

Structural Basis of Lipid Binding for the Membrane-embedded Tetraacyldisaccharide-1-phosphate 4'-Kinase LpxK*

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Background: LpxK is an essential membrane-bound kinase in the lipid A biosynthetic pathway.

Results: Structural and kinetic studies reveal the molecular basis of lipid binding.

Conclusion: The LpxK active site recognizes the lipid's glucosamine/phosphate headgroups and only accommodates disaccharides.

Significance: These studies provide a structural template for designing novel antibiotics and a glimpse of catalysis at the membrane interface.

The membrane-bound tetraacyldisaccharide-1-phosphate 4'-kinase, LpxK, catalyzes the sixth step of the lipid A (Raetz) biosynthetic pathway and is a viable antibiotic target against emerging Gram-negative pathogens. We report the crystal structure of lipid IV_A, the LpxK product, bound to the enzyme, providing a rare glimpse into interfacial catalysis and the surface scanning strategy by which many poorly understood lipid modification enzymes operate. Unlike the few previously structurally characterized proteins that bind lipid A or its precursors, LpxK binds almost exclusively to the glucosamine/phosphate moieties of the lipid molecule. Steady-state kinetic analysis of multiple point mutants of the lipid-binding pocket pinpoints critical residues involved in substrate binding, and characterization of N-terminal helix truncation mutants uncovers the role of this substructure as a hydrophobic membrane anchor. These studies make critical contributions to the limited knowledge surrounding membrane-bound enzymes that act upon lipid substrates and provide a structural template for designing small molecule inhibitors targeting this essential kinase.

Lipid molecules that reside in the membrane are acted upon by enzymes that either reside in the membrane themselves or are recruited there by other effectors. Despite the countless critical cellular processes that occur at the membrane interface, how enzymatic catalysis proceeds in such a heterogeneous environment remains a poorly understood phenomenon. In particular, due to the complex nature of biological membranes, the strategies with which these enzymes find and act upon their

lipid substrates are yet to be elucidated. The bacterial lipid A (Raetz) biosynthetic pathway provides a useful template for understanding how cytosolic precursors become fully embedded membrane lipids and illustrates the various stages of membrane catalysis that accompany this transition (1).

In Gram-negative bacteria, lipid A is the hydrophobic anchor of lipopolysaccharide (LPS), which makes up the outer leaflet of the asymmetric outer membrane of these organisms. This acylated disaccharide of glucosamine plays an important role in eliciting an immunogenic response to bacterial pathogens and is essential to the survival of the vast majority of these microbes (2, 3). The sixth step of the lipid A biosynthetic pathway in *Escherichia coli* is the phosphorylation at the 4'-position of the tetraacyldisaccharide-1-phosphate intermediate (DSMP)² by the cytosol-facing inner membrane kinase LpxK to produce lipid IV_A (Fig. 1A) (4, 5). Crystal structures of *Aquifex aeolicus* LpxK in apo-, nucleotide-bound, and nucleotide analog-bound forms have been reported (6, 7), revealing LpxK as a distinct member of the P-loop-containing nucleoside triphosphate hydrolase superfamily that acts upon a lipid substrate at the membrane surface (8–10). The LpxK C-terminal domain, a substructure unique to this enzyme, connects to the N-terminal domain by a long two-stranded β -sheet linker and serves as the "lid" to trap the nucleotide substrate between the two domains, forming the "closed" conformation (Fig. 1B). In the apo-form, the C-terminal lid domain is rotated away from the active site by a $\sim 25^\circ$ angle, resulting in an "open" conformation of the ATP-binding pocket (6).

Much less is known regarding recognition by LpxK of its lipid ligands DSMP and lipid IV_A. Kinetic studies have revealed that DSMP and ATP/Mg²⁺ must concurrently bind to the enzyme in order for catalysis to occur, and the kinase activity is a surface-dependent process (7). *A. aeolicus* LpxK has an apparent K_m for *Escherichia coli*-derived DSMP of $7.0 \pm 0.3 \mu\text{M}$ and shows activity *in vitro* despite molecular differences between

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The atomic coordinates and structure factors (code 4LKV) have been deposited in the Protein Data Bank (<http://www.pdb.org/>).

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² The abbreviations used are: DSMP, tetraacyldisaccharide-1-phosphate; AMP-PCP, 5'-adenylyl (β , γ -methylene)diphosphonate; Walker B motif, carboxylate-containing metal-binding motif of P-loop kinases.

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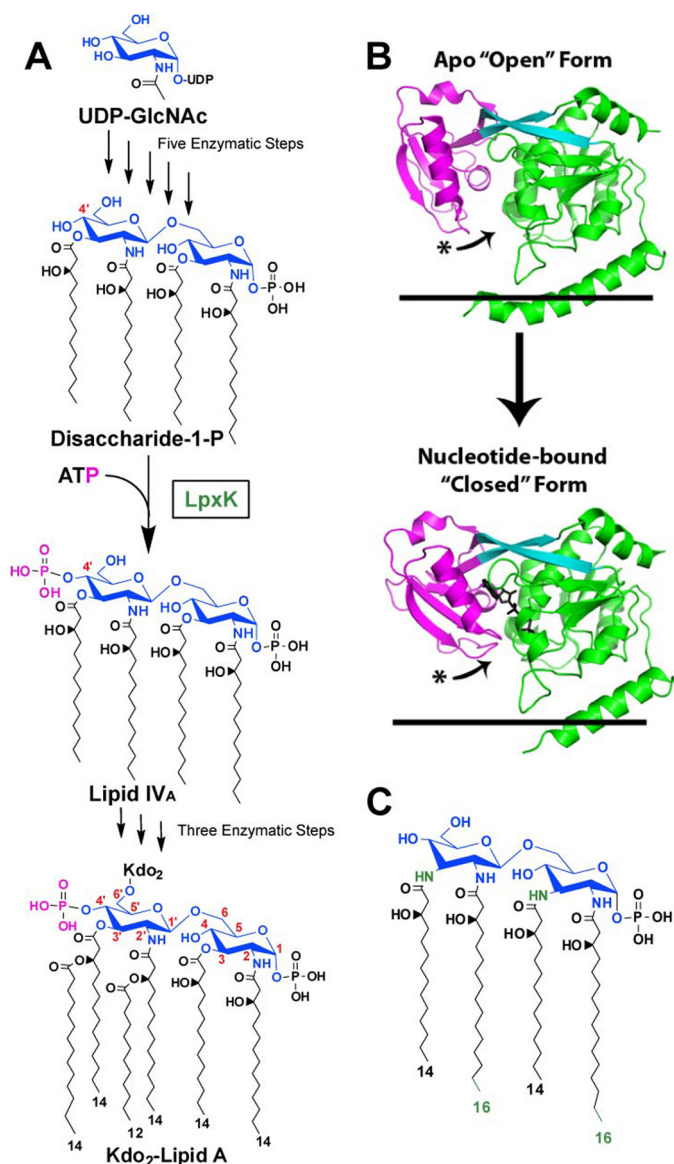


FIGURE 1. Role of LpxK in the Raetz pathway of lipid A biosynthesis. *A*, the biosynthesis of lipid A begins with uridine diphosphate-*N*-acetyl-D-glucosamine (UDP-GlcNAc). The pathway consists of nine steps, the sixth being the phosphorylation (magenta) of DSMP at the 4'-hydroxyl by LpxK, a membrane-bound kinase. Atoms of the glucosamine headgroups of lipid A are highlighted (blue), and positions on these sugars are indicated (red). Acyl chain length in Kdo₂-lipid A is also indicated. Kdo, 3-deoxy-D-manno-oct-2-ulosonic acid. *B*, LpxK consists of an N-terminal domain (green), a linker region (cyan), and a C-terminal domain (magenta). The kinase undergoes a conformational transition upon catalysis in which its C-terminal "lid" domain closes around its nucleotide ligand (black sticks). The curved arrows indicate relative movement of C-terminal loops undergoing this hinge motion. Black lines represent the cytoplasmic inner membrane surface (Protein Data Bank entries 4EHX and 4ITL). *C*, predicted structure of *A. aeolicus* DSMP based on the structure of *Aquifex* lipid A (11, 12). *A. aeolicus* DSMP contains longer acyl chains at the 2- and 2'-positions on the glucosamine backbone as well as amide linkages for the 3- and 3'-acyl chains, which differentiate it from the *E. coli* DSMP. Differences are colored in green.

