot of the fraction of remaining substrate against time.

Steady state kinetic studies

Values of V_{max} and K_m for the XI-catalyzed isomerization of D-xylose to give D-xylulose in D_2O at pD 7.0 or 8.0 in the presence of 24 mM imidazole and 10 mM MgCl_2 at 25 °C and $I = 0.10$ (NaCl) were determined from the nonlinear least squares fit of the initial velocity data to the Michaelis-Menten equation. The value of K_i for competitive inhibition of XI by DGA was determined from the nonlinear least squares fit of the initial velocities v to eq. 5. The quoted errors are standard deviations. The initial velocities determined in these experiments were reproducible to better than $\pm 5\%$.

$$\frac{v}{V_{\text{max}}} = \frac{[S]}{[S] + K_m \left(1 + \frac{[I]}{K_i}\right)} \quad (5)$$

RESULTS

Kinetic and product studies

D-xylose isomerase from *Streptomyces rubiginosus* was assayed in D_2O by coupling the enzyme-catalyzed isomerization of D-xylose to form D-xylulose to the reduction of D-xylulose by NADH using sorbitol dehydrogenase. Table 1 reports the values of k_{cat} and K_m determined for the XI-catalyzed isomerization of D-xylose in D_2O at pD 7.0 in the presence of 24 mM imidazole buffer and 10 mM MgCl_2 at 25 °C and $I = 0.1$ (NaCl). Values of $k_{\text{cat}} = 3.3 \text{ s}^{-1}$ and $K_m = 5.0 \text{ mM}$ at pD 8.0 were determined for this isomerization reaction under otherwise similar conditions. By comparison, the XI-catalyzed (*Streptomyces violaceoruber*) isomerization of D-xylose in H_2O at 35 °C exhibits a broad pH optimum at neutral pH, with $k_{\text{cat}} = 10 \text{ s}^{-1}$ and $K_m = 2.8 \text{ mM}$ (23).

The inhibition of the XI-catalyzed isomerization of D-xylose by DGA at pD 7.0 was examined using six concentrations of D-xylose between 2.5 mM and 30 mM and fixed concentrations of 3.2 and 6.3 mM DGA (data not shown). The presence of DGA does not change the maximum velocity for the reaction in the presence of saturating D-xylose. This shows that inhibition by DGA is competitive. An inhibition constant for DGA of $K_i = 3.0 \pm 0.1 \text{ mM}$ was determined from the nonlinear least-squares fit of the initial velocity data to eq 5. The initial velocities determined for the XI-catalyzed isomerization of 21 mM D-xylose in the presence of 5.3 mM and 10.3 mM DGA at pD 8.0 are also consistent with $K_i = 3 \text{ mM}$ for DGA which shows that there is no large change in K_i for the enzyme-catalyzed reaction on moving to the higher pD.

The nonenzymatic reaction of DGA in D_2O at pD 7.0 in the presence of 24 mM imidazole buffer and 10 mM MgCl_2 at 25 °C and $I = 0.1$ (NaCl) was monitored by ^1H NMR for three weeks, during which time there was ca. 25% disappearance of the starting DGA. The major reaction products were formate and glycolaldehyde from the oxidative cleavage of DGA. There was no detectable formation of DHA from a nonenzymatic aldose-ketose isomerization reaction. These data gave $(k_N)_{\text{app}} \approx 1.4 \times 10^{-7} \text{ s}^{-1}$ as the net observed rate constant for all of the nonenzymatic reactions of DGA (Scheme 3).

The XI-catalyzed isomerization of DGA in D₂O at pD 7.0 or 8.0 in the presence of 24 mM imidazole buffer and 10 mM MgCl₂ at 25 °C and *I* = 0.1 (NaCl) was monitored by ¹H NMR spectroscopy for ca. 9 h during which time there was disappearance of ca. 10% of the starting DGA. Figure 1A shows the linear initial rate time courses, with slope *k*_{obsd}, for the fractional conversion of 11 mM DGA to give *h*-DHA and *d*-DHA, (*f*_{DHA})_{tot} (eq 1), catalyzed by XI at pD 7.0 ([E] = 0.30 mM) or pD 8.0 ([E] = 0.20 mM). Values of the *apparent* first-order rate constants *k*_{obsd} = (2.2 ± 0.1) × 10⁻⁶ s⁻¹ and (1.5 ± 0.1) × 10⁻⁶ s⁻¹ at pD 7.0 and 8.0, respectively, were determined from the slopes of the linear correlations in Figure 1A (eq 2), where the quoted error is the standard deviation in this slope. The values of *k*_{cat} and *k*_{cat}/*K*_m (Table 1) for the XI-catalyzed isomerization of DGA in D₂O (Scheme 3) were obtained from the *apparent* first-order rate constants using eq. 6, with the assumption that *K*_i = 3 mM determined for competitive inhibition by DGA of the XI-catalyzed reaction of D-xylose is equal to *K*_m for the XI-catalyzed reaction of DGA.

$$k_{\text{obsd}} = \frac{k_{\text{cat}}[E]}{K_m + [DGA]_0} \quad (6)$$

Figure 2 shows representative partial ¹H NMR spectra of in the region of the free keto form of DHA, obtained during the XI-catalyzed isomerization of DGA in D₂O. The singlet at 4.300 ppm with an area *A*_{CH₂} is due to the α-CH₂OD groups of both *h*-DHA and *d*-DHA, and the 0.024 ppm upfield-shifted triplet with an area *A*_{CHD} is due to the α-CHDOD group of *d*-DHA (Scheme 3). In these spectra the area *A*_{CHD} is ca. 2.5% and 1.2%, respectively, that of the area *A*_{CH₂} for reactions at pD 7.0 (Figure 2A) and pD 8.0 (Figure 2B). Figure 1B shows the change with time in the fraction of the product DHA that is labeled with deuterium at pD 7.0 and 8.0, determined from the peak areas *A*_{CH₂} and *A*_{CHD} using eqs 3 and 4. The increase in *f*_{*d*-DHA}/(*f*_{DHA})_{tot} with time results from the slow XI-catalyzed deuterium exchange of the methylene protons of the product *h*-DHA to give additional *d*-DHA. The *initial* fractions of the product DHA that contains deuterium, 0.095 and 0.05 at pD 7.0 and 8.0, respectively, were determined by making a short linear extrapolation of the data in Figure 1B to zero reaction time. Table 1 reports the values of *k*_{cat}/*K*_m for the XI-catalyzed isomerization of DGA with intramolecular hydride transfer to give *h*-DHA and with proton transfer with incorporation of deuterium from solvent D₂O to give *d*-DHA, calculated from the kinetic parameters for the overall XI-catalyzed isomerization of DGA and these fractional product yields.

No products of XI-catalyzed isomerization were detected during the reaction of 11 mM DGA for 9 h in D₂O at pD 7.0 in the presence of 24 mM imidazole buffer, 10 mM MgCl₂, 0.27 mM XI and 0.5 mM (5,000 *K*_i) of the competitive inhibitor D-threonoxyhydroxamic acid (28) at 25 °C and *I* = 0.1 (NaCl).

