Multidisciplinary Interrogation of a Crucial Protein Interface in the Type II Secretion System

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28 **ABSTRACT** The type IV filament superfamily comprises widespread membrane-associated
 29 polymers in prokaryotes. The Type II secretion system (T2SS), a significant virulence pathway in
 30 many pathogens, belongs to this superfamily. A knowledge gap in the understanding of the T2SS
 31 is the molecular role of a small ‘pseudopilin’ protein. Using multiple biophysical techniques, we
 32 have deciphered how this missing component of the Xcp T2SS architecture is structurally
 33 integrated, and thereby also unlocked its function. We demonstrate that the low abundance XcpH
 34 is the adapter that bridges a trimeric initiating tip complex XcpIJK with a periplasmic filament of
 35 XcpG subunits. Our model reveals that each pseudopilin protein caps an XcpG protofilament in
 36 an overall pseudopilus compatible with the dimensions of the periplasm and the outer membrane-
 37 spanning secretin through which substrates of the T2SS pass. Unexpectedly, to fulfill its adapter
 38 function, the XcpH N-terminal helix must be unwound, a property shared with the XcpG subunits.
 39 We provide the first complete structural model of a type IV filament, a result immediately
 40 transferable to understanding of other T2SS and the type IV pili.

INTRODUCTION

Bacteria have sophisticated secretory nanomachines that evolved to deliver exoproteins to the bacterial cell surface, into the surrounding medium or directly into host cells. Among these, the type 2 secretion system (T2SS) is a trans-envelope apparatus specialized for secretion of folded proteins from Gram negative bacteria¹. In many cases, these secreted substrates are virulence factors: examples include the cholera toxin of *Vibrio cholera*, the exotoxin A of *Pseudomonas aeruginosa*, and the heat labile toxin of enterotoxigenic *Escherichia coli*². The conformational constraint of transporting folded substrates is shared by the Type VII, or ESX, system³ and the Type IX secretion system⁴, but is in clear contrast to the linear export of unfolded polypeptides by the T3SS⁵ or the T4SS⁶. The substrates of the T2SS are not known to share any sequence or 3D structural motifs⁷, and thus an open question is how substrates are individually recognized and ushered out. Dynamic and changing interactions, potentially involving nucleation of structure in intrinsically disordered regions, must be at the heart of this selection process⁸.

The T2SS accomplishes its task with a distinctive mode of transport involving a pilus-like structure (the pseudopilus) in the periplasmic space, at the interface between an inner membrane assembly platform and a large outer membrane pore, the secretin⁹. Structural, functional and evolutionary commonalities to the extracellular Type IV pili critically inform the T2SS field and *vice versa*¹⁰. Thanks to recent spectacular developments in cryo-electron microscopy, the 3D structures of T2SS secretins^{11–14} and the filament formed upon overexpression of the pseudopilin subunit PulG¹⁵ have been solved, and reveal important structural features. While secretins form a tightly gated giant double β -barreled pore of 80 Å diameter in the outer membrane, PulG adopts a right-handed homopolymeric helix with a diameter of 70 Å, compatible with observations that a filament extends through the secretin when the pseudopilin is overexpressed^{16,17}. In this pseudopilus, inter-subunit contacts occur along hydrophobic N-terminal α helices of the pseudopilin, with a special role for the negatively charged glutamic acid at position 5 in both recruiting and stabilizing interactions within the filament^{18–21}. It has long been assumed that under native expression levels the pseudopilin forms a short and transient thread entirely within the periplasm during secretion²², a model which has been difficult to verify experimentally until recently, when a cryo-electron

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tomography reconstruction of the *Legionella pneumophila* T2SS revealed pseudopilus density in ~20% of the complexes²³.

In addition to the major pseudopilin, additional low abundance (or *minor*) pseudopilins have been identified in the T2SS, based on being encoded within the same operon and sharing amino acid sequence with the pseudopilin, plus undergoing maturation by the same prepilin peptidase^{24–26}. The four minor pseudopilins are generally designated as the H, I, J, and K proteins (variously prefixed with Gsp-, Pul-, Xcp-, depending on the particular T2SS, please see on line Materials and Methods for nomenclature notes). While none of these has been directly observed in a T2SS pseudopilus, the periplasmic domains of minor pseudopilins GspI, GspJ, and GspK have the ability to assemble into a ternary complex whose crystal structure has been solved. The four periplasmic domains of GspH, GspI, GspJ and GspK, form a quaternary complex^{27,28}, but its structure is unknown. As a group, the minor pseudopilins are known to play a role in initiation and control of pseudopilus assembly and potentially retraction. The complex furthermore interacts with secretion substrates³² and with the periplasmic domains of the secretin³³. Although there is strong evidence that the quaternary complex is found, at least transiently, at the tip of the pseudopilus, its structure has been elusive. Minor pseudopilin subunits could not be resolved in the *L. pneumophila* T2SS cryo electron tomograms²³. In a recent T4P cryo electron tomography study, minor pilins were attributed to density blobs but no conclusions could be drawn about which protein belonged to which density nor about the molecular nature of their interactions³⁴.

While many reports have established the importance of the minor pseudopilins and pilins as a group for initiation and/or assembly of (pseudo)pili, fewer have analyzed their individual contributions to the secretion process. Nonetheless, specific roles have been assigned or proposed for all four minor pseudopilins: GspJ primes filament assembly through the recruitment of other minor pseudopilins³⁷, GspI serves as a protein-protein interaction hub²⁷, and GspK controls pseudopilus length and/or number^{27,37}. GspH interacts with GspJ through its globular domain and with the structurally similar major pseudopilin via its N-terminal hydrophobic α -helices^{18,27,39,40}, thus it is tempting to suggest GspH plays an adapter function between the tip and the body of the pseudopilus. However, the essentiality of GspH for secretion has been called into question because

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its absence can be overcome by overproduction of other minor pseudopilins³⁷. There are no molecular data on how GspH is embedded in the quaternary complex.

