Structural insights into lipoprotein N-acylation by *Escherichia coli* apolipoprotein N-acyltransferase

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Gram-negative bacteria express a diverse array of lipoproteins that are essential for various aspects of cell growth and virulence, including nutrient uptake, signal transduction, adhesion, conjugation, sporulation, and outer membrane protein folding. Lipoprotein maturation requires the sequential activity of three enzymes that are embedded in the cytoplasmic membrane. First, phosphatidyglycerol: proteinprotein diacylglycerol transferase (Lgt) recognizes a conserved lipobox motif within the prolipoprotein signal sequence and catalyzes the addition of diacylglycerol to an invariant cysteine. This S-diacylglycerol lipidobox motif with the consensus sequence [LVI]N-terminal hydrophobic signal peptide that contains a conserved nitrogen hydrolase domain containing a Cysgl2–Lys catalytic triad. Based on these structures and complementary cell-based, biochemical, and molecular dynamics approaches, we propose a mechanism for substrate engagement and catalysis by *E. coli* Lnt.

Significance

Lipoprotein biosynthesis is crucial for Gram-negative bacterial viability and involves the activities of three essential integral membrane proteins embedded in the inner membrane (Lgt, LspA, and Lnt). These enzymes function sequentially to produce mature triacylated lipoproteins, many of which are then transported to the outer membrane. Lnt is responsible for catalyzing the addition of palmitate to the N terminus of diacylated apolipoproteins. Despite a number of studies that have biochemically characterized *Escherichia coli* Lnt, the structural basis for substrate engagement and catalysis remains unclear. Here we present the crystal structures of wild-type *E. coli* Lnt and a C387S active-site mutant. These structures provide insights into the molecular mechanisms of apolipoprotein N-acylation by Lnt and shed further light on the mechanism of lipoprotein biosynthesis by these essential bacterial enzymes.


The authors declare no conflict of interest.

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Data deposition: The crystallography, atomic coordinates, and structure factors have been deposited in the Protein Data Bank, www.pdb.org (PDB ID codes SVRG and SVRH).

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Fig. 1. Lnt is required for in vitro and in vivo growth and is enzymatically active in vitro. (A) Schematic of the lipoprotein biosynthetic pathway in Gram-negative bacteria. (B) Lack of in vitro growth of CFT073 lnt mutant after depletion of Lnt. WT CFT073 (black) and CFT073 lnt (red) were incubated in 2% arabinose (filled symbols) or 0.2% glucose (open symbols), and cfus were enumerated at various times posttreatment. The gray dashed line indicates the limit of detection for the assay. These data are representative of two independent experiments. (C) Transmission electron microscopy of WT CFT073 and CFT073 lnt mutant after treatment for 2 or 4 h with 0.2% glucose. Bars represent 0.5 μm. (D) The Pal substrate peptide<sub>substrate</sub> (Pam2Cys-SSNKNGK-Biotin, molecular mass = 1,713 Da) and product peptide<sub>product</sub> (Pam3Cys-SSNKNGK-Biotin, molecular mass = 1,951 Da), as detected by SAMDI mass spectrometry. A total of 50 nM recombinant Lnt was incubated with 10 μM Pal peptide and 100 μM POPE. Reactions were either prequenched (Left) or allowed to react for 21.5 h at room temperature (Right). Inset shows a Western blot of a reaction that has been allowed to go to completion. Substrate and product peaks were monitored to determine the peptide fraction conversion [(AUC<sub>product</sub>/AUC<sub>product</sub> + AUC<sub>substrate</sub>)] and product concentration [fraction conversion × (peptide<sub>total</sub>)]. (E) SAMDI mass spectrometry was used to monitor wild-type Lnt activity over 180 min while simultaneously varying the concentrations of POPE from 12.5–100 μM and Pal peptide<sub>substrate</sub> from 0.3125–5 μM. Global fitting of the initial velocities with a two-substrate ping-pong model was used to calculate the intrinsic K<sub>m(POPE)</sub>, K<sub>m(Pal peptide)</sub>, and k<sub>cat</sub> (55 μM, 14 μM, and 0.0093 s<sup>−1</sup>, respectively). Lineweaver–Burk double reciprocal plot was generated from the global fitting results, and the data represent velocities determined from average of duplicate samples.
of enzymes involved in diverse activities including nitrilase, amidase, carboxymethylase, and N-acetyltransferase catalysis. These enzymes are often multimeric and adopt a characteristic α-β-α sandwich fold that contains a canonical Lys–Cys–Glu catalytic triad (23). The Lnt active site is contained within its carbon-nitrogen hydrolase domain, with a catalytic triad comprised of E267, K335, and C387, each of which is essential for Lnt activity (24). A general two-step ping-pong mechanism for N-acetylation by Lnt has been proposed in which a thioester acyl-envelope intermediate is first formed at the catalytic cysteine through nucleophilic attack on the carbonyl of the sn-1-glycerolphospholipid of PE, resulting in S-palmitoylated Lnt. In the second step, the acyl-envelope intermediate is resolved by nucleophilic attack by the α-amino group of an apolipoprotein, resulting in a mature triacylated lipoprotein (24–26). Lnt has been shown to exist primarily in the acylated form in vivo, priming it for the more efficient second step of the N-acetylation reaction (25). Other residues have been reported to be essential for the activity of Lnt, however their function remains unclear due to a lack of an atomic structure (24, 25, 27).

