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# Lipoprotein SmpA is a component of the YaeT complex that assembles outer membrane proteins in *Escherichia coli*

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A major role of the outer membrane (OM) of Gram-negative bacteria is to provide a protective permeability barrier for the cell, and proper maintenance of the OM is required for cellular viability. OM biogenesis requires the coordinated assembly of constituent lipids and proteins via dedicated OM assembly machineries. We have previously shown that, in *Escherichia coli*, the multicomponent YaeT complex is responsible for the assembly of OM  $\beta$ -barrel proteins (OMPs). This complex contains the OMP YaeT and three OM lipoproteins. Here, we report another component of the YaeT complex, the OM lipoprotein small protein A (SmpA). Strains carrying loss-of-function mutations in *smpA* are viable but exhibit defects in OMP assembly. Biochemical experiments show that SmpA is involved in maintaining complex stability. Taken together, these experiments establish an important role for SmpA in both the structure and function of the YaeT complex.

The outer membrane (OM) of Gram-negative bacteria, such as *Escherichia coli*, constitutes the outermost compartment of the cell and contributes to overall cellular integrity. The barrier function of the OM requires stable maintenance of a unique lipid and protein composition. It is an asymmetric bilayer with phospholipids in the inner leaflet and glycolipids, primarily LPS, in the outer leaflet (1). There are two major types of proteins in the OM, OM  $\beta$ -barrel proteins (OMPs) and OM lipoproteins. The OMPs consist of antiparallel  $\beta$ -strands, organized into a cylindrical shape, producing a  $\beta$ -barrel, whereas OM lipoproteins are tethered to the inner leaflet of the OM via amino-terminal lipid moieties.

All of the components of the OM are synthesized within the cytoplasm and must be subsequently targeted to the OM. Transport of these amphipathic molecules across the hydrophilic environment of the periplasm requires dedicated transport and assembly machineries. Significant progress has been made in identifying various proteins required for OM biogenesis in *E. coli* (2). Lipoproteins are targeted by components of the essential Lol pathway (3). The essential OM YaeT complex is required for OMP assembly (4), whereas yet another essential OM complex, Imp/RlpB, catalyzes LPS assembly (5–7). It is likely that the targeting and insertion of LPS and OMPs are in some way coordinated to maintain a proper lipid–protein ratio and preserve the integrity of the OM. Genetic evidence supporting such a homeostatic mechanism exists, because mutations in genes specifying members of the YaeT complex can suppress specific OM defects conferred by a particular mutation in *imp* (8–10).

Information concerning the interactions between the members of the YaeT complex has been obtained through both genetic and biochemical analysis. This complex contains the OMP YaeT and three associated OM lipoproteins, YfgL, NlpB, and YfiO (4). Cells lacking either NlpB or YfgL are viable and exhibit only minor OMP assembly defects (9, 11). The other members of the complex, YaeT and YfiO, are essential, and depletion of either of these proteins from the cell results in severe OMP targeting defects (4, 11, 12).

Here, we report an additional, nonessential component of the YaeT complex, the OM lipoprotein small protein A (SmpA). We show that SmpA plays an important role in both the stability and function of the YaeT complex.

## Results

**SmpA Copurifies with the YaeT Complex.** To gain insight into the structure of the YaeT complex, we have been probing the interactions between the different protein components (4, 11). Biochemical studies using wild-type and mutant proteins raised the possibility that the complex might contain an additional component(s). To address this possibility, we used a variety of different methods to purify the complex before and after cross-linking with formaldehyde. Potential interacting proteins were excised from the gel and identified by tandem mass spectrometry. One of the proteins identified was SmpA (molecular weight, 12,162.5 Da) (13). This protein was of particular interest, because a homolog of SmpA in *Pseudomonas aeruginosa*, OmlA, has been characterized as an OM lipoprotein that functions in maintaining the cell envelope integrity (14).

