Crystal structure analysis reveals *Pseudomonas* PilY1 as an essential calcium-dependent regulator of bacterial surface motility

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Edited by John J. Mekalanos, Harvard Medical School, Boston, MA, and approved December 3, 2009 (received for review October 7, 2009)

Several bacterial pathogens require the “twitching” motility produced by filamentous type IV pili (T4P) to establish and maintain human infections. Two cytoplasmic ATPases function as an oscillatory motor that powers twitching motility via cycles of pilus extension and retraction. The regulation of this motor, however, has remained a mystery. We present the 2.1 Å resolution crystal structure of the *Pseudomonas aeruginosa* pilus-biogenesis factor PilY1, and identify a single site on this protein required for bacterial translocation. The structure reveals a modified β -propeller fold and a distinct EF-hand-like calcium-binding site conserved in pathogens with retractile T4P. We show that preventing calcium binding by PilY1 using either an exogenous calcium chelator or mutation of a single residue disrupts *Pseudomonas* twitching motility by eliminating surface pili. In contrast, placing a lysine in this site to mimic the charge of a bound calcium interferes with motility in the opposite manner—by producing an abundance of nonfunctional surface pili. Our data indicate that calcium binding and release by the unique loop identified in the PilY1 crystal structure controls the opposing forces of pilus extension and retraction. Thus, PilY1 is an essential, calcium-dependent regulator of bacterial twitching motility.

calcium binding | human pathogen | microbiology | pilus biogenesis | structural biology

The pathogenesis of bacterial infections typically depends on microbial adherence to host tissue, aggregation of the pathogen at the site of infection, and subsequent dissemination to other anatomical sites within the host. Type IV pili (T4P) are thin surface filaments involved in biofilm formation, bacterial aggregation, and microbial adherence to biotic and abiotic surfaces. T4P biogenesis requires more than a dozen proteins conserved across a range of bacterial species (1). A subset of bacteria, including the opportunistic human pathogen *Pseudomonas aeruginosa*, cyclically extend and retract their T4P, facilitating a form of surface movement termed twitching motility (2–4). Retractable T4P are required for *P. aeruginosa* adherence to host tissue, virulence, and dissemination of infection (5–7).

The pilin subunits that form T4P are incapable of self-assembly. Instead, pilus extension and retraction requires repeated rounds of pilin polymerization and depolymerization, both of which demand ATP hydrolysis (8, 9). In *P. aeruginosa*, the cytoplasmic ATPase PilB drives the assembly of pilin monomers into mature T4P (8, 10), while the structurally related cytoplasmic ATPase PilT disassembles T4P back into pilin monomers (8, 11, 12). Loss-of-function mutations in PilT result in the loss of twitching motility due to the inability of formed pilus fibers to retract (12). The C-terminal half of the 117-kDa *P. aeruginosa* PilY1 protein shares sequence homology with the C-terminal domain (CTD) of the PilC proteins of pathogenic *Neisseria* species; in contrast, the N-terminal regions of the PilCs and PilY1 are divergent in sequence (13). Although the PilC proteins from

Neisseria have been implicated in antagonizing pilus retraction by the pilus-biogenesis ATPase PilT, the N-terminal domains of these proteins appear critical for host cell attachment (14, 15). Thus, we hypothesized that the conserved CTDs of PilY1 and PilC may play a common role in T4P biogenesis rather than adhesion, and pursued a structural and functional analysis of the CTD from *Pseudomonas aeruginosa* PilY1.