Despite considerable biochemical insights, the molecular mechanisms that underlie lipid binding and apolipoprotein N-acetylation by Lnt remain unknown. Here we present the high-resolution crystal structures of wild type and a C387S mutant of E. coli Lnt. These structures reveal an eight-transmembrane helix fold and shed light on the role of conserved residues in the two-step reaction mechanism. We use a variety of approaches to confirm the mechanism of lipid binding and acyl-envelope intermediate formation, postulating a mechanism for subsequent engagement of diacylated apolipoprotein substrates and N-acetylation.

**Results**

**In Vitro and in Vivo Characterization of E. coli Lnt.** To determine the role of Lnt in clinical isolates, we generated an lnt mutant in the uropathogenic E. coli strain CFT073 (CFT073 lnt) (see SI Materials and Methods and Table S1 for bacterial strains and plasmids) that allowed Lnt protein levels to be induced or repressed (28). Briefly, in the presence of arabinose, Lnt is expressed under the control of the BβA promoter of the araBAD operon, whereas expression is repressed in the presence of glucose, a negative regulator of this operon. We then confirmed that Lnt is essential for growth in vitro, as CFT073 lnt did not grow when Lnt transcription is repressed (Fig. 1B). Further, Lnt was shown to be essential for virulence in mice, as no viable CFT073 lnt colony-forming units (cfus) were detected in the liver and spleen 24 h after i.v. infection compared with wild-type CFT073 (Fig. S1A). On a cellular level, depletion of Lnt in the CFT073 lnt mutant initially led to swelling of the periplasmic space at the bacterial poles (CFT073 lnt, 2 h), as measured by transmission electron microscopy (Fig. 1C). At later times, we noted cellular swelling and loss of density in the cytoplasm surrounding the nucleoid. Depletion of Lnt to levels that still support bacterial growth in vitro resulted in increased sensitivity to human serum killing (Fig. S1B) as well as to vancomycin (Fig. S1C), a Gram-positive antibiotic that is inactive against E. coli due to its impermeable OM. These data suggest that Lnt is essential for bacterial viability in vitro and in vivo and that depletion of Lnt results in increased OM permeability.

In addition to these cell-based approaches to examining Lnt activity, we assessed the in vitro biochemical activity of recombinant E. coli Lnt using self-assembled monolayer desorption/ionization (SAMDI) mass spectrometry to measure accumulation of a modified peptide product (29). Following biochemical reactions, biotinylated, diacylated peptide substrate and product were captured on a SAMDI surface, and all other reaction constituents were washed away before analysis by mass spectrometry. Enzymatic activity was monitored by ratiometrically assessing the fraction conversion of the substrate to product. In the absence of Lnt, only the substrate peptide was observed. Both substrate and product were observed following an incubation with recombinantly purified Lnt at room temperature (Fig. 1D). Reactions containing wild-type Lnt were linear with regard to time and enzyme concentration in the presence of 1-palmitoyl-2-oleoyl-sn-glycero-3-phosphoethanolamine (POPE) with both a short and long version (two versions were used due to limited availability of either peptide) of diacylated Peptidoglycan-associated lipoprotein (Pal) peptide substrates (Fig. S2). Similar to previous studies, these data suggested that wild-type Lnt functions in two steps via a ping-pong mechanism (Fig. 1E) (26). The kinetics of wild-type Lnt reacting with PE in this assay are in line with results reported previously, with a kcat/Km value of 674 M−1 s−1 (26).

**Overall Structure of Wild-Type E. coli Lnt.** To gain mechanistic insights into lipoprotein N-acetylation by E. coli Lnt, we crystallized a wild-type, selenomethionine (SeMet)-labeled Lnt by vapor diffusion and also obtained crystals of native Lnt using in meso methods. We determined the crystal structure of the SeMet-labeled Lnt at a resolution of 3.29 Å by single-wavelength anomalous dispersion (SAD) phasing. This structure was then used to phase the higher resolution (2.52 Å) native wild-type dataset (Fig. 2A and Fig. S3). Data collection and refinement statistics are shown in Table 1. Lnt crystallized as a monomer and is comprised of eight transmembrane helices (TM1–8; Fig. 2B) with both the N and C termini located in the cytoplasm.
Table 1. Crystallographic data and refinement statistics