We cloned the *smpA* gene from the *E. coli* genome and modified it to express SmpA with a C-terminal His tag. This construct produced a tagged but functional SmpA protein that complemented the *smpA*-null phenotype discussed below (data not shown). We used this His-tagged SmpA protein for affinity purification experiments to verify its interaction with the YaeT complex. As shown in Fig. 1A, SmpA-His successfully pulled down all four previously known members of the YaeT complex. Complementary affinity purifications verified that SmpA copurifies with the YaeT complex regardless of which lipoprotein member was tagged (Fig. 1B). These experiments demonstrate that SmpA interacts directly with the YaeT complex *in vivo*.

**SmpA Is Not Essential for Viability in *E. coli*.** The YaeT complex is required for the assembly of OMPs into the OM (4), and two of its members, YaeT and YfiO, are essential in *E. coli* (4, 12). To determine whether SmpA performs an essential cellular function, we replaced the last 110 codons of the 113 codons of *smpA* with a gene coding for either chloramphenicol or kanamycin resistance by using previously described recombining methods (15). Both of these mutant *smpA* strains were phenotypically identical in terms of growth on LB medium. Because there is no gene immediately downstream of *smpA* (16) and we are able to complement *smpA* mutant phenotypes with a functional copy of *smpA* on a plasmid (data not shown), any phenotype arising from an *smpA*-null strain would not be a consequence of polarity. We monitored the growth of strains lacking SmpA at 37°C in rich media and observed no apparent growth defect (Fig. 2A). Strains lacking SmpA grew normally at temperatures ranging from 23°C to 42°C in both rich

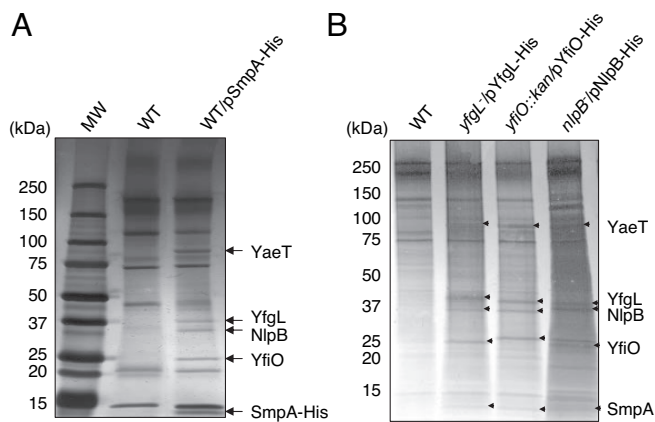
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The authors declare no conflict of interest.

Abbreviations: OM, outer membrane; OMP, outer membrane  $\beta$ -barrel protein; SmpA, small protein A; MBP, maltose-binding protein; TBS, Tris-buffered saline.

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**Fig. 1.** SmpA copurifies with the YaeT complex. (A) Silver-stained SDS/polyacrylamide gel of samples from cell lysates of wild-type (WT), *wt/pSmpA-His* strains immunoprecipitated with anti-His tag antibody. (B) Silver-stained SDS/polyacrylamide gel of samples from cell lysates of WT, *yfgL-1pYfgL-His*, *yfiO::kan/pYfiO-His*, and *nlpB-1pNlpB-His* strains immunoprecipitated with anti-His tag antibody. MW, molecular weight.

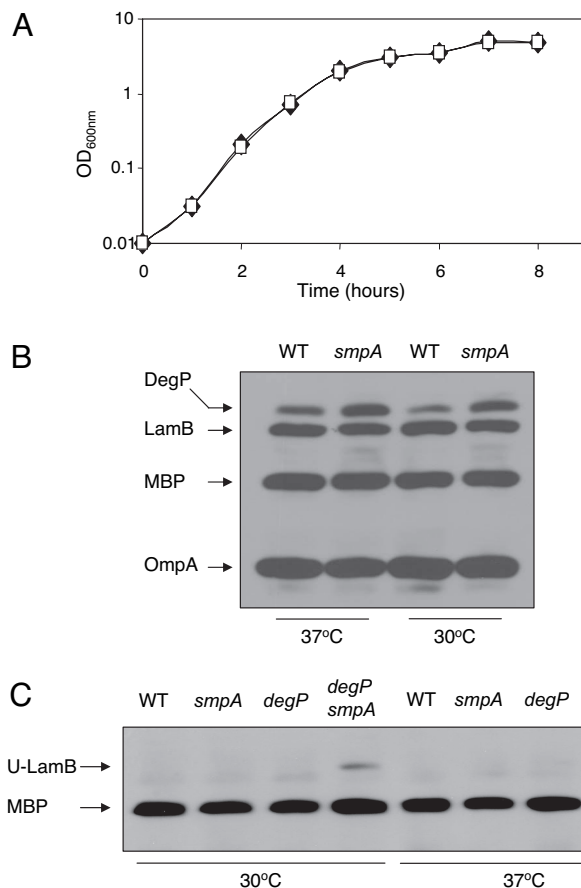
and minimal media (data not shown). Therefore, SmpA is a nonessential member of the YaeT complex.