E. coli DSMP and the predicted structure of *A. aeolicus* DSMP, which include longer acyl chains at the 2- and 2'-positions along with amide-linked acyl chains at the 3- and 3'-positions as opposed to ester linkages (Fig. 1C) (11, 12). Based on the directionality of the nucleotide phosphates in the AMP-PCP-bound LpxK structure, DSMP is predicted to bind to a shallow

basic cleft on the N-terminal domain formed between the N-terminal helix and core of the domain (7). The size and location of this cleft suggest that LpxK binds a smaller portion of its lipid substrate when compared with the few other proteins that have been crystallized in complex with lipid A or its precursors and imply a distinct strategy for finding its lipid substrate.

Knowledge of how LpxK interacts with the membrane and its lipid substrate will directly facilitate the development of small molecule antibiotics targeting this essential lipid A enzyme in Gram-negative bacteria and provide molecular insight into the activity of membrane-bound lipid modification enzymes, about which there exist precious little structural or kinetic data to date.

EXPERIMENTAL PROCEDURES

Complementation of an E. coli lpxK Knockout with A. aeolicus LpxK—Methods for manipulating DNA were adapted from previously described techniques (13). The *E. coli lpxK* gene was amplified from the plasmid pJK2 (5) via PCR by primers designed to incorporate 5' NdeI and 3' XhoI restriction sites into the PCR product. PCR was performed using the KOD Hot Start kit (EMD Millipore, Billerica, MA) on a Mastercycler gradient thermocycler (Eppendorf, Hamburg, Germany). This oligonucleotide, along with the pET21b vector (EMD Millipore, Billerica, MA), was digested with NdeI and XhoI (New England Biolabs, Ipswich, MA), and the digested PCR product containing the *E. coli lpxK* gene was ligated into the expression vector by T4 ligase (Invitrogen), creating the plasmid pRPE10, and transformed into chemically competent DH5(α) cells (14) (Invitrogen). In order to transfer both the *E. coli* and *A. aeolicus lpxK* genes into the pBAD30 vector (15), the plasmids pRPE7 (*A. aeolicus lpxK*) and pRPE10 (*E. coli lpxK*) were digested with XbaI and XhoI in order to maintain the ribosome-binding site from pET21b. The inserts were gel-purified and ligated into the pBAD30 vector, which had been digested by XbaI and SalI (SalI and XhoI have compatible sticky ends). The pBAD30 vectors containing the pET21b ribosome binding site and *E. coli* and *A. aeolicus lpxK* were designated pRPE43 and pRPE44, respectively, and transformed into electrocompetent TG1/pTAG1 cells (16). The presence of the correct construction was confirmed by sequencing.

TG1/pTAG1 is an *E. coli* strain with a chromosomal knockout of *lpxK* replaced by a kanamycin cassette. The knockout is complemented by the *lpxK* gene on a pMAK705 plasmid, which is temperature-sensitive and stops being replicated at 42 °C (17). TG1/pTAG1 was transformed with the pRPE43, pRPE44, or empty pBAD30 vector and grown on LB-agar plates supplemented with 50 μg/ml chloramphenicol, 25 μg/ml kanamycin, and 50 μg/ml ampicillin at 30 °C. One colony from each plate was used to inoculate 6 ml of LB overnight supplemented with chloramphenicol, kanamycin, and ampicillin. The overnight cultures grew to an OD of ~2, at which point they were diluted in LB to an OD of 1 and serially diluted 10-fold eight successive times. 3 μl of each dilution was spotted onto duplicate LB-agar plates supplemented with kanamycin and ampicillin. One plate was incubated at 30 °C and the other at 42 °C for 24 h, at which point relative colony growth between the three strains was assessed for the two temperatures.

In order to further confirm the ability of *A. aeolicus* LpxK to complement the *E. coli* enzyme *in vivo*, W3110 cells (18) that had been chemically transformed with the plasmids pBAD30 (vector control), pRPE43 (*EcLpxK* in pBAD30), or pRPE44 (*AaLpxK* in pBAD30) were transduced by P1_{vir} bacterial lysates of the TG1/pTAG1 strain, transferring the chromosomal knockout of the *lpxK* gene. Upon outgrowth of the transductions onto fresh LB-agar plates supplemented with kanamycin/ampicillin, colonies were only observed for the strains of W3110 that harbored plasmid pRPE43 or pRPE44 but not the vector control. To confirm the correct placement of the kanamycin cassette disrupting the endogenous *lpxK* gene, colony PCR was performed on each of the surviving strains with a forward primer complementary to a sequence on the upstream *msbA* gene (*prmsbA*F) and reverse primers complementary to either the downstream *ycaR* gene (*prycar*R) or the transduced kanamycin cassette (*prkan*R). Results of colony PCR were analyzed on a 1% (w/v) agarose gel.

Crystal Structure of LpxK Bound to Lipid IV_A—Wild type *A. aeolicus* LpxK was generated by growth of C41(DE3) (19) cultures expressing the construct pRPE7 and purified as described previously (6). Crystals were grown in 24-well sitting drop vapor diffusion trays from Hampton (Aliso Viejo, CA) at 20 °C. All conditions included a 700- μ l well volume and 10- μ l drop volume, and crystals were immediately flash-frozen in liquid nitrogen after being looped. Small cube-shaped crystals (0.03 \times 0.03 \times 0.03 mm) were obtained when LpxK was incubated with lipid IV_A, ADP, and Mg²⁺. Lipid IV_A was a gift from Dr. Hak Suk Chung, which was purified from the CMR300 *E. coli* strain, as has been previously described (20). The lipid IV_A stock solution contained 0.1% (w/v) Triton X-100 and 25 mM Tris, pH 6.8, along with 2 mM lipid. The drop included seven parts reservoir solution containing 60% (v/v) 2-methyl-2,4-propanediol and 0.1 M HEPES, pH 7.5, and three parts protein solution containing 7.6 mg/ml LpxK, 1 mM lipid IV_A, 12 mM Tris, pH 6.8, 0.05% (w/v) Triton X-100, 0.28% (w/v) dodecyl maltoside, 8.5% (v/v) glycerol, 21 mM HEPES, pH 8, 570 mM NaCl, 2.2 mM MgCl₂, and 2.2 mM ADP. Initial crystals were harvested after 5 months of incubation and diffracted poorly. A microcrystalline seed stock was then made with three of these crystals homogenized in a 500 μ l of 60% (v/v) 2-methyl-2,4-propanediol and 0.1 M HEPES, pH 7.5, solution.