<table>
<thead>
<tr>
<th>Data collection</th>
<th>Lnt wild type, SeMet</th>
<th>Lnt wild type, native</th>
<th>Lnt C387S</th>
</tr>
</thead>
<tbody>
<tr>
<td>Beamline</td>
<td>ALS 5.0.2</td>
<td>APS 22ID</td>
<td>APS 22ID</td>
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<tr>
<td>Wavelength</td>
<td>0.97957</td>
<td>0.9792</td>
<td>0.9792</td>
</tr>
<tr>
<td>Resolution range(^a)</td>
<td>43.82–3.29 (3.41–3.29)</td>
<td>45.16–2.52 (2.61–2.52)</td>
<td>42.04–2.14 (2.21–2.14)</td>
</tr>
<tr>
<td>Space group</td>
<td>P 3(_2_1)</td>
<td>P 2(_1_2_2)</td>
<td>P 1_2_1</td>
</tr>
<tr>
<td>Unit cell</td>
<td>153.54, 153.54, 89.51</td>
<td>87.76, 158.03, 44.76</td>
<td>52.67, 72.59, 75.61</td>
</tr>
<tr>
<td>Total reflections</td>
<td>36,544 (3,700)</td>
<td>43,030 (4,220)</td>
<td>60,043 (6,032)</td>
</tr>
<tr>
<td>Unique reflections</td>
<td>18,294 (1,184)</td>
<td>21,533 (1,937)</td>
<td>30,212 (3,039)</td>
</tr>
<tr>
<td>Multiplicity</td>
<td>2.0 (2.0)</td>
<td>2.0 (2.0)</td>
<td>2.0 (2.0)</td>
</tr>
<tr>
<td>Completeness, %</td>
<td>93.08 (63.76)</td>
<td>98.03 (91.71)</td>
<td>97.36 (98.99)</td>
</tr>
<tr>
<td>Mean (\text{I} / \text{sigma}(\text{I})</td>
<td>10.42 (2.06)</td>
<td>9.79 (2.01)</td>
<td></td>
</tr>
<tr>
<td>Wilson B factor</td>
<td>55.38</td>
<td>32.82</td>
<td>27.61</td>
</tr>
<tr>
<td>R-merge</td>
<td>0.06087 (0.3954)</td>
<td>0.06562 (0.3566)</td>
<td>0.06904 (0.4642)</td>
</tr>
<tr>
<td>CC1/2</td>
<td>0.999 (0.751)</td>
<td>0.995 (0.773)</td>
<td>0.974 (0.715)</td>
</tr>
</tbody>
</table>

Refinement

| Reflections used in refinement | 21,380 (1,937) | 30,207 (3,039) |
| Reflections used for R-free   | 1,099 (83)     | 1,461 (153)    |
| R-work                       | 0.2265 (0.2729) | 0.2068 (0.2927) |
| R-free                       | 0.2674 (0.3364) | 0.2502 (0.2976) |
| Number of nonhydrogen atoms  | 4,143          | 4,247           |
| Macromolecules               | 3,878          | 3,851           |
| Ligands                      | 119            | 138             |
| Solvent                      | 146            | 258             |
| Protein residues             | 494            | 490             |
| rmsd bonds (Å)               | 0.01           | 0.003           |
| rmsd angles (\(^\circ\))    | 1.14           | 0.69            |
| Ramachandran favored, %      | 94.69          | 96.71           |
| Ramachandran allowed, %      | 4.69           | 3.09            |
| Ramachandran outliers, %     | 0.61           | 0.21            |
| Rotamer outliers, %          | 1.71           | 1.72            |
| Clashscore                   | 11.07          | 5.67            |
| Average B factor             | 34.28          | 33.91           |
| Macromolecules               | 33.91          | 32.45           |
| Ligands                      | 44.05          | 57.45           |
| Solvent                      | 36.05          | 43.05           |

\(^a\)Numbers in parentheses represent the high-resolution shell.

TM3 contains a pronounced kink at G71 such that the C-terminal portion of the helix extends away from the rest of the transmembrane helical bundle at an \(\sim70^\circ\) angle. Side chain interactions contribute to similar angles through the membrane for TM4 and TM5, which are slightly splayed near the outer leaflet of the inner membrane to form a groove leading up into the periplasmic domain of the enzyme. TMs 5 and 6 comprise a portion of the protein that was previously presumed to form two partially membrane-embedded helices (27). The large loop between these helices (W141–V169) is primarily embedded in the membrane except for a short, amphipathic helix (K161–G168) that lies against the periplasmic side of the membrane before TM6. These helices and the intervening loop form a small, aqueous cavity that appears to allow solvent to penetrate into the membrane.

Lnt has a large periplasmic carbon–nitrogen hydrolase domain with a characteristic \(\alpha\)-\(\beta\)-\(\beta\)-\(\alpha\) sandwitch fold between TMs 7 and 8, as expected from sequence homology analyses. The two helices near the N terminus of this domain are located fully in the periplasm, whereas the helices C-terminal to the carbon–nitrogen hydrolase beta sheets pack against the membrane. Within this domain, the catalytic residues K335, E267, and C387 exist as a preformed active site, with E267 properly positioned to act as the general base that will abstract a hydrogen from C387 and initiate the first step of catalysis (Fig. 2C). Interestingly, these residues are located in a cavity that is raised above the outer leaflet of the inner membrane, with the \(C_\text{a}\) of C387 located \(\sim10\ \text{Å}\) above the predicted membrane interface. A small channel formed by highly conserved hydrophobic residues leads from the active site toward the cytoplasmic membrane (Fig. 2D). This hydrophobic cleft is enclosed on one side by a loop comprised of residues V339–V345 (loop 1) and on the other side by another loop (S78–G87; loop 2) directly preceding TM4. Both loops have high crystallographic B factors, suggesting that they are dynamic relative to the rest of the molecule. FS2 is poorly ordered and extends across the hydrophobic groove.