**SmpA Mutants Exhibit OMP Assembly Defects.** Although SmpA is not required for cell viability, it is still possible that it plays a nonessential role in OM biogenesis. If this is true, strains lacking SmpA may exhibit increased sensitivity to various toxic small molecules (2). Indeed, *smgA* strains exhibited 4-fold increased sensitivity to rifampin and 2-fold increased sensitivity to cholerae relative to the wild-type strain. Furthermore, *smgA* strains were unable to grow on medium containing 0.5% SDS and 1 mM EDTA. Therefore, the barrier function of the OM is compromised in *smgA* mutants.

Given that SmpA associates with the YaeT complex (Fig. 1), strains that lack this protein may be unable to efficiently assemble OMPs in the OM. For example, strains that lack YfgL, another nonessential lipoprotein member of the YaeT complex, exhibit OMP assembly defects as evidenced by decreased steady-state levels of two OMPs, LamB and OmpA (9). Similarly, we found a slight decrease in the levels of OmpA and LamB in an *smgA* mutant strain, whereas the levels of the periplasmic protein maltose-binding protein (MBP) were unaffected (Fig. 2B).

The slightly lower levels of OMPs in an *smgA* mutant may result from the degradation of mistargeted OMPs by periplasmic proteases such as DegP. Activation of the  $\sigma^E$  stress response, which is induced by the presence of misfolded OMPs in the periplasm, increases the levels of DegP (17, 18). If the loss of SmpA leads to an OMP assembly defect, DegP levels should increase. Steady-state levels of DegP were elevated 1.5- to 2-fold in an *smgA* mutant (Fig. 2B), reflecting the stress caused by defective assembly of OMPs in this strain.

To directly observe the folding status of OMPs in various mutants, we used a gentle-lysis protocol, which preserves the native conformation of OMPs (19). Normally, processed LamB monomer assumes a folded structure that subsequently forms trimers in the OM (20). When OMP assembly is compromised, the accumulation of steady-state levels of unfolded LamB monomer represents a population of LamB molecules that is unable to assemble properly into trimers. However, unfolded LamB is often hard to detect because it is rapidly degraded by the DegP protease (19). Indeed, although we were unable to detect any unfolded LamB monomer in an *smgA* mutant, we observed a significant amount of unfolded LamB in an *smgA degP* double mutant at both 30°C and 37°C (Fig. 2C). Based on these results, we suggest that unfolded LamB