Based on x-ray crystallographic structures of constructs representing periplasmic domains of GspH, GspI, GspJ, and GspK from several microbial species solved alone and/or in dimeric or trimeric subcomplexes^{28,39–43}, each pseudopilin fold resembles major T2SS pseudopilins and Type IV pilus (T4P) major pilins characterized by an extended N-terminal α -helix, the first half of which (α 1N) is not included in the soluble crystallography targets but is expected to be exposed to the membrane environment, and the second half of which (α 1C) is partially buried in a globular α/β domain. Despite this common overall topology, it is important to bear in mind that the periplasmic domains of the minor pseudopilins interact to form the soluble quaternary complex in the absence of α 1N, and since these minor subunits are at the tip of the pseudopilus they cannot depend on vertical interactions with major subunits for upward assembly interactions. Within this general framework, each minor pseudopilin has distinctive sequence and structural hallmarks. GspH is most similar in size, sequence and structure to the major pseudopilin. GspI is the smallest of the four. GspJ is larger, and displays a pronounced groove on its surface. Finally, GspK carries an additional structural domain of over 100 amino acids. GspH, GspI, and GspJ share the glutamate at position 5 with GspG, whereas GspK does not.

Based on two crystal structures of ternary complexes of minor pseudopilin periplasmic domains (3CI0 from the enterotoxigenic *E. coli* Gsp T2SS and 5VTM from the *P. aeruginosa* Xcp T2SS equivalent minor pilins XcpIJK^{28,40}), we know the stabilizing interactions among the periplasmic domains (designated here with a subscript “p”) GspI_p, GspJ_p, and GspK_p include buried salt bridges between α 1C residues for all three pairwise interactions: XcpI_p Asp51 to XcpK_p Arg45, XcpJ_p Glu58 to XcpK_p Arg45, and XcpI_p Asp51 to XcpJ_p Glu58. The interaction is also stabilized by salt bridges involving amino acids of XcpI_p α 1 and XcpJ_p β 11, including XcpI_p Asp51 and XcpK_p Arg195. Contacts at the bottom of the bundle of helices further lock the ternary complex in place. Double involvement of XcpI_{D51} in XcpI_p:J_p & XcpI_p:XcpK_p interfaces is evidence to bolster the original finding that XcpI_p supports the linear organization of XcpJ_p:I_p:K_p interactions in the complex²⁷. Somewhat perplexingly, the largest buried surface area between any two of the soluble proteins in both ternary complexes is between XcpJ_p (GspJ_p) and XcpK_p (GspK_p), with a total

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buried surface area of $\sim 3600 \text{ \AA}^2$ in 5VTM. And yet, these two subunits do not interact with each other in solution in the absence of XcpI_p²⁷.

Several lines of evidence support downward addition of XcpH to the base of the XcpJ:I:K complex through an interaction between the globular domains of XcpH and XcpJ. XcpH_p completes the quaternary complex by interacting with XcpJ_p²⁷. Additionally, $\alpha 1$ of XcpH is essential in this interaction and XcpH_p can be positioned most convincingly in a small angle x-ray scattering envelope of the quaternary complex below XcpJ_p²⁸. The structural compatibility of the ternary tip complex GspI_pJ_pK_p with downward addition of GspH below GspJ was proposed by Korotkov *et al.* in agreement with their previous suggestion that GspH could be positioned at the top of a GspG fiber but not at the bottom^{39,40}. The lack of Glu5 in GspK can also be taken as evidence that GspK does not require helix:helix packing stabilization via a salt bridge with a higher subunit in the filament and thus is located at the top of the pseudopilus³¹.

In this study we set out to validate the biological importance of XcpH and to decipher the molecular details of the XcpH:I:J:K interaction. Because all available data to date suggest an unstable, dynamic and/or small interaction interface with XcpH, we turned to a collection of compatible techniques suited to interrogation of such interactions. We show that XcpH is indeed critical for efficient Type II secretion in *P. aeruginosa*. We demonstrate it interacts specifically with the $\alpha 1$ helix of XcpJ, and we narrow the interaction residues to a well-defined interface that is highly conserved among T2SSs. Our experimentally determined constraints allow us to synthesize a molecular model of the XcpHIJK quaternary structure, and to place this in the context of the complete pseudopilus filament, with biological implications for the role of the GspH/XcpH low abundance pseudopilin. Moreover, the molecular organization involving an inner membrane platform, priming by minor pilins, and a helical subunit assembly that is remarkably well conserved both evolutionarily and structurally in other type IV filamentous nanomachines^{10,44} including the canonical type IV pilus (T4P)^{19,31,38}, competence pilus^{45,46}, and to some extent the archaellum⁴⁷⁻⁴⁹. Thus our work has wide-ranging implications for understanding assembly of several fundamental microbial organelles.

RESULTS

The four minor pilins are essential for secretion.

In our quest to understand the structure-function relationship of the minor pseudopilin complex, we endeavored to systematically assess the independent requirement of each of the four minor pseudopilins of the *P. aeruginosa* Xcp T2SS during the secretion process. Individual in-frame deletion mutants were constructed in the PA01 reference strain for *xcpH*, *xcpI*, *xcpJ* and *xcpK* (see online Materials and Methods for nomenclature). Each mutant was assessed for its ability to secrete the major Xcp T2SS effector LasB (**Figure 1a**) and for the ability of secreted LasB to degrade casein on milk plates (**Figure 1b**). All four individual deletion strains were similar in their inability to secrete LasB and lacked protease activity, like the negative control strain in which the entire T2SS operon is missing, thus demonstrating the absolute requirement of each minor pseudopilin in the Xcp T2SS secretion process. This finding is supported by complementation of each secretion deficient phenotype by introduction of the corresponding wild type gene on a plasmid. Our data are apparently in conflict with a 2018 publication suggesting that XcpH and XcpK are not required for T2S in *P. aeruginosa*²⁸. Our finding that XcpH is indispensable for secretion prompted us to further explore the 3D integration of XcpH into the XcpHIJK quaternary complex.

Chemical shift perturbation studies of labeled XcpH_p in the presence of XcpJ_p.

XcpH_p solution NMR assignments

To discover sites of interaction between XcpH and XcpJ, we performed solution NMR experiments. We initially isotopically labeled the ~16 kDa XcpH_p, a feasible NMR target, with ¹⁵N and ¹³C, and carried out amino acid assignments. Regions heavily crowded in the HSQC spectrum were excluded, leading to 85% coverage of the XcpH_p amino acid sequence (**Figure 2a**). The high signal dispersion of the HSQC spectrum indicates an overall well-folded protein, while the area in the spectrum with high resonance overlap and strong signal (around 8.2 ppm in the ¹H dimension and 121 ppm in the ¹⁵N dimension) indicates a highly dynamic region of the protein.

We employed these assignments to experimentally determine the secondary structure elements within XcpH_p (**Figure 2b**). Furthermore, we generated two 3D homology models of XcpH_p using

PDB 2KNQ and 2QV8³⁹. Each 3D model matched our experimental secondary structure assignments well (not shown).