Seed stock (see above) was used to nucleate a second batch of crystals. These crystals were grown in 24-well sitting drop trays as above with 0.5 μ l of the seed stock per well. The drop consisted of seven parts reservoir solution containing 70% (v/v) 2-methyl-2,4-propanediol and 0.1 M HEPES, pH 7.5, and three parts protein solution containing 6.7 mg/ml LpxK, 1 mM lipid IV_A, 12 mM Tris, pH 6.8, 0.05% (w/v) Triton X-100, 0.42% (w/v) dodecyl maltoside, 8.5% (v/v) glycerol, 21 mM HEPES, pH 8, 570 mM NaCl, 1 mM MgCl₂, and 1 mM ADP. Cube-shaped crystals similar to the original seed stock were obtained after 1 month. Data were collected at a single wavelength (1.0 Å) at the SER-CAT (Southeast Regional Collaborative Access Team) 22-ID line at the Advanced Photon Source (Argonne National Laboratory) to a resolution of 3.5 Å in the C22₁ space group with unit cell dimensions of $a = 136.6$ Å, $b = 140.7$ Å, $c = 173.8$ Å. Data were reduced and scaled using HKL-3000, and the struc-

ture was solved using the PHASER molecular replacement module within the PHENIX suite (21, 22). Apo-LpxK was used as the search model (Protein Data Bank code 4EHX). Iterative structure refinement was accomplished using the PHENIX software suite, included optimized atomic displacement parameters and stereochemistry weights, and used the search model as a restraint reference model. Riding hydrogen atoms were included throughout the refinement. There were four molecules in the asymmetric unit, and ligands (lipid IV_A and HEPES) were added to the model using an electron density acceptance criterion of $d \geq 3 \sigma$ (d) in the $F_o - F_c$ difference electron density map. The final model was validated via MOLPROBITY (23). Data collection and refinement statistics can be found in Table 1. A simulated annealing composite omit map for the lipid ligands was generated using CNS (24).

Point Mutagenesis of Lipid-binding LpxK Residues—Quik-Change mutagenesis (Stratagene, La Jolla, CA) was employed to generate point mutants N43A, R72A, R119A, Q142A, H143A, R171A, and E172A. Previously constructed point mutants Y74A, E100A, D138A, D138N, D139A, D139N, and H261A (6, 7) were also obtained. To generate partially purified LpxK point mutants, the plasmids were transformed into C41(DE3), expressed, and solubilized from membranes as described (7).

Solubilized membranes of each of the point mutants were assayed in the standard assay condition but with varied DSMP concentrations (1.56–100 μ M) and a fixed concentration of ATP and MgCl₂ of 5 mM. The other assay components were 50 mM Tris, pH 8.5, 0.5% (w/v) Triton X-100 (Thermo Scientific, Rockford, IL), 1 mg/ml BSA (Sigma-Aldrich), 0.1 M NaCl, and enzyme. The TLC-based assay was performed as described previously (6, 7), and the data were fit to the Michaelis-Menten equation using KaleidaGraph (Synergy Software, Reading, PA) in order to determine apparent K_m and k_{cat} with respect to DSMP. Enzyme concentration was varied to capture the linear range of activity for each mutant.

N-terminal Helix Truncation Studies—Two truncated forms of *A. aeolicus* LpxK were generated, one in which amino acids 2–12 were removed (referred to as Δ 12LpxK) and another in which amino acids 2–29 were removed (referred to as Δ 29LpxK). This corresponds to deleting roughly half of the N-terminal helix and the full N-terminal helix, respectively. Primers were used to amplify the Δ 12LpxK or Δ 29LpxK PCR product from pRPE7 and bestow 5' NdeI and 3' XhoI restriction sites. PCR was performed on a T3000 Thermocycler (Biometra, Göttingen, Germany) with the KOD Hot Start kit (EMD Millipore, Billerica, MA). The PCR products were digested with the respective enzymes, ligated with T4 ligase (Invitrogen) into the pET21b expression vector (EMD Millipore, Billerica, MA), and transformed into chemically competent DH5(α) cells (Invitrogen). Sequencing confirmed the correct construction of the Δ 12LpxK (pRPE72) and Δ 29LpxK (pRPE73) mutants. These plasmids were transformed into C41(DE3) cells, and membranes expressing these constructs were generated as with wild type LpxK (6, 19). All constructs expressed to comparable levels as judged by SDS-PAGE. Cell-free extract from cells expressing these constructs along with wild type LpxK and empty vector were assayed using the standard assay conditions (6, 7).

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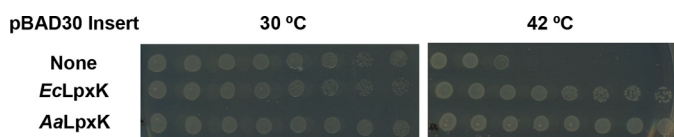


FIGURE 2. Complementation of *E. coli* LpxK with *A. aeolicus* LpxK. The TG1/pTAG1 strain of *E. coli*, containing a chromosomal knockout of *lpxK* complemented with *E. coli* LpxK on a pMAK plasmid with a temperature-sensitive replicon, was transformed with pBAD vectors containing no insert, *E. coli* *lpxK*, or *A. aeolicus* *lpxK*. 3 μ l of eight successive 10-fold serial dilutions (from left to right) of cultures starting at an A_{600} of 1 were spotted and grown at 30 or 42 °C. Cells containing the pBAD vectors with either LpxK ortholog, but not with an empty insert, were able to support growth at 42 °C upon loss of the pMAK vector, indicating that they could complement the chromosomal knockout of the *lpxK* gene.

Membranes derived from C41(DE3) overexpressing the empty vector, full-length LpxK, Δ 12LpxK, and Δ 29LpxK were obtained and suspended in 50 mM HEPES, pH 8.0, to \sim 9 mg/ml. Each of these membrane samples was diluted to 1.5 mg/ml into 0.5 ml of either 50 mM HEPES, pH 8.0, 1 M NaCl, 0.2 M sodium carbonate, pH 11.5, or the normal solubilization buffer consisting of 0.5 M NaCl, 1% (w/v) Triton X-100, and 20% (v/v) glycerol. Samples were incubated for 2 h at 4 °C in ultracentrifuge-safe microtubes (Beckman Coulter, Brea, CA) and centrifuged at 150,000 \times *g* for 1 h at 4 °C. The supernatant was collected, and the pellet was homogenized in 0.5 ml of 50 mM HEPES, pH 8.0. Both soluble and insoluble fractions were analyzed using SDS-PAGE with 20 μ l of each fraction per lane.

RESULTS

Complementation of an *E. coli* *lpxK* Knockout with *A. aeolicus* LpxK—To determine whether the LpxK ortholog from *A. aeolicus* could complement an *E. coli* knockout of the *lpxK* gene, pBAD30 vectors harboring the genes from both *E. coli* and *A. aeolicus* were transformed into the TG1/pTAG1 strain (16). When grown at 30 °C, TG1/pTAG1 transformed with *E. coli* *lpxK*, *A. aeolicus* *lpxK*, or the empty pBAD30 vector all grew robustly (15) (Fig. 2). In these cells, the temperature-sensitive replicon of the pMAK705 plasmid is able to function, and a sufficient amount of LpxK is expressed for the cells to survive (17). However, on identical plates that had been incubated at 42 °C, TG1/pTAG1 cells transformed with pBAD30 containing either *E. coli* *lpxK* or *A. aeolicus* *lpxK* were able to grow, whereas the empty vector could not (Fig. 2), indicating that *A. aeolicus* LpxK can perform the function of *E. coli* LpxK sufficiently well to complement a chromosomal knockout of the gene.

In a second set of experiments designed to further confirm the complementation of *E. coli* LpxK with the *A. aeolicus* ortholog *in vivo*, a chromosomal knockout of the *lpxK* gene in *E. coli* (16) was transduced into W3110 strains (18) expressing *E. coli* or *A. aeolicus* LpxK on a pBAD30 vector. Both strains were able to survive upon transduction, whereas vector control was not. Colony PCR confirmed the presence of the *lpxK::kan* insertion at the correct location on the chromosome. These results indicate that the molecular differences between *E. coli* and *A. aeolicus* DSMP do not prevent *A. aeolicus* LpxK from utilizing the *E. coli*-derived substrate and validate our pursuit of structural and kinetic studies involving *A. aeolicus* LpxK with the more accessible *E. coli*-derived reaction product lipid IV_A.