The XI-catalyzed isomerization of DHA (11 mM) in D₂O at pD 7.0 in the presence of 24 mM imidazole buffer and 10 mM MgCl₂ and 0.30 mM XI at 25 °C and *I* = 0.1 (NaCl) was monitored by ¹H NMR spectroscopy for ca. 25 h during which time ca. 6% of the starting *h*-DHA had undergone deuterium exchange with solvent to form *d*-DHA. Small amounts of *h*-DGA and *d*-DGA (ca. 3% of total DHA) from thermodynamically unfavorable isomerization were also detected. No products of the enzyme-catalyzed deuterium exchange or isomerization reactions of DHA were detected by ¹H NMR after a 25 h reaction time under the same conditions, but in the presence of 0.5 mM D-threonoxyhydroxamic acid.

X-Ray crystallographic analyses

XI crystallized in space group I222 with unit cell dimensions $a = 92.6$, $b = 98.1$, and $c = 102.6$. XI from *S. olivochromogenes* has been shown to crystallize in space group $P2_12_12$ with pseudo I222 symmetry (15). However, these data were not collected from crystals cooled to 100 K. The first XI structure determined from cryogenic data displayed true I222 symmetry (16). Given that attempts at refinement of the present structures in several orthorhombic space groups failed, we conclude that I222 is also the true space group for these crystals. Data collection and refinement statistics for the structures of XI crystals obtained after soaking for nine h in a solution that contained 50 mM DGA are presented in Table 2. The structure at 0.97 Å resolution (Figure 3) showed electron density that was modeled as two ligands bound at the enzyme active site: the hydrate of DGA (DGAH, 49% occupancy) and formate ion (51% occupancy). The C3 hydroxyl of DGAH interacts with the protein through a hydrogen bond to N ϵ of the imidazole side chain of His54 (2.6 Å) and to a bridging water molecule, with the γ -oxygen of Thr90 (2.8 Å). XI shows good catalytic activity with either Mg^{2+} or Mn^{2+} bound at the two metal sites (22). We have worked with the Mg^{2+} enzyme. The C2 hydroxyl is coordinated to M1 (2.1 Å). The two C1 hydroxyls of DGAH, Oa and Ob, interact with M1 (2.2 Å) and with an ordered solvent molecule. One oxygen atom of formate ion is found to bridge M1 and M2a (2.2 Å and 2.5 Å, respectively). The second oxygen is coordinated to M2a (2.3 Å) and interacts by hydrogen bonding with the ζ -ammonium ion of Lys183 (3.1 Å). The water/hydroxide ion ligand of M2a is 2.0 Å from the Mg^{2+} atom. In this arrangement the binding of DGAH and formate appears to be mutually exclusive. A second structure (not shown) of XI soaked with DGA for a period of 15 minutes is nearly identical to the structure DGAH despite the much shorter soaking time.

DISCUSSION

There are two pathways for the slow TIM-catalyzed reactions of the neutral truncated substrates DGA and glycolaldehyde in D_2O : (1) a reaction at the enzyme active site that is shut down by binding of the competitive inhibitor 2-phosphoglycolate (PGA); and, (2) a reaction that is insensitive to PGA and/or to mutations that cripple the reaction of the natural substrate DGAP at the enzyme active site (8). This second nonspecific reaction pathway has also been documented for the reaction of glycolaldehyde catalyzed by bovine serum albumin (39). By contrast, the observed XI-catalyzed isomerization of DGA to give DHA is completely abolished by the tight-binding competitive inhibitor D-threono-hydroxamic acid (28), which shows that this isomerization occurs exclusively at the enzyme active site. Similarly, the very slow XI-catalyzed deuterium exchange reaction of DHA is also completely inhibited by the binding of D-threono-hydroxamic acid.

XI-catalyzed isomerization of DGA

The XI-catalyzed isomerization of $[2-^3H]$ -D-glucose in water at pH 7 proceeds with transfer of tritium from substrate to solvent about once every 10^9 substrate turnovers to form product (13). This is consistent with intramolecular hydride transfer in conversion of the α -hydroxy aldehyde to the α -hydroxy ketone, and a very rare competing enzyme-catalyzed proton transfer reaction. The improbable alternative proton transfer mechanism would require an extraordinary level of shielding of a putative catalytic side-chain that shuttles the proton between the two reacting carbon atoms (13). By contrast, the XI-catalyzed isomerization of DGA to give DHA in D_2O reported here proceeds with incorporation of 0.10 and 0.05 mole fraction of deuterium for the reactions at pD 7.0 and 8.0, respectively (Figure 1B), so that at least 5 – 10% of the reaction of the truncated substrate proceeds by a proton transfer mechanism (Scheme 4). We also find that XI catalyzes the slow transfer of deuterium from solvent D_2O into DHA to give *d*-DHA (k_{ex} , Scheme 4). The XI-catalyzed reactions of both

DGA and DHA are abolished by the binding of the competitive inhibitor D-threono-hydroxamic acid (28).

We conclude that turnover of the minimal substrate DGA by XI proceeds by the competing hydride and proton transfer pathways (Scheme 4) previously observed for the nonenzymatic isomerization of DGA in aqueous solution (19, 20). Therefore, the interaction between the terminal CH(OH)CH₂OH fragment of D-xylose and XI appears to provide a specific stabilization of the transition state for enzyme-catalyzed isomerization with intramolecular hydride transfer, so that the truncation of this fragment results in a large increase in the relative rate of the reaction with proton transfer. We suggest that the very rare tritium exchange reaction of [2-³H]-D-glucose and the isomerization of DGA with transfer of deuterium from solvent to product DHA catalyzed by XI proceed by adventitious deprotonation of bound substrate by a basic residue at the enzyme active site. There is ample precedent for nonspecific protein-catalyzed deprotonation of α -carbonyl carbon with second-order rate constants similar to that reported here for XI. The value of $k_{\text{cat}}/K_{\text{m}} = 3 \times 10^{-3} \text{ M}^{-1} \text{ s}^{-1}$ for the XI-catalyzed isomerization of DGA by proton transfer to give *d*-DHA (Table 1) is 10-fold larger than $k_{\text{N}} = 3.2 \times 10^{-4} \text{ M}^{-1} \text{ s}^{-1}$ for nonenzymatic deprotonation of DGA by the tertiary amine 3-quinuclidinone (Table 3) (26), but it is similar to $k_{\text{cat}}/K_{\text{m}} = 0.013$ and $3 \times 10^{-3} \text{ M}^{-1} \text{ s}^{-1}$ determined for the deuterium exchange reactions of glycolaldehyde in D₂O catalyzed by bovine serum albumin and the crippled K12G mutant of yeast TIM, respectively (8, 39).

There is a decrease in the fractional yield of *d*-DHA from the XI-catalyzed isomerization of DGA in D₂O from 9.5% to 5% on moving from pD 7.0 to 8.0 (Figure 1B), but this increase in pD results in no significant change in the value of $k_{\text{cat}}/K_{\text{m}}$ for the overall reaction that gives *h*-DHA as the major product (Table 1). Therefore, the rate-determining step for the overall isomerization reaction is pD-independent in this range, but the dependence on pD of the steps that determine the relative yields of *h*-DHA and *d*-DHA are different. We are unable to provide a simple explanation for these data.