A number of lipids are bound to Lnt in the wild-type structure. Interestingly, although intact mass spectrometry analysis indicated that a large portion of the purified protein was covalently bound to palmitate (Fig. S4), no electron density was observed that was consistent with a covalent modification of the catalytic cysteine. This lack of covalent density is likely due to hydrolysis of the lipid during crystallization, as we noted an \(\sim238\)-Da mass shift in the predominant peak following a prolonged incubation in crystallization buffer (Fig. S4A, Inset). Despite this lack of a covalent modification, the structures of wild-type Lnt showed F\(_c\) = 2·F\(_o\) density likely corresponding to two aliphatic lipid tails leading from the active-site cavity down the hydrophobic cleft mentioned above. The presence of this density provides evidence that this hydrophobic groove represents the binding site for substrate lipids on Lnt. Due to low occupancy of lipid and poor electron density, we were unable to unambiguously identify the lipids bound in the
active site of the crystal structures, and they have been built as monoolein molecules that are derived from the lipid phase of the LCP matrix.

Structure of a Lipid-Bound Lnt C387S Mutant. Previous studies have demonstrated that mutating the active-site cysteine of Lnt to a serine leads to the formation of an acyl-enzyme intermediate that is resistant to reducing agents (25). Such mutants are also unable to efficiently resolve the acyl enzyme intermediate in the second step of N-acylation, likely owing to the substantially higher pKₐ of the serine side chain, which makes it a poor leaving group compared with the cysteine in the wild-type enzyme. In an attempt to obtain a structure of Lnt that is stably and covalently bound to palmitate, we purified a C387S mutant and solved its structure in the presence of POPE to 2.14 Å resolution (Fig. 3A). In terms of overall architecture, this structure is highly similar to the wild-type Lnt structure, with an overall average Cα rmsd of 0.268 Å.

We were surprised to find that no discernable electron density corresponded to a covalently bound lipid in the C387S mutant structure. This lack of covalent modification was again in contrast to mass spectrometry analysis performed before protein crystallization (Fig. S4B). However, the crystal structure revealed the presence of two monoolein lipids (lipid 1 and lipid 2) bound at two sites (site 1 and site 2, respectively) within the hydrophobic channel that connects the active site to the membrane (Fig. 3B). Both lipid acyl chains are partially embedded in the membrane. The structure suggests that lipid 1 binds in the site that would normally be occupied by the acyl chain of the S-palmitoyl cysteine (based on its proximity to C387), representing the acyl donor. Lipid 2 binds further from the C387, likely in the site that would normally contain the lyso-PE byproduct of the first half of the reaction.

The poorly ordered loop following TM3 in the wild-type Lnt structure (loop 2) is well ordered in the C387S structure, with F82 positioned across the lipid-binding groove (Fig. 3C). This positioning has the result that both bound lipids are completely encircled by protein, indicating that Lnt may bind lipids via an induced-fit mechanism before nucleophilic attack on the sn-1-acyl chain by C387. To probe the possibility of this induced-fit mechanism further, we performed a 10-ns molecular dynamics simulation using the crystal structure of wild-type apo Lnt embedded in a POPE lipid bilayer. Indeed, this simulation shows F82 and the rest of loop 2 to be highly flexible with regard to the rest of the protein, leading us to propose that this residue may serve as a gate for lipid binding and release (Fig. 3D and Movie S1).

Lnt Active Site and a Putative Phosphate-Binding Site. The active-site geometry in our lipid-bound C387S mutant structure supports a mechanism wherein E267 acts as a general base, abstracting a hydrogen to activate C387 for nucleophilic attack on the carbonyl group of the phospholipid sn-1-acyl chain. K335 is positioned to stabilize the oxyanion of the tetrahedral intermediate formed in both steps of the N-acylation reaction (Fig. 4A and B). Using our structure to model a full POPE substrate places the carbonyl oxygen of the sn-1-acyl chain within hydrogen-bonding distance of K335, an interaction that would likely strengthen the electrophilic nature of the carbonyl carbon before nucleophilic attack by the catalytic cysteine. Nucleophilic attack would then form a thiocysteine-enzyme intermediate, which we have modeled in Fig. S5.

In our crystal structures, a chloride ion is bound in a cavity surrounded by the side chains of W237, W415, N412, Q233, and K236, each with coordination distances of ~3.1–3.6 Å (Fig. 4C). Although it is possible that this ion-binding site is simply a crystallization artifact, we are intrigued that the chloride ion binds where we expect the negatively charged phosphate portion
of the POPE headgroup to bind to Lnt based on our model of this substrate bound to Lnt (Fig. 4D). Additionally, it is well-established that W237 is essential for Lnt activity (24, 25, 27), although the reason behind this requirement has been elusive. Based on our structure, we therefore propose this structural element to be a potential phosphate-recognition site.