Results

PilY1 Exhibits a Modified β -Propeller Fold. Examination of the amino acid sequence of PilY1 from *P. aeruginosa* revealed conservation between its C-terminal 550 residues and the same regions of the PilC proteins from *Neisseria*, and a predicted structural homology to the eight-bladed β -propeller fold exhibited by a range of proteins (16, 17). Thus, we overexpressed, purified, and crystallized the PilY1 CTD (amino acids 6145–1163) for examination by x-ray diffraction. Native diffraction data were collected to 2.1 Å resolution (Table 1); however, attempts to determine the structure using existing β -propeller search models or by isomorphous replacement methods were not successful. PilY1 CTD contains only six methionine residues, which upon selenomethionine substitution failed to produce an adequate signal to support structure determination by anomalous dispersion (18). To overcome this problem, a series of single leucine-to-methionine (L-M) substitutions were placed in the PilY1 CTD to provide additional sites for single-wavelength anomalous dispersion (SAD) phasing (Figs. S1 and S2). After several attempts, three L-M substitutions (at positions 712, 812, 823) were successfully combined, crystallized, and generated clear selenomethionine fluorescence signal upon examination in the x-ray beam. The structure was determined by SAD phasing using crystals containing nine total selenomethionine residues per protein monomer (Table 1).

The structured 505 residues of the PilY1 CTD exhibit a seven-bladed modified β -propeller fold composed of 31 β -strands and 9 α -helices (Fig. 1A and Figs. S1 and S2; residues at the termini, as well as 713–721 and 1061–1068 were disordered and are not in the refined model). Blades I through IV are four-stranded antiparallel β -sheets and align well with blades of a canonical β -propeller enzyme, quinoximoprotein alcohol dehydrogenase from *Comamonas testosteroni* [Protein Data Bank (PDB)

Author contributions: J.O., M.D.J., K.A.C., M.C.W., and M.R.R. designed research; J.O., M.D.J., K.A.C., J.R.S., and R.W.H. performed research; J.O., M.D.J., K.A.C., J.R.S., R.W.H., M.C.W., and M.R.R. analyzed data; and J.O., M.D.J., K.A.C., M.C.W., and M.R.R. wrote the paper.

The authors declare no conflict of interest.

This article is a PNAS Direct Submission.

Data deposition: The atomic coordinates and structure factors have been deposited in the Protein Data Bank, www.pdb.org (PDB ID code 3HX6).

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This article contains supporting information online at www.pnas.org/cgi/content/full/0911616107/DCSupplemental.

Table 1. Crystallographic data collection, phasing, and refinement

PilY1 C-terminal domain		
Data set	Native	SeMet
Unit cell, <i>abc</i> , Å; $\alpha\beta\gamma$, °	<i>a</i> = 64.4, <i>b</i> = 108.3, <i>c</i> = 159.0	<i>a</i> = 169.8, <i>b</i> = 64.9, <i>c</i> = 116.6, β = 108.7
Space group	P2 ₁ 2 ₁ 2 ₁	C2
Wavelength, Å	1.0718	0.9792
X-ray source	APS SER-CAT 22-BM	APS SBC-CAT 19-ID
Resolution, Å (highest shell)	50-2.10 (2.18-2.10)	50-2.80 (2.90-2.80)
No. of reflections, unique	65477	29869
Completeness, %	99.7 (98.8)	99.9 (100.0)
<i>R</i> _{sym} *, %	8.5 (43.5)	7.1 (24.4)
<i>I</i> / σ	39.7 (6.7)	48.6 (9.8)
Redundancy	13.9 (13.3)	7.8 (7.9)
SAD phasing statistics to 2.8 Å		
Se atom sites in AU (observed/total)		14/18
Mean figure of merit		
MLPHARE (highest shell)		0.65 (0.60)
After DM (highest shell)		0.82 (0.41)
Refinement summary		
Resolution, Å (highest shell)	50-2.10 (2.18-2.10)	
Molecules per AU	2	
No. of protein atoms/AU	7,361	
No. of waters/AU	818	
<i>R</i> _{working} [†] , %	19.6 (23.7)	
<i>R</i> _{free} [‡] , %	23.5 (28.1)	
Average B factor, Å ²		
Protein	29.5	
Water	42.3	
rms deviations		
Bonds, Å	0.005	
Angles, °	1.37	
Coordinate error estimates, Å		
Luzzati	0.23	
SigmaA	0.15	
Ramachandran, %		
Allowed, generous, disallowed	98.6, 1.0, 0.4	
Protein Data Bank ID code	3HX6	

AU, asymmetric unit; APS, Advanced Photon Source; SER, Southeast Regional; CAT, Collaborative Access Team; SBC, Structural Biology Center; BM, bending magnet; ID, insertion device; SAD, single-wavelength anomalous dispersion; MLPHARE, maximum likelihood heavy atom refinement and phase calculation; DM, density modification.