A subset of XcpH_p amino acids, in particular lysines, could not be assigned in our NMR spectra. The inability to assign these lysines implies a flexible region of the protein backbone. We tested the role of these residues in secretion by creating an in-frame deletion of amino acids 108-117, which form a long excursion between β3 and β4, resembling an old-fashioned mouse trap spring in one homology model (**Supplemental Figure 1**). We found that trans complementation with the *xcpH*_{Δ34} deletion restores LasB secretion and activity to the Δ*xcpH* strain (**Figure 1b, c**). Thus residues 108-117 are not required for LasB secretion, and therefore an disorder to order transition of this region cannot be required for secretion of LasB either.

Specific sites of spectral changes along the XcpH_p helix suggest binding interface: Chemical Shift Perturbation

With XcpH_p chemical shifts in hand, we performed chemical shift perturbation (CSP) analysis to identify structural regions impacted in XcpH_p upon binding of its partner XcpJ_p. It is expected that resonances from residues close to the protein-protein interface will be shifted by the presence of the binding partner. Thus, a sample of 100 μM ¹⁵N uniformly labeled XcpH_p was titrated with increasing amounts of XcpJ_p and the 2D ¹H-¹⁵N HSQC spectrum was collected. XcpJ_p concentrations above 80 μM could not be used since at that concentration there was a significant signal intensity loss due to the large size of the complex. Significant CSPs were observed in multiple places in the amino acid sequence (**Figure 3a**). By mapping these CSPs on a structural model of XcpH_p, we observed one subset of amino acid residues located close to the tip of the alpha helical spine of XcpH_p (residues G42 to G59 in the helix, and G130 and G131 on the nearby β4-β5 loop) that are perturbed by the presence of XcpJ_p (**Figure 3b**). In addition, there is a second set of residues with significant CSP located on the C-terminal β sheet of XcpH_p (residues F135 to R138 in β5, L149 and S153 in β6 and E162 and A164 in β7). These additional perturbed sites may indicate a secondary point of interaction with XcpJ_p, or changes in XcpH_p propagated through its structure after binding to XcpJ_p. Global smaller amplitude changes across much of XcpH_p suggest that XcpH_p:J_p interaction subtly affects dynamics or the structure of the entire XcpH_p molecule.

Site specific cysteine incorporation in XcpJ_p provides XcpH_p:J_p interaction sites.

Cysteine cross-linking

The knowledge that a significant structural interaction interface for the major pseudopilins is the packing of their $\alpha 1$ helices¹⁵, combined with our CSP results implicating the XcpH $\alpha 1C$ in the XcpH_p:J_p interaction, led us to hypothesize that $\alpha 1C$: $\alpha 1C$ packing forms a major interaction between these two minor pseudopilins. We introduced cysteines at surface-exposed $\alpha 1C$ positions in XcpH_p (V46, D49, L53) and XcpJ_p (R46, R53) and mixed purified proteins under oxidizing conditions. In every instance, homodimers are formed (**Figure 4**). Heterodimers between XcpH_p and XcpJ_p were also observed in some prominent cases, most notably for XcpJ_{R46C}:H_{L53C} (**Figure 4**). Heterodimers were weak or absent when XcpH_{pV46C} was one of the partners. Although these results provide no quantitative measure of affinity or specificity, they served as motivation to use XcpJ_p $\alpha 1C$ cysteine variants as probes of specific sites of interaction in our NMR platform.

Paramagnetic relaxation enhancement

To identify local interactions and specific amino acid involvements in the XcpH:XcpJ interface, we performed paramagnetic relaxation enhancement (PRE) experiments guided by cysteine cross-linking results. In this approach, the effect of a specifically incorporated spin-label (1-Oxyl-2,2,5,5-tetramethylpyrroline-3-methyl) methanethiosulfonate (MTSL) within XcpJ_p on the ¹⁵N XcpH_p spectrum is observed; in particular, XcpH_p residues in proximity to the spin label on XcpJ_p will experience a decrease in signal intensity. XcpJ_p cysteine variants R46C and R53C were spin-labeled, as these positions in XcpJ_p were implicated in cysteine cross-linking experiments described above to participate in an XcpH $\alpha 1C$ - XcpJ $\alpha 1C$ packing interaction (**Figure 4**). In addition, XcpH_{pT178C} and XcpH_{pE180C} were created and used for spin-labeling; these amino acids are on the face of XcpJ opposite $\alpha 1C$, where we did not expect any change in signal from spin label incorporation. We chose to perform PRE experiments using a 10:3 molar ratio of ¹⁵N-U XcpH_p to spin-labeled XcpJ_p variants. Although a 1:1 ratio would in theory have provided more quantitative results, for this protein pair, it would have led to signal loss as seen in the previously described titration.

The spin label MTSL at positions 46 or 53 within XcpJ_p has a marked effect on XcpH_p signal intensities (**Figure 5a**). XcpJ_{pR46C}-MTSL induced a signal reduction on XcpH_p $\alpha 1C$ (residues 42

to 54), with a maximum effect on D49. In addition, residues located in the loop between XcpH_p β4 and β5 are also affected (G130 and E131) (**Figure 5**), in agreement with CSP results. XcpJ_pR53C-MTSL had an effect on similar amino acid residues. Within XcpH_p α1C the overall signal intensity decrease was slightly less compared to the C46 label, but interestingly the effect was shifted up the helix with the residue most greatly affected by the paramagnetic label, XcpH_p-N55, located at the tip of XcpH_p (**Figure 5**). No significant effect was observed on XcpH_p when XcpJ_p was labeled at positions T178C or E180C (**Supplemental Figure 2**). Thus, the PRE experiment clearly supports that the interface between XcpH_p and XcpJ_p is mediated by α1C interactions with involvement of residues in the β4-β5 loop of XcpH. In addition, the difference in signal intensity decrease and the shift on the most affected residue towards the tip of XcpH_p when the MTSL label is at XcpJ position 53 as opposed to 46 suggest that the helical register is organized with XcpJ_p above XcpH_p in the heterodimer.

Cross-linking Mass Spectrometry using acidic cross links *via* the DMTMM activating reagent

In order to extend our site-specific information from engineered disulfide bonds and PRE measurements, we applied a recently developed methodology for crosslinking of acidic side chains within a defined distance, followed by mass spectrometry identification of cross-linked peptides⁵⁰. We established our pipeline using a 1:1 molar ratio of XcpH_p and XcpJ_p (**Supplemental Methods** and **Supplemental Figure 3**). Previous work has established that XcpJ_p is indispensable for XcpH_p involvement in the quaternary complex, which was the motivation for using this pair in our NMR experiments also^{27,28}. Subsequently, we interrogated the complete quaternary complex of XcpH_pI_pJ_pK_p, known already to form *in vitro* a quaternary complex with a molar ratio of 1:1:1:1²⁸. These latter experiments confirmed the set of acidic crosslinks between XcpH_p and XcpJ_p are the same in the heterodimer as in the heterotetramer, and thus we focus here on the quaternary complex results.

