

Structural and mechanistic insight into covalent substrate binding by *Escherichia coli* dihydroxyacetone kinase

Rong Shi^a, Laura McDonald^b, Qizhi Cui^c, Allan Matte^c, Miroslaw Cygler^{a,c}, and Irena Ekiel^{b,c,1}

^aDepartment of Biochemistry, McGill University, Montréal, QC, Canada H3G 1Y6; ^bDepartment of Chemistry and Biochemistry, Concordia University, Montréal, QC, Canada H4B 1R6; and ^cBiotechnology Research Institute, National Research Council of Canada, 6100 Royalmount Avenue, Montréal, QC, Canada H4P 2R2

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The *Escherichia coli* dihydroxyacetone (Dha) kinase is an unusual kinase because (i) it uses the phosphoenolpyruvate carbohydrate: phosphotransferase system (PTS) as the source of high-energy phosphate, (ii) the active site is formed by two subunits, and (iii) the substrate is covalently bound to His218^{K*} of the DhaK subunit. The PTS transfers phosphate to DhaM, which in turn phosphorylates the permanently bound ADP coenzyme of DhaL. This phosphoryl group is subsequently transferred to the Dha substrate bound to DhaK. Here we report the crystal structure of the *E. coli* Dha kinase complex, DhaK–DhaL. The structure of the complex reveals that DhaK undergoes significant conformational changes to accommodate binding of DhaL. Combined mutagenesis and enzymatic activity studies of kinase mutants allow us to propose a catalytic mechanism for covalent Dha binding, phosphorylation, and release of the Dha-phosphate product. Our results show that His56^K is involved in formation of the covalent hemiaminal bond with Dha. The structure of H56N^K with noncovalently bound substrate reveals a somewhat different positioning of Dha in the binding pocket as compared to covalently bound Dha, showing that the covalent attachment to His218^K orients the substrate optimally for phosphoryl transfer. Asp109^K is critical for activity, likely acting as a general base activating the γ -OH of Dha. Our results provide a comprehensive picture of the roles of the highly conserved active site residues of dihydroxyacetone kinases.

DhaK–DhaL complex | kinase mechanism

Dihydroxyacetone phosphate (Dha-P), an intermediate for the synthesis of pyruvate, can be generated in bacteria through glycerol fermentation by the sequential actions of glycerol dehydrogenase forming dihydroxyacetone and Dha kinase generating Dha-P. Dha kinases are divided into two classes, those using ATP as the phosphoryl donor or those using the PTS-dependent DhaM protein to provide the phosphoryl group (1). The ATP-dependent Dha kinases are present in animals, plants, and some bacteria and consist of a single chain containing two domains, whereas the PTS-dependent Dha kinases exist only in bacteria and are composed of two subunits. The most studied representative of the PTS-dependent family is the *Escherichia coli* Dha kinase, which is composed of DhaK and DhaL subunits. DhaM, the phosphotransferase subunit of Dha kinase, is phosphorylated by the small phospho-carrier protein HPr of the PTS (1). Phosphorylated DhaM then forms a complex with the ADP-bound DhaL subunit and transiently phosphorylates ADP to ATP (1). The ATP-loaded DhaL subsequently associates with the DhaK subunit containing the Dha substrate covalently bound to His218^{K*} through a hemiaminal bond (2). Finally, ATP bound to DhaL, serving as a coenzyme (3), transfers phosphate to Dha yielding Dha-P.

Crystal structures for the two individual subunits of the *E. coli* PTS-dependent Dha kinase—DhaK (2) and DhaL (4)—have been previously determined, as well as the structure of an ATP-dependent single chain, two-domain Dha kinase from *Citrobacter*

freundii (5), which has 30% sequence identity to the *E. coli* kinase. Both ATP- and PTS-dependent Dha kinases display high structural similarity of their corresponding domains/subunits and are structurally distinct from other kinases. The active site residues are highly conserved between bacteria and eukaryotes reflecting the unique phosphotransfer mechanism. The structure of the homodimeric, ATP-dependent *C. freundii* Dha kinase shows swapped K- and L-domains, with bound Dha and an ATP analogue separated by approximately 14 Å (5). It is therefore unlikely that this structure represents a phosphotransfer-competent state of this enzyme, leaving the question of the mechanism unanswered. In addition to the structures of DhaK and DhaL, the structure of the *Lactococcus lactis* DhaM domain has been reported as well as its complex with DhaL, providing insight into phosphotransfer from phosphohistidine to ADP (6).