**R*_{sym} = $\sum |I - \langle I \rangle| / \sum I$, where *I* is the observed intensity and $\langle I \rangle$ is the average intensity of several symmetry-related observations.

[†]*R*_{working} = $\sum ||F_o| - |F_c|| / \sum |F_o|$, where *F*_o and *F*_c are the observed and calculated structure factors, respectively.

[‡]*R*_{free} = $\sum ||F_o| - |F_c|| / \sum |F_o|$ for 5% of the data not used at any stage of the structural refinement.

1KB0; overall rmsd of 3.6 Å over 290 equivalent Cα positions with 15% sequence identity; Fig. 1B] (16). This β-propeller enzyme was identified by Phyre to align with the PilY1 predicted structure (*E*-value 0.014, 95% certainty) (16). Blades V through VII of the PilY1 CTD, however, deviate from the standard β-propeller fold and align poorly with established β-propellers. Blade V contains a fifth strand (β20) that is shared with blade VI. Blade VI is composed of six strands, including β20, and maintains an insert containing a short helix and β-turn-β motif (α9, β26–27). Finally, blade VII is truncated and composed of three relatively short strands.

WD40 β-propeller proteins mediate protein–protein interactions via surface contacts at the propeller center (19). Superposition of the PilY1 CTD structure onto that of the seven-bladed β-propeller WDR5, complexed with a lysine 4-containing histone H3 peptide, revealed an rmsd of 3.9 Å over 261 equivalent Cα positions sharing only 12% sequence identity (20). Blades I–IV of the PilY1 CTD align well with the first four blades of WDR5, whereas the last three blades of each protein deviate more significantly in structure (Fig. S3). However, helices four and seven of PilY1 appear to block the surface site used by WDR5 to interact with peptide (19). Both helices are predicted to be conserved in the C-terminal domains of the related *Neisseria* PilC proteins (Fig. S2) (17). These observations suggest that PilY1 and related pilus-biogenesis factors do not mediate pro-

tein–protein interactions using the canonical WD40 surface-binding site.

PilY1 Contains a Unique Calcium-Binding Loop. A strong $|F_o| - |F_c|$ electron density peak in the PilY1 CTD structure was interpreted as a calcium ion based on the observed coordination by three aspartic acid side chains (851, 855, 859), one asparagine side chain (853), one main-chain carbonyl oxygen of valine 857, and one water (Fig. 2A and Fig. S1). The calcium binding observed in PilY1 is similar to that seen in the canonical EF-hand Ca-binding motifs (Fig. 2B and Fig. S4) (21). PilY1 and EF hands both employ seven contacts to the bound calcium ion, including one from a water molecule and one bidentate contact via an acidic side chain. PilY1 is unique, however, in that it achieves this calcium chelation using a stretch of only nine amino acids (851–859) in a loop between two β-strands (β13 and β14) rather than the twelve residues between two α-helices typically observed in the EF hands (21). In addition, the carboxylic acid group of the bidentate Asp-859 residue is rotated nearly 90° with respect to the equivalent residue in EF-hand proteins (21). Asp-859 forms a hydrogen bond with His-797, which is conserved in PilY1 proteins of known sequence. The calcium-chelating residues observed in the structure are conserved in the PilY1s of known sequence and in the related PilCs, with the exception that Asn-853 is replaced with an aspartic acid in the *Neisseria* proteins (Fig. 2C and Fig. S2). This

Fig. 1. Crystal structure of the *P. aeruginosa* PilY1 CTD. (A) 2.1 Å resolution crystal structure of the *P. aeruginosa* PilY1 C-terminal domain, which begins at residue 644 and ends at 1148 and encompasses seven β -blades (I–VII; colored red, orange, yellow, green, blue, magenta, and violet). (B) Superposition of the seven-bladed PilY1 CTD β -propeller (red through violet) on the eight-bladed β -propeller of the quinoximinoalcohol alcohol dehydrogenase from *Comamonas testosteroni* (1KB0; gray).

modified EF-hand-like motif is also present in PilY1 orthologues found in other bacteria that utilize T4P, including the human pathogen *Kingella kingae*. Thus, a unique calcium-binding site appears to be conserved in the C-terminal domains of pilus-biogenesis proteins related to PilY1.