The four minor pseudopilin soluble constructs XcpH_p, XcpI_p, XcpJ_p and XcpK_p contain 25, 14, 36 and 42 acidic residues, respectively, which are distributed along their amino acid primary sequences. Thus the use of adipic acid dihydrazide (ADH) coupled to 4-(4,6-dimethoxy-1,3,5-triazin-2-yl)-4-methyl-morpholinium chloride (DMTMM) as a coupling reagent appeared

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promising. DMTMM activates carboxylic acid functionalities, usually poorly reactive, allowing coupling of the crosslinking reagent ADH to the extended amino acid side chains with high efficiency⁵⁰. When added in large excess to a freshly prepared aqueous solution containing a 1:1:1:1 molar ratio of XcpH_p, XcpI_p, XcpJ_p and XcpK_p, DMTMM/ADH reagents formed covalent bonds (**Figure 6a**), which connect proximal acidic functionalities with a maximum linking distance of 20-25 Å⁵¹. The most likely tetramer band observed on SDS-PAGE of this mixture was excised and analyzed by a classical proteomic bottom-up approach to validate the presence of all four proteins. Each protein was identified unambiguously, with a sequence coverage of 92% for XcpH_p (61 peptides), 78% for XcpI_p (31 peptides), 86% for XcpJ_p (71 peptides) and 91% for XcpK_p (146 peptides).

Due to the long length of the linkers, these cross links provide complex and valuable information to constrain a model of the quaternary structure of the assembled XcpH_pI_pJ_pK_p complex. Most revealing for our goals of understanding the placement of XcpH within the pseudopilus quaternary complex were crosslinks between glutamate or aspartate side chains of XcpH_p with any of the other three soluble pseudopilin subunits in the complex. Many inter-subunit crosslinks between XcpH_p and XcpJ_p were identified between α1C of XcpH_p, in particular within the peptide LAGLIGVLTDEAVLDNR, and the several peptides that together span the analogous α1C of XcpJ_p and the β hairpin that follows it (**Figures 6b, 6c**). An example MS/MS spectrum reveals intense peaks defining the connection between the two peptides (**Figure 6b**), with both chains unambiguously characterized by many fragment ions distributed along the peptide sequences. Moreover, the large mass difference observed between y₆α⁺ and y₇α⁺ characterizes precisely the position of crosslinking. The same α1C region of XcpH_p also cross links with two regions of XcpK_p and one region of XcpI_p therefore suggesting they are in physical proximity (**Figure 6c, Table S3**). Interestingly, an acidic residue located in the XcpH_p β₄β₅ loop shown by PRE analysis to be in close proximity with the α1C region of XcpJ_p (**Figure 5b**) is also involved in cross links with XcpJ_p, XcpI_p and XcpK_p (**Figure 6c, Table S3**).

Crosslinks between XcpI_p and XcpJ_p as well as a single crosslink identified between XcpI_p and XcpK_p were all in agreement with the crystallographically determined structures of GspI_pJ_pK_p and XcpI_pJ_pK_p. Moreover, intrasubunit crosslinks agreed with known protein 3D structures (data not

shown). This rich library of crosslinks between amino acid side chains within the quaternary complex provides strong constraints for modeling of a three-dimensional structure of the XcpH_pI_pJ_pK_p complex.

Integrating experimental data to build quaternary complex model

XcpH_pI_pJ_pK_p complex

Data from spin labelling studies and acidic cross linking mass spectrometry were used as restraints on the interaction of XcpH_p with XcpI_pJ_pK_p, to model the quaternary complex using HADDOCK software for protein-protein docking. Docking used the structure of the XcpI_pJ_pK_p ternary complex (PDB: 5VTM), the XcpH_p homology model, PRE experimental results and acidic cross-linking interactions as input (**Supplemental Tables 3 & 4**). The best model obtained is in good agreement with these experimental restraints (**Figure 7**). In this model, the α1C of XcpH_p is in direct contact with that of XcpJ_p, and runs approximately parallel to the helical bundle in the XcpI_pJ_pK_p complex (**Figure 7**). XcpH_p interaction with the ternary complex is maintained by a large network of electrostatic interactions and hydrogen bonds (**Figure 7**). One group of contacts corresponds to an electrostatic interaction between helices, in particular including a salt bridge between side chain XcpH_p D49 and XcpJ_p R42 (possibly also to XcpJ_p R46, given the rotamers available to Arg side chains and the limitations of resolution of this model) as well as numerous main chain and side chain H-bonds. A second group comprises β-strand to β-strand contacts. In this interaction, XcpH_p loop β6-β7 G155 and F156 backbone carboxyl groups form hydrogen bonds with XcpJ_p β7 R131 side chain and Q133 main chain, respectively. Other contacts observed between XcpH_p and XcpJ_p correspond to: 1) XcpH_p loop α1-β1 and XcpJ_p loop β4-β5 (N55-R92), 2) XcpH_p loop β4-β5 and XcpJ_p loop β4-β5 (S129-R92) (**Figure 7**). Remarkably, although no data from the chemical shift perturbation experiments were explicitly included in the HADDOCK restraints, many of the interactions in the model are recapitulated as chemical shift perturbations, for example the very high signal interaction within XcpH_p β7 is explained by its close approach to XcpJ_p.

The use of acidic cross-linking restraints for XcpH_p with the two other subunits from the quaternary complex disambiguates the XcpH_p position *vis a vis* these proteins, for example the crosslinks to XcpK_p help to position the tip of XcpH_p α1C close to the XcpK_p β-sheet, allowing additional contacts that stabilize the complex. These include XcpH_p-XcpK_p residues E50-R283,

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D54-R283, R56-D282, R56-R284 and N55-G309 (**Figure 7**). We were able to further support this quaternary structure by analyzing the co-evolution of residues in XcpH_p and XcpJ_p. (**Supplemental Figure 4**). The analysis similarly points to XcpH polar amino acids at the tip of α 1C (D49, E57) and in the β 4- β 5 loop (S128, S129, E131) and pairs them with two sets of XcpJ co-evolved amino acids located within a large flap at the tip of XcpJ (R92 and R99, forming electrostatic interactions), or in the vicinity of α 1C (V45, R189 and W107, potentially contributing to packing geometry) (**Supplemental Figure 4**). This co-evolution result adds credence to the robustness and physiological relevance of our XcpH:XcpJ interaction model. Altogether, we present a model of the XcpH_pI_pJ_pK_p quaternary complex supported by experimental evidence, which shows the interaction of XcpH with XcpJ and XcpK is maintained by several electrostatic interactions.