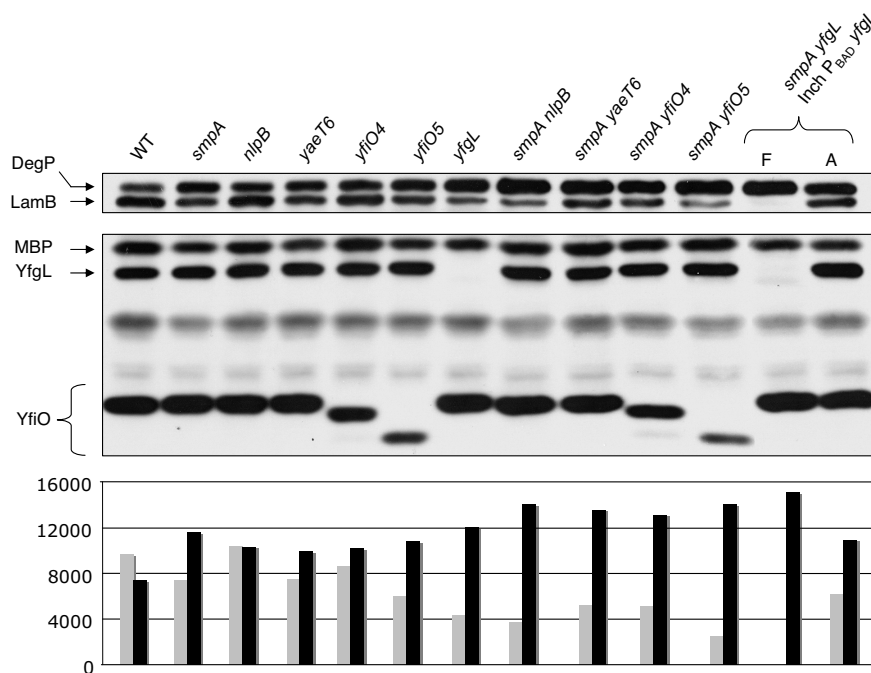


**Fig. 2.** SmpA is not essential for growth in *E. coli*. (A) Growth curves for wild-type cells ( $\blacklozenge$ ) or *smgA* cells ( $\square$ ) at 37°C are shown. Growth was monitored by measuring the optical density at 600 nm every hour for 8 h. The results of a representative experiment are shown above. (B) Levels of envelope proteins in an *smgA* mutant during midlogarithmic growth at 37°C and 30°C. Western blot analysis shows a very small reduction in the levels of OmpA and LamB in an *smgA* mutant strain when compared with a wild-type (WT) strain. Levels of the periplasmic chaperone/protease DegP are elevated 1.5- to 2-fold in an *smgA* mutant. (C) Unfolded LamB accumulates in an *smgA degP* double mutant. Cells were grown to midlogarithmic growth at 37°C and 30°C and harvested by centrifugation. These samples were gently lysed to preserve the native conformation of OMPs. Western blot analysis by using LamB antiserum was used to visualize unfolded LamB (U-LamB). Levels of MBP are shown as a loading control.

accumulates in cells lacking SmpA and is rapidly degraded by DegP. Thus, *smgA* mutants exhibit minor defects in the assembly of OMPs.

**SmpA Affects the Stability of the Multiprotein YaeT Complex.** We have previously shown that affinity purification of a particular His-tagged member of the complex in a wild-type background allows copurification of all other complex members (4) (Fig. 1). When these same experiments are performed in backgrounds defective in any one component of the complex, these interactions are weakened to varying degrees. The absence of nonessential components or the presence of defective mutant proteins destabilize the interaction between the His-tagged protein and other components of the complex, which can be judged by the amount of proteins that copurifies with the His-tagged protein in each mutant strain. This type of analysis allows us to determine the relevant contributions of each protein to the stability of the complex as a whole. By using this method, we have previously shown that the YaeT6 mutation does not affect the stability of the complex (8), that YaeT interacts with YfgL independently of YfiO and NlpB, and





**Fig. 4.** *smpA* genetic interactions. Western blot analysis shows the levels of DegP, LamB, MBP, YfgL, and YfiO proteins in strains carrying mutations that affect different YaeT complex members and in *smpA* double mutants during midlogarithmic growth at 30°C. The *smpA yfgL* depletion strain was grown in the presence of L-arabinose (A) to induce expression of *yfgL* or in the presence of D-fucose (F) to prevent its expression. Below the Western blots is a chart that shows the quantification of LamB (gray bars) and DegP (black bars). The values of the y axis represent the areas of the bands as quantified by using ImageJ image-processing software.

biochemical results demonstrating that SmpA is a structurally and functionally important component of the YaeT complex.