Measurement of calcium-binding affinity using a fluorescence competition assay established that purified PilY1 CTD exhibits a K_d of 2.6 μM for calcium (Fig. 2D), which is similar to the affinities reported for traditional EF-hand proteins (22). Titration with increasing concentrations of magnesium chloride reveals that this cation does not bind to PilY1. Mutation of the calcium-chelating residue Asp-859 to alanine eliminates specific calcium binding by the protein (Fig. 2D). Removal of calcium either by Chelex 100 or by mutating a chelating residue (e.g.,

Fig. 2D and Fig. S5) did not change the overall structure (as measured by circular dichroism spectropolarimetry; Fig. S6) or melting temperature (Fig. S7) of the PiY1 CTD (23). Thus, calcium binding appears not to play a role in the overall structure of the PiY1 C-terminal domain.

Calcium Binding and Release by PilY1 are Essential for Functional Pili.

Expression of mutant versions of PilY1 in *P. aeruginosa* revealed that calcium binding and release are essential for pilus production and twitching motility. To examine the phenotypic consequences of altering the PilY1 calcium-binding site on T4P production and function, we engineered plasmid-encoded versions of full-length wild-type PilY1 and PilY1 carrying the D859A substitution. Both proteins were expressed to native levels in a *P. aeruginosa* strain in which the chromosomal *pilY1* gene was deleted (Fig. S8). The PilY1^{minus} strain is devoid of surface T4P, lacks measurable twitching motility, and is phenotypically indistinguishable from a nonpilated PilA^{minus} strain, which does not produce the major T4P subunit, pilin (Fig. 3A and B). Complementation of the PilY1^{minus} strain with plasmid-expressed PilY1 restored twitching motility and pilus production to wild-type levels (Fig. 3A and B). In contrast, complementation of the PilY1^{minus} strain with full-length PilY1 D859A resulted in significantly less twitching motility ($p < 0.001$) (Fig. 3A). The level of surface T4P produced by the PilY1 D859A expressing strain was also dramatically reduced compared to wild type and likely accounts for the observed defect in twitching motility (Fig. 3B).

PilY1 was previously shown to traffic to the surface T4P fraction (13) (Fig. S8), where it presumably functions through a direct association with the pilus fiber. To determine whether the defect in twitching observed with the D859A form of PilY1 was caused by disrupted pilus assembly/extension or increased pilus retraction, the PilY1 D859A mutant was expressed in a mutant strain lacking the PilT-retraction ATPase (PilT^{minus}, PilY1^{minus} strain). These bacteria regained the ability to produce normal levels of T4P, and the mutant PilY1 protein was capable of trafficking to the sheared pilus fraction (Fig. 3C–F). This result indicates that the loss of surface pili observed with the PilY1 D859A mutant is caused by PilT-mediated pilus retraction. In the presence of PilT, the D859A form of PilY1 is incapable of antagonizing PilT-mediated pilus retraction despite normal trafficking of the protein. We conclude that the calcium-binding site in the PilY1 CTD is necessary for normal pilus function by antagonizing the retractile activity of PilT.