DGA is a competitive inhibitor of the XI-catalyzed isomerization of D-xylose in D₂O with $K_{\text{i}} = 3.0 \text{ mM}$ that is similar to $K_{\text{m}} = 4.9 \text{ mM}$ for isomerization of D-xylose under the same conditions (Table 1). Therefore, the addition of an ethylene glycol fragment to DGA to give D-xylose does not change the *apparent* stability of the complex with XI. The observation that the addition of this fragment to DGA results in a large increase in $k_{\text{cat}}/K_{\text{m}}$ for XI-catalyzed isomerization by hydride transfer from 0.034 to 490 $\text{M}^{-1} \text{ s}^{-1}$ (Table 1) shows that its interactions with XI are expressed only in the transition state and are therefore utilized to stabilize the linear sugar in the transition state relative to the cyclic sugar in the Michaelis complex.

The data in Table 3 show that at pD 7.0 TIM provides only a 50-fold rate acceleration of the reference nonenzymatic deprotonation of DGA by the small Brønsted base catalyst 3-quinuclidinone in D₂O (26), while XI provides a somewhat larger 2100-fold rate acceleration of the reference nonenzymatic Zn²⁺-catalyzed isomerization reaction of DGA that is promoted by lyoxide ion shown in Scheme 5 (20). The similar observed catalytic activities of XI ($k_{\text{cat}}/K_{\text{m}} = 0.034 \text{ M}^{-1} \text{ s}^{-1}$) and TIM ($k_{\text{cat}}/K_{\text{m}} = 0.017 \text{ M}^{-1} \text{ s}^{-1}$) towards isomerization of the common minimal substrate DGA by hydride transfer and proton transfer, respectively (Table 3), therefore requires that the difference in the kinetic parameters for catalysis of the isomerization of the whole substrates D-xylose and DGAP, respectively, by these enzymes results from different contributions of the intrinsic binding energy of the nonreacting portions of the substrate to stabilization of the transition state (40).

Insights from Crystallography

TIM shows a large selectivity for the binding and reaction of the reactive carbonyl form of DGAP (27). This probably reflects the snug fit of ligands at the active loop-closed enzyme (41, 42) and the exclusion of the sterically demanding hydrated substrate. By contrast, we report here that the X-ray crystal structure of the complex obtained by soaking XI in a solution that contains 50 mM total DGA shows that the ligand is bound as DGA hydrate (DGAH, Figure 3). Since XI forms an initial Michaelis complex with the cyclic form of glucose/xylose (14, 43, 44), there is no obvious requirement that the enzyme show selectivity towards binding the carbonyl form of the truncated substrate DGA. It would appear that this selectivity for the reactive carbonyl form of substrate is provided by the subsequent development of strong interactions of the enzyme with the C-4 and C-5 hydroxyls of the linear form of the full substrate D-xylose in the transition state (see above). While the binding mode observed herein for the DGAH/formate complex is strikingly similar to that of the cyclic form of glucose (PDB ID 1XIF; Figure 4a), it is a closer match to that of the linear xylulose (PDB ID 1XII; Figure 4b). Since these small fragments can adopt multiple binding orientations in the active site, this finding highlights a preference of XI for binding the linear forms of pentose/hexose sugars.

The structure of XI with bound hydrated triose does not represent a productive complex due to both the presence of the hydrate form and the orientation of C1/C2 with respect to the catalytic metals (C2 hydroxyl group does not bridge M1 and M2 as expected for the hydride transfer mechanism). However, the DGAH structure does show that binding of the triose substrate can induce conformational changes in active site residues that are remarkably similar to those observed in structures with bound pentose and hexose substrates (Figure 4). Specifically, the X-ray structures demonstrate that in addition to mobility of the M2 metal center, the catalytic residues Glu181, Glu186, Asp255, and Asp257 adopt alternate conformations upon substrate binding compared to those observed in unliganded XI. This finding is also consistent with EPR spectroscopy studies on XI which show that binding of the cyclic form of substrates or substrate analogue inhibitors is sufficient to induce the shift of the metal ion in this site (45). A recent joint X-ray and neutron diffraction study shows that the catalytic metal, M2, occupies two sites after ring-opening and before isomerization (46). Thus, the metal ion movement may well be independent and not necessarily coupled to the isomerization reaction hypothesized to take place from the linear form of the sugar.

Rate acceleration and intrinsic binding energies for XI-catalyzed isomerization

The linear open form of D-xylose and the free carbonyl form of DGA are expected to show a similar *intrinsic* chemical reactivity in nonenzymatic Zn^{2+} -catalyzed isomerization by hydride transfer in aqueous solution. However, the apparent rate constant for Zn^{2+} -catalyzed isomerization of D-xylose should be ca. 500-fold smaller than the observed rate constant for isomerization of DGA at the same pD (Scheme 5). This is because the fraction of D-xylose present as the reactive linear sugar (0.01%, Scheme 6, (47)) is 500-fold smaller than the fraction of DGA present as the reactive free aldehyde (5% (26)). A 500-fold correction of the apparent second-order rate constant for nonenzymatic Zn^{2+} -catalyzed isomerization of DGA by hydride transfer at pD 7.0, $k_{\text{Zn}} = k_{\text{T}}[\text{DO}^-] = 1.6 \times 10^{-5} \text{ M}^{-1} \text{ s}^{-1}$ (Scheme 5), gives an estimated rate constant of $k_{\text{Zn}} = 3.2 \times 10^{-8} \text{ M}^{-1} \text{ s}^{-1}$ for the Zn^{2+} -catalyzed isomerization of D-xylose under the same conditions (Table 3). This can then be combined with $k_{\text{cat}}/K_{\text{m}} = 490 \text{ M}^{-1} \text{ s}^{-1}$ for the XI-catalyzed isomerization of D-xylose to give an estimated enzymatic rate acceleration for isomerization of D-xylose of 1.5×10^{10} -fold (Table 3).

The data for the XI-catalyzed reaction of DGA allow us to portion the 1.5×10^{10} -fold rate acceleration for the XI-catalyzed isomerization of D-xylose into a smaller 2100-fold rate acceleration for reaction of the truncated substrate DGA (Table 3) and a much larger 7 x

10^6 -fold rate acceleration that is due to *specific* stabilization of the transition state for isomerization by interactions between XI and the terminal ethylene glycol fragment of D-xylose. This corresponds to an apparent 9.3 kcal/mol intrinsic binding energy for this substrate fragment. By comparison, a larger ca. 12 kcal/mol intrinsic binding energy is estimated for the nonreacting substrate phosphodianion group in the TIM-catalyzed isomerization of DGAP (26, 48). The larger phosphodianion binding energy underscores the greater strength of electrostatic interactions between fully charged species, such as between the phosphodianion of DGAP and the alkylammonium side chain of Lys-12 at TIM (8, 49), compared with interactions with neutral hydrogen bond donors/acceptors.