Critical Residues for Lnt Activity and E. coli Growth. To probe the importance of key active-site, lipid-binding groove, and phosphate-binding site residues of Lnt, we examined the ability of mutations in these regions to functionally complement E. coli in the absence of wild-type Lnt expression. In addition to these cell-based assays, a subset of mutants were recombiantly expressed and purified for in vitro biochemical activity assays using SAMDI mass spectrometry.

As expected, no growth of CFT073 lnt cells was detected in the absence of wild-type Lnt expression. This growth defect was rescued by a plasmid expressing wild-type Lnt but not by the empty plasmid (Fig. 5A). The first panel of mutants we tested included residues at or near the active site that have previously been demonstrated to be essential for Lnt function in E. coli (24, 25, 27): namely, C387S, C387A, K335A, E267A, E267D, E343A, E343D, and E389A. Although previous studies have not examined N314, we also included a N314A mutant in our panel, given the proximity of this residue to the active site. This mutant was unable to rescue growth of the CFT073 lnt mutant (Fig. 5B) and likely plays an important role in stabilizing the active site due to the fact that it hydrogen bonds with both E267 and E343. In accordance with previous studies, alanine mutations for all of the other residues listed above also did not rescue growth of the CFT073 lnt mutant or exhibit significant biochemical activity (Fig. 5B and Fig. S6A). All of these mutant proteins were, however, expressed in the CFT073 lnt strain (Fig. S6B).

Although the C387S mutant protein showed modest in vitro activity after an overnight reaction, this was not sufficient to rescue bacterial growth of the Lnt-depleted CFT073 lnt mutant (Fig. 5B and Fig. S6A). Even the more conservative E267D and E343D mutations did not rescue CFT073 lnt growth in vitro, underscoring the importance of a properly positioned general base at position 267. In the absence of structural information, the specific role of E343 was previously unknown. In our structures, this residue hydrogen bonds with K335, likely ensuring its proper positioning for stabilizing the transition state of catalysis. Additionally, it is possible that E343 plays the dual role of substrate binding given that it makes a second hydrogen bond with the monoolein molecule that binds in site 2 in our C387S structure (this lipid represents the sn-2-acyl chain of PE). It is unclear if this hydrogen bond would also occur on a native lipid substrate or if it is simply an artifact of the crystallization system. E389 potentially serves two roles. This residue appears to stabilize the active-site conformation through a network of hydrogen bonds with the peptide backbone and a hydrogen bond with the side-chain of Y333. Beyond these interactions, this residue also likely creates a sterically block that further stabilizes the positioning of K335.

We also sought to test the importance of the putative phosphate-binding residues that coordinate a chloride ion in our C387S mutant structure (Fig. 5C). W237 has already been shown to be critical for Lnt activity (24, 25, 27). Indeed, the W237A mutant enzyme was completely inactive biochemically, and both this mutant and a W237E mutant failed to rescue growth of the Lnt-depleted CFT073 lnt mutant despite showing protein expression in this strain (Fig. 5C and Fig. S6A and C). The other putative phosphate-binding residues (Q233A and W415A) also failed to rescue growth of the CFT073 lnt mutant (Fig. 5C). It should be noted that in addition to participating in ion coordination, W415 also contacts the aliphatic chain of lipid 2. Interestingly, expression of the W415A Lnt protein was undetectable in CFT073 lnt cells, as detected by Western blot (Fig. S6C). These findings underscore the importance of the Lnt ion-coordinating residues, which are distant from the active site, in Lnt function, bolstering the hypothesis that they may be involved in phosphate recognition, although we cannot rule out the possibility that this site is in fact an ion coordination element that is important for overall Lnt protein stabilization.

Finally, we probed the importance of several residues located along the Lnt lipid-binding groove. Lnt-inducible knockout strains harboring F82A, F341A, and Y388F mutants all grew to levels similar to wild type, whereas F146A, W148A, F365, Y388A, and F416A mutants all could not rescue growth of the CFT073 lnt mutant despite protein being expressed (Fig. 5D and Fig. S6D). We were somewhat surprised to see that the F82 and F341 mutants exhibited normal growth, given that they are involved in enclosing lipid substrates in the lipid-binding groove of Lnt. One possible explanation for this normal growth could be that the affinity of the site for POPE is still high enough to maintain sufficient N-acylation for normal bacterial cell viability. Alternatively, it is possible that the removal of this gating mechanism may have actually increased the inherent substrate turnover, yielding a more efficient enzyme. Y388 contributes to the hydrophobic nature of the lipid-binding groove and likely makes Van der Waals interactions with what would be the acyl tail of the acyl enzyme intermediate. This residue is also located near the active site and makes backbone hydrogen-bonding interactions, but these interactions are not likely to be important for catalysis given that a Y388F mutant showed growth similar to wild type. Overall, our data confirm that the lipid-binding groove observed in our structures is important for Lnt function biochemically and in bacterial cells. The results of our mutagenesis assays are mapped onto the Lnt structure in Fig. 5E.