In *E. coli*, defects in OMP folding as well as changes in OM composition trigger both the  $\sigma^E$  and Cpx stress responses (21, 22). These stress responses help to alleviate problems in the cell envelope by up-regulating the expression of genes encoding periplasmic folding factors, proteases, and proteins involved in biosynthesis and assembly of cell envelope components. Approximately 100 genes have been predicted to be members of the  $\sigma^E$  and Cpx regulons in *E. coli* (23, 24). The genes that encode all of the components of the YaeT complex are all members of the  $\sigma^E$  regulon (12, 24–27). Therefore, like the genes encoding other members of the YaeT complex, *smpA* expression is up-regulated by the  $\sigma^E$  regulon when the OM biogenesis is compromised. In a genome-wide profiling study to look for genes that might be regulated by the CpxA/R envelope stress response in *E. coli*, multiple potential CpxR-P binding sites were identified in the promoter region of *smpA* (23). However, additional experiments are needed to verify the physiological significance of these observations.

Mutations in members of the YaeT complex show increased sensitivity to hydrophobic antibiotics, bile salts, and detergents (8, 9). The *Pseudomonas aeruginosa* homolog of *smpA*, *omlA* (39% sequence identity), was reported to be an OM lipoprotein (14), and *omlA* mutants showed hypersensitivity to anionic detergents such as SDS and increased sensitivity to hydrophobic antibiotics such as rifampin (14). Based on these phenotypes, it is safe to assume that the cell envelope of *P. aeruginosa* is compromised in *omlA* mutants (14). This is also true in *E. coli*, because we have observed increased sensitivity to cholate, rifampin, and SDS/EDTA in *smpA* mutants in addition to slight defects in OMP biogenesis (Fig. 4).

Affinity purification of the YaeT complex in strains that carry mutations in different lipoprotein components of the complex provide us with information about how the complex is organized and stabilized. Based on this type of analysis, we proposed that YaeT contacts both YfiO and YfgL directly and that the C terminus

of YfiO is required to maintain a stable interaction between NlpB and the rest of the complex (11). Here, we have shown that SmpA, a new component in the multiprotein YaeT complex, associates with YfiO and NlpB independent of YfgL (Fig. 3). The identification of this additional component may explain why OM defects are worse in doubly mutant *nlpB<sup>-</sup> yfiO $\Delta$ C-term* cells than in either of the single mutants because there are more interdependent interactions than we originally detected.

Clearly, three lipoproteins in the YaeT complex, YfgL, NlpB, and SmpA, perform nonessential function(s) because strains lacking each one of these three proteins individually are viable and exhibit rather modest OM defects. It follows therefore that the function(s) performed by these three lipoproteins serve instead to increase the rate and/or the efficiency of OMP assembly, either directly or indirectly. Indeed, strains lacking YfgL exhibit defects in the rate and/or the efficiency at which OMP assembly intermediates are delivered to the YaeT complex, but once delivered, OMP assembly appears to occur at normal rates (28). As noted above, YfgL interacts with YaeT in a manner that is independent of NlpB, SmpA, and YfiO and vice versa. In light of these structural differences, it will be of interest to determine whether NlpB and SmpA function at different steps in the complex process of OMP assembly.

## Materials and Methods

**Strains and Plasmids.** Strains are listed in Table 1. All strains were constructed by using standard microbiological techniques. The chromosomal disruption of the *smpA* gene was generated according to the procedure described by Datsenko and Wanner (15). PCR products were generated by using the following primers (5'-TGAGCCACGTA CTGCTCGGGCCCGAAAAGGAATC-AAATCACTATGCGCTGTTTGTAGGCTGGAGCTGCTTCG-3' and 5'-CAGCGCAGGTTTGTATCAATATTGGTCAACACACCGCTACTGTTAAAGGTGCATATGAATATCC-TCTTAG-3') from the template plasmids pKD3 and pKD4. The PCR products were gel-purified and resuspended in distilled water.

