Aerobactin, a citrily-hydroxamate siderophore, is produced by a number of pathogenic Gram-negative bacteria to aid in iron assimilation. Interest in this well-known siderophore was reignited by recent investigations suggesting that it plays a key role in mediating the enhanced virulence of a hypervirulent pathotype of *Klebsiella pneumoniae* (hvKP). In contrast to classical opportunistic strains of *K. pneumoniae*, hvKP causes serious life-threatening infections in previously healthy individuals in the community. Multiple contemporary reports have confirmed fears that the convergence of multidrug-resistant and hvKP pathotypes has led to the evolution of a highly transmissible, drug-resistant, and virulent “super bug.” Despite hvKP harboring four distinct siderophore operons, knocking out production of only aerobactin led to a significant attenuation of virulence. Herein, we continue our structural and functional studies on the biosynthesis of this crucial virulence factor. *In vivo* heterologous production and *in vitro* reconstitution of aerobactin biosynthesis from hvKP was carried out, demonstrating the specificity, stereoselectivity, and kinetic throughput of the complete pathway. Additionally, we present a steady-state kinetic analysis and the X-ray crystal structure of the second aerobactin synthetase IucC, as well as describe a surface entropy reduction strategy that was employed for structure determination. Finally, we show solution X-ray scattering data that support a unique dimeric quaternary structure for IucC. These new insights into aerobactin assembly will help inform potential antivirulence strategies and advance our understanding of siderophore biosynthesis.

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This article contains Figs. S1–S7, Tables S1 and S2, Schemes S1–S3, and Data S10–S25.

The atomic coordinates and structure factors (code 6CN7) have been deposited in the Protein Data Bank (http://wwpdb.org).

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3 The abbreviations used are: KP, *Klebsiella pneumoniae*; hvKP, hypervirulent pathotype KP; NRPS, nonribosomal peptide synthetase; NIS, NRPS-independent siderophore; RMSD, root mean square displacement; DL, disordered loop; SER, surface entropy reduction; hLys, N^6^-hydroxy-L-lysine; ahLys, N^6^-acetyl-N^4^-hydroxy-L-lysine; SAXS, small-angle X-ray scattering.

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![Figure 1. Aerobactin biosynthetic pathway.](image)

Bacillus subtilis and other sporulating bacteria contain a gene cluster encoding the bactin siderophore, which is utilized by the synthetases IucA and IucC. This work further our understanding of the catalytic function of these closely-related enzymes.

**Results**

**Heterologous in vivo aerobactin production**

Much of the prior investigation on aerobactin biosynthesis originated from a pathogenic strain of *E. coli* harboring the aerobactin operon in the ColV-K30 plasmid (29). Although Neilands and co-workers (31) studied aerobactin biosynthesis by knocking out and re-complementing aerobactin genes originating from this plasmid, we employed a parallel approach in which an aerobactin-deficient *E. coli* strain was engineered to harbor the aerobactin gene cluster from a single pathway to be structurally and functionally characterized. This work furthers our understanding of NIS biosynthesis and provides the biochemical and structural foundation to further explore this potential antivirulence target.
However, as predicted, when expressing only the synthetase genes iucA and iucC, none of the aerobactin intermediates were detected (Fig. 2B). Finally, when the E. coli were transformed and induced to express all four iucA–D genes, a pronounced peak corresponding to aerobactin was observed alongside secondary peaks corresponding to the intermediates ahLys and citryl-ahLys (Fig. 2C). In addition to demonstrating aerobactin biosynthesis, this engineered system served as a relatively convenient/accessible source for isolating in vivo biosynthesized aerobactin and citryl-ahLys without using a pathogenic bacterial strain or requiring iron-deficient culture media. Although these metabolites were detected and isolated from the culture media supernatant, it is unclear whether they were actively secreted or if they accumulated via cell lysis or passive diffusion.

**In vitro aerobactin reconstitution**

After demonstrating expression of the four iucA–D genes was sufficient for aerobactin production in vivo, the pathway was investigated in vitro using purified enzymes (Fig. 3, Table S1 and Fig. S1). Despite a tendency to aggregate during expression, IucD, a putative membrane-associated enzyme, was purified as a monodisperse sample with the help of detergent. IucB, expressed as a cleavable chaperone chimera, had a broader size distribution and also had a tendency to aggregate at elevated concentrations and temperatures. However, when incubated together with substrates L-lysine, NADPH, acetyl-CoA, ATP, citrate, and the co-factor FAD, they were able to biosynthesize ahLys (Fig. 3B). When IucD, IucB, and IucA were combined, all of the ahLys was ligated to citrate, forming the citryl-ahLys intermediate, but no aerobactin was detected (Fig. 3C).