Discussion

We have determined the crystal structures of both wild-type and a C387S mutant of E. coli Lnt with two lipids bound in the active site. These structures provide a detailed view of lipid substrate binding and the catalytic mechanism, which is supported by our mutational analysis. One question that remains unanswered is how a lipid or diacylated substrate peptide is extracted from the lipid bilayer and transported ~10 Å into the periplasmic catalytic site of Lnt. Our molecular dynamics simulations suggest that the helices and loops that make up the walls of the lipid-binding groove are quite mobile. Given that this is a highly dynamic region of the protein, it is reasonable to assume that Brownian motion and appropriate gating by the protein may be the primary driver of lipid movement into the Lnt active site.

The Lnt crystal structures allow us to suggest a mechanism by which the protein regulates the entry of substrates to the active-site cavity and the exit of products (Fig. 6). We propose that in the crystal structure of the C387S mutant, lipid 2 occupies the site that would normally contain either the lyso-PE product of the first half of the reaction or one of the substrate peptide acyl groups that will help direct the α-amino group of the lipoprotein to the thioester carbonyl of the acyl-enzyme intermediate for nucleophilic attack. Based on our studies, initial substrate binding likely involves an induced-fit mechanism wherein loop 1 and loop 2 open to allow the noncovalent product to exit site 2 (Fig. 6A). This would then enable entry of one of the acyl chains of the diacylated protein substrate into site 2. Based on modeling this interaction into the C387S mutant crystal structure (Fig. S7), it is possible that the Phe clamp would remain open during this step to accommodate the presence of a third acyl chain. Binding would likely be mediated by the diacylglycerol moiety of the substrate peptide rather than the peptide

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**Fig. 5.** Critical residues for Lnt function in *E. coli.* (A) Rescue of CFT073 lnt mutant growth in vitro with wild-type Lnt but not with empty plasmid. CFT073 lnt mutant was grown under Lnt depleting conditions (media containing glucose; open gray squares) containing either an empty plasmid (open gray circles) or a plasmid expressing wild-type Lnt (filled gray squares). As a control, CFT073 lnt mutant was grown under Lnt inducing conditions (media containing arabinose; filled black squares). (B) Active-site residues, (C) phosphate-binding residues, and (D) lipid-binding groove residues critical for *E. coli* growth. CFT073 lnt mutants lacking wild-type Lnt expression were complemented with plasmids expressing various Lnt mutants. Lnt mutants that rescue growth of the Lnt-depleted CFT073 lnt cells are shown in blue, whereas those that could not rescue growth are shown in red. The gray dashed line indicates the limit of detection for the assay. (E) Map of Lnt mutational analysis onto a model of the Lnt crystal structure with POPE bound. Mutants that were unable to rescue wild-type growth of CFT073 lnt in glucose-containing media are shown with a red sphere at the Cα position. Mutants that showed growth similar to wild type in glucose-containing media are denoted with a blue sphere at the Cα position.
itself, given that *E. coli* lipoproteins have diverse sequences downstream of the conserved cysteine and are all substrates of Lnt. It is possible, however, that Lnt makes hydrogen-bonding interactions with the main chain of the substrate peptide. Binding would position the α-amino group of the lipoprotein substrate for nucleophilic attack on the carbonyl of the S-palmitoyl cysteine intermediate. The mature, triacylated lipoprotein product would then be released, and the now vacant lipid-binding groove would be available to bind another PE substrate (Fig. 6B).

Overall, our structures provide a more complete picture of the lipoprotein biosynthetic pathway in Gram-negative bacteria. One question that remains to be answered is how lipoproteins are transferred from one enzyme to the next during this biosynthetic process. It is unclear if this is achieved by simple 2D diffusion through the membrane or if these proteins may exist as a higher order lipoprotein biosynthetic complex to facilitate more efficient lipidation. Further studies into this possibility will certainly deepen our understanding of this important biochemical pathway in Gram-negative bacteria.

**Materials and Methods**

**Lnt Expression and Purification.** *E. coli* Lnt constructs containing a noncleavable C-terminal 6xHis tag were expressed recombinantly in *E. coli* BL21 (DE3) cells by 64 h autoinduction at 16 °C. Cells were harvested by centrifugation at 4,668 × g for 20 min at 4 °C, resuspended in lysis buffer (20 mM Tris, pH 8.0, 500 mM NaCl, 10% glycerol, 1x complete protease inhibitor mixture; Roche), and lysed by three passages through a microfluidizer at 10,000 psi. Cell debris was removed by centrifugation at 18,000 g for 15 min at 4 °C. The membrane fraction was then isolated by ultracentrifugation at 125,000 g for 1 h at 4 °C. The membrane pellet was resuspended in buffer A (20 mM Tris, pH 8.0, 300 mM NaCl, 5 mM imidazole, 10% glycerol, 1% DDM) supplemented with 1x complete protease inhibitor mixture and agitated overnight at 4 °C using a magnetic stirrer. Insoluble material was separated by ultracentrifugation at 125,000 g for 1 h at 4 °C. The supernatant was then incubated with cobalt affinity resin for 1 h at 4 °C on a nutator. The resin was added to a gravity flow column and washed with five column volumes (CVs) of buffer A followed by five CVs buffer B (20 mM Tris, pH 8.0, 300 mM NaCl, 5 mM imidazole, 10% glycerol, 0.02% DDM). Bound protein was eluted with buffer B containing 250 mM imidazole. Pure fractions were combined and applied to a Superdex 200 16/60 column that had been pre-equilibrated with buffer B containing 250 mM imidazole. Pure fractions were combined and concentrated to 25–40 mg/mL.