**Table 1. Strains and plasmids**

Strains or plasmids	Genotype and relevant features	Ref.
<b>Bacterial strains</b>		
MC4100	F <sup>-</sup> <i>araD139</i> Δ( <i>argF-lac</i> ) <i>U169 rpsL150 relA1 flb5301 deoC1 ptsF25 thi</i>	30
DY378	W3110 λcl857 Δ( <i>cro-bioA</i> )	29
JG5212	MC4100 <i>smpA::cam</i>	This study
JG5213	MC4100 <i>smpA::kan</i>	This study
JCM158	MC4100 <i>ara</i> <sup>fl</sup> , spontaneous arabinose-resistant	11
JCM175	JCM158 <i>yfgL::kan</i>	This study
JCM252	JCM158 <i>yfgL::kan</i> Δ( <i>latt-lom</i> ):: <i>bla</i> P <sub>BAD</sub> <i>yfgL araC</i>	This study
JCM304	JCM158 <i>nlpB::kan</i>	11
JCM344	JCM158 <i>yfiO4::cam</i>	11
JCM345	JCM158 <i>yfiO5::cam</i>	11
JCM375	JCM158 <i>smpA::cam</i>	This study
JCM456	JCM158 <i>yfiO4::cam smpA::cam</i>	This study
JCM457	JCM158 <i>yfiO5::cam smpA::cam</i>	This study
JCM473	JCM252 <i>smpA::cam</i>	This study
JCM474	JCM158 <i>smpA::cam nlpB::kan</i>	This study
JCM480	JCM158 <i>yaeT6 yadG::cam</i>	This study
JCM494	JCM158 <i>smpA::cam yaeT6 yadG::cam</i>	This study
<b>Plasmids</b>		
pBAD18	Cloning vector; P <sub>BAD</sub> -dependent expression	Invitrogen
pKD4	Contains a kanamycin-resistance cassette	15
pKD3	Contains a chloramphenicol-resistance cassette	15
pET42a(+)	Cloning vector with His tag	Novagen
pET22b(+)	Cloning vector	Novagen
pET22-42	Cloning vector derived from pET22b(+) and pET42a(+)	This study
<i>pYfgL-His</i>	Encodes C-terminal His-tagged YfgL	4
<i>pYfiO-His</i>	Encodes C-terminal His-tagged YfiO	4
<i>pNlpB-His</i>	Encodes C-terminal His-tagged NlpB	4
<i>pSmpA-His</i>	Encodes C-terminal His <sub>6</sub> -tagged SmpA	This study
<i>pYaeT-His</i>	Encodes N-terminal His <sub>6</sub> -tagged YaeT	This study
pBAD18:: <i>yfgL</i>	P <sub>BAD</sub> - <i>yfgL</i> on pBAD18 vector	This study

One hundred microliters of cells and 10–50 ng of PCR product were used to transform the lambda-red strain DY378 (29). Mutations were verified by using PCR with primers external to the ORF of *smpA*. Sequences are available upon request. All *smpA* mutants and double mutants were constructed by moving either *smpA::cam* or *smpA::kan* into the recipient strain by using P1 transduction.

The multiple cloning site of the pET42a(+) vector (Novagen, Madison, WI) was moved into the pET22b(+) vector (Novagen) by using XbaI and Bpu1102I restriction sites to create pET22-42. Primers 5'-ATGACATATGCGCTGTAAAACGCTGACTGC-3' and 5'-ACGTCTCGAGGTTACCACTCAGCGCAGGTTTGTATCAATATTG-3' were used to amplify *smpA* from MC4100 genomic DNA by using PCR. The fragment was introduced into pET22-42 by using XhoI and NdeI restriction enzymes. The resulting construct, *pSmpA-His*, expresses C-terminally tagged SmpA that complements the *smpA* chromosomal disruption described above.

To construct a strain carrying a second, arabinose-inducible copy of *yfgL*, we first cloned *yfgL* into pBAD18 by using primers 5'-TTGAATTCGATTACATTTTGAGGA-3' and 5'-TTCTAGATTATGTATTGCTGCTGT-3' to amplify the gene from the MC4100 chromosome and by using EcoRI and XbaI restriction enzymes for cloning. Next, the portion of the plasmid containing *araC* P<sub>BAD</sub>-*yfgL* *bla* was integrated into the λ-attachment site by using the protocol established by Boyd *et al.* (30) to create JCM252 (Table 1).