Filament model

The quaternary complex presented here reveals the interaction of the minor pseudopilin soluble domains, and in particular places XcpH_p in this ensemble. However, these proteins also interact via their missing hydrophobic α helices (α 1N) in the biological context of the pseudopilus. Thus, we used available experimental data and chemically reasonable restraints to model the complete Type II secretion system pseudopilus. For that purpose, hydrophobic α 1N were added to the XcpH_pI_pJ_pK_p proteins in the soluble quaternary complex model. In addition, the major pseudopilin XcpG filament was modeled using as template the structure of PulG filament (PDB: 5WDA)¹⁵. Then, the quaternary complex was added to the tip of the XcpG filament by aligning XcpH to the first XcpG unit. Finally, the whole model was minimized using PyRosetta⁵². The best model was selected based on the lowest RMSD with respect to the HADDOCK model (**Figure 8**). An important feature included as a restraint in calculating this model was the salt bridge between N-terminal amino groups and the carboxylic acid of E5 of each preceding unit in the filament, including XcpHIJK at the tip of the pseudopilus. This contact neutralizes these charges in the transmembrane helices as seen in other filament structures such as *P. aeruginosa* PAK pilus (PDB 5VXY). As expected, E5-F1 contacts continue into the minor pseudopilin complex in the following sequence: XcpG-XcpH-XcpJ-XcpI-XcpK. This arrangement maintains the right-handed nature of the major pseudopilin filament structure. In the filament model, the average distance between E5 carboxyl group and F1 amino group is 3.58 ± 0.34 Å between adjacent subunits. In addition,

Cisneros *et al.*³⁷ showed the close contact between the residues 16 and 10 of neighboring subunits (PulJ-PulI and PulI-PulK). Even though these contacts were not used as restraints during modelling, the current model reproduces them between the minor and major pseudopilins throughout the filament (**Figure 8** and data not shown). Thus, the C α -C α distance from residue 10 to 16 of the neighbouring subunits in the PulG filament is 8.5 Å, in the model that distance is on average 8.9 ± 1.4 Å. Strikingly, to maintain the restraint contacts, in particular the E5 to F1 salt bridge between XcpH and XcpJ, it was necessary to allow XcpH α 1N between residues 20 and 26 to unravel. Although there is no other experimental evidence that the XcpH helix adopts this melted secondary structure, the equivalent amino acids in the major T2SS pseudopilin and the major T4P pilin are strikingly extended in high resolution filament models^{15,53}. Additionally, the presence of a glycine at XcpH position 25 supports this conclusion, since glycine and proline residues destabilize α helices.

Another feature observed in the filament model, specifically related to the minor pseudopilin tip, is the capping of each XcpG protofilament by a minor pseudopilin and the off-center placement of XcpK. If the major pseudopilin filament is separated into 4 protofilaments, each one is capped by a minor pseudopilin (**Figure 8**). The exception to this is XcpJ; although the XcpJ globular domain is directly above one protofilament, a small gap exists between XcpJ and the XcpG unit directly below it. An additional consequence of capping one XcpG protofilament with XcpH is that the XcpK globular domain protrudes beyond the major pseudopilin filament diameter, giving the appearance of a hook-like structure to the filament tip (**Figure 8**).

While our model cannot be considered an atomic structure of the Type II secretion pseudopilus, it provides a valuable path to testable predictions such as the importance of XcpH helix unravelling, possible substrate binding sites available on the filament tip, the presence of dynamic contacts between XcpG and minor pseudopilins, and the importance of the XcpK globular domain for secretion.

DISCUSSION

Study of dynamic interactions prioritized by binding affinity is challenging but we have used chemical, structural, microbiological, and computational approaches to describe the complete

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ultrastructure of the T2SS pseudopilus and in particular to propose a specific and testable model for the location of the adapter unit XcpH (GspH) between XcpIJK and the filament formed by XcpG. Addition of XcpH would be the next step in a dynamic chronological pathway following XcpIJK association at the periplasmic membrane. Subsequently, XcpG addition absolutely requires the α 1N helices, as there is no interaction between the soluble domains of XcpH and XcpG (**Supplemental Figure 5**), nor indeed between the soluble domain of XcpG and the soluble domain of any of the quaternary complex soluble domains individually²⁷. We have amassed independent and complementary data from several techniques in this integrative study. In particular, the pioneering methodologies in acidic crosslinking represent an important technical breakthrough in the experimental approach to studying protein:protein interactions in multipartner complexes.

This study provides a rationale for the observation that at least in the *P. aeruginosa* Xcp T2SS, XcpH is required for secretion. It serves as the adapter that connects the regular helical assembly of XcpG subunits to the asymmetric complex that catalyzes initiation of the filament and interacts with substrates and other components of the secretion system. It may be possible that XcpH serves other as yet unverified roles in the system as well. For example, given its very strong structural homology to the major XcpG subunit, XcpH might occasionally integrate into the filament and thereby destabilize it, possibly initiating disassembly of the pseudopilus³¹.

Previous interaction studies using surface plasmon resonance²⁷ testing pairwise interactions of the soluble domains of XcpG, H, I, J and K as well as our current NMR results (**Supplemental Figure 5**) reveal that XcpH interacts exclusively with XcpJ. In addition, the binding of XcpH to XcpJ is more efficient when XcpJ is engaged in the XcpIJK ternary complex (Figures 2 & 3 in ²⁷). These data suggested that conformational changes in XcpJ upon binding to XcpI and XcpK create a more favorable docking platform for XcpH and/or favor the downward addition of XcpH to the ternary complex over that of XcpG when both are available within the inner membrane. Our current data supporting many pairwise interactions among the quaternary complex proteins, including between XcpH and each of the other three, fit well with the hypothesis that alone the soluble domains have weak and transient interactions with each other which in some cases are below detection but that within the full ensemble, these interactions are revealed. Our model furthermore accentuates that

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the central α helices are, not surprisingly, fundamentally important for more stable interactions. Future experiments will rely on the α 1N helical tails being part of the interactions.