TABLE 1

Data collection and refinement statistics for LpxK-lipid IV_A complex

The data set was collected from two crystals.

Parameters	Values
Data collection	
Space group	C222 ₁
Unit cell <i>a</i> , <i>b</i> , <i>c</i> (Å)	136.6, 140.7, 173.8
Wavelength (Å)	1.0
Resolution (Å)	50.0–3.5 (3.56–3.50) ^a
<i>R</i> _{merge}	0.107 (0.687)
<i>I</i> / σ	21.2 (2.0)
Completeness (%)	94.2 (74.2)
Redundancy	12.8 (5.7)
Reflections/unique	257,205/20,053
Refinement	
<i>R</i> _{work} / <i>R</i> _{free} (%)	26.7/33.1
No. of atoms (average <i>B</i> -factor, Å ²)	
Protein	10,272 (88.3)
Lipid IV _A	94 (74.5)
Ramachandran plot	
Favored/allowed/outlier (%)	96.7/3.0/0.3
Root mean square deviations	
Bond length (Å)	0.01
Bond angles (degrees)	0.82

^a Values in parenthesis are for the highest resolution shell.

Crystal Structure of LpxK Bound to Its Lipid Product—In order to identify the specific molecular interactions between LpxK and its lipid ligands, *A. aeolicus* LpxK was crystallized in the presence of its product, lipid IV_A. The resulting crystals diffracted to 3.5 Å with four molecules in the asymmetric unit (Table 1 and Fig. 3A). Each of the LpxK protomers adopted the “open” form of the enzyme with respect to nucleotide binding (6) (Figs. 1B and 3B). Interpretable density was found in two of the four putative lipid-binding pockets (chains A and C), and lipid IV_A was modeled into both (Fig. 3A). The fact that the shape of the electron density at these two sites within the asymmetric unit is similar enough to allow for the placement of lipid IV_A in the same orientation (Fig. 3B) further supports the inclusion of these ligands within the overall structure. Because the electron density is best defined in the pocket of chain A and the B-factors are considerably lower for these atoms (average B-factor of 59.4 Å² for the chain A ligand *versus* 90.3 Å² for the chain C ligand), further analysis regarding lipid IV_A binding will be in reference to this protomer. A simulated annealing composite omit map for the lipid ligand of chain A was generated using CNS, and the corresponding difference density showed a strong signal for the lipid IV_A headgroups (24) (Fig. 3C).

Lipid IV_A is bound in the putative lipid-binding pocket on the underside of the N-terminal domain (Figs. 3D and 4A). Electron density for the glucosamine/phosphate headgroups of the lipid is strong, whereas density for the acyl chains is comparatively poor (Fig. 3C). Conserved residues of the L1 (purple), L4 (cyan), and L5 (orange) loops participate in lipid binding, as hinted at in previous studies (6) (Fig. 4). Loop L1, containing RGY from the highly conserved SRGY motif among LpxK orthologs, runs along the internal face of the lipid-binding pocket. The side chain of Arg⁷² is positioned to assist in binding the distal lipid IV_A headgroup via a hydrogen bond with the 6'-oxygen and the 1-phosphate, whereas the Tyr⁷⁴ side chain contributes a potential van der Waals interaction (Fig. 4, B and C). The putative hydrogen bond formed between the Arg⁷²

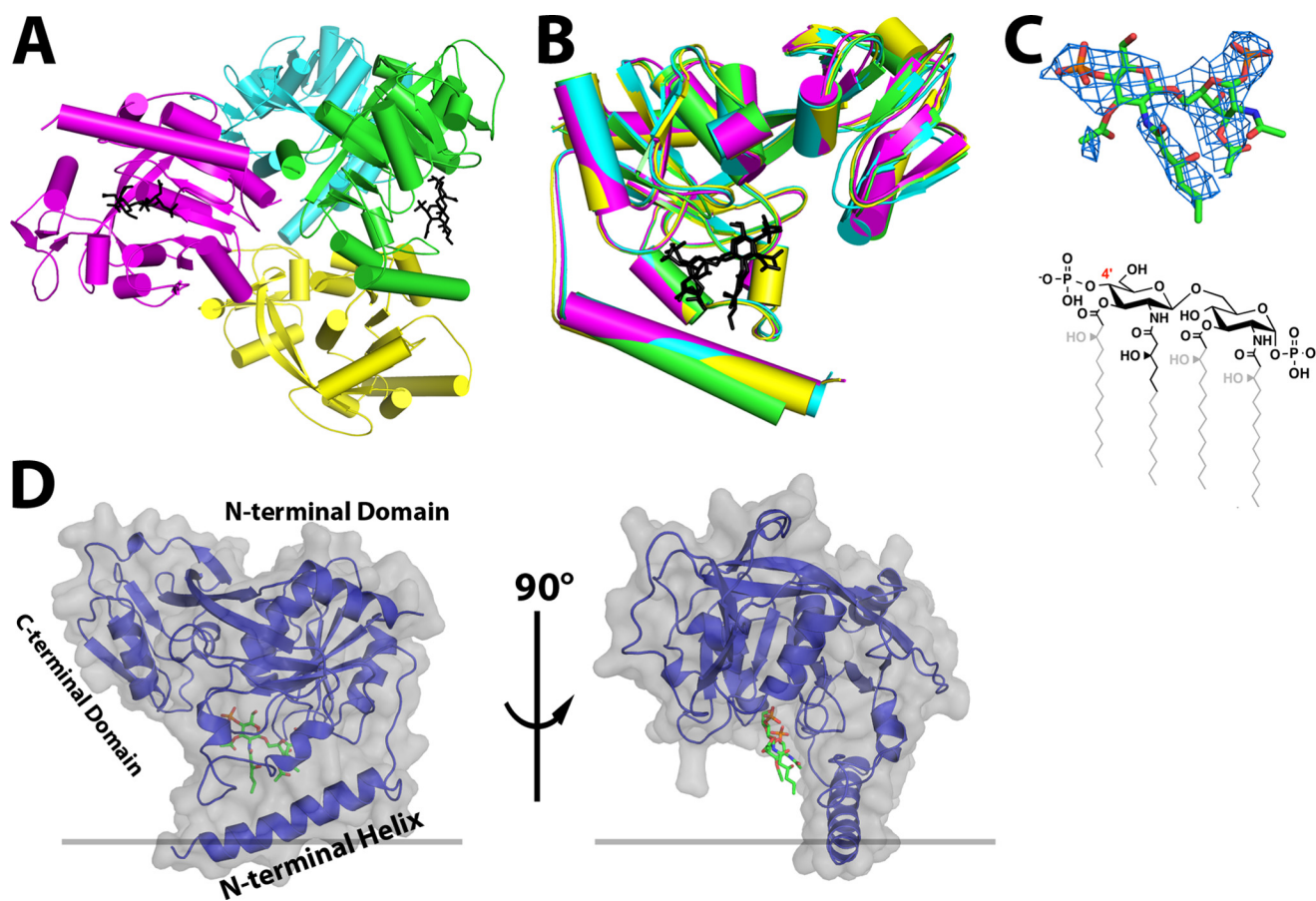


FIGURE 3. **Crystal structure of LpxK bound to lipid IV_A.** *A*, the asymmetric unit contains four LpxK molecules (green, cyan, magenta, and yellow for chains A, B, C, and D, respectively) and two molecules of lipid IV_A (black sticks). *B*, alignment of polypeptide chains A–D illustrates that lipid IV_A is bound to the “open” form of the enzyme in this crystal. *C*, stick representation of lipid IV_A in the crystal structure (from chain A) with the corresponding simulated annealing composite omit electron density (blue mesh) calculated with coefficients $2F_o - F_c$ contoured at 1.5σ . For comparison, the chemical structure of lipid IV_A is included, with atoms omitted due to poor electron density shown in gray. The 4′-position is indicated in red for reference. *D*, the crystal structure of the LpxK–lipid IV_A complex (chain A) reveals that the lipid is bound in a cleft at the base of the N-terminal domain between the N-terminal helix and core domain. Polypeptide backbone (purple diagram) is traced through the surface representation of LpxK (gray) with the lipid displayed as green sticks.

guanidinium and the 6′-hydroxyl of the distal glucosamine may protect this oxygen from being erroneously phosphorylated.