We have determined the crystal structure of the *E. coli* DhaK–DhaL complex bound to ADP and Dha at 2.2-Å resolution, providing a snapshot of the final step of phosphoryl transfer from ATP to Dha. Site-directed mutagenesis, activity measurements, and structural data indicate that His56^K is important for formation of the covalent hemiaminal bond. We show further that the covalent attachment of Dha to His218^K optimally positions the substrate for phosphoryl transfer but is likely dispensable for enzyme activity. Together, these data allow us to propose a general catalytic mechanism for dihydroxyacetone kinases, consistent with the available structural and biochemical data.

Results and Discussion

Structure of the DhaK–DhaL Complex Reveals Conformational Rearrangements upon Complex Formation. Each asymmetric unit contains one molecule each of DhaK and DhaL with all residues well defined in the electron density map. The biological unit of the complex is a heterotetramer containing a central DhaK dimer and two molecules of DhaL (2:2 stoichiometry). Each DhaL subunit interacts with only one DhaK subunit of the dimer, with the two DhaL subunits being approximately 37 Å apart (Fig. 1A). An ADP molecule and two magnesium ions are bound to DhaL, whereas DhaK contains a Dha molecule covalently bound to His218^K (Fig. 1B).

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Data deposition: The atomic coordinates and structure factors have been deposited in the Protein Data Bank, www.pdb.org [PDB ID codes 3PNK (DhaK), 3PNL (DhaK–DhaL complex), 3PNM (DhaK H56A), 3PNO (DhaK H56N), and 3PNQ (DhaK H56N–Dha)].

*Superscript letter ^K (or ^L) denotes kinase subunit.

¹To whom correspondence should be addressed. E-mail: irena.ekiel@nrc.ca.

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domain swapping in *C. freundii* (Fig. S1A). Importantly, and in contrast to the *C. freundii* Dha kinase structure, in the *E. coli* structure, the orientation of DhaL relative to DhaK is consistent with a direct transfer of the γ -phosphoryl moiety from ATP to the γ -OH group of Dha (Fig. 1B).

We have also crystallized *E. coli* DhaK in a different crystal form (Table S1) than that previously reported (2). Interestingly, in our free DhaK structure, the loop $\beta 6^K/\alpha 4^K$ (residues 138–145) displays a conformation intermediate between that of free DhaK (PDB ID code 1O12) and that of DhaL-bound DhaK (see above), providing a snapshot of the movement of this loop (Fig. S2). The flexibility of the loop $\beta 6^K/\alpha 4^K$ is pivotal for the ability of DhaK to bind DhaL, because conformations of this loop in both free DhaK structures are not compatible with complex formation. The difference between the two structures of free DhaK is likely caused by changes in packing environment.

The Dha Kinase Active Site Is Formed at the Interface of DhaL-DhaK Subunits. In order to bring DhaL-bound ADP in proximity to DhaK-bound Dha, the tip of the DhaL helical barrel containing loop $\alpha 7^L/\alpha 8^L$ (residues 175–185) that caps the nucleotide-binding site, dips into the cavity formed between the $\beta 6^K/\alpha 4^K$ and $\beta 8^K/\beta 9^K$ loops, displacing both loops from their locations observed in free DhaK (Fig. 2A). The distance between the β -phosphate oxygen of ADP and the γ -OH group of Dha is 4.8 Å, sufficient to accommodate the missing ATP γ -phosphate in our DhaK–DhaL(ADP) complex as shown by our model of DhaK–DhaL(ATP) (Fig. 2B).

Both subunits contribute residues to form the active site. The key residues in the Dha binding pocket, including His218^K covalently bound to Dha, Gly53^K, His56^K, Lys104^K, and Asp109^K, are in almost identical positions as in the free DhaK structure. Dha is further anchored by Arg178^L, whose guanidinium group hydrogen bonds to the γ -OH of Dha and likely stabilizes the trigonal bipyramidal transition state during phosphoryl transfer. All of these residues are highly conserved in bacterial Dha kinases. Mutations of His218^K in *E. coli* DhaK or Arg161^L in *L. lactis* DhaL (corresponding to Arg178^L in *E. coli*) result in inactive enzymes (2, 6).