Based on the near atomic resolution of the XcpQ secretin¹² and taking into account that the pseudopilus tip physically interacts with the secretin³³, a satisfying outcome of our data-driven modeling of the complete pseudopilus is the observation that the filament (~6 nm) docks readily into the periplasm-facing vestibule (~8 nm)¹² of the XcpD secretin (**Supplemental Figure 6**). Moreover, the positioning of the Xcp pseudopilus plus secretin complex into the cryo-tomographic map of the complete T2SS in its natural context, the bacterial envelope²³, indicates that 8 to 16 pseudopilin subunits in addition to the tip complex are required to span the periplasm between the IM and the entry into, or the internal periplasmic gate of, the secretin interior, respectively (**Supplemental Figure 6**).

The results presented here provide insights into not only the *P. aeruginosa* T2SS but indeed into the dozens of T2SS recovered in proteobacteria with similar genome content of five pre-pseudopilins equivalent to XcpG, H, I, J, and K². The conservation of amino acid sequences of these proteins across species implies their protein:protein interaction interfaces are also conserved, and coevolved as shown here for the XcpH_p:XcpJ_p interface. By extension, both the minor pseudopilins of the T2SS and the minor pilins of the T4P can be expected to form a similar quaternary complex in which an adapter minor (pseudo)pilin subunit sets the stage for downward addition of the major (pseudo)pilin subunit to form the extended filament by interacting via its periplasmic domain with the three upper minor subunits and via its unraveled α 1C with the first major subunit. Correspondingly, the set of *E. coli* T4P minor pilins restore assembly of the major T2SS pseudopilin PulG in absence of minor pseudopilins PulHJK³⁷. An open question is whether the secretion ATPase motor protein, GspE, is required for insertion and/or α 1C unraveling in this adapter minor pilin at the junction between the tip complex and the filament itself.

ONLINE MATERIALS AND METHODS

Gene and protein nomenclature

Here we have extended the general T2SS nomenclature to the *P. aeruginosa* Xcp T2SS components. Thus, former XcpT, U, V, W and X proteins are now labelled XcpG, H, I, J and K in agreement with their homologs in other T2SSs. All pseudopilin soluble domains (deleted for their N-terminal hydrophobic domains ($\alpha 1N$)) used in this study are designated with a subscript p indicating periplasmic. According to the standard in the field, polypeptides are numbered with amino acid 1 as the first residue in the mature pseudopilin following prepilin peptidase cleavage.

Bacterial strains and plasmids

Escherichia coli K-12 DH5 α (laboratory collection) and BL21(DE3) pLysS (laboratory collection) were used for cloning procedures and soluble protein production, respectively. *Pseudomonas aeruginosa* PAO1 wild type (laboratory collection), PAO1 $\Delta xcpH$ (this study), $\Delta xcpI$ (this study), $\Delta xcpJ^{43}$, and $\Delta xcpK^{26}$ or PAO1 Δxcp (also called DZQ40)⁵⁴ strains were used for *in vivo* complementation assay. Construction of the *xcpH* and *xcpI* *P. aeruginosa* PAO1 deletion strains was performed as described previously¹⁶ using the pKN- ΔH and pKN- ΔI mutator plasmids (**Table S1**).

Plasmids and oligonucleotides used in this study are listed in **Table S1** and **Table S2**, respectively. Site directed mutagenesis was performed using quick change technology (Stratagen) or inverse PCR. The gene encoding the XcpH_p protein deleted for its mouse trap domain was cloned into the pJN105 vector using SLIC technology⁵⁵. Plasmid pET-XcpH_p was constructed following the strategy used by Durand et al.¹⁶ to construct pET-XcpG_p. All constructs have been verified by DNA sequencing.

LasB secretion and protease activity on plates.

Preparation of culture supernatants from *P. aeruginosa* for analysis of secreted proteins has been described⁵⁶. Gel analysis was standardized so that the volume of supernatant equivalent to 2 OD₆₀₀ units of culture was used for each sample (**Figures 1a, 1c**). Protease activity was tested by spotting 5 μ l of bacterial culture grown in LB to early stationary phase on TSA plates containing 1.5% lyophilized milk and appropriate antibiotics, followed by 14 h incubation at 30°C.

Protein production and purification

Production and purification of XcpH_p and XcpJ_p with or without cysteine substitutions for NMR purposes (except XcpG_p/H_p NMR CSP, see Supplemental Material) was carried out with minor modifications from published procedures³². Expression was performed in BL21(DE3) pLysS *E. coli* carrying the plasmids pETG-20A-XcpH_p or pETG-20A-XcpJ_p. For ¹³C and ¹⁵N uniform protein labeling, cells were grown in 3 L of LB at 37°C to OD₆₀₀ ~0.6. Cells were collected and washed with M9 media without glucose or ammonium chloride to remove remaining LB media. After washing, cells were transferred to 1 L M9 media supplemented with 1 g of ¹⁵N-U ammonium chloride and 2 g of ¹³C-U glucose (Cambridge Isotope Laboratories) and incubated at 25°C for 2 hours before inducing with 1 mM IPTG overnight. Purification of XcpH_p proceeded using a combination of nickel affinity and size exclusion chromatography. The cell pellet was resuspended in Buffer A (50 mM tris-HCl buffer pH 8.0 containing 300 mM NaCl) and lysed using a French press. Supernatant from lysate clarified by centrifugation at 25,000 x G for 30 min. was loaded into a 5 mL Ni-NTA column (Quiagen) and washed with 100 mL of Buffer A containing 50 mM imidazole. Protein was eluted from the column using Buffer A with 500 mM imidazole. Peak fractions containing target protein were pooled and treated with TEV protease (purified in house) added to a final concentration of ~40 µg/mL and the sample was dialyzed overnight at 4°C against Buffer A to remove imidazole. Removal of TEV protease proceeded by loading samples into a 1 mL Ni-NTA column which was washed with the same buffer. Fractions containing XcpH_p were collected and concentrated for size exclusion chromatography in a Sephacryl S-100 HiPrep 16/60 column (GE). During this step, buffer was exchanged to NMR buffer (25 mM sodium phosphate buffer pH 6.5 containing 25 mM NaCl). XcpJ_p was expressed and purified using the identical protocol but without ¹³C and ¹⁵N labeling.