The 9.3 kcal/mol intrinsic binding energy of the terminal ethylene glycol fragment of D-xylose is similar to that observed for other enzymatic reactions of sugars. For example, intrinsic transition state binding energies of 4 – 6 kcal/mol per sugar hydroxyl group have been reported for enzyme-catalyzed glycosyl transfer reactions (50). An unusually large 10 kcal/mol intrinsic binding energy was reported for the galactosyl C-2 hydroxyl in the breakdown of the covalent β -D-galactopyranosyl intermediate of β -galactosidase-catalyzed glycosyl transfer reactions (51).

The structure of XI determined by the time-of-flight neutron Laue method has been used to identify the location of the hydrogen atoms of enzyme-bound D-xylulose (52, 53). The neutron diffraction and X-ray crystal structures (15, 16, 25, 54) of XI show that the C-4 hydroxyl of substrate interacts with the metal ion designated as M1, and that the C-5 hydroxyl is hydrogen bonded to N_{ε2} of His54. The neutron diffraction structure requires a surprising perturbation in the relative pK_as for the C-5 sugar hydroxyl and N_{ε2} of His54 that results in formal proton transfer from the substrate oxygen to the protein imidazole. This suggests that the complex of the O-ionized sugar is stabilized by a strong, single potential minimum hydrogen bond (55, 56) to the cationic N_{ε2} of His54 (53).

Ground state and transition state effects

The value of $K_m = 4.9$ mM for the XI-catalyzed isomerization of D-xylose at pD 7.0 is not significantly different from $K_m \approx K_i = 3.0$ mM for DGA (Table 1). This shows that the binding interactions between XI and the terminal ethylene glycol fragment of D-xylose are *not* expressed in the Michaelis complex with D-xylose. Rather, these interactions develop on proceeding from the Michaelis complex between XI and the cyclic sugar to the transition state for sugar isomerization, and are expressed in the much larger value of $k_{cat} = 2.4$ s⁻¹ for D-xylose compared with $k_{cat} = 9.0 \times 10^{-5}$ s⁻¹ for XI-catalyzed isomerization of DGA by hydride transfer (Table 1).

There is a normal primary kinetic deuterium isotope effect on the XI-catalyzed isomerization of [2-²H]-D-glucose which shows that the ring-opening of glucose is a fast and reversible step that precedes rate-determining intramolecular hydride transfer (57, 58). A simple explanation for the observed substituent effects on k_{cat} and K_m is that the interactions between XI and the C-4 hydroxyl and the C-5 ring oxygen of D-xylose are minimal in the ground state Michaelis complex with the cyclic sugar, and develop only after ring opening of D-xylose to form the linear sugar. This must be the case for the hydrogen bond between the C-5 hydroxyl of the linear sugar and the cationic N_{ε2} of His54 (53). We propose that the interaction between the C-4 hydroxyl of D-xylose and the metal ion designated as M1 is also weak or absent at the *productive* complex with the cyclic sugar. If the interactions between XI and the C-4 and C-5 hydroxyl develop only upon conversion of cyclic D-xylose to the linear form, then the affinity of the linear sugar for XI will be much higher than that of the cyclic form of D-xylose. In fact, linear D-xylitol typically shows a higher affinity for XI than does D-xylose (22, 59). For example, the value of $K_i = 0.026$ mM for inhibition of XI-catalyzed [*Streptomyces* sp. (A.T.C.C. 21132)] isomerization by xylitol

at pH 7.1 is more than 100-fold smaller than $K_m = 3.1$ mM for the isomerization of D-xylose (22).

XI and TIM use different strategies to obtain an increase in k_{cat} through the utilization of enzyme-ligand binding interactions to obtain specific transition state stabilization relative to the Michaelis complex. In the case of XI, the substrate ring-opening is a step on the pathway from the Michaelis complex to the transition state, and the new binding interactions that develop with the open linear sugar are expressed as an increase in k_{cat} . Similarly, there is good evidence that a tightening of the electrostatic interactions between the cationic side chain of Lys-12 and the bound substrate DGAP on proceeding from the Michaelis complex (substrate dianion) to the transition state (enediolate-like trianion) result in a ca. 10^5 -fold increase in k_{cat} for TIM-catalyzed isomerization (8).

SUMMARY AND CONCLUSIONS

XI achieves its catalytic power through the summation of several stabilizing interactions between the protein catalyst and the substrate pieces. The enzyme provides a 2100-fold increase in the second-order rate constant for metal-ion-catalyzed isomerization by hydride transfer in D₂O. The K_m of 3 mM for the XI-catalyzed isomerization of DGA is consistent with a modest stabilization of the Michaelis complex. However, the X-ray crystal structure of this complex shows the triose bound as the unreactive hydrate (DGAH). If the dominant XI•DGAH complex is nonproductive, then the binding of DGA as the productive Michaelis complex must be relatively weak, and the weak binding compensated by effective protein activation of catalysis by the metal cation. The transition state for aldose-ketose isomerization of D-xylose is stabilized by 9.3 kcal/mol by interactions of XI with the C-4 and C-5 hydroxyls of substrate. This significant, but not extraordinarily large, interaction can be rationalized by an examination of the neutron diffraction (52, 53) and X-ray crystal structures (15, 16, 25, 54) of XI•sugar complexes. The larger kinetic parameters for catalysis by TIM compared with XI are due to: (i) The smaller catalytic burden borne by TIM, which must overcome only a 1.7 kcal/mol barrier to dehydration of DGAP to give the free carbonyl form, compared with XI, which must overcome a 5.4 kcal/mol barrier for ring-opening of D-xylose. (ii) The larger ca. 12 kcal/mol intrinsic binding energy for the phosphodianion fragment of DGAP, compared with the 9.3 kcal/mol binding energy of the terminal ethylene glycol fragment of D-xylose (Table 3).

The major impediment to perfection in enzymatic catalysis of isomerization of D-xylose is the evolution of an enzyme active site capable of accommodating the very different structures of the cyclic (99.99%) and linear (0.01%) forms of D-xylose (47). The higher affinity of XI for linear xylitol compared with the cyclic sugars suggests that this catalyst has evolved to optimize binding interactions with linear sugars, while the large K_m of 4.9 mM for D-xylose in D₂O (Table 1) shows that cyclic sugars bind relatively weakly. It is interesting that the apparent $k_{cat}/K_m = 490 \text{ M}^{-1} \text{ s}^{-1}$ for XI-catalyzed isomerization of D-xylose in D₂O (Table 1) and the fraction of sugar present in the linear form (10^{-4}) are consistent with a value of $k_{cat}/K_m = 4.9 \times 10^6 \text{ M}^{-1} \text{ s}^{-1}$ for direct catalysis of the isomerization of *linear* D-xylose. This rate constant is not far removed from that for a diffusion-controlled reaction, such as that carried out by TIM.