Despite the lack of conformational changes in DhaL upon binding to DhaK, the adenine ring of ADP undergoes a *syn* to *anti* conversion (Fig. S3) when compared with the free DhaL structure (4). Our molecular dynamics (MD) simulations indicate that in the *syn* conformation, ADP has a potential steric clash of its adenine ring with Thr107^K (Fig. S4). Flipping of the adenine ring is accompanied by formation of three additional water-mediated H bonds, two between its purine N1 and the carbonyl group of Val117^L and the amide group of Ala123^L and the third between its purine N6 and the OH group of Thr107^K. In addition, one of the two Mg²⁺ ions (M2) in the ADP binding site of DhaL moves by approximately 1.0 Å compared to free DhaL and becomes a bridging element at the DhaK–DhaL interface (Fig. S3). Its coordination number increases from three to five with the addition of the carbonyl of Phe78^K and a water molecule.

His56^K and Asp109^K Are Important for Dha Kinase Activity. In order to evaluate the contributions of residues in the active site for activity, we performed site-directed mutagenesis and activity measurements for several mutant enzymes, focusing on the conserved residues His56^K, Asp109^K, His218^K, and Arg178^L. Gel filtration chromatography revealed that all of the DhaK mutant enzymes (H56A^K, H56N^K, D109A^K, D109N^K, and H218K^K) were able to form complexes with DhaL, while no complex formation between DhaK and R178E^L was observed (see *SI Text*). Two variants of His56^K, H56A^K and H56N^K, show a moderate decrease in $k_{\text{cat}}^{\text{app}}$ but at least a 40- to 300-fold increase in K_m^{app} (Table 1). Both D109A^K and D109N^K mutant enzymes showed no measurable activity (Fig. S5). Mutation of His218^K to either lysine (Fig. S5) (2)

Table 1. Kinetic constants of wild-type DhaK and His56^K mutants for Dha substrate

	WT	H56A ^K	H56N ^K
$k_{\text{cat}}^{\text{app}}$ (min ⁻¹)	507 ± 25	96.1 ± 2.0	223 ± 4.3
K_m^{app} (mM)	<0.008*	0.302 ± 0.005	2.20 ± 0.15
$k_{\text{cat}}^{\text{app}}/K_m^{\text{app}}$ (mM ⁻¹ min ⁻¹)	>6.3 × 10 ⁴	318	101

Standard deviations were calculated from three independent experiments. *Due to the limited sensitivity of the glycerol-3-phosphate dehydrogenase coupled assay for WT at low substrate concentrations the K_m^{app} could not be accurately determined.

or alanine (2) also abolishes activity, consistent with its role in binding and orienting the substrate. In addition to the coupled enzyme assay, similar results were obtained by NMR spectroscopy (Fig. S6).

Structures of DhaK H56N^K and H56A^K Reveal the Ability of Dha to Bind Noncovalently. To gain further insight into the reduced activity exhibited by the H56N^K and H56A^K mutant enzymes we determined their crystal structures. The overall structure of either the H56N^K or H56A^K enzyme is almost identical to that of the previously reported structure of wild-type DhaK (2), although they crystallized in a different packing environment (Table S1) with four subunits in the asymmetric unit. However, in contrast to the structures of wild-type DhaK, no electron density corresponding to the Dha substrate was observed in any of the subunits in the structures of these mutant enzymes (Fig. S7). These structures were immediately suggestive of a critical role for His56^K in hemiaminal bond formation with Dha. The H56N^K and H56A^K structures reveal very small conformational changes in the active site as compared to wild-type DhaK with covalently bound Dha. These differences include an approximately 35° rotation of the imidazole ring of His218^K, possibly resulting from the shorter Asn56^K and Ala56^K side chains in the mutant enzyme structures. The carboxylate group of Asp109^K also rotates approximately 35°, possibly due to the loss of its hydrogen bonding interaction with the hydroxyl groups of Dha.

To better understand how the H56N^K mutant enzyme could retain some activity in the absence of covalent substrate binding, we determined the crystal structure of H56N^K in the presence of Dha by supplementing the crystallization buffer with 10 mM Dha. These crystals contained four molecules in the asymmetric unit (Table S1). In one subunit there is clear electron density showing the presence of Dha bound in the active site (Fig. 3A). Although this non-covalently bound Dha molecule occupies the same binding pocket as in the covalently bound structure, its C2 and O2 atoms are displaced by approximately 2 Å and approximately 3 Å, respectively, as compared with covalently bound Dha. The absence of the covalent bond results in somewhat different interactions between Dha and the surrounding residues. The Dha carbonyl group is anchored via hydrogen bonds with the main chain amide group of Gly53^K; the hydroxyl group (α -OH) buried in the binding pocket is within H-bonding distance of Lys104^K(NZ) and His218^K(NE2), while the other exposed hydroxyl group (γ -OH) forms a hydrogen bond to Asp109^K(OD1) and Ser80^K(OG) (Fig. 3A). Although this γ -OH is still orientated toward the β -phosphate group of ADP, it is shifted approximately 1.7 Å away from its position in the covalently bound substrate (Fig. 3B). We surmise that this rearrangement contributes to the observed reduced catalytic efficiency of the mutants (Table 1). Another contribution to the lower activity of the H56N^K and H56A^K mutants is their weaker affinity for the substrate, as reflected by a significant increase in the K_m^{app} (Table 1). Although our upper estimate of K_m^{app} for Dha in wild-type DhaK is 8 μ M, the corresponding K_m^{app} is 302 μ M and 2.2 mM for H56A^K and H56N^K, respectively (Table 1). Despite the different positioning of Dha in the H56N^K