The L4 loop, which includes the Walker B motif, also appears to contain important residues for binding the lipid substrate. These residues make up the top of the lipid-binding pocket and are involved in the interaction with the glucosamine/phosphate moieties of lipid IV_A (Fig. 4). Asp¹³⁸, a residue implicated in magnesium binding according to previous structural data of both LpxK and other P-loop kinases (7, 9, 10), faces away from the lipid-binding site and points toward that of ATP/Mg²⁺. Asp¹³⁹ of the Walker B motif appears to form a hydrogen bond with both the Arg¹⁷¹ and Arg⁷² guanidinium moieties, probably positioning these residues for substrate binding. Gln¹⁴² also interacts with Arg¹⁷¹ and may also contribute to van der Waals interactions with the lipid IV_A 1-phosphate. His¹⁴³, as hinted at upon analysis of HEPES sulfate binding to the “apo” structure (6), forms a hydrogen bond to the 1-position phosphate of the lipid.

The L5 loop (Leu¹⁶⁵, Arg¹⁷¹, and Glu¹⁷²) makes up a portion of the external wall of the lipid-binding pocket (Fig. 4). Leu¹⁶⁵ is located in a favorable position to make hydrophobic contacts with the distal acyl chains of the lipid ligand, although a lack of acyl chain density prevents a more specific assignment. As previously mentioned, the Arg¹⁷¹ guani-

dinium group interacts with the Asp¹³⁹ carboxylate and may also help LpxK bind to DSMP through extensive van der Waals contact with the lipid headgroups and C2′-acyl chain (Fig. 4). A slight shift in conformation of this residue could conceivably see this guanidinium side chain interacting with glucosamine oxygens or with the carboxamide oxygen of the 2′-acyl chain. Glu¹⁷², another highly conserved residue of the L5 loop, faces away from the lipid-binding site, forming a hydrogen bond with Gln¹⁴².

Other residues outside these three loops also appear to have potential roles in lipid binding. Asn⁴³, a conserved residue of the N-terminal domain, may help position residues such as Arg¹⁷¹ and the catalytic Lys⁵¹ (Fig. 4). Val⁴⁶, a residue that precedes the P-loop motif, is less conserved and may contribute van der Waals interactions to the distal glucosamine of the 4′-phosphate. Glu¹¹⁷, a less conserved N-terminal domain β-sheet residue, is hydrogen-bonded to the 4-position oxygen of the proximal glucosamine moiety (Fig. 4B). Tyr¹³ of the N-terminal helix is well positioned to interact with the acyl chains of lipid IV_A, especially the 2′-hydroxymyristoyl group, given the extended electron density for this acyl chain compared with that of the other hydrocarbons (Figs. 3C and 4). Residues Glu¹⁰⁰ and His²⁶¹, previously implicated in ATP/

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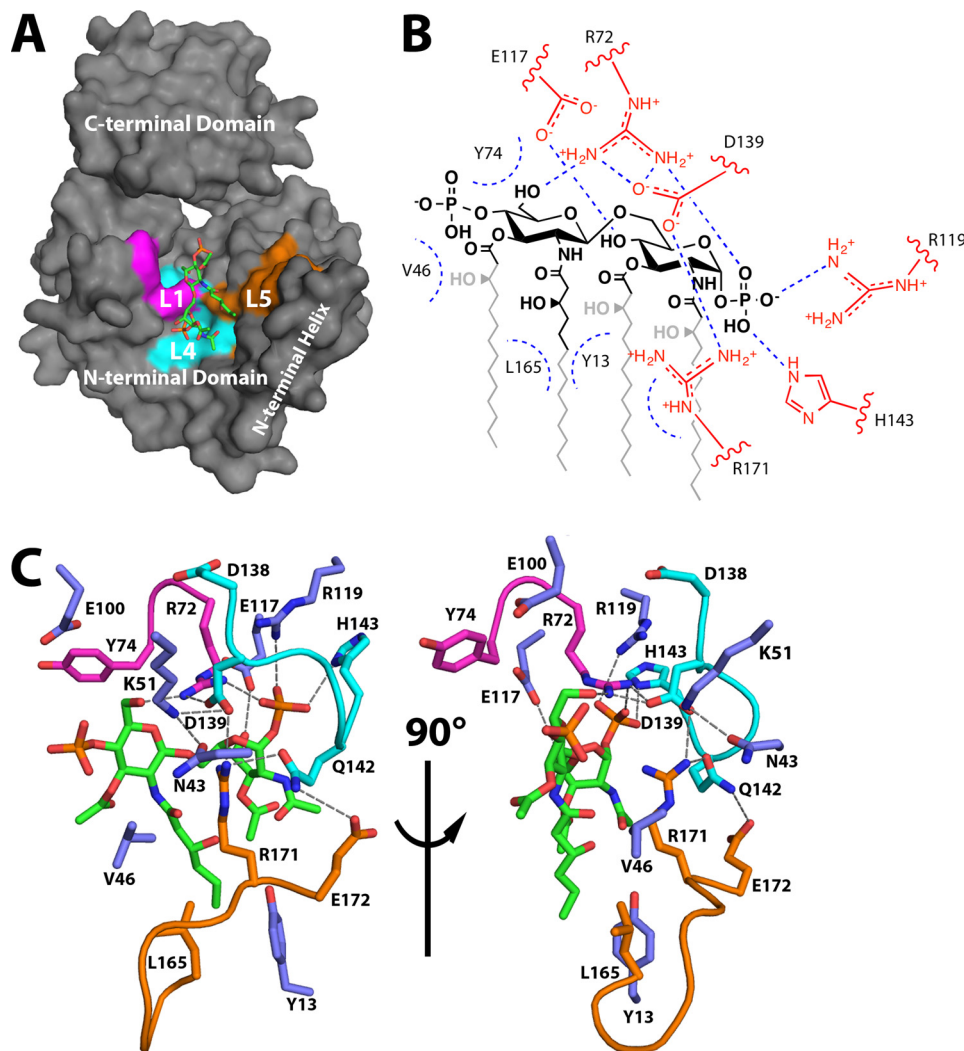


FIGURE 4. LpxK residues involved in lipid binding. *A*, surface representation of LpxK from the point of view of the inner membrane, with the N-terminal domain, N-terminal helix, and C-terminal domain substructures indicated. The surface created by residues of the conserved loops L1, L4, and L5 (magenta, cyan, and orange, respectively) makes up the left, back, and right sides of the lipid-binding pocket. *B*, schematic overview of lipid-binding residues of LpxK. Side chain atoms are highlighted in red. Coloring for the lipid is conserved from Fig. 3C. Straight blue dotted lines indicate putative hydrogen bonds, whereas curved blue dotted lines represent potential van der Waals contacts. *C*, orthogonal view of LpxK residues involved in lipid binding. Orientation of the lipid is conserved from Fig. 3D with the 4'-phosphate group oriented to the left (left) and front (right), respectively. Nearby side chains are represented as sticks with coloring conserved from *A*. Purple sticks indicate side chains that do not belong to the three highlighted motifs.