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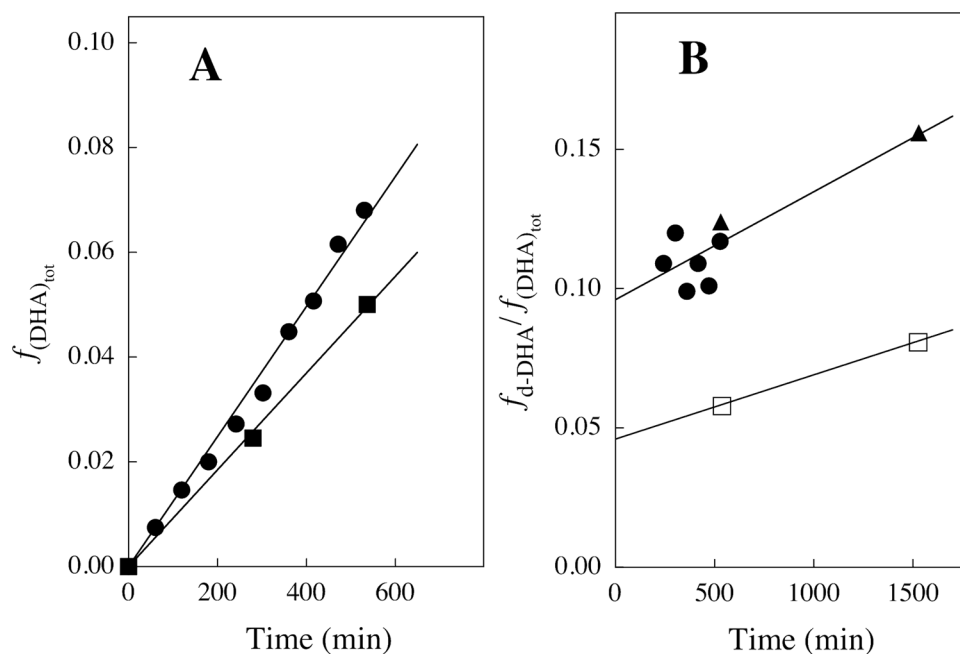


Figure 1.

Data for the XI-catalyzed isomerization reaction of 11 mM DGA to give DHA in D_2O in the presence of 24 mM imidazole buffer and 10 mM MgCl_2 at 25 °C and $I = 0.1$ (NaCl) monitored by ^1H NMR spectroscopy. A. The time course for the fractional formation of *total* DHA where $(f_{\text{DHA}})_{\text{tot}}$ was determined from the area of the signals for the products *h*-DHA and *d*-DHA and the remaining substrate using eq 1: (●) Reaction catalyzed by 0.30 mM XI at pD 7.0; (■) Reaction catalyzed by 0.20 mM XI at pD 8.0. B. The change with time in the observed fractional yield of *d*-DHA (see eq 4): (●) Reaction catalyzed by 0.30 mM XI at pD 7.0; (▲) Reaction catalyzed by 0.27 mM XI at pD 7.0; (□) Reaction catalyzed by 0.20 mM XI at pD 8.0.

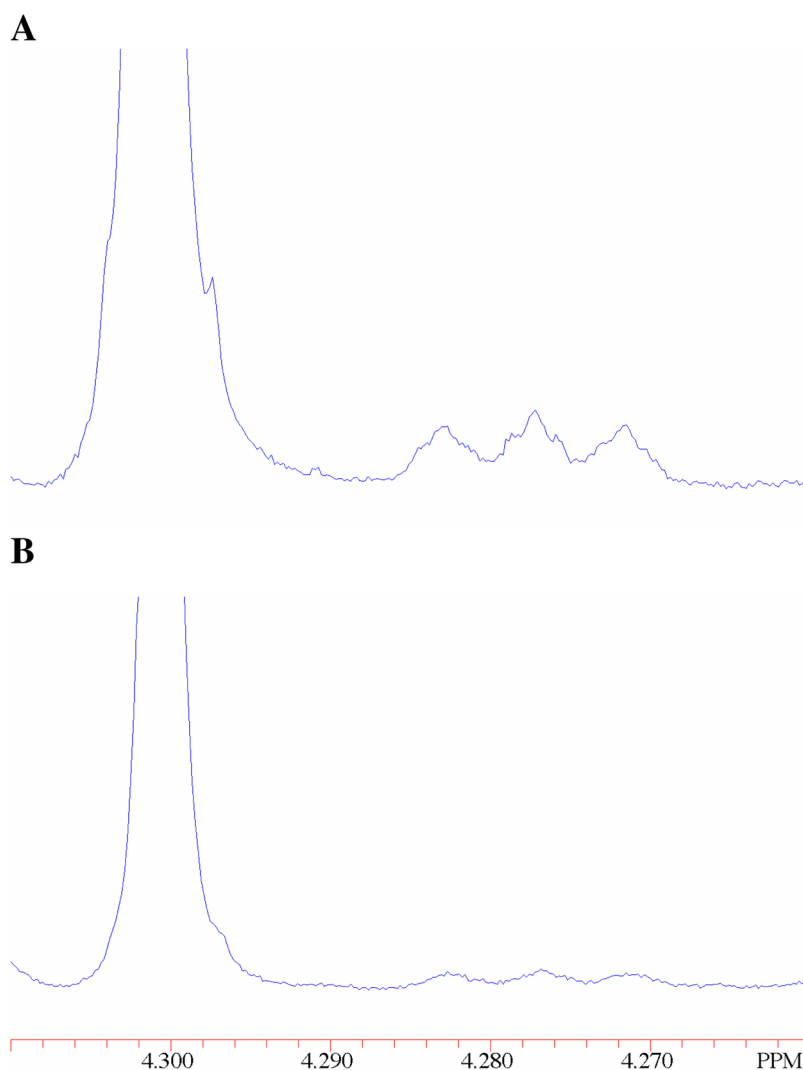
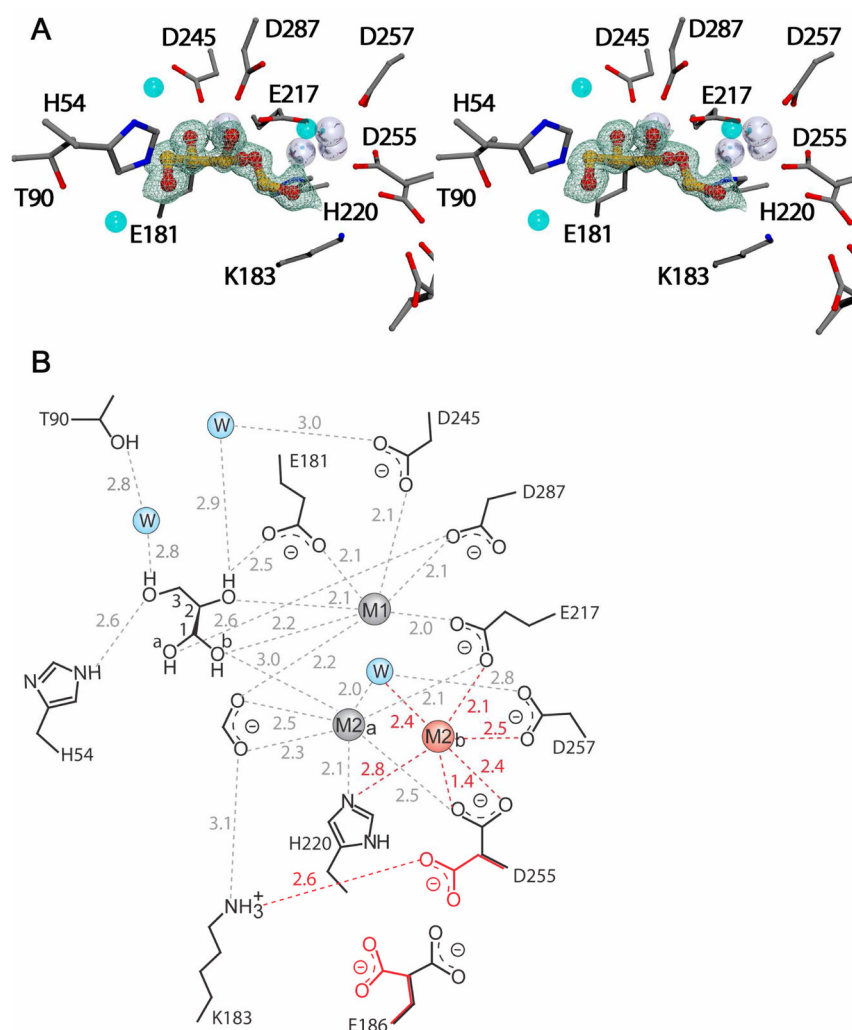