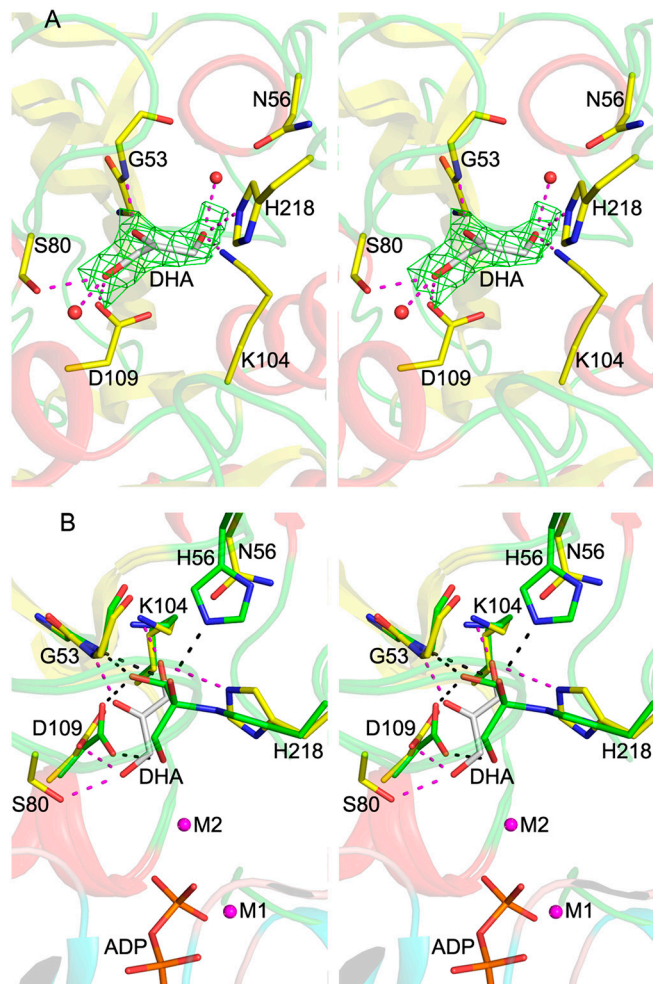


Fig. 3. Crystal structure of the DhaK H56N^K mutant. (A) Active site of the DhaK H56N^K mutant bound noncovalently to Dha. The DhaK molecule is shown in cartoon representation. Asn56^K and the residues involved in substrate binding are shown in stick mode. Hydrogen bonding interactions involving Dha are indicated by magenta dashed lines. Red spheres represent two water molecules. The difference ($F_o - F_c$) electron density map for Dha (omitted from refinement) contoured at 2.5σ is shown in green. (B) Substrate binding pocket of the DhaK H56N^K-Dha complex superimposed onto that of the DhaK-DhaL complex. The carbon atoms of the labeled residues in the DhaK H56N^K mutant and in the DhaK-DhaL complex are colored in yellow and green, respectively. The phosphate moieties of ADP as well as the two magnesium ions in the DhaK-DhaL complex are also shown. Hydrogen bonds involving the noncovalently and covalently bound Dha molecules are represented by magenta and black dash lines, respectively.

active site, the γ -OH remains within H-bonding distance (~ 2.9 Å) of Asp109^K(OD) therefore retaining the ability of Asp109^K to act as a catalytic base by deprotonating Dha.