We then created a mutation in full-length PilY1 designed to mimic calcium binding and to potentially render the protein calcium insensitive. Specifically, a lysine residue was substituted for the bidentate calcium-chelating Asp-859 residue (D859K). Although a lysine residue is not equivalent to a divalent metal ion, modeling of the D859K substitution mutation indicates that the ϵ -amino group of lysine would be expected to form a salt bridge with the two proximal aspartic acids at 851 and 855 (Fig. S9). Purified D859K PilY1 CTD does not bind calcium and is not distinct in structure or melting temperature from the wild-type CTD (Figs. S5–S7). When expressed to native levels in the *P. aeruginosa* PilY1^{minus} strain (Fig. S8), the full-length PilY1 D859K mutant was significantly reduced for bacterial twitching motility ($p < 0.001$) compared to the same strain expressing wild-type PilY1 (Fig. 3A). Interestingly, the level of surface pilus production in the PilY1 D859K expressing strain was equivalent to that of the wild-type strain (Fig. 3B), indicating that reduced fiber production does not account for the twitching motility defect. PilY1 D859K was capable of trafficking to the surface T4P fraction when expressed in both a *pilY1* mutant background (Fig. S8) and a PilT^{minus}, PilY1^{minus} strain (Fig. 3C–F). Thus, using the crystal structure as a guide, we produced a single-site mutation that dramatically alters pilus function without affecting pilus production. Based on these data, we conclude that

Twitching motility mediated by PilY1's influence on the PilB/PilT oscillatory motor would be expected to be coordinated with the significant >100 pN retractile motion exerted by the system once target cell adhesion occurs (25, 26). Surface translocation by twitching motility likely involves relatively moderate affinity contacts between a surface and T4P (26). For example, a disulfide loop in pilin, the major pilin subunit of the pilus fiber, has been proposed to be a point of contact between surfaces and T4P (27). We propose that the cycles of pilus extension and retraction actuated by PilY1's C-terminal domain may control twitching motility involving these moderate affinity contacts. Once target host cell contact has been achieved, however, the T4P system would be expected to convert to a rapidly retracting, PilT-dominated state in which nearly nanonewton forces are exerted (28, 29). The function of this mode would be to quickly draw the bacterial cell into close contact with host tissue cells. It is tempting to speculate that

Fig. 3. PilY1 calcium-binding site is essential for T4P production and function. (A) Twitching motility of *P. aeruginosa* (WT) or a *pilY1* mutant expressing either empty vector (–), PilY1, or mutant versions of PilY1 (D859A or D859K). The twitching motility zones produced by bacteria expressing mutant PilY1 were significantly (*, $p < 0.001$) reduced compared to those of bacteria expressing wild-type PilY1. (B) Coomassie blue-stained gel of pilin isolated from the bacterial surface. (C–F) Abolishing T4P retraction allows conditional localization of PilY1 to the bacterial surface. (C) Coomassie-stained gel of pilin isolated from the bacterial surface. (D) Western blot of T4P-containing surface fractions probed with PilY1-specific antiserum. (E) Western blot of whole cell lysates probed with a pilin-specific antibody. (F) Western blot of whole cell lysates probed with a PilY1-specific antiserum.

PilY1 may serve as the switch between twitching and rapid retraction. The sequence divergent N-terminal domains of PilY1 orthologues are proposed to perform target cell-specific adhesion; once that is achieved, perhaps the conserved C-terminal

Fig. 4. Calcium chelation impacts T4P production. (A) Pilus preparations from a *pilT* mutant or a *pilY1* mutant expressing either PilY1, or mutant versions of PilY1 grown in the presence (+) or absence (–) of the calcium chelator EGTA. (B) Summary of the impact specific mutants have on pilus state. Wild-type PilY1 appears to balance extension and retraction to produce twitching motility (boxed), whereas gene deletions, PilY1 site mutants, and the calcium chelator EGTA all significantly disrupt the equilibrium between pilus extension and retraction.

domains switch to the calcium-free state to facilitate rapid PilT-mediated retraction (14, 30, 31).

The C-terminal domain of *P. aeruginosa* PilY1 shares significant sequence identity with the equivalent regions of the *Neisseria* PilC's (Fig. 2C) and similar levels of identity in PilY1 homologues from other human, plant, and marine infectious bacteria (17). We find that each PilY1 homologue contains a predicted calcium-binding site highly similar in sequence to that observed in the PilY1 C-terminal domain (Fig. S2). Thus, calcium binding and release may play an equally important role in pilus biogenesis and motility in organisms that maintain a PilY1 homologue. Taken together, the structural and functional data we outline provide unique insights into the complex functional regulation of the T4P essential to a range of human pathogenic bacteria.