Figure 2.

Representative partial ^1H NMR spectra at 500 MHz obtained during the XI-catalyzed isomerization reaction of 11 mM DGA in D_2O in the presence of 24 mM imidazole buffer and 10 mM MgCl_2 at 25 °C and $I = 0.1$ (NaCl). The large singlet at 4.300 ppm is the signal for the two $\alpha\text{-CH}_2\text{OD}$ groups of *h*-DHA and the single $\alpha\text{-CH}_2\text{OD}$ group of *d*-DHA. The upfield-shifted triplet at 4.276 ppm is the signal for the single $\alpha\text{-CHDOD}$ group of *d*-DHA. A. Reaction catalyzed by 0.30 mM XI at pD 7.0. The area of the triplet is 2.5 % that of the singlet. B. Reaction catalyzed by 0.20 mM XI at pD 8.0. The area of the triplet is 1.2 % that of the singlet.

**Figure 3.**

The structure of the XI active site complexed with DGAH. A. Shown as a stereo-view with the [Fo]-[Fc] simulated annealing omit electron density map shown as grey wireframe, the metal ions as silver spheres, and the water molecules as blue spheres. These figures were generated with the POVSCRIPT+ modification of the MOLSCRIPT program (60) and POV-Ray (61) (<http://www.povray.org/>). B) A diagram of the structure which shows the waters as blue spheres and the hydrogen bonds shown as dashed lines labeled with distances in Å. The atoms and corresponding distances in red indicate those with partial occupancy.

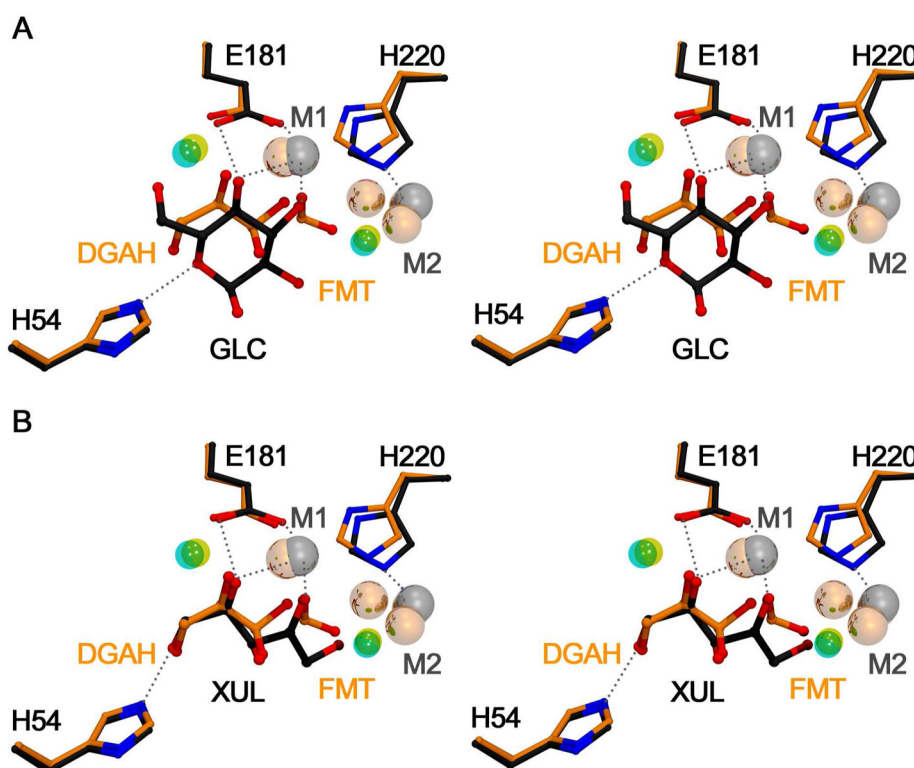
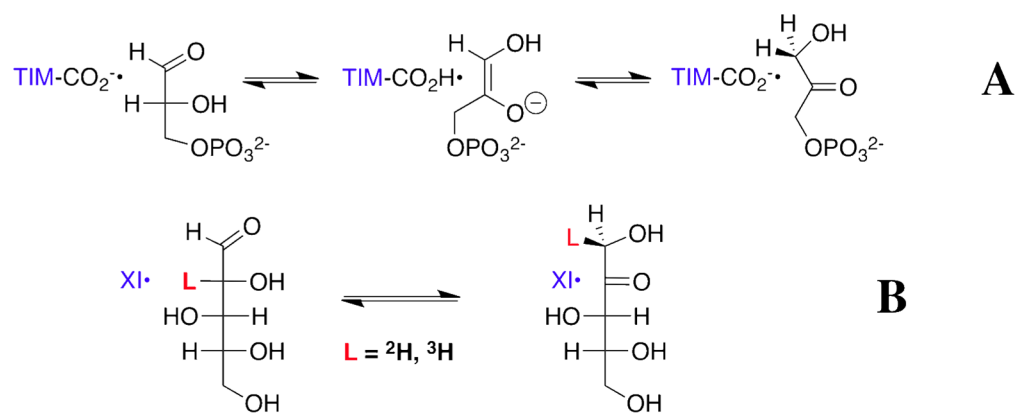
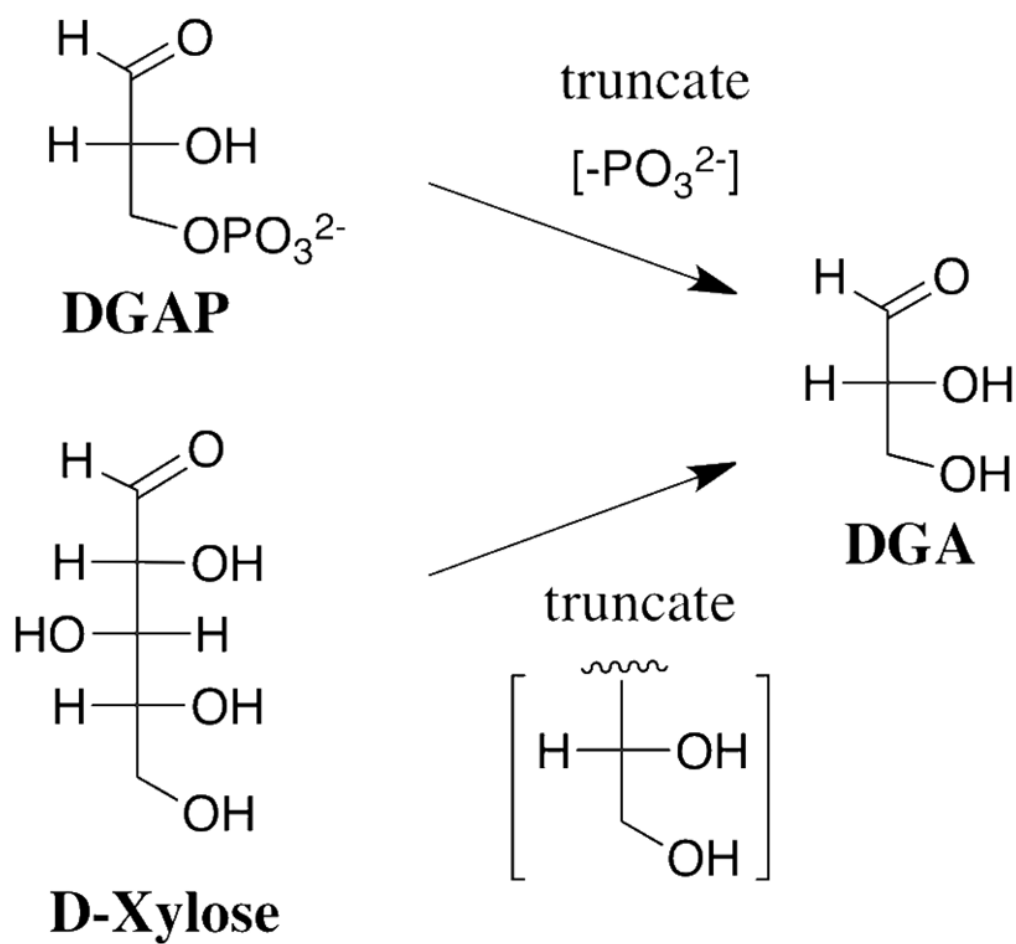