Mechanistic Insight into Phosphotransfer. The chemical mechanisms of phosphoryl transfer in general, and with respect to kinases in particular, have been well studied (reviewed in ref. 7). However, little has been proposed regarding how DhaK forms the covalent adduct, catalyzes phosphoryl transfer, and ultimately releases its product. We divide the overall reaction into three stages: (i) binding of Dha to DhaK and formation of the covalent hemiaminal linkage; (ii) phosphoryl transfer from ATP to the covalently bound Dha, generating Dha-P and ADP; and (iii) release of Dha-P from DhaK, setting the stage for a new catalytic cycle. We propose here a plausible chemical mechanism for the overall reaction (Fig. 4). With the exception of the demonstrated roles for His218^K of DhaK (2) and Arg178^L of DhaL (6), the remain-

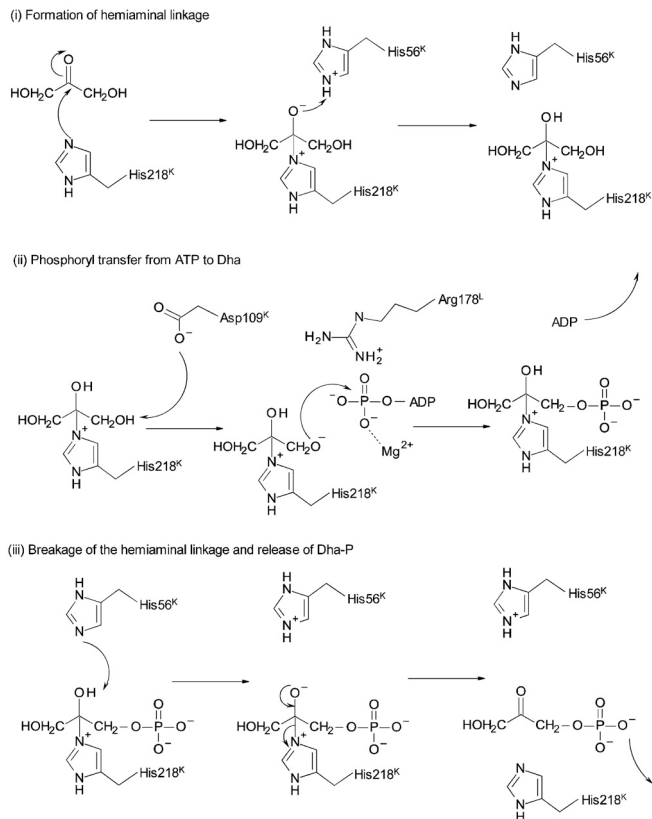


Fig. 4. Proposed catalytic mechanism for the DhaK-DhaL kinase, consisting of (i) initial formation of the hemiaminal linkage between His218^K and Dha; (ii) phosphoryl transfer from ATP to Dha, yielding Dha-P; and (iii) breaking of the hemiaminal linkage and release of Dha-P.

ing residues in the active site have not been previously implicated as having a role in catalysis.

Consistent with our structural data, we propose that formation of the initial covalent complex with Dha (Fig. 4*i*) involves nucleophilic attack of His218^K on the electrophilic carbonyl C β atom of Dha. For this, the NE2 atom of His218^K needs to be deprotonated. The proximal protonated His56^K acts subsequently as a general acid, donating its proton to form the Dha β -OH moiety (Fig. 4*i*). Although in general a water molecule could also play the role of general acid, there is no suitably located water molecule in any of the available crystal structures of wild-type DhaK. Formation of the covalent enzyme-substrate complex allows for precise orientation of the substrate with respect to other active site residues, including Gly53^K, His56^K, Lys104^K, and Asp109^K.

Following formation of the covalent DhaK-Dha complex, the next step is transfer of the γ -phosphoryl group of ATP to the γ -OH of Dha (Fig. 4*ii*). This reaction requires activation of the γ -OH through removal of a proton, in order to generate an oxyanion suitable for attack at the ATP γ -phosphorous atom. This activation requires a residue acting as a general base, which we identify as Asp109^K, within H-bonding distance of Dha γ -OH. This role is supported by the loss of activity observed for the Asp109^K mutant enzymes (Fig. S5). In the DhaK-DhaL complex, the leaving oxygen (O2B of ADP) and the entering oxygen (O γ of Dha) are separated by 4.8 Å (Fig. 1*B*). Based on molecular dynamics simulations on the DhaK-DhaL-ATP model, the Py-O γ (Dha) distance in the Michaelis complex is estimated to be 3.9 Å (Fig. 2*B*). The locations of negative charges developing during phosphoryl transfer may be inferred from the location of stabilizing, positively charged groups provided by the enzyme. An invariant residue, Arg178^L, is located at the active site of the DhaK-DhaL complex and has been suggested to be essential