Methods

Structure Determination. PilY1 residues 615–1163 were cloned from genomic DNA from the PAK strain of *P. aeruginosa* (GenBank #EU234515), and over-expressed recombinantly in *E. coli*. The protein was purified using Ni-affinity and size-exclusion chromatography steps to >95% purity, and crystallized in 0.3 M sodium malonate (pH 6.5), 20% (wt/vol) PEG 3350, 50 mM DTT, 30 mM ammonium phosphate, and 4–8% trifluoroethanol. Residues 6145–1163 of PilY1 contained only five methionine residues; as such, crystals containing selenomethionine-substituted protein failed to produce sufficient signal to allow structure determination. L712M, L812M, and L823M mutations were successfully generated, combined, expressed, purified, and crystallized in PilY1 specimen sufficient for x-ray data collected. Diffraction data to 2.8 Å resolution were collected at the peak wavelength (0.979 Å) for selenium SAD phasing (Table 1) and indexed and scaled in space group C2 using HKL3000 (32, 33). Automated model building correctly built 30% of the final structure; complete building and refinement was accomplished using the programs COOT and CNS, respectively (34, 35). Once the selenomethionine-substituted structure was refined at 2.8 Å resolution, it was used in molecular replacement to solve the 2.1 Å native structure, which was refined using COOT and CNS to the final statistics shown in Table 1 (34, 35).

Calcium Binding and Biophysical Studies. Calcium was removed from purified PilY1 using Chelex 100 (BioRad) prior to biophysical studies (23). A binding curve for Oregon Green® 488 BAPTA-5N (Invitrogen) was measured on a PHERAstar (BMGLabtech) at 488 nm and the K_d established at 10.1 μM. WT, D859A, and D859K PilY1 CTD was serially diluted 2-fold to calculate PilY1 CTD K_d for calcium binding, if applicable. For biophysical characterizations via circular dichroism spectropolarimetry, a wavelength scan from 195–260 was performed at 22 °C. Thermal denaturation protein samples were measured 210 nm at a range of 10–85 °C.

Isolation of Surface T4P. *P. aeruginosa* strains were spread on LB agar plates and grown until confluent. Bacteria were collected and suspended in 1 ml 0.15 M NaCl/0.2% formaldehyde and vortexed vigorously for 1 min to release surface T4P. Bacterial cells were removed by centrifugation at 12,000 × g for 5 min. T4P fractions were separated by SDS-PAGE (18% polyacrylamide) and pilin visualized by GelCode blue stain (Pierce).

Twitching Motility. Bacteria were stab inoculated to the bottom of 100 mm tissue culture-treated plates containing 5 mL LB agar (1% wt/vol). Plates were incubated for 40 h at 37 °C in a humidified incubator. Twitching motility was quantified by measuring the diameter of the subsurface zone of bacterial spread.

Western Blots. Whole cell lysates and T4P fractions were separated on 18% (pilin) or 7.5% (PilY1) SDS-polyacrylamide gels and transferred to nitrocellulose or PVDF membranes, respectively. Membranes were probed with PKL1 antipilin monoclonal antibody or anti-PilY1 CTD rabbit serum. Horseradish peroxidase-conjugated secondary antibodies were used for the detection of specific antibody-antigen complexes. Blots were developed with chemiluminescence reagents and visualized via autoradiography.

ACKNOWLEDGMENTS. We thank Wladik Minor and Zbyszek Otwinowski for HKL3000 guidance, Joseph Lomino and Lisa Charlton of the Redinbo Laboratory for help with CD experiments, and Rebekah Nash and members of the Redinbo and Wolfgang groups for assistance with manuscript preparation. Supported by National Institutes of Health Grants AI078924 (to M.R.R.) and AI069116 (to M.C.V.), and a Howard Hughes Medical Institute Med-into-Grad Initiative Training Grant (K.A.C.).

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