Mg²⁺ binding and catalysis, respectively, appear to be uninvolved in lipid binding in this open conformation. Glu¹⁰⁰ remains out of bonding range to the lipid (Fig. 4C), and His²⁶¹ resides ~15 Å away from the lipid-binding pocket on the C-terminal domain (Fig. 7A).

Point Mutagenesis of Lipid Binding Residues in LpxK—In order to further characterize lipid binding on a residue-by-residue basis, various point mutants of conserved residues of the DSMP binding pocket were generated and evaluated through steady-state kinetics. Apparent K_m and k_{cat} values with respect to DSMP are displayed in Table 2. The alanine mutant of Asn⁴³, a residue that precedes the P-loop and appears to help position Arg¹⁷¹ for lipid binding, has little effect on K_m for DSMP and decreases k_{cat} a modest 6.7-fold (Table 2; Fig. 5 visually maps the K_m data onto the structure). Alanine mutants of Glu¹⁰⁰ and His²⁶¹, conserved residues that do not directly bind the lipid according to the lipid IV_A complex, only increase K_m for DSMP 2–3-fold. The corresponding k_{cat}

TABLE 2
Kinetic parameters of LpxK point mutants with respect to DSMP

LpxK mutant	Apparent K_m	K_m increase	Apparent k_{cat}	k_{cat} decrease
WT	7 ± 1	1	2.4 ± 0.1	1
N43A	9 ± 2	1	(3.6 ± 0.3) × 10 ⁻¹	6.7
R72A	28 ± 3	4	(3.9 ± 0.1) × 10 ⁻¹	6.2
Y74A	48 ± 7	7	(3.1 ± 0.2) × 10 ⁻²	77
E100A	17.4 ± 0.8	3	(1.23 ± 0.02) × 10 ⁻³	2,000
R119A	9.7 ± 0.6	1	(3.60 ± 0.07) × 10 ⁻²	67
D138A	47 ± 5	7	(1.06 ± 0.05) × 10 ⁻³	2,300
D138N	7 ± 2	1	(2.8 ± 0.2) × 10 ⁻²	87
D139A	46 ± 8	7	(5.1 ± 0.4) × 10 ⁻⁴	4,700
D139N	14 ± 2	2	(6.3 ± 0.2) × 10 ⁻³	380
Q142A	28 ± 3	4	(3.5 ± 0.1) × 10 ⁻¹	6.8
H143A	50 ± 10	7	(1.6 ± 0.2) × 10 ⁻¹	15
R171A	128 ± 9	20	(1.38 ± 0.06) × 10 ⁻²	170
E172A	18 ± 3	3	(7.0 ± 0.1) × 10 ⁻¹	3.4
H261A	15 ± 1	2	(3.1 ± 0.9) × 10 ⁻³	770

decrease (2,000- and 770-fold, respectively) is much more significant, consistent with their primarily catalytic roles, as demonstrated previously (7).

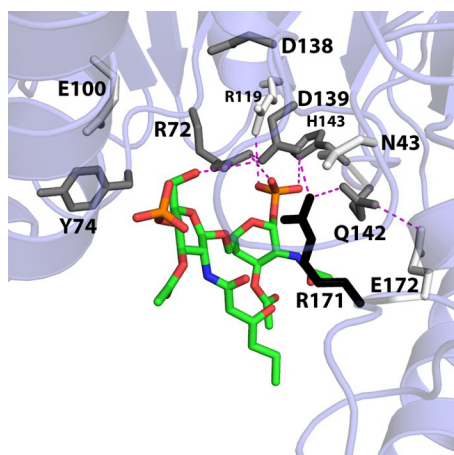


FIGURE 5. Apparent DSMP K_m perturbations of lipid-binding pocket alanine mutants mapped onto the LpxK-lipid IV_A coordinates. In order to visualize the effects of alanine mutants of conserved residues within the lipid-binding pocket on the apparent DSMP K_m , these residues are shaded in the model based on how much this parameter is increased when mutated to alanine using the following criteria: 0–4-fold K_m increase, white; 4–8-fold K_m increase, gray; >8-fold K_m increase, black. With the exception of Arg¹¹⁹, the alanine mutants of residues that appear to contact the lipid increase the K_m at least 4-fold. Arg¹⁷¹ appears to be the most critical residue for lipid substrate binding, with an increased DSMP K_m 20-fold higher than wild type. Potential hydrogen bonding interactions are shown as magenta dashes. Font size is indicative of depth.

Side chains of the conserved L1 loop SRGY motif (Arg⁷² and Tyr⁷⁴) are positioned to extend into the putative lipid-binding pocket and interact with the lipid substrate. Alanine mutants of Arg⁷² and Tyr⁷⁴ have a more significant effect on K_m (4- and 7-fold increase, respectively) (Table 2), whereas their k_{cat} values differ by an order of magnitude (6.2- and 77-fold decrease, respectively). Side chains of the L4 loop, which immediately follows the Walker B motif, also show a significant increase in DSMP K_m . The Q142A and H143A mutants increase this parameter 4- and 7-fold, respectively (Table 2) while decreasing k_{cat} a modest 6.8- and 15-fold. Another residue in the same locale, Arg¹¹⁹, has little effect on K_m and decreases k_{cat} by 67-fold when mutated to alanine, despite its proximity to the 1-phosphate of lipid IV_A in the crystal structure (Fig. 4). Such a modest effect may reflect a lack of conservation of Arg¹¹⁹ among LpxK orthologs compared with other lipid-binding residues (6).

Another substructure thought to be involved in DSMP binding, based on residue conservation and the lipid IV_A crystal structure, is the L5 loop (Fig. 4). Arg¹⁷¹, held in place through a strong interaction with Asp¹³⁹, appears to provide extensive van der Waals contacts for the lipid. This residue, when mutated to alanine, results in a 20-fold increase in apparent K_m for the DSMP substrate and a 170-fold decrease in apparent k_{cat} (Table 2). Glu¹⁷², another residue of this loop, which appears to be indirectly involved in lipid binding through an interaction with Gln¹⁴², was also mutated to alanine, resulting in a 3-fold K_m increase and a 3.4-fold k_{cat} decrease.

Point mutants of the Walker B motif carboxylates (common to P-loop kinases) were also analyzed for their effect on the kinetic parameters of LpxK with respect to the lipid substrate. The D138A and D139A mutants both increase the apparent DSMP K_m 7-fold (Table 2) while also having a significant impact on k_{cat} (2,300- and 4,700-fold decrease, respectively).

Asparagine mutants D138N and D139N result in rescued DSMP K_m values and partially rescue the k_{cat} value (87- and 380-fold, respectively) (Table 2). Surprisingly, Asp¹³⁸, a residue that is hypothesized to bind to the catalytic Mg²⁺ ion, had a non-trivial effect on the DSMP K_m . This residue, besides its catalytic role, may be critical for maintaining the correct orientation of the L4 loop. The mutation of Asp¹³⁸ may directly affect the position of adjacent residue Asp¹³⁹, which, according to the lipid-bound crystal structure and point mutagenesis, is involved in lipid binding (Fig. 4). The interaction of Asp¹³⁹ with Arg¹⁷¹ appears to be highly relevant because effects on K_m and k_{cat} are much more drastic for the D139A mutant compared with the D139N mutant, which at least preserves some bonding characteristics. This result may indicate that the D139A point mutant alters the conformation of the crucial Arg¹⁷¹ side chain so that this moiety can no longer position the lipid substrate for efficient phosphate transfer.