**Crystallization.** Lnt at 40 mg/mL was reconstituted into the lipidic cubic phase (LCP) by mixing with monooolein at a 2:3 (wt/vol) protein-to-lipid ratio using two coupled syringes. A Mosquito LCP robot (TTP Labtech) was used to dispense 50 nL boluses of protein-laden mesophase overlayed with 800 nL precipitant in 96-well glass sandwich plates. Crystals with P2_1_2_1 symmetry were obtained at 20 °C using a precipitant solution containing 50 mM ADA,
ph 6.5, and 24% PEG 400. Crystals were harvested using MTEGen cryoloops and flash-cooled in liquid nitrogen without added cryoprotectant.

The C387S mutant of Lnt was also crystallized in LCP and grew in a condition containing 0.1 M MES, pH 5.7–6.3, 27% PEG 500 DME, 0.1 M sodium chloride, 0.1 M magnesium chloride, and 0.01 M copper(II) chloride dihydrate as the precipitant. The LCP host lipid contained a monolaurin mixture containing 0.25 mol% POPE. These crystals grew at 20 °C, had P2_12_1 symmetry, and were harvested as above.

For phasing efforts, aliquots of SeMet-labeled Lnt at 25 mg/mL were adjusted to 10 mg/mL with buffer C. Twenty-five microliter aliquots of SeMet-labeled Lnt were then supplemented with 300 μg E. coli polar lipid extract (Avanti Polar Lipids) and incubated overnight at 4 °C on a nutator. Aggregates were removed from the lipidated sample by centrifugation at 20,000 g for 90 min at 4 °C. Crystallization was carried out at 18 °C using the hanging drop vapor diffusion method combined with streak seeding using a seed stock containing native Lnt crystals obtained by the same method. Crystals with P2_12_1 symmetry were obtained in a precipitant solution containing 50 mM sodium acetate, pH 5.0, 50 mM magnesium acetate, and 28–36% PEG 200. Crystals were cryoprotected in reservoir solution containing 40% PEG 200 and flash-cooled in liquid nitrogen.

Structure Determination. Initial phases of wild-type Lnt were obtained by SAD from data collected using beamline 5.0.2 at the Advanced Light Source (ALS) on SeMet derivative crystals. The heavy atom substructure of 12 selenium atoms was determined by the Hybrid Substructure Search submodule in the Phenix package (30), and initial phases were obtained using AutoBuild (31). Automatic chain tracing using AutoBuid (32) yielded an interpretable electron density map, and this was used as an initial guide for building an intermediate model. The intermediate model was used as a search model in Phaser (33) against the native dataset containing Lnt on beamline 22-ID of the Advanced Photon Source (APS). Iterative rounds of refinement and manual rebuilding using PHENIX (34) and COOT (35) resulted in an almost complete chain of Lnt. The wild-type model was used to phase the Lnt C387S datasets.

Molecular Dynamics Simulations. Molecular dynamics simulations of wild-type Lnt embedded in a POPE lipid bilayer was performed using Desmond Molecular Dynamics submodule from the Schrödinger suite of programs (36). Briefly, the crystal structure of wild-type apo Lnt was prepared by building missing loops using Prime (37), bond orders assigned, and energy minimized with hydrogen bond optimization within Maestro (Schrödinger). The mini-
mimized protein was then placed in a simulation box that satisfied the periodic boundary conditions, embedded in a POPE membrane bilayer, and solvent and counterions were added to neutralize the system. The position of the membrane was adjusted based on the Orientation of Proteins in Membranes database (38). The membrane-embedded protein system was relaxed before running the 10-ns molecular dynamics simulation. The trajectory was ana-
lized using the tools in Desmond and exported to PyMOL to generate the movie of the molecular dynamics simulation.