for activity by mutagenesis (6). This residue has been postulated to stabilize the γ -phosphoryl group during phosphoryl transfer. In the crystal structure, the guanidinium group of Arg178^L is within H-bonding distance of the γ -OH group of Dha while at the same time does not directly interact with the β -phosphate of ADP (3.8 Å away). As previously stated, the second magnesium ion (M2) in the complex is coordinated by five oxygens, one of which is from a water molecule located between the β -phosphate of ADP and the substrate γ -OH group and likely mimics one of the oxygen atoms in the γ -phosphate group of ATP. Such an arrangement indicates that both Arg178^L and M2 are important for stabilizing the negative charges accumulated on the γ -phosphoryl group (γ -phosphate) during phosphoryl transfer. In the DhaK–DhaL complex, no positively charged residue other than Arg178^L could be found at the interface.

The final step in catalysis involves dissociation of Dha-P from DhaK, requiring breaking the hemiaminal linkage. Based on the structure of the active site, His56^K is well positioned to act as a general base, abstracting the proton from the β -OH of Dha-P, with concomitant formation of the carbonyl group at C2 and breakage of the Dha-P-His218^{NE2} bond (Fig. 4*iii*). The only other amino acid that could potentially act as a general base, Asp109^K, which interacts directly with the primary hydroxyl groups of Dha via its γ -carboxyl oxygens, is located too far to interact with the β -OH of Dha. Here, the oxygen–oxygen distance is 5.4 Å as opposed to 2.6 Å between NE2 of His56^K and the β -OH of Dha. As suggested previously (8), the γ -carboxyl oxygen of Asp109^K loses the ability to form an H bond with the primary hydroxyl group of Dha after the phosphoryl-ester is formed. Unfavorable interactions between the phosphate group of Dha-P and the carboxylate group of Asp109^K may help release the product from the enzyme. Together our data allow us to propose a chemical mechanism for DhaK supported by several structural and biochemical studies, consistent with involvement of His56^K and Asp109^K in addition to the previously characterized His218^K (2) and Arg178^L (4). These residues are highly sequence conserved in both bacterial and eukaryotic Dha kinases suggesting their common mechanism.

Our results indicating some activity for the H56A^K and H56N^K mutants require explanation. The first possibility is that covalently linked substrate is not essential for catalysis. Noncovalent binding of Dha, as reflected by our crystal structure, leads to reduced affinity as indicated by increased K_m^{app} as well as to a suboptimal orientation of γ -OH, affecting $k_{\text{cat}}^{\text{app}}$. If the covalent linking is not essential for activity, why are H218K^K and H218A^K mutants dead? We believe that because the lysine mutant introduces a more bulky side chain it does not allow for the binding of Dha. Furthermore, the small Dha molecule could not be well positioned in the absence of the histidine imidazole ring in the H218A^K mutant. The second possibility is that catalysis in H56^K mutants is achieved through the covalently bound Dha, assuming that His56^K determines the equilibrium between the covalently and noncovalently bound states of Dha. The covalent bond might be formed without participation of His56^K; for example, a water molecule could replace His56^K as general acid in the H56A^K and H56N^K mutants. The fact that K_m^{app} rather than $k_{\text{cat}}^{\text{app}}$ is more affected by the His56^K mutation provides some support for this idea. However, we could not identify appropriate water molecule in any of the His56^K mutant structures. Moreover, despite having analyzed 40 independent subunits of the crystal structures of DhaK H56A^K or H56N^K mutants, we were not capable of capturing the covalently bound substrate even at 40 mM Dha. Therefore, it is more likely that covalent binding of Dha is not essential for activity.

Functional Implications of Covalent vs. Noncovalent Substrate Binding by Dha Kinases. Formation of a transient, covalent bond between substrate and enzyme usually directly contributes to catalysis

(9, 10, 11); however, covalent substrate binding by dihydroxyacetone kinases is distinctively different. Previous studies have suggested that the primary role of the hemiaminal bond formed between DhaK and Dha or Dha-P is to provide the enzyme with a way of specifically selecting for short chain ketoses and aldoses vs. polyalcohols, because the enzyme displays remarkable selectivity for Dha in comparison with the closely related molecule, glycerol (8). This is in contrast to glycerol kinase, which is able to effectively utilize both Dha and glycerol as substrates for phosphorylation (12, 13). Quantum-mechanical calculations did not provide any evidence for an influence of the hemiaminal bond on activation of the Dha γ -O atom involved in nucleophilic attack of the γ -phosphate of ATP (8). In our hands, addition of glycerol (10 mM–1 M) to phosphotransferase reactions showed no decrease in kinase activity for wild-type and both H56^K mutant enzymes, indicating that the specificity of DhaK for Dha is not a consequence of hemiaminal bond formation per se. The apparent inability of glycerol to compete with Dha for binding to the DhaK active site may be a consequence of subtle structural differences between the two molecules, due to the sp^2 β -C in Dha vs. the sp^3 β -C of glycerol, resulting in steric conflicts of the latter within the quite rigid DhaK active site (based on comparison of all available crystal structures).