The plasmid expressing His-tagged YaeT, *pYaeT-His*, was constructed such that an in-frame hexahistidine tag was incorporated near the N terminus of the mature sequence. Two PCR products

were amplified from the MC4100 genome, corresponding to the N-terminal plus 6x-His and C-terminal sequences, respectively, ligated together, and reamplified to obtain the full-length *yaeT-His* DNA fragment. The N-terminal fragment of *yaeT* corresponding to the sequence encoding the first 21 amino acids of the protein was amplified by using primers 5'-GTCCTAGAGCATATGGCGATGAAAAAGTTGC-3' (NdeI) and 5'-ACACCTGCAGCATGGTGATGGTGATGGTGAGCACCGTATACGGTGG-3' (PstI). The C-terminal primer for this reaction incorporated sequence coding for the amino acids HHHHHHAA. The C-terminal fragment of *yaeT-His* was amplified by using primers 5'-ACACGCTGCAGAAAGGGTTCGTAGTGAAAGATATTC-3' (PstI) and 5'-ACACGCGGCCGCTTACCAGGTTTACCAGTGTAAA-CTG-3' (NotI). The two PCR fragments were digested with PstI, ligated together, and reamplified by using the external primers described above. This PCR product was digested with NdeI and NotI and ligated into pET22b(+) treated with the same enzymes. The resulting construct encodes the His-tagged YaeT with an insertion of eight residues, HHHHHHAA, between Ala-21 and Glu-22. Strains carrying this plasmid can be crossed via a P1 lysate with a *yaeT* null allele (*yaeT::kan*) (4), demonstrating that the His-YaeT protein expressed from *pYaeT-His* is functional.

**Media and Growth Conditions.** LB was prepared as previously described (31). When necessary, media were supplemented with 20 μg/ml chloramphenicol, 25 μg/ml kanamycin, 5 μg/ml rifampin, or 25 μg/ml tetracycline. All bacterial cultures were grown under aerobic conditions at 30°C unless otherwise noted. Minimum inhibitory concentration experiments for cholera and rifampin were performed as described previously (8).

**Western Blot Analysis.** Cultures were grown overnight in LB medium, with strain JCM473 supplemented with 0.025% L-arabinose. Cultures were diluted the following day into fresh media 1:100, with strain JCM473 subcultured at the same dilution into medium containing 0.025% of either L-arabinose or the competitive inhibitor of arabinose, D-fucose (32). After growth to an OD<sub>600</sub> of 0.4–0.45, 1-ml samples were harvested by centrifugation and immediately resuspended in SDS loading buffer in a volume in milliliters equal to the OD<sub>600</sub> ÷ 6. After boiling for 10 min, equal volumes were loaded onto 12% polyacrylamide gels. Gentle lysis was performed on relevant samples as described previously (19). After electrophoresis, gels were transferred to a nitrocellulose membrane and probed with antibodies used to detect LamB and OmpA (1:15,000 dilution) (18), MBP (1:15,000 dilution) (19), DegP (1:30,000 dilution) (33), YfgL (1:10,000 dilution) (4), YaeT (1:3,000) (this study), and YfiO (1:20,000 dilution) (this study). The YaeT antibody was generated against the first 15 residues of the mature protein (AEGFVVKDIHFEGQL) (Rockland Immunochemicals, Gilbertsville, PA). Donkey anti-rabbit IgG horseradish peroxidase (Amersham Biosciences, Piscataway, NJ) conjugate was used as the secondary antibody at a concentration of 1:8,000. ECL (Amersham Biosciences) and XAR film (Kodak, Rochester, NY) were used to visualize the protein bands. Protein bands were analyzed with gel image analysis software (ImageJ Software).

To generate the YfiO antibody, His-tagged cells expressing signal sequenceless YfiO were lysed by French press and centrifuged at 10,000 × g for 20 min. The soluble fraction was subjected to Ni-nitrilotriacetic acid affinity purification as described below. The eluted solution was dialyzed against 20 mM Tris buffer (pH 8) and 150 mM NaCl. The resulting protein preparation was used to generate a rabbit polyclonal antiserum (Princeton Animal Facility, Princeton University).