Characterization of N-terminal Helix Truncation Mutants—According to the crystal structure of lipid IV_A-bound LpxK, the only density-supported interaction between the lipid substrate and the N-terminal helix is a potential van der Waals interaction with Tyr¹³ (Fig. 4). Intrigued by the lack of visible interaction between the hydrophobic acyl chains and hydrophobic residues of the N-terminal helix, we endeavored to more concretely assign a role to this substructure. Based on surface charge and hydrophobicity analysis, we previously predicted that this helix is the primary mediator of membrane binding (6). To test this hypothesis, truncation mutants missing the first half of the N-terminal helix and the full helix were generated (designated Δ 12LpxK and Δ 29LpxK, respectively) and expressed on pET21b vectors in the C41(DE3) strain (19) (Fig. 6A). The cell-free extracts of Δ 12LpxK, Δ 29LpxK, and WT *A. aeolicus* LpxK were assayed for specific activity (7). The Δ 12LpxK truncation mutant decreased activity 81-fold, whereas the Δ 29LpxK truncation mutant decreased activity 16,000-fold to near background levels.

In order to assess the nature of the interaction of the N-terminal helix with the membrane, membrane fractions of the C41(DE3) strain expressing empty vector, WT LpxK, Δ 12LpxK, or Δ 29LpxK were isolated from the cell-free extract by ultracentrifugation. Upon SDS-PAGE analysis, it was revealed that the vast majority of the protein for each construct continued to partition with the membrane. These membrane fractions were then treated with various buffer conditions, ultracentrifuged in order to separate solubilized protein from the remaining membrane-bound fraction, and analyzed by SDS-PAGE (Fig. 6B). In a simple HEPES buffer condition, all constructs remained bound to the membrane fraction. In a high salt condition designed to disrupt the charge-charge interactions between LpxK and the phospholipids of the membrane, wild type enzyme remained bound to the membrane. However, the Δ 12 mutant displayed significant solubilization of the enzyme in this salt condition, whereas the Δ 29 mutant showed nearly full solubilization. In another condition designed to disrupt charge interactions, which involved membrane fraction treatment with 0.2 M carbonate at pH 11.5, a similar trend emerged, in which elimination of half or all of the N-terminal helix resulted in partial or full solubilization, respectively. Carbonate was also able to solubilize a portion of the wild type enzyme. As a posi-

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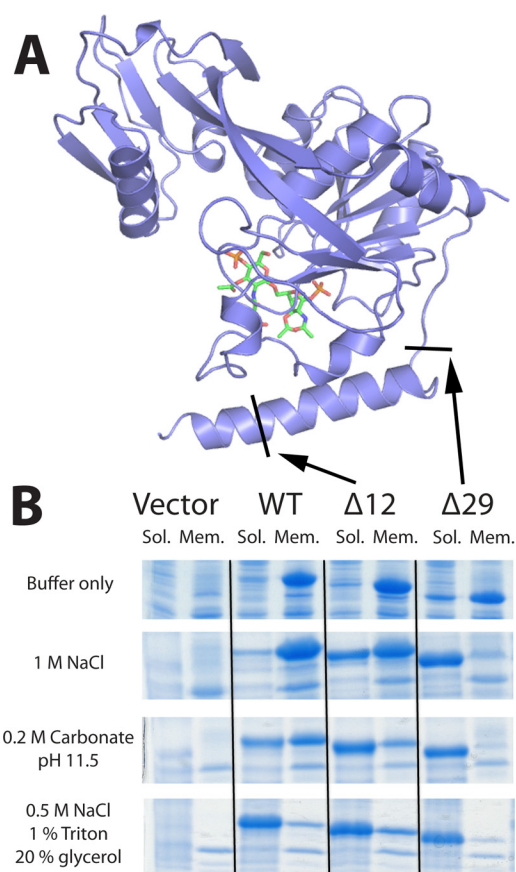


FIGURE 6. N-terminal truncation mutants reduce hydrophobic contact with the membrane. *A*, schematic representation of LpxK with arrows indicating truncation points for the $\Delta 12$ and $\Delta 29$ LpxK mutants. Coloring is conserved from Fig. 3*D*. *B*, SDS-polyacrylamide gels revealing the differential partitioning of the LpxK protein between insoluble membrane fractions and soluble fractions upon ultracentrifugation based on different solubilization buffer conditions. Truncation mutants are better solubilized than WT protein in conditions that disrupt the charge-charge interactions with the membrane, such as high salt or high pH. Band migration of the WT, $\Delta 12$, and $\Delta 29$ peptides was comparatively consistent with their predicted molecular masses of 36.5, 35.2, and 33.1 kDa, respectively.

tive control, the solubilization condition of 0.5 M NaCl, 1% (w/v) Triton X-100, and 20% (v/v) glycerol successfully solubilized every construct, combining disruption of both charge and hydrophobic contacts with the membrane.

This result indicates that the primary function of the N-terminal helix is hydrophobic interaction with the membrane because when the N-terminal helix was partially or fully removed, the enzyme primarily relied on charge-charge interactions to remain membrane-bound, and conditions that disrupt these charge-charge interactions were sufficient to fully solubilize these constructs. This result also provides a plausible explanation for why the truncation mutants perform so poorly in the standard assay. Because the enzymes are performing their function on Triton X-100 micelles, which have uncharged (tetramethylbutyl)-phenyl headgroups, LpxK in the assay condition relies on the hydrophobicity of the N-terminal helix to remain bound to the micelles at the location of the DSMP substrate. Without this helix, LpxK probably disassociates from the membrane and is unable to access its lipid substrate, thus precluding efficient catalysis.

DISCUSSION

The crystal structure of lipid IV_A -bound LpxK reveals almost exclusive interaction with the glucosamine/phosphate headgroups of the lipid, consistent with the notion that LpxK resides at the membrane interface and scans the surface of the inner membrane for its substrate DSMP (6). Such a lipid recognition mode is distinct from structurally characterized lipid A or lipid A precursor interactions with receptors and enzymes. For example, the β -sandwich fold of myeloid differentiation factors MD-1 and MD-2 form a hydrophobic pocket for the acyl chains of their lipid ligands both alone (25, 26) and in complex with Toll-like receptor 4 (TLR4) (27, 28), whereas the glucosamine/phosphate moieties are not extensively coordinated. LpxI of *Caulobacter crescentus*, the UPD-2,3-diacylglucosamine hydrolase that resides two steps upstream of LpxK in the lipid A biosynthetic pathway of this organism, also makes extensive van der Waals contacts with the acyl chain of its diacylated lipid ligands but comparatively few contacts with the glucosamine headgroup (29). Furthermore, structurally characterized ligand binding modes for lipid kinases, such as sphingosine kinase, display extensive acyl chain burial unlike LpxK (30).

In contrast to the aforementioned lipid-binding proteins, a headgroup-centric recognition mode, as seen in LpxK, has been noted in eukaryotic phosphatidylinositol-binding domains (31–33). Unlike LpxK, however, these effectors are transient peripheral membrane proteins and do not possess enzymatic activity on their own. The binding of soluble phosphoinositide analogs in complex with polyphosphate 5-phosphatase has been described recently (34), but this enzyme is not a *bona fide* membrane protein like LpxK, which possesses a hydrophobic binding component contributing to membrane affinity.