Our structural and biochemical results show that covalent bond formation does contribute, in part, to orientation of the substrate for optimal phosphoryl transfer. An additional potential role for hemiaminal bond formation to that in catalysis is for it to serve as an on/off switch, thereby exerting a regulatory function. A covalent bond between Dha and DhaK would support the enzyme existing in either a “Dha-loaded” or “Dha-unloaded” form, as dictated by (i) the inherent affinity of the enzyme for substrate and (ii) the effective concentration of Dha within the cell. The hemiaminal bond, once formed, commits DhaK to the “loaded” state, regardless of further changes in intracellular Dha concentration. It also provides a form of DhaK already primed for catalysis, once an ATP-loaded form of DhaL is available. In this manner, Dha binding and phosphoryl transfer are effectively uncoupled, allowing, for example, variations in the intracellular PEP pool independent of the particular state of DhaK. In a similar manner, the covalent bond would allow for an on/off switch with regards to transcriptional activation or repression that has been shown to occur via interactions with the transcription regulator DhaR (14). In either case, one would predict that a DhaK enzyme incapable of forming the hemiaminal bond in vivo would exhibit an attenuated response as opposed to an on/off behavior, given that the substrate could bind/unbind to the enzyme active site. Importantly, His56^K is as strongly sequence conserved as is the catalytic Asp109^K in Dha kinases across various species, including higher eukaryotes. This suggests a physiological importance for covalent substrate binding for this class of kinase in both PEP and ATP-dependent branches. The precise nature of the physiological consequences on cellular Dha metabolism resulting from loss of hemiaminal bond formation is the subject of future studies.

Materials and Methods

Protein Expression, Purification, and Characterization. Details for cloning, mutagenesis, expression, and purification are available in *SI Text*. Briefly, DhaK and DhaL N-terminal His₈-fusion proteins were purified by nickel affinity chromatography and the tags on DhaK removed by cleavage with tobacco etch virus (TEV) protease. Enzyme I and DhaM GST-fusion proteins were purified by glutathione affinity chromatography and cleaved from the column by TEV. HPr was purified by solubilizing inclusion bodies in urea and refolding the protein on nickel resin. For crystallization, the His₈-tag was cleaved from DhaL and the DhaK–DhaL complex was further purified on a Superdex 200 size exclusion column equilibrated with 50 mM Tris pH 8, 10 mM NaCl, 1 mM magnesium acetate, 1 μ M AMP-PNP, and 1 mM DTT. Complex formation of kinase mutants was analyzed by size exclusion chromatography in the same manner except that 1 μ M ADP was substituted for AMP-PNP.

Crystallization. Crystals of the DhaK–DhaL complex were obtained using the Anion Suite screen (Qiagen). Optimization of conditions led to the best crystals from hanging drop vapor diffusion at 19 °C by equilibrating 2 μ L of protein (10 mg/ml) with 1 μ L reservoir solution (0.1 M HEPES pH 7.5, 3.5 M sodium formate) over 1 mL of reservoir solution. Crystals belong to space group $P4_21_2$ with unit cell dimensions $a = b = 74.6$, $c = 268.8$ Å. Crystals of wild-type DhaK and mutant enzymes were also obtained by hanging drop vapor diffusion by equilibrating 1 μ L of protein (2.5 mg/ml of DhaK and 1.7 mg/ml of DhaL-his) with 1 μ L reservoir solution (0.1 M sodium citrate pH 5.6, 20% [w/v] PEG 8000) over 1 mL of reservoir solution. Crystals of the wild-type DhaK belong to space group $P2_1$ with $a = 49.8$, $b = 91.5$, $c = 73.2$ Å, $\beta = 89.9^\circ$. Crystals of H56A^K and H56N^K belong to space group $P1$ with unit cell dimensions $a = 59.7$, $b = 82.6$, $c = 92.9$ Å, $\alpha = 77.9$, $\beta = 78.1$, $\gamma = 71.1^\circ$, whereas that of the DhaK–H56N^K–Dha complex form in space group $P2_1$ with $a = 82.2$, $b = 101.1$, $c = 99.3$ Å, $\beta = 89.95^\circ$. For data collection, crystals were transferred to reservoir solution supplemented with 12% (v/v) ethylene glycol and flash cooled in a nitrogen stream at 100 K (Oxford Cryosystems).