In contrast, when IucD, IucB, and IucC were incubated together, no appreciable citryl-ahLys or aerobactin was detected (Fig. 3D). Finally, when substrates and all four purified IucA–D enzymes were included in the reaction mixture, a prominent peak corresponding to aerobactin was observed alongside lesser amounts of ahLys and citryl-ahLys remaining in the mixture (Fig. 3E). This in vitro reconstitution of aerobactin biosynthesis using purified enzymes provides direct functional evidence supporting the reported aerobactin pathway (30).

**Aerobactin pathway turnover frequency**

In addition to establishing the competency of the in vitro aerobactin biosynthetic system, the turnover of the pathway was also examined quantitatively. Aerobactin was purified from the culture supernatant of the engineered E. coli strain (Fig. S2, D–F) and used to construct a standard curve (Fig. 4A). The reconstitution reaction was sampled along its time course to monitor how aerobactin evolved over time (Fig. 4B). The linear time points on the reaction progress curves were fit with regres-
Aerobactin synthetase stereoselectivity

The stereoselectivity of the aerobactin synthetases IucA and IucC has not been determined. Two distinct stereocenters are relevant in aerobactin biosynthesis. *E. coli* IucD was previously shown to only be capable of hydroxylating L-lysine, and had no activity with D-lysine (32). Therefore, it follows that the stereocenters derived from the α-carbon of lysine have the 2′S configuration in aerobactin and its precursors. Condensation of ahLys with one of the two pro-chiral carboxymethyl groups of citrate breaks the symmetry of the molecule and results in a second stereocenter at the citrate-derived C3 position in citryl-ahLys (Fig. 1). Citryl-ahLys purified from *iucA–D* co-expression eluted as a single peak by HPLC (Fig. 5A). Similarly, citryl-ahLys isolated from the *in vitro* IucA-catalyzed condensation of ahLys and citrate also eluted as a single peak (Fig. 5B). In contrast, a synthetic epimeric mixture of citryl-ahLys at the C3 position eluted as two distinct peaks, presumed to represent the two diastereomers (Fig. 5C). Elution of authentic 3R,2′S- and 3S,2′S-enriched citryl-ahLys produced through total synthesis, featuring enantioenrichment of a citrate intermediate by fractional recrystallization, supported this presumption and allowed the absolute configuration of the two peaks to be assigned (Fig. 5, D and E). The correlation of the retention times of the biosynthesized samples and the synthetic 3S,2′S-enriched sample indicated that only the 3S,2′S-diastereomer was appreciably produced by IucA. When biosynthesized citryl-ahLys was the substrate of IucC, all of it was turned over to make aerobactin (Fig. 5, A and B). Conversely, when the synthetic citryl-ahLys samples were incubated with IucC, only the peak corresponding to the 3S,2′S-diastereomer diminished, whereas the peak corresponding to the 3R,2′S-diastereomer remained unperturbed (Fig. 5, C–E). Taken together, these results demonstrate that the aerobactin synthetases are highly stereoselective; only 3S,2′S-citryl-ahLys is produced by IucA, and only 3S,2′S-citryl-ahLys is consumed by IucC in the production of aerobactin (Figs. 1 and 5).
X-ray crystal structure of IucC

Crystals of WT IucC were grown, but despite extensive optimization efforts, they were of insufficient quality to solve its X-ray crystal structure. To overcome this hurdle and grow better diffracting crystals, a surface entropy reduction strategy was employed (Fig. S4). Using one of 12 engineered IucC surface mutants (designated M1–M12), the X-ray crystal structure of IucC surface mutant M5 (containing five substitutions, S182G, E183S, D185T, Q187G, and Q188T) was solved to 2.45-Å resolution by molecular replacement (Table 2, Fig. 6).

The IucC monomeric tertiary structure is described using a cupped hand analogy (Fig. 6A), following the convention employed for other structurally-characterized NIS synthetases (38, 40). The thumb domain (residues 2–113) consists of a 3-helix bundle adjoined by a distorted 4-stranded antiparallel β-sheet. The extended loop region (114–163) contains a loop that is directed into the interior of the protein followed by four short α-helices. The fingers domain (164–362) comprises a 6-stranded and a 3-stranded anti-parallel β-sheet decorated with loops and α-helices. This domain also includes the mutated surface patch (182–188) that helped mediate crystallization. Finally, the palm domain (363–576) contains a prominent 3-helix bundle and forms the base of the central active site depression. The tertiary fold is homologous to the other structurally characterized NIS synthetases, including Type C AlcC (RMSD = 1.5 Å, 392 Ca), and AsbB (RMSD = 2.3 Å, 386 Ca), as well as Type A IucA (r.m.s. displacement = 4.1 Å, 361 Ca), and AcsD (RMSD = 5.1 Å, 315 Ca).