In Vitro Growth and Serum-Killing Assays. For in vitro growth curves, overnight cultures of WT CFT073 and CFT073 Int grown in Luria–Bertani (LB) medium containing 2% arabinose were back-diluted 1:100, grown to midexponential phase (OD_{600} = 0.6), and then diluted 1:100 in media containing either 2% arabinose or 0.2% glucose to initiate growth curves. For complementation studies in the CFT073 Int mutant, Lnt mutants were generated using the QuikChange Lightning Site-Directed Mutagenesis kit (Agilent Technologies) as per the manufacturer recommendations. Primers used to generate the mutant Lnt-expressing plasmids are shown in Table S2. Wild-type or mutant Lnt coding regions were cloned into the pLMG188tet plasmid. The cfs were counted by plating on LB + kanamycin (50 μg/mL) agar. Serum sensitivity assays were carried out using normal human sera (NHS) pooled from six donors, at 50% serum con-
centration. A Checkerboard microdilution minimal inhibition concentration (MIC) assay was performed to determine if decreasing Lnt expression (by decreasing arabinose concentrations in the growth medium) from the pBAD plasmid in- creases sensitivity to vancomycin or human serum killing. WT CFT073 or CFT073 Int colonies were picked from an overnight LB plate, and MIC assays were performed according to Clinical and Laboratory Standards Institute (CLSI) guidelines.

Biochemical Assays. The enzymatic activity of wild-type and mutant E. coli Lnt was quantified using SAMDI mass spectrometry (SAMDI Tech, Inc.) by moni-
toring the ratio of substrate (diacylated) and product (triaoylated) pep-
tides and calculating the fraction conversion as AUC_{triacylated} / (AUC_{triacylated} + AUC_{diacylated}). The amount of product peptide formed was determined by multiplying the fraction conversion by the total peptide concentration. Two peptide substrates were synthesized based on the Pal lipoprotein, Pal peptide_short (Pam2Cys-SSKNGK-Gbiotin; CPC Scientific, Inc.) and Pal peptide_long (Pam2Cys-SSKNASDDGEMGLGAGTGMGD-Kbiotin; Anspec, Inc.). Reactions were carried out at room temperature in Lnt assay buffer (50 mM Tris, pH 7.5; 150 mM NaCl, 0.05% DDM (Anatrace); 0.05% bovine skin gelatin (Sigma); and 1 mM TCEP in a 384-well polypropylene plate (Greiner)). Two lipid substrates were used, POPE (Avanti Polar Lipids) formulated as POPE/DM mixed mi-
celles, and E. coli i–ω-PE in chloroform (Avanti Polar Lipids). Enzymatic activity was monitored in the presence of PE or POPE and peptide_{short} or peptide_{long}. Reactions were quenched by the addition of a final concentration of 0.5% formic acid. The kinetic mechanism of wild-type Lnt was assessed by simulta-
nuously varying the concentration of the peptide_{short} and POPE substrates and monitoring enzyme activity over 3 h. Intrinsic K_{m} and V_{max} values for each substrate were determined by global fitting the initial, linear enzymatic rates as a function of concentration of both substrates according to two sub-
strate ping-pong (v = v_{max}[A][B] / (K_{m} [A] + [B] + [A][B]) or ternary mechanisms (v = v_{max}[A][B] / [K_{m} + [A][B]] + [A][B]), where A is the free substrate and B is the peptide_{short} substrate. Reaction rates were determined using Prism software (GraphPad Software), and global fitting was done using Grafit soft-
ware (Erithacus Software, Ltd.). Global fitting was also used to distinguish between these two general mechanisms. Lineweaver–Burk plots of the re-
ciropral initial velocity (1/v_{b}) versus the reciprocal of the POPE concentration (1/[A]) at each fixed peptide substrate concentration were generated for vi-
sualization purposes using the global fitting results.

Transmission Electron Microscopy. Bacteria were grown to midexponential phase, washed with PBS, and incubated in modified Karnovsky’s fixative (0.1 M sodium cacodylate, pH 7.2, 2% paraformaldehyde, and 2.5% glutaraldehyde). Samples were then postfixed in 1% aqueous osmium tetroxide (EM Sciences) for 1 h followed by overnight incubation in 0.5% uranyl acet-
ae. The samples were then washed through a series of ethanol followed by propylene oxide and embedded in Eponate 12 (Ted Pella, Inc.). Ultrathin sections (80 nm) were cut with an Ultracut microtome (Leica), stained with 0.2% lead citrate, and examined in a JEOI JEM-1400 transmission electron microscope at 80 kV. Digital images were cap-
tured with a GATAN Ultrascan 1000 CCD camera.

Mouse Infection Model. Overnight bacterial cultures of WT CFT073 and CFT073 Int grown in medium containing 2% arabinose were back diluted 1:100 in M9 media and grown to an OD_{600} of 0.8–1 at 37 °C. Cells were harvested, washed once with PBS, and resuspended in PBS containing 10% glycerol. Cells were frozen in aliquots, and thawed aliquots were measured for cfu before mouse infections. Virulence of WT CFT073 and CFT073 Int was measured using the neutropenic E. coli infection model (39). Briefly, 7 wk-old C57BL/6 J mice (Jackson Laboratory) were rendered neutropenic by peritoneal injection of two doses of cyclophosphamide (150 mg/kg on day −1 and 100 mg/kg on day −1). On day 0, mice were infected with 5 × 10^5 cfu mid-exponential phase bacteria diluted in PBS by i.v. injection through the tail vein. At 30 min and 24 h post infection, bacterial burdens in the liver and spleen were determined by serial dilutions of tissue homogenates on LB plates.

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