X-ray Data Collection, Structure Solution, and Refinement. Diffraction data for free DhaK and the DhaK–DhaL complex were collected to 2.2 Å at the 31-ID beamline (LRL–CAT), Advanced Photon Source, Argonne National Laboratory. Data integration and scaling were performed with the program HKL2000 (15). Structure determination was performed by molecular replacement using the program Phaser (16) from the CCP4 suite with the previously reported *E. coli* DhaK (PDB 1O12) (2) and DhaL (PDB ID code 2BTD) (4) structures as the search models. Refinement was carried out with the program Refmac5 (17) with final $R_{\text{work}}/R_{\text{free}}$ of 0.172/0.210 and 0.189/0.225 for DhaK and DhaK–DhaL, respectively. Diffraction data for crystals of DhaK mutant enzymes were collected to 2.55 Å (H56A^K), 1.97 Å (H56N^K) and 2.2 Å (H56N^K–Dha) at the CMCF1 beamline at the Canadian Light Source. Data processing and structure determination was done similarly as described above. All models have good geometry as analyzed with PROCHECK (18). Final data collection and refinement statistics are shown in Table S1.

Molecular Dynamics Simulations. The DhaK–DhaL–ATP, DhaK–DhaL–*syn*–ADP, and DhaK–DhaL–*anti*–ADP complexes were sampled by 5-ns MD simulations using the AMBER10 suite of programs (19) together with the AMBER ff03 force field for the proteins and a modified force field for ATP and ADP (20). Details of the calculations are available in SI Text.

Phosphotransferase Assay. Phosphorylation of Dha was measured in a coupled assay similar to that described previously (1). To remove residual endogenous DhaK that could copurify with recombinant DhaL, the latter

was further purified by size exclusion (Superdex 200) in buffer containing 20 mM Tris pH 8, 300 mM NaCl, 1 mM DTT, 1 mM MgCl₂, and 0.01 mM ADP. Phosphotransferase reactions for specific activity measurements contained 1 μ M Enzyme I, 1 μ M HPr, 0.5 μ M DhaM, 0.5 μ M DhaL, 0.5 μ M DhaK, 50 mM potassium phosphate pH 7.5, 16 mM Dha, 2.5 mM DTT, 2.5 mM MgCl₂, 2 units glycerol-3-phosphate dehydrogenase (Sigma Chemical Co.), and 1 mM NADH and were initiated by the addition of 2 mM PEP. The glycerol-3-phosphate dehydrogenase coupling enzyme uses NADH to convert Dha-P to glycerol-3-phosphate. The production of Dha-P in the reaction was followed by monitoring oxidation of NADH at 340 nm in a Spectramax 250 plate reader at room temperature. The effect of glycerol on phosphotransferase activity for wild-type, H56A^K and H56N^K was examined for reactions in the presence of 10 mM, 100 mM and 1 M glycerol. $k_{\text{cat}}^{\text{app}}$ was measured for wild-type enzyme from coupled phosphotransferase reactions (0–1.6 mM Dha) containing 0.5 μ M Enzyme I, 0.5 μ M HPr, 0.25 μ M DhaM, 0.25 μ M DhaL, 0.25 μ M DhaK, 50 mM potassium phosphate pH 7.5, 2.5 mM DTT, 2.5 mM MgCl₂, 2 units glycerol-3-phosphate dehydrogenase (Sigma Chemical Co.), and 1 mM NADH, initiated by addition of 2 mM PEP. These conditions are similar to those reported previously (8), and kinetic constants were derived by varying concentration of Dha only, leading to apparent values of kinetic parameters. Due to the limited sensitivity of the assay for wild-type enzyme seen at low substrate concentrations an accurate value for K_m^{app} could not be determined. For H56A^K and H56N^K, $k_{\text{cat}}^{\text{app}}$ and K_m^{app} were determined from reactions containing 1 μ M Enzyme I, 1 μ M HPr, 0.5 μ M DhaM, 0.5 μ M DhaL, and 0.5 μ M DhaK at a range from 0–16 mM and 0–64 mM Dha, respectively. Values were calculated by nonlinear regression fit to a hyperbola.

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