The IucC model has three gaps where disordered loops (DL1–3) were unresolved in the electron density map (Fig. 6A). In Fig. 6B, the three disordered loops were modeled onto the crystal structure to better appreciate their proximity to the putative active site. The first gap (DL1, ~279–288) in the fingers domain encompasses a lid loop extending over the putative active site and has been shown to be important in binding ATP and citrate in the other structurally-characterized NIS synthetases. The second gap (DL2, ~331–341) in the fingers domain is a surface loop connecting a β-sheet and a short α-helix. Finally, the third disordered loop (DL3, ~551–558) occurs proximally contacting the phosphate groups of ATP (Fig. S6). The quaternary structure of IucC active site and has been shown to be important in binding ATP and citrate in the other structurally-characterized NIS synthetases.

The third disordered loop (DL3, ~551–558) was modeled into the putative active site to highlight the proximally-located disordered loops limit our ability to infer how ahLys and citryl-ahLys bind in the active site, as well as how IucC catalyzes their condensation.

The quaternary structure of IucC

Features from the crystal lattice were combined with biochemical and solution-phase small-angle X-ray scattering (SAXS) data to investigate the oligomeric state and quaternary structure of IucC. Despite migrating a bit lower than expected by SDS-PAGE (Fig. 5A), MS indicated that the full construct was expressed, and had a monomeric mass of 66.1 kDa (Table 3).
assemblies revealed both plausible tetrameric and dimeric unit can be,
A tetramer (Fig. 7 in crystallo consistent with the general size and shape of the molecular envelope dimeric (Fig. 7 inset) assemblies are highlighted using the same color scheme as depicted as sticks. crystal structure of IucA (PDB code 5JM8) are also modeled into the active site with IucC the side chains of several predicted binding residues labeled and were collected on mass (264 kDa) (Table S2). However, when SAXS data (230–240 kDa) slightly underestimated its theoretical tetrameric mass was reduced significantly to 155 kDa, a value marginally overestimating a dimer (132 kDa). Size exclusion chromatography also illustrated the propensity of IucC to transition to a dimer upon exposure to its nucleotide substrate (Fig. S1). During purification, IucC eluted from size exclusion chromatography primarily as a tetramer, partially overlapping a secondary dimer peak (Fig. S1D). After freezing, storing, and thawing tetrameric fractions, the tetrameric oligomer was observed to be stable (Fig. S1, B and C) and appeared the most crystallizable oligomer in screening experiments. Molecular weight calculations from SAXS data of IucC in standard storage buffer (≈230–240 kDa) slightly underestimated its theoretical tetrameric mass (264 kDa) (Table S2). However, when SAXS data were collected on IucC in the presence of ATP, the calculated molecular mass was reduced significantly to ≈155 kDa, a value marginally overestimating a dimer (132 kDa). Size exclusion chromatography also illustrated the propensity of IucC to transition to a dimer upon exposure to its nucleotide substrate (Fig. S1E).

Inspection of the asymmetric unit containing eight IucC molecules revealed both plausible tetrameric and dimeric assemblies in crystallo (Fig. 6C). The crystalline asymmetric unit can be reduced into two abutting tetramers adopting a circular configuration. This tetramer can be further divided into two identical dimers bearing resemblance to the dimeric assembly reported for the Type C NIS synthetases AlcC and AsbB (Fig. S7) (40). Comparison of the experimental IucC scattering curve to scattering curves calculated from potential crystalline assemblies revealed the strongest correlation with the tetramer (Fig. 7A). Likewise, the size and shape of an ab initio molecular envelope calculated from the SAXS data were consistent with the general size and shape of the in crystallo tetramer (Fig. 7A, inset). SAXS data collected on IucC in the presence of 1 mM ATP and MgCl₂ showed a marked reduction in the near zero-angle scattering intensity, indicating a reduction in particle mass (Fig. 7B). In the conditions containing ATP, the SAXS data were most closely approximated by the curve calculated from the dimer, particularly in the low q region. Again, there is reasonable correlation between the crystalline dimer and an ab initio molecular envelope calculated from IucC in the presence of ATP (Fig. 7B, inset). Nevertheless, inspection of the molecular envelopes in Fig. 7 reveals that they are a bit too “snug” for the tetramer and a bit too “loose” for the dimer. Moreover, the molecular weight estimations seemed to underestimate a tetramer and overestimate a dimer (Table S2). Both of these observations suggest IucC likely exists as a mixture of both tetramer and dimer under both conditions. Modeling the scattering curves as a mixture of the two oligomers revealed that the experimental data were best fit by a 91.9% mixture of tetramer:dimer in storage buffer, whereas it was best approximated as a 28.72% tetramer:dimer ratio in the presence of its substrate ATP.