The crystal structure of LpxK bound to lipid is perhaps most comparable with the *Staphylococcus aureus* membrane transglycosylase structure with a lipid II analog, where lipid recognition also appears to occur primarily at the headgroup and involves two carbohydrate moieties (35). Although there are multiple arginine residues contributing to lipid binding as with LpxK, they are not nearly as critical for catalytic activity as Arg¹⁷¹ of LpxK. Furthermore, the transglycosylase is anchored in the membrane through a transmembrane helix in contrast to the amphiphilic helix of LpxK, and its lipid substrate contains a single polyprenol chain, unlike the tetraacylated lipid IV_A product captured in the LpxK structure.

The complex presented in the work involves the “open” form of LpxK, in which the catalytic dyad of Asp⁹⁹-His²⁶¹ is not formed, and this histidine lies ~ 12 Å away from the aspartate (6, 7) (Figs. 1*B* and 7*A*). Because the bound lipid is the phosphorylated LpxK product, this crystal structure most likely represents a postcatalytic complex in which ADP/Mg²⁺ has already left its binding site. Although the order of binding of the two substrates is not known, the fact that crystal complexes of LpxK bound to either the nucleotide or lipid alone can be attained may suggest a random order. Alignment of AMP-PCP-bound LpxK (7) and lipid IV_A -bound LpxK provides a good model for the arrangement of the substrates in the putative LpxK-ATP/Mg²⁺-DSMP ternary complex (Fig. 7*A*). The N-terminal domains of the two structures are nearly superimposable, with

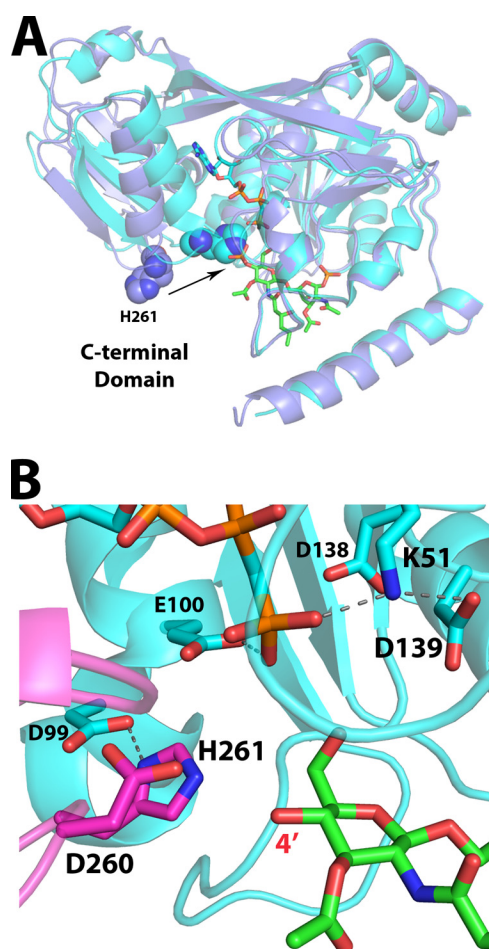


FIGURE 7. Overlay of AMP-PCP and lipid IV_A-bound structures as a model for the catalytic ATP-DSMP-LpxK ternary complex. *A*, alignment of the LpxK-lipid IV_A complex (purple schematic) and the nucleotide-bound complex in the “closed” form (Protein Data Bank code 4ITL) (cyan schematic) illustrates how the motion of the C-terminal domain closing around the nucleotide assembles the active site. The movement of the catalytic histidine His²⁶¹ (spheres) upon closure of the C-terminal domain to assemble the active site is indicated by an arrow. *B*, visualization of the active site of the AMP-PCP bound LpxK structure (cyan schematic and sticks) with DSMP modeled (green sticks, with atoms for the 4′-phosphate removed) based on the alignment of the N-terminal domains from *A*. The C-terminal domain (magenta) closure moves the α-carbon of His²⁶¹, the putative catalytic base, over 7 Å so that it may interact with Asp⁹⁹ to be activated. Among the plausible catalytic bases Glu¹⁰⁰, Asp¹³⁹, and His²⁶¹, the histidine resides nearest to the 4′-hydroxyl position (red) at 3.3 Å in this model, giving further credence to the hypothesis that this residue is the catalytic base. Hydrogen bonds within the AMP-PCP-bound LpxK structure are indicated with gray dashes. Font size is indicative of depth.

the most noticeable structural differences being C-terminal domain “closure” around AMP-PCP, a motion that effectively assembles the catalytic residues in the active site (Fig. 7A). Among the possible catalytic bases outlined in previous studies (7), His²⁶¹ appears closest to the 4′-hydroxyl of the DSMP substrate (3.3 Å) according to the model (Fig. 7B). Alanine mutants of residues Glu¹⁰⁰ and His²⁶¹, which are of catalytic importance according previous kinetic analysis (7), have a small effect on the apparent K_m for the DSMP substrate while decreasing k_{cat} significantly, confirming their primary role in catalysis (Table 2). These residues do not appear to participate in DSMP binding and are brought into play only upon domain closure with nucleotide-bound LpxK.

Several additional key residues were identified both structurally and kinetically as critical for LpxK lipid binding. His¹⁴³ anchors the 1-phosphate in position for correct binding orientation (Fig. 4) and ensures that molecules like lipid X (2,3-diacetylglucosamine-1-phosphate), whose headgroup consists of the 1-phosphate with only a single glucosamine, are not phosphorylated (4). As demonstrated previously, lipids that contain both glucosamine moieties but with altered acyl chain number are still substrates for the *E. coli* LpxK kinase activity (5), giving further support for the hypothesis that substrate recognition occurs primarily at the DSMP headgroup. Of all of the residues studied through point mutagenesis, Arg¹⁷¹ of the L5 loop reveals itself to be the most crucial to lipid binding, increasing the DSMP K_m 20-fold when mutated to alanine (Table 2). Arg¹⁷¹ appears to provide a critical anchor point for residues of the lipid-binding pocket, may maintain van der Waals interactions with lipid IV_A along its side chain, and may “lure” the negatively charged phosphates of the lipid substrate into this pocket using its guanidinium functionality.

The extensive characterization of the interaction between LpxK, its lipid ligands, and the cell membrane have provided the underpinnings for further study of membrane-embedded lipid modification enzymes. LpxK has differentiated itself from other P-loop kinases significantly through the development of a hydrophobic N terminus and surrounding basic residues that direct LpxK to the membrane and allow it to recognize the glucosamine/phosphate headgroups of its lipid substrate. Furthermore, this lipid-bound crystal structure makes LpxK a highly attractive target for the development of novel antimicrobials for three reasons. First, the dynamic conformational switch (Fig. 7A) that LpxK undergoes upon substrate binding makes for a specific high-affinity binding site which, if probed with an inhibitor, could trap LpxK in the closed state and prevent catalysis. Second, the large surface area of the lipid-binding site is naturally primed for a conjugated glucosamine sugar scaffold, providing an additional pathway toward the development of potent inhibitors. Finally, the existence of an LpxK crystal structure in complex with either nucleotide or lipid ligand of this bisubstrate enzyme presents the possibility of targeting both binding sites with a single drug. Such a strategy has been employed to engineer highly specific and highly potent inhibitory compounds for both protein kinases and other P-loop kinases, such as adenylate kinase (36, 37), providing a plausible template with which to combat the spread of Gram-negative bacteria with resistance to commercial antimicrobials.

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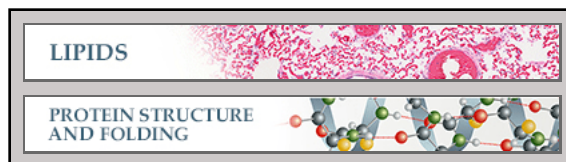
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Lipids:

**Structural Basis of Lipid Binding for the
Membrane-embedded
Tetraacyldisaccharide-1-phosphate 4'
-Kinase LpxK**



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