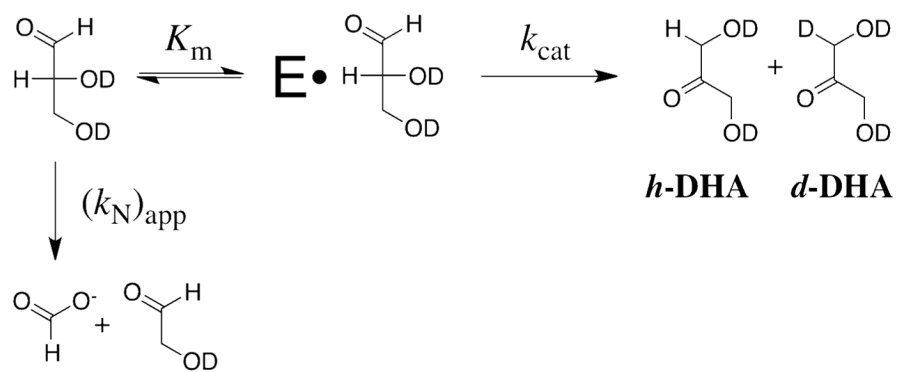
Figure 4. Overlays of the structure of XI complexed with DGAH/formate/ Mg^{2+} (DGAH and FMT, yellow backbone, Mg^{2+} as light grey spheres, H_2O as cyan spheres) with those of: (A) the cyclic form of D-glucose (GLC, grey backbone, Mg^{2+} as dark grey spheres, H_2O as yellow sphere, PDB ID 1XIF); and, (B) the linear form of D-xylulose (XUL, grey backbone, Mg^{2+} as dark grey spheres, H_2O as yellow sphere, PDB ID 1XII). The hydrogen and coordination bonds are shown for the D-glucose and D-xylulose ligands as grey dashed lines. Models were superimposed with the SSM algorithm as implemented in COOT. The Figures prepared using MOLSCRIPT (60).



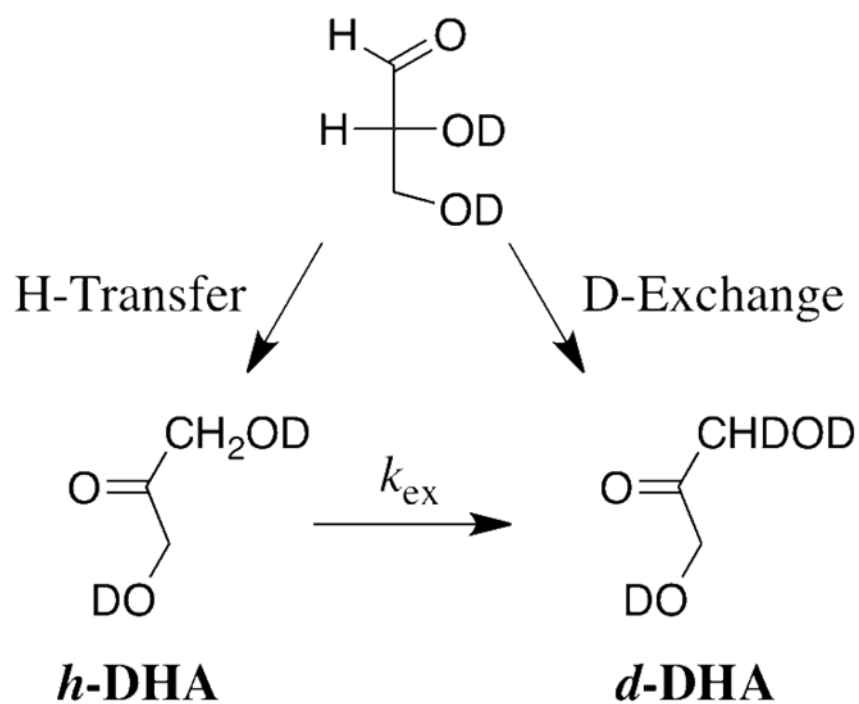
Scheme 1.