Computational analysis of the crystal lattice (PISA (41) and EPPIC (42)) predicted the dimer was the only biologically relevant oligomeric assembly, burying ≈1,080 Å² of surface area between the two protomers. This prediction is also supported by the observation that all four dimers within the asymmetric unit are identical, whereas the structure of the two tetramers are marginally different (Fig. S7, A and B). Taken together, it appears IucC is capable of forming both a tetramer and dimer in solution. However, although the tetrameric assembly appears to promote crystallization, it is more likely an artifactual consequence of heterologous expression and purification, with the...
Structural and functional aerobactin biosynthesis

The four purified iucA–D enzymes were also able to reconstitute aerobactin biosynthesis in vitro. Although hLys was not detected in the assay when only IucD was included, the three downstream metabolites ahLys, citryl-ahLys, and aerobactin were all clearly identified when additional luc enzymes were added. Enzyme combinations in which either IucD or IucB were omitted did not lead to the identification of nonhydroxylated or nonacetylated aerobactin analogues, respectively (data not shown). This observation demonstrates the aerobactin secondary metabolic pathway is engrained with a high-degree of substrate selectivity to limit production of nonfunctional analogues. Both of the synthetases are required to synthesize aerobactin; neither was capable of compensating for the activity of the other to any appreciable extent. This is in contrast to the Type C NIS synthetase AsbB in petrobactin biosynthesis, which was reported to partially compensate for the absence of its Type A counterpart AsbA (40). Although having one synthetase capable of catalyzing both condensations would be ideal from a metabolic efficiency perspective, the presence of two distinct synthetases in aerobactin biosynthesis, as well as many of the other NIS pathways, likely implies a difficult evolutionary challenge of designing a single stereoselective active site to accommodate both sets of substrates. One unique way of addressing this challenge would be by combining two synthetase active sites into a single polypeptide, an approach that may be adopted by the putative NIS synthetase CiuE in the biosynthesis of corynebactin, a lysine and citrate-based siderophore closely-related to aerobactin (45, 46).

Functional characterization of the Type A NIS synthetase AcsD described a highly stereoselective mechanism for the condensation of citrate and L-serine (38). Furthermore, structural models of the AcsD and IucA active sites predicted a mechanism in which only one of the two pro-chiral carboxymethyl groups of citrate is activated by ATP to form a citryl-adenylate intermediate (36, 38). Experiments presented herein confirmed the structural predictions by functionally demonstrating the stereoselective mechanisms of IucA and IucC. Although the final symmetry of aerobactin renders the citrate- and serine-derived C3 position achiral, stereochemistry at the C3 position has been shown to play a critical role in influencing siderophore potency and iron affinity in the NIS staphyloferrin B (43).

In addition to demonstrating aerobactin biosynthesis in vitro, the velocity of the pathway was determined to more quantitatively evaluate aerobactin production and compare it to other reconstituted pathways. The aerobactin turnover frequency ($6 \pm 2$ min$^{-1}$) is similar to values reported for the in vitro production of the siderophores pyochelin (2 min$^{-1}$) (47) and yersiniabactin (1.4 min$^{-1}$) (48), despite employing contrasting NRPS and hybrid NRPS/polyketide synthetase enzymatic mechanisms, respectively. The apparent steady-state kinetics of the individual aerobactin synthetase IucC was also examined quantitatively. IucC showed an affinity for ATP similar to IucA ($K_m = 130 \pm 30$ $\mu$M) (36), a result supporting the common ATP binding mechanism that NIS synthetases are biologically relevant dimer predominating in the presence of its substrate ATP.

**Discussion**

Heterologous expression of the four iucA–D genes in an aerobactin-deficient strain of *E. coli* enabled the production of the siderophore. This implies that all the necessary substrates can be harvested from the cellular metabolic pool, not requiring any additional enzymatic activity to supply the pathway. It is interesting to note that the operon of the closely related, and virulence-promoting, NIS staphyloferrin B contains specialized enzymes dedicated to citrate and amino acid precursor production (43). The relatively low burden of the 5-gene, 8-kb aerobactin operon likely contributes to its relatively wide distribution in pathogenic members of the Enterobacteriaceae family, including *Escherichia*, *Klebsiella*, *Salmonella*, *Yersinia*, *Enterobacter*, *Shigella*, *Serratia*, and *Citrobacter*, as well as marine *Vibrio* (44) species.