**Immunoprecipitation.** Cells were grown in the 50 ml of LB media to an OD<sub>600</sub> of ≈0.6 and harvested by centrifugation at 5,000 × g for 10 min. The cells were resuspended in 1 ml of Tris-buffered saline (TBS) with 1% ZW3-14 and supplemented with 100 μg/ml lysozyme, 50 μg/ml DNase I, and 50 μg/ml RNase. Cells were lysed by incubating the mixture at room temperature for 20 min on a shaker. The lysate was then centrifuged at 18,000 × g for 5 min, and the supernatant was transferred into a new Eppendorf tube. A total of 4 μg of anti-5His monoclonal antibody (Qiagen, Valencia, CA) was added into the lysate, and the mixture was shaken at 4°C for 1 h. A total of 60 μl of protein G-Sepharose beads (Sigma–Aldrich, St. Louis, MO) was added, and the mixture was shaken for another hour at 4°C. This mixture was then loaded into a micro bio-spin column (Bio-Rad, Hercules, CA), and the beads were washed with 3 ml of 1× immunoprecipitation buffer (Sigma–Aldrich) and

then eluted with 50 μl of boiling SDS-sample buffer without 2-mercaptoethanol. A total of 25 μl of eluate was used for SDS-PAGE analysis. Silver stain was conducted by using the Silver Stain Plus kit from Bio-Rad and by following the protocol therein.

**Affinity Purification.** A total of 250 ml of culture was grown in LB media to midlogarithmic growth (OD<sub>600</sub>, 0.4–0.6). Cells were harvested by centrifugation at 5,000 × g for 10 min. Cells were lysed in 2 ml of TBS (20 mM Tris·HCl, pH 7.4, 150 mM NaCl), 2% Triton X-100, and 10 mM EDTA, supplemented with 100 μg/ml lysozyme and 50 μg/ml DNase I and RNase by shaking at room temperature for 10 min. MgCl<sub>2</sub> solution was added to a final concentration of 10 mM, and the suspension was shaken for another 10 min. The cell lysate was centrifuged at 10,000 × g for 10 min to remove unbroken cells and cell debris. The clear lysate was exchanged into buffer TBS, 0.1% Triton X-100, and 20 mM imidazole by using a PD-10 column (GE Healthcare, Piscataway, NJ). The lysate was loaded twice into a column packed with 0.5 ml of Ni-nitrilotriacetic acid resin (Qiagen), which had been preequilibrated with the previous buffer. The column was washed with 10 ml of TBS, 0.05% N-dodecyl-β-maltoside, and 20 mM imidazole, and eluted with 5 ml of TBS, 0.05% N-dodecyl-β-maltoside, and 200 mM imidazole. The eluate was concentrated in an ultrafiltration device (Amicon Ultra; Millipore, Beverly, MA; MWCO 5K) by centrifugation at 5,000 × g for ≈2 h to a volume of ≈50 μl. Samples were mixed 1:1 with SDS loading buffer and boiled for 10 min, and equal volumes were loaded onto 4–20% gradient polyacrylamide gels. After electrophoresis, gels were transferred to a PVDF membrane and probed with antibodies as described above. YfiO, YfgL, YaeT, and His antibodies were all used at a dilution of 1:3,000.

**Cellular Protein Levels in Different YaeT-His and SmpA-His Strains.** To adequately detect cellular levels of YaeT-His, YfgL, YfiO, and SmpA-His, the membrane fraction of cell cultures was isolated. A total of 250 ml of culture was grown in LB media to OD<sub>600</sub> ≈ 0.6. Cells were harvested by centrifugation at 5,000 × g for 10 min. The cells were lysed in 10 ml of TBS (20 mM Tris·HCl, pH 7.4, 150 mM NaCl) and supplemented with 100 μg/ml lysozyme and 50 μg/ml DNase I and RNase by passage through a French pressure cell. Unlysed cells were pelleted at 5,000 × g for 10 min. The membrane fraction was pelleted by ultracentrifugation at 100,000 × g for 30 min. The membrane was resuspended in 250 μl of TBS, 2% Triton X-100, and 10 mM EDTA, and insoluble lipids were again pelleted at 100,000 × g for 30 min. The soluble fraction was mixed 1:1 with SDS loading buffer and boiled for 10 min. Equal volumes were loaded onto 4–20% gradient polyacrylamide gels. After electrophoresis, gels were transferred to a PVDF membrane and probed with antibodies as described above.

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