Production and purification of XcpH_p, XcpI_p, XcpJ_p, XcpK_p, XcpH_pV46C, XcpH_pD49C, XcpH_pL53C, XcpJ_pR46C and XcpJ_pR53C for cysteine cross-linking (and cross-linking MS experiments was performed in BL21(DE3) pLysS *E. coli* carrying the corresponding pETG-20A plasmid (**Table S1**) as published.²⁷

In vitro cysteine cross-linking

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40 μ M of a single cysteine variant or two variant proteins in 1:1 molar ratio were incubated in Tris-HCl 50 mM, NaCl 100mM, pH 8 in a total volume of 500 μ L. The mixture was supplemented with DTT (20 mM) to allow reduction of intra-chain disulfide bands formed during purification steps. The mixture was dialyzed against 300 mL of Tris-HCl 50 mM, NaCl 100 mM, pH 8 buffer for 2 h at RT to allow DTT removal and cysteine oxidation. To analyze disulfide bond formation, 75 μ L of each reaction was mixed with 25 μ L of Leammli loading buffer with or without reductant as appropriate. The samples were heated for 5 min at 95°C and then visualized by 12% Coomassie blue stained SDS-PAGE.

Acidic Crosslinking of XcpH_p + XcpJ_p and XcpH_p + XcpI_p + XcpJ_p + XcpK_p

Preparation of multimers For assembling the dimer, 17 μ L of XcpH_p at 4 mg/mL were mixed to 25 μ L of XcpJ_p at 4 mg/ml, leading to a mixture of 4 nmol of each protein (molar ratio 1:1). For the tetramer, 17 μ L of XcpH_p (4 mg/mL), 12 μ L of XcpI_p (4 mg/mL), 25 μ L of XcpJ_p (4 mg/mL) and 34 μ L of XcpK_p (4 mg/mL) were mixed to yield a 4 nmol solution of each protein (molar ratio 1:1:1:1). Samples were dried under vacuum, and dissolved in 100 μ L of PBS buffer pH 7.4 to achieve a final concentration of each protein of 40 μ M. Proteins were kept at 25°C for 1 h to let the association of the dimer or tetramer occur.

Cross-linking reaction To crosslink the dimer, 10 μ L of ADH (adipic acid dihydrazide) and 16 μ L of DMTMM ((4-(4,6-dimethoxy-1,3,5-triazin-2-yl)-4-methyl-morpholinium chloride) were added to the solution of XcpH_p:J_p complex, reaching final concentrations of ADH and DMTMM equal to 46 mM (8 and 12.7 mg/mL, respectively). For the tetramer, 23 μ L of ADH and 36 μ L of DMTMM were added to the solution of XcpH_p:I_p:J_p:K_p complex, reaching final concentrations of ADH and DMTMM equal to 81 mM (14.5 and 22.6 mg/mL, respectively). These concentrations were chosen to impose a molar excess of crosslinking reagent of more than 1,000. The mixture was incubated 2 h at 37°C under agitation (750 rpm) to allow intensive crosslinking reaction. The reaction was quenched by reagent removal using a ZebaSpinDesalting Column (0.5mL, 7k, Pierce), employed according to the manufacturer recommendations. The resulting sample was dried using a Speedvac system.

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SDS-PAGE Cross-linked species were imaged by classical SDS-PAGE analysis. The protein mixture was suspended in 170µL of 8M urea, leading to a global protein concentration of 48µM. 20µg of proteins (=10µL) were dried before being solubilized in 20µL of Laemmli buffer (Tris HCl pH 6.8 (65mM), SDS 2%, glycerol 20% and DTT 350mM) and a spatula tip of bromophenol blue. The sample was heated at 100°C for 5 min, and 10µL (~10µg) were loaded on the gel (NuPage™ 4-12% Bis-Tris-Gel, 1.0 mm x 10 wells). The migration starts with a voltage of 200V applied for 40min (stacking), and then 150V for the separation. The electrophoresis system is switched-off when the migration blue line is localized at the extremity of the gel. The gel is fixed 3h in a bath composed of 50% ethanol, 47% water and 3% phosphoric acid, washed three times with ultrapure water, and finally incubated in water containing 34% methanol, 17% ammonium sulfate and 3% phosphoric acid. The coloration of the bands was made during 3 days with Coomassie blue G250 added at 360 mg/l in the solution. The gel was finally washed several times in pure water to remove the excess of Coomassie blue.

In-gel digestion The bands corresponding to the XcpH_p:XcpJ_p dimer or XcpH_p:XcpI_p:XcpJ_p:XcpK_p tetramer, respectively, were excised, and cut into small pieces. Resulting pieces of gel were then washed by 50µL of 50mM NH₄HCO₃ then centrifuged (5min, 600 rpm) to remove the supernatant. The same step was repeated with 50µL of 50/50 (v/v) 50mM NH₄HCO₃/acetonitrile. The two previous steps were repeated twice. To ensure a good penetration of the enzymes into the pieces of gel, the latter were dehydrated twice with 50µL of pure acetonitrile removed by centrifugation (5 min, 600 rpm). The gel was then rehydrated at 0°C with 3µL of a solution containing 1/100 of Lys-C and 1/50 of trypsin in 50mM NH₄HCO₃. The digestion was performed at 37°C for 4 hr. Enzymatic activity was quenched by acidifying the medium using 25µL of 1% trifluoroacetic acid. Resulting peptides were eluted from the gel by incubating the sample overnight at 20°C under 600 rpm.

LC-MS/MS 1µg of the digested material was analysed using a UPLC nanoACQUITY (Waters, UK) coupled to a Q-Exactive Plus Hybrid Quadrupole-Orbitrap Mass Spectrometer (Thermo Scientific, USA). The chromatographic system is equipped with two columns. The first one, dedicated to the trapping of peptides, is a Symmetry C18 (5 µm, 180 µm x 20 mm, Waters, UK). The second one which performs the analytical separation is a HSST3 C18 (1.8 µm, 75 µm x 250

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mm, Waters). The dimensions given for the columns are in the order: particle diameters, internal diameters and column lengths. Solvent A was water, acidified with 0.1% formic acid and solvent B was acetonitrile, also acidified with 0.1% formic acid. Cross-linked peptides were first trapped for 3min (98/2, v/v, A/B) at a flow rate of 20 μ L/min before being eluted with a gradient of 57 min at a flow rate of 700 nL/min. The elution started with a linear gradient of B from 2% to 7% in 5min, followed by an increase from 7% to 40% in 25min, then to 85% in 3min. This composition A/B 15/85 is kept for 5min before changing to 98/2 for reconditioning the analytical column. The Q-Exactive Plus spectrometer was set in a nanoESI positive mode acquisition for 57 minutes. The acquisition was recorded in full scan MS and data dependent MS/MS, in a mass range m/z 400-1,750. For the MS stage, resolving power was set at 70,000 @m/z 200, with an automatic gain control (AGC) target at 1e6 (or 50 ms as a maximum injection time). For MS/MS, a “Top 12” experiments was applied, meaning that the twelve most intense ions of each MS scan were selected for fragmentation. Singly charged ions, ions with undetermined charge (for example, electronic noise) and ions with signal intensities below the AGC threshold set at 1e3 were excluded from this selection. For precursor ions, the selection window was 2.0 m/z, the AGC target was 1e5 (or 50ms as a maximum injection time) and the resolving power of 17,500 @m/z 200. Normalized collision energy was 25. A dynamic exclusion of 10s was also applied to avoid the redundancy of MS/MS spectra of the same ions.