Figure 7. iucC SAXS. A, the experimental scattering curve of IucC in storage buffer extrapolated to infinite dilution overlaid with scattering curves calculated from potential oligomeric assemblies found in the crystalline asymmetric unit. Inset, a crystallographic tetramer docked inside an *ab initio* molecular envelope (wireframe) calculated from the experimental curve. B, the experimental scattering curve of IucC in storage buffer plus 1 mM ATP and MgCl$_2$ extrapolated to infinite dilution overlaid with scattering curves calculated from potential oligomeric assemblies found in the crystal structure. Inset, a crystallographic dimer docked inside an *ab initio* molecular envelope (wireframe) calculated from the experimental curve.
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predicted to share. The relatively low catalytic efficiency of IucC also appeared comparable with values reported for the Type C NIS synthetase homologue AsbB (240 m⁻¹ s⁻¹) (40). These seemingly modest pathway turnover frequencies and enzyme catalytic efficiencies support the premise that high production may not be essential for these siderophore pathways to meet the demand for iron.

Although WT hvKP IucC was a stable enzyme and expressed robustly, a surface entropy reduction mutant (M5) was required to grow high-quality crystals. Computational analysis of the resulting crystal lattice using EPPIC (42) revealed that residues within the mutated surface patch (182–188) contributed to one-third (5 of 15) of the distinct crystal lattice interfaces, while not disturbing the predicted biologically relevant dimer interface or catalytic competency (Fig. S5). This significant contribution to crystal lattice contacts helps rationalize how this particular mutant patch was able to promote higher quality crystal growth. Additionally, this most successful surface mutant had the highest melting temperature off all the evaluated IucC constructs, consistent with the commonly-held assertion that maximizing protein stability promotes successful crystallization (Fig. S4C) (49).

Attempts to obtain a co-crystal structure with substrates, especially with ATP, were surprisingly unsuccessful, given the co-crystallization success of the other NIS synthetases. Although the underlying reasons behind this phenomenon are uncertain, it was observed that the disordered loops (DL1–3) in the asymmetric unit. Given the presumed active site flexibility required to bind the relatively complex substrates ahLys and asymmetric unit. Given the presumed active site flexibility required to bind the relatively complex substrates ahLys and citryl-ahLys, it is possible that inter-particle interactions from the asymmetric unit. Given the presumed active site flexibility required to bind the relatively complex substrates ahLys and citryl-ahLys, it is possible that inter-particle interactions from the asymmetric unit. Given the presumed active site flexibility required to bind the relatively complex substrates ahLys and citryl-ahLys, it is possible that inter-particle interactions from the asymmetric unit. Given the presumed active site flexibility required to bind the relatively complex substrates ahLys and citryl-ahLys, it is possible that inter-particle interactions from the asymmetric unit. Given the presumed active site flexibility required to bind the relatively complex substrates ahLys and citryl-ahLys, it is possible that inter-particle interactions from the asymmetric unit. Given the presumed active site flexibility required to bind the relatively complex substrates ahLys and citryl-ahLys, it is possible that inter-particle interactions from the asymmetric unit. Given the presumed active site flexibility required to bind the relatively complex substrates ahLys and citryl-ahLys, it is possible that inter-particle interactions from the asymmetric unit. Given the presumed active site flexibility required to bind the relatively complex substrates ahLys and citryl-ahLys, it is possible that inter-particle interactions from the asymmetric unit. Given the presumed active site flexibility required to bind the relatively complex substrates ahLys and citryl-ahLys, it is possible that inter-particle interactions from the asymmetric unit. Given the presumed active site flexibility required to bind the relatively complex substrates ahLys and citryl-ahLys, it is possible that inter-particle interactions from the asymmetric unit. Given the presumed active site flexibility required to bind the relatively complex substrates ahLys and citryl-ahLys, it is possible that inter-particle interactions from the asymmetric unit. Given the presumed active site flexibility required to bind the relatively complex substrates ahLys and citryl-ahLys, it is possible that inter-particle interactions from the asymmetric unit. Given the presumed active site flexibility required to bind the relatively complex substrates ahLys and citryl-ahLys, it is possible that inter-particle interactions from the asymmetric unit. Given the presumed active site flexibility required to bind the relatively complex substrates ahLys and citryl-ahLys, it is possible that inter-particle interactions from

**Heterologous in vivo aerobactin production**

The four *iucA–D* genes were subcloned into duet expression vectors for simultaneous co-expression. The *iucB* and *iucD* genes were subcloned into MCS1 (Ncol and BamHI) and MCS2 (NdeI and Xhol) of the CDFDuet-1 expression plasmid (StrepR/Spec³), respectively. Similarly, the *iucA* and *iucC* genes were subcloned into MCS1 (Ncol and BamHI) and MCS2 (NdeI and BglII) of the pETDuet-1 expression plasmid (Amp³), respectively. The *iucA* gene construct included a N-terminal His tag, whereas the other three enzymes were untagged. *E. coli* BL21(DE3) were then transformed with either CDFDuet-1, pETDuet-1, or both of the plasmids. The transformed *E. coli* strains were grown in 60 ml of M9 minimal media with appropriate antibiotics to an *A₆₀₀ nm* of ≈0.6–0.8 (37 °C, 250 rpm). Protein expression was induced with the addition of 750 μM isopropyl 1-thio-β-D-galactopyranoside, and cells were incubated for ≈18 h overnight. The cells were pelleted by centrifugation at 4,500 rpm for 10 min, and the culture supernatant was decanted and frozen at −20 °C for subsequent analysis.