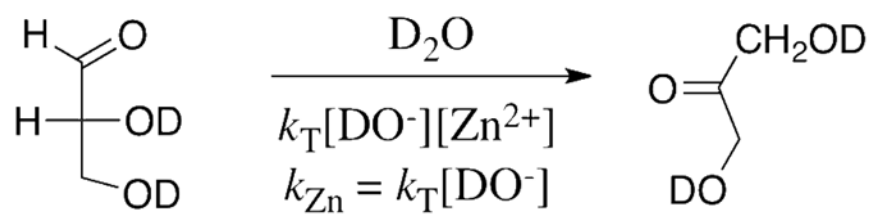
Scheme 2.



Scheme 3.



Scheme 4.



Scheme 5.

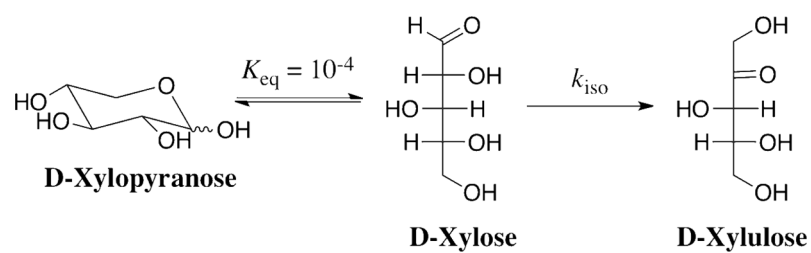
**Scheme 6.**

Table 1

Kinetic Parameters for Isomerization of D-Glyceraldehyde (DGA) and D-Xylose Catalyzed by D-Xylose Isomerase from *Streptomyces rubiginosus* in D₂O at 25 °C.^a

Substrate	Pathway	k_{cat} (s ⁻¹)	K_{m} (M)	$k_{\text{cat}}/K_{\text{m}}$ (M ⁻¹ s ⁻¹)
DGA	Total	1.0×10^{-4}	3×10^{-3}	0.034 (0.035) ^b
DGA ^c	Hydride Transfer	9×10^{-5}	3×10^{-3}	0.031
DGA ^d	Proton Transfer	1×10^{-5}	3×10^{-3}	3×10^{-3}
D-Xylose	Hydride Transfer	2.4	4.9×10^{-3}	490

^a At pD 7.0 in the presence of 24 mM imidazole buffer and 10 mM MgCl₂ at $I = 0.1$ (NaCl), unless noted otherwise. Rate constants are reported for the reaction of *total* DGA or D-xylose.

^b Data for reaction at pD 8.0.

^c Kinetic parameters for the reaction with intramolecular transfer of hydrogen to form *h*-DHA, calculated from the observed kinetic parameters and the 90% fractional yield of *h*-DHA.

^d Kinetic parameters reaction with transfer of deuterium from solvent to product to form *d*-DHA, calculated from the observed kinetic parameters and the 10% fractional yield of *d*-DHA.

Table 2

Statistics from X-Ray Data Collection and Refinement.

Data Collection Statistics	DGAH
Resolution (Å) (last shell) ^a	20.00-0.97 (1.00-0.97)
Wavelength (Å)	0.9399
No. of reflections	
Observed	7,151,163
Unique	272,713
Completeness (%) ^a	99.7 (96.7)
R _{merge} (%) ^{a,b}	5.5 (39.4)
Redundancy	8.5 (5.4)
I/σ(I) ^a	39.5(3.4)
Refinement Statistics	
No. of reflections in working set	262,140
No. of reflections in test set	13,001
R _{cryst} (R _{free})	0.116 (0.125)
No. of non-hydrogen atoms	3,206
No. of solvent atoms	523
Average B-factor (Å ²)	
Protein atoms	11.0
DHA/Formate	8.7/8.7
Mg/Mn(II) ions	12.6
Solvent	25.1
RMS deviations	
Bond lengths (Å)	0.007
Bond angles (°)	1.329

^aData for the highest resolution shell are given in parenthesis.^b $R_{\text{merge}} = \frac{\sum_{\text{hkl}} \sum_i |I_{\text{hkl},i} - \langle I_{\text{hkl}} \rangle|}{\sum_{\text{hkl}} \sum_i I_{\text{hkl},i}}$, where $\langle I_{\text{hkl}} \rangle$ is the mean intensity of the multiple $I_{\text{hkl},i}$ observations for symmetry-related reflections.

Enzymatic Rate Accelerations for Turnover of the Whole Substrates and the Common Substrate Fragment D-Glyceraldehyde by D-Xylose Isomerase and Triosephosphate Isomerase.

Table 3

Catalyst	Substrate	Pathway	k_N^a ($M^{-1} s^{-1}$)	k_{cat}/K_m ($M^{-1} s^{-1}$)	Rate Effect ^b	Fragment Binding Energy
Zn^{2+}	DGA		$1.6 \times 10^{-5} c$			
Zn^{2+}	D-Xylose		$3.2 \times 10^{-8} d$			
XI	DGA	Hydride Transfer		0.034 ^e	2100	9.3 kcal/mol ^f
XI	D-Xylose			490 ^e	1.5×10^{10}	
Buffer ^g	DGA		$3.2 \times 10^{-4} h$			
TIM	DGA	Proton Transfer		0.017 ^h	50	
TIM	DGAP			$9.6 \times 10^6 i$	3×10^{10}	11.9 kcal/mol ^j

^a Second-order rate constants for the reference *nonenzymatic* isomerization reactions of *total* DGA or D-xylose in D₂O at pD 7.0 and 25 °C.

^b The enzymatic rate acceleration calculated as the ratio of the second-order rate constants for the enzyme-catalyzed reaction and the reference nonenzymatic reaction catalyzed by Zn^{2+} or the Brønsted base 3-quinuclidinone in aqueous solution.

^c Calculated as $k_{Zn} = k_T[DO^-]$ where $k_T = 950 M^{-2} s^{-1}$ is the third-order rate constant for the lyoxide-ion promoted Zn^{2+} -catalyzed isomerization of *total* DGA and $[DO^-] = 1.7 \times 10^{-8} M$ at pD 7.0 (Data from Ref. 20).

^d Calculated from k_{Zn} for isomerization of DGA with a 500-fold correction for the smaller fraction of D-xylose than of DGA present in the reactive carbonyl form. We make the assumption that the free carbonyl forms of DGA and D-xylose have similar reactivities in Zn^{2+} -catalyzed isomerization by hydride transfer (see text).

^e Data at pD 7.0 from Table 1.

^f The intrinsic binding energy of the terminal CH(OH)CH₂OH fragment of D-xylose, calculated from the ratio of the enzymatic rate accelerations for the XI-catalyzed reactions of D-xylose and DGA.

^g Data for nonenzymatic deprotonation of DGA by the tertiary amine base 3-quinuclidinone. We make the assumption that DGA and DGAP have similar reactivities in nonenzymatic proton transfer.

^h Data from Ref. 26.

ⁱ Calculated from $k_{cat} = 4300 s^{-1}$ and $K_M = 0.45 mM$ for turnover of *total* DGAP by rabbit muscle TIM in H₂O at pH 7.5 and 25 °C (Ref. 26).

^j The intrinsic binding energy of the phosphodianion group of DGA, calculated from the ratio of the enzymatic rate accelerations for the TIM-catalyzed reactions of DGAP and DGA.