The supernatant was injected over a 3 × 50-mm Agilent InfinityLab Poroshell 120 EC-C18 2.7 μm column held at 30 °C with a flow rate of 0.4 ml/min. The mobile phases consisted of 0.1% formic acid in water (A) and 0.1% formic acid in acetonitrile (B). The elution

**Experimental procedures**

**Cloning, expression, and purification of hvKP luc enzymes**

Details of Luc enzyme expression and purification are outlined in Table S1; a general expression and purification procedure is described below. The four *iucA–D* genes were individually amplified from the hvKP1 clinical isolate (50) genomic DNA using primers to incorporate restriction sites at the 5′ and 3′ ends of the gene. The genes were subcloned into modified expression vectors with N-terminal His tags and protease recognition sites for tag removal. The vectors were transformed into *E. coli* BL21(DE3) for protein production. Cells were grown in LB media (250 rpm) to an *A₆₀₀ nm* of ≈0.5–0.7. Expression was induced with the addition of isopropyl 1-thio-β-D-galactopyranoside, followed by incubation for ≈18 h overnight at 16 °C. Cells were harvested by centrifugation at 6,000 × g for 15 min at 4 °C. After decanting off the supernatant media, the cell pellets were flash frozen in liquid nitrogen and stored at −80 °C for later use.

Frozen cell pellets were re-suspended in lysis buffer and lysed by sonication. The resulting slurry was separated by ultracentrifugation at 185,000 × g. The supernatant was filtered over a 0.45-μm polysulfone membrane before being subjected to immobilized metal affinity chromatography. The lysate supernatant was passed over a 5-ml Ni²⁺-nitritriacetic acid column. Bound proteins were eluted from the column using lysis buffer plus 300 mM imidazole. Fractions that contained His-tagged luc enzymes by SDS–PAGE were combined and dialyzed overnight at 4 °C or room temperature with the appropriate protease in dialysis buffer for His/chaperone tag removal. After spiking with imidazole to 20 mM, the dialyzed sample was passed over the Ni²⁺-nitritriacetic acid column for a second time. The flow-through fractions containing the desired protein without the His tag were combined and concentrated using an appropriate centrifugal filter before being subjected to size exclusion chromatography (HiLoad™ 16/60 Superdex™ 200, GE Healthcare Life Sciences). The desired fractions were combined, concentrated, and flash frozen in liquid nitrogen before being stored at −80 °C for subsequent use.

**Heterologous in vivo aerobactin production**

The four *iucA–D* genes were subcloned into duet expression vectors for simultaneous co-expression. The *iucB* and *iucD* genes were subcloned into MCS1 (Ncol and BamHI) and MCS2 (NdeI and Xhol) of the CDFDuet-1 expression plasmid (StrepR/Spec³), respectively. Similarly, the *iucA* and *iucC* genes were subcloned into MCS1 (Ncol and BamHI) and MCS2 (NdeI and BglII) of the pETDuet-1 expression plasmid (Amp³), respectively. The *iucA* gene construct included a N-terminal His tag, whereas the other three enzymes were untagged. *E. coli* BL21(DE3) were then transformed with either CDFDuet-1, pETDuet-1, or both of the plasmids. The transformed *E. coli* strains were grown in 60 ml of M9 minimal media with appropriate antibiotics to an *A₆₀₀ nm* of ≈0.6–0.8 (37 °C, 250 rpm). Protein expression was induced with the addition of 750 μM isopropyl 1-thio-β-D-galactopyranoside, and cells were incubated for ≈18 h overnight. The cells were pelleted by centrifugation at 4,500 rpm for 10 min, and the culture supernatant was decanted and frozen at −20 °C for subsequent analysis.

The supernatant was injected over a 3 × 50-mm Agilent InfinityLab Poroshell 120 EC-C18 2.7 μm column held at 30 °C with a flow rate of 0.4 ml/min. The mobile phases consisted of 0.1% formic acid in water (A) and 0.1% formic acid in acetonitrile (B). The elution
gradient started with isocratic 5% B for 2 min, followed by a linear gradient to 20% B over 10 min. The eluant was monitored for UV absorbance at 218 nm and by an in-line Adviron Expression 
compact mass spectrometer configured for positive mode electrospray ionization. The mass spectrometer was further configured with the following parameters: scan range: 100–1,000 m/z; capillary voltage: 150 V; ESI voltage +3,500 V; capillary temperature: 150 °C; gas temperature: 350 °C; gas flow: 4 liter/min.

**In vitro aerobactin reconstitution**

A 100-μl reaction mixture was prepared containing 50 mM HEPES pH 7.5, 5 mM MgCl₂, 50 mM NaCl, 0.2 mM tris-(2-carboxyethyl)phosphine, 5 mM L-lysine, 5 mM NADPH, 0.5 mM FAD, 5 mM acetyl-CoA, 5 mM ATP, 5 mM citrate, and various combinations of the luc enzymes at 10 μM concentration. After incubating at 37 °C, 10-μl aliquots of the reaction mixture were removed at specified time points (up to 1 h) and quenched by the addition of 40 μl of 0.1% formic acid and 5% acetonitrile in water before being eluted with 100% methanol. Fractions evidenced from the supernatant and rinsed with 30 ml of water. This crude residue was re-dissolved in 5% acetonitrile in water before being eluted with 100% methanol. Fractions evidenced from the supernatant and rinsed with 30 ml of water. The general purification strategy was informed by the work of Butler and colleagues (44, 51). After acidifying to pH 2.5 using concentrated HCl, 250 ml of M9 minimal media culture supernatant from iucA–D co-expression was stirred with 10 g of AmberLite XAD-4 resin overnight at 4 °C. The resin was filtered from the supernatant and rinsed with 30 ml of water before being eluted with 100% methanol. Fractions evidenced to contain iron chelators by the liquid chrome azurol S assay and analytical HPLC-MS analysis were combined and then concentrated to a residue (~120 mg) before being purified by the identical semi-preparative HPLC procedure outlined above. The combined yield of the reaction and purification was 65%.

**Preparation of synthetic ahLys and citryl-ahLys**

The detailed synthesis of aerobactin precursors ahLys and citryl-ahLys is presented in the supporting information (Data S10–S25). Briefly, the N⁰-oxidized and acetylated L-lysine derivative, ahLys was synthesized in accordance with previously described methods reported by Hu and Miller (53). Late stage divergence from the reported route allowed for functionalization of the α-amino moiety of the fully protected ahLys derivative (Fig. 8). Enantiopure (S)-1,3-dimethyl citrate was prepared by co-crystallization with (R)-methylbenzylamine (54, 55). Subsequent activation to the corresponding citrylchloride and coupling to the α-amino of the protected ahLys derivative yielded methyl-N⁰-acetoxy-N⁰-acetyl-N⁰-[S]-1,3-dimethylcitril]-L-lysine. Global saponification with lithium hydroxide afforded the penultimate aerobactin intermediate (3S,2'S)-citryl-ahLys with minor epimerization.

**IucC steady-state kinetics**

A coupled NADH oxidation assay was employed to evaluate the steady-state kinetics of IucC (56). Reaction mixtures (50 μl) were prepared containing 50 mM HEPES pH 7.5, 15 mM MgCl₂, 3 mM PEP, 500 μM NADH, 10 units/ml of pyruvate kinase, lactate dehydrogenase, and adenylate kinase, 2 μM IucC, and varied concentrations of substrates. The concentration of a single substrate was varied while the other two were at saturating/near saturating concentrations. When not being varied, ATP and citryl-ahLys were at 5 and 6.5 mM, respectively. When hydroxylamine was employed as a surrogate nucleophile, it was at 75 mM. Reactions were carried out in triplicate in 96-well half-area clear polystyrene plates at 37 °C; the absorbance (340 nm, ε = 6,220 M⁻¹ cm⁻¹) was measured every 10–15 s using a Biotek Synergy 4 plate reader for 10 min. The maximum initial absorbance slopes were converted to enzyme activity and plotted versus varying substrate concentrations. The plots were

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**Figure 8. Synthesis of aerobactin precursor (3S,2’S)-citryl-ahLys.** Reaction conditions: (a) Pd/C, H₂, EtOH, room temperature, 1 h; (b) SOCl₂, 85 °C, 1 h; (c) Et₃N, 0 °C to room temperature, 2 h; (d) LiOH, THF, H₂O.
then fit with nonlinear Michaelis-Menten regression in GraphPad Prism to calculate \( V_{\text{max}} \) and \( K_m \) values.

**lucC surface entropy reduction (SER) and crystallization**

The general SER approach was informed by numerous literature reports, two of which are cited here (57, 58). A homology model of hvKP lucC was created based on the crystal structures of the two Type C NIS synthetases AsbB and AlcC using YASARA software. The homology model was visually inspected to identify sequences on the surface of the enzyme enriched in high-entropy charged/polar residues, including Lys, Glu, Gln, Arg, Asp, and Asn. Mutations were then designed to introduce lower entropy side chains (Ala, Gly, Ser, Thr, Tyr, and His) to promote the formation of stable crystal lattice contacts, while trying to avoid making the surface of the enzyme too hydrophobic. In total, 12 constructs were designed with a single mutated surface patch (Fig. S4B). The hvKP lucC plasmid was sent to GENEWIZ, Inc. (South Plainfield, NJ) to synthesize and sequence the 12 mutant genes. *E. coli* BL21(DE3) were transformed with the vectors containing the mutant lucC genes, and protein expression and purification was carried out using the same general procedure described for WT lucC.

Of the 12 surface mutant proteins designed, seven were successfully purified in sufficient quantity for crystallization screening using a 120-condition in-house screen and the 1,536-condition high-throughput crystallization screening service available at the Hauptman-Woodward Medical Research Institute (59). The thermal stability of WT lucC and the seven successfully purified SER mutants was assayed using a fluorescence-based thermal shift assay (60). Although crystallization leads were identified for all seven of the surface mutants, initial crystals from surface mutant M5 appeared most promising. Conditions were optimized to \( \sim 5.5 \text{ mg/ml of lucC M5 combined 1:1 with a mixture of 10–11% PEG 20,000 and 100 mM MES, pH 6.5, in hanging-drop vapor-diffusion format at 20 °C. Microseeding was employed to control nucleation and promote the growth of single crystals. Crystals were cryo-protected by serial transfer through mixture plus 8, 16, and 24% ethylene glycol before being flash frozen in liquid nitrogen.**

**X-ray data collection, structure determination, and refinement**

Frozen crystals were shipped to GM/CA ID-B at the Advanced Photon Source (APS) at the Argonne National Laboratory (Lemont, IL) for remote X-ray diffraction data collection. A total of 400 images were collected on a single crystal using a 100 × 45 \( \mu \text{m} \) beam (no attenuation) over 360° (0.9°, 0.5 s) on a Dectris Eiger-16m detector at a distance of 350 mm. The diffraction images were processed using iMosflm (61) and the structure was solved by molecular replacement using Phaser (62) software and employing monomeric AlcC (PDB code 3XOQ, 29.9% identity) as the search model. Automated model building was performed using PHENIX.Autobuild (63), manual model building and refinement was carried out using Coot (64), and automated structure refinement was executed using PHENIX.Refine (65). Data collection and refinement statistics are presented in Table 2. In addition to the disordered loops discussed in the text, the protein chain was modeled for residues 542–545 despite their weak electron density. Structure factors and coordinates for hvKP lucC surface mutant M5 have been deposited in the Protein Data Bank (code 6CN7).

**SAXS**

SAXS data on WT hvKP lucC were collected at Stanford Synchrotron Radiation Lightsource (SSRL) beamline 4—2 (66, 67). Protein samples (40 \( \mu \text{l} \)) were prepared in a dilution series of 5.3, 4.0, 2.7, and 1.3 mg/ml in a buffer of 50 mM HEPES, 150 mM NaCl, and 0.2 mM tris(2-carboxyethyl)phosphine. To investigate the effect of ATP on the conformation of lucC, a second dilution series (5.1, 3.8, 2.5, and 1.3 mg/ml) was prepared in an identical buffer system plus 1 mM ATP and MgCl₂. Each sample was held at 10 °C and data were collected with 20 exposures of 1-s duration. Scattering from a buffer blank was subtracted from the data. The scattering curves were produced using SAS-Tool software (67) and the scattering curves were analyzed using software from the ATSAS program package (68): PRIMUSQT, DAMMIF, DAMAVER, CRYSOL, SUPCOMB, OLIGOMER, and DATVC. Guinier analysis (0.4 ≤ qR ≤ 1.3) was used to determine the zero-angle scattering intensity (\( I_0 \)). This value was then used to calculate the molecular mass by employing lysozyme (14.3 kDa, 16.3 mg/ml) as a reference standard. For molecular weight calculations based on Porod volume, the pair distance distribution (\( P(r) \)) function (\( q ≤ 0.3 \)) was employed to calculate the Porod volume, which was then divided by 1.66 to yield the molecular mass in Da (69). The DATVC software was used to calculate the molecular weight by the volume of correlation (\( V_c \)) including data for \( q ≤ 0.3 \). The lucC ab initio molecular envelopes were calculated using P222 (without ATP) and P2 (with ATP) symmetry restraints in DAMMIF and including data for \( q ≤ 0.3 \). The CRYSOL scattering curves were calculated using the solvent electron density of water (0.334 e⁻/Å³). To correct for a slight concentration dependence (minor interparticle repulsion) in the SAXS data, PRIMUSQT was used to generate scattering curves extrapolated to infinite dilution.

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