- Data Analysis: Spectrum Identification Machine SIM-XL version 1.5.0.14⁵⁷ was used for identification of cross-linked peptides. ADH-DMTMM crosslinker was set up in the software to create a mass shift of 138.0905 Da for each crosslink and 156.1012 for hydrolyzed monolinks. The possible reaction sites were restricted to C-terminal extremities and acidic side chains of Glu and Asp amino acids. Accuracy on mass measurements was 2 ppm for the precursor and 10 ppm for the fragment ions, to reduce to their maximum the false positives and to increase the reliability of the results. Oxidation of methionine, due to experimental conditions, has also been considered as a variable modification. Only the crosslinked peptides characterized by a SIM-XL internal score of 2.5, 3.0, 2.3, 3.0 or 2.5 and higher have been retained for IK, IJ, HJ, HI, or HK pairs, respectively. All the detected crosslinked peptides have been manually validated by analyzing their corresponding MS/MS spectra for exact masses, numbers of identified fragments, and signal intensity (e.g. **Figure 6b**, **Supplemental Figure 3b**).

NMR experiments.

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All solution NMR experiments (except XcpH_p/J_p chemical shift perturbation) were run at 37°C in a Bruker 900 MHz NMR spectrometer at the National Magnetic Resonance Facility at Madison. Sample condition used was NMR buffer containing 0.01% sodium azide, 50 μM 2,2-dimethyl-2-silapentane-5-sulfonate (DSS) (Sigma) and 8% D₂O.

Backbone sequential assignments were carried out using standard 3D NMR experiments including HNCO, HN(CA)CO, HNCACB, CACB(CO)NH. These experiments were collected using non-uniform sampling (NUS). Data processing of NUS data was performed using NMRPipe⁵⁸. Chemical shift referencing was done using DSS as reference. Backbone assignments were facilitated using NMRFAM-Sparky software⁵⁹. Calculation of secondary chemical shifts for secondary structure estimation was done using random coil chemical shifts calculated for the XcpH amino acid sequence using ncIDP library⁶⁰. Secondary chemical shifts were calculated as $\Delta\delta\text{Ca} - \Delta\delta\text{Cb}$, where $\Delta\delta\text{Ca} = (\delta\text{Ca}_{\text{XcpH}} - \delta\text{Ca}_{\text{random coil}})$ and $\Delta\delta\text{Cb} = (\delta\text{Cb}_{\text{XcpH}} - \delta\text{Cb}_{\text{random coil}})$. This step effectively removes any error in chemical shift referencing.

Chemical shift perturbation assays were carried out using 100 μM ¹⁵N-U XcpH and increasing amounts of unlabeled XcpJ (20 μM, 50 μM, and 80 μM). An independent ¹H-¹⁵N HSQC spectrum was collected for each sample. Total change of the amide proton and nitrogen chemical shift ($\Delta\delta\text{NH}$) was calculated for each residue as $((\Delta\delta\text{H})^2 - 1/6 (\Delta\delta\text{N})^2)^{1/2}$, where $\Delta\delta\text{H}$ and $\Delta\delta\text{N}$ correspond to the difference in chemical shift of proton and nitrogen respectively. $\Delta\delta\text{NH}$ was calculated using the control spectrum as the initial point and the spectrum with 80 μM XcpJ as the endpoint.

To perform paramagnetic relaxation enhancement (PRE) experiments, four different XcpJ_p cysteine mutant constructs (R46C, R53C, T178C, E180C) were used to attach the spin label (1-Oxyl-2,2,5,5-tetramethylpyrroline-3-methyl) methanethiosulfonate (MTSL) (Santa Cruz Biotechnology). XcpJ_p variants were purified as above in Buffer A containing 1 mM DTT. MTSL labeling of XcpJ_p was performed in NMR buffer and 0.5 mM DTT. XcpJ_p samples were incubated with a 20x molar excess of MTSL for 4 h at room temperature. Additional 20x molar excess of MTSL was added to the sample which was further incubated at room temperature overnight. Excess MTSL was removed by dialysis against the NMR buffer. The NMR sample contained 80 μM ¹⁵N-U XcpH_p and 24 μM of MTSL labeled XcpJ_p. The ratio of XcpH to XcpJ_p was selected based on the chemical shift perturbation assay to avoid signal loss due to complex formation. A 2D ¹H-¹⁵N HSQC spectrum was acquired for each sample. After collecting the HSQC experiment of the oxidized or paramagnetic form of MTSL, 2 mM sodium ascorbate (Sigma-Aldrich) was

added to the sample and incubated at room temperature for at least one hour before collecting the spectrum of the reduced or diamagnetic MTSL form. Effects of the paramagnetic label were quantified as the ratio of signal intensity of the spectrum collected with MTSL in paramagnetic versus diamagnetic form ($I_{\text{para}}/I_{\text{dia}}$).

Quaternary complex computational modeling

Modeling of the XcpH_pI_pJ_pK_p quaternary complex was achieved using high ambiguity driven protein-protein Docking (HADDOCK)⁶¹. Structures used for HADDOCK corresponded to the XcpI_pJ_pK_p crystal structure (PDB: 5VTM) and an XcpH_p model prepared based on the structure of EpsH_p from *Vibrio cholerae* (PDB: 2QV8) using the Phyre2 server⁶². Allowed distances for acidic crosslinking restraints between C_α carbons were set between 8-18 Å, while 6-18 Å were used for PRE restraints (**Supplemental Tables 3 & 4**). Docking the XcpH_p model onto the XcpI_pJ_pK_p structure was performed using the HADDOCK 2.2 server⁶¹. The best structure obtained from HADDOCK was selected for further analysis of the XcpH_pI_pJ_pK_p complex.

Biological filament modeling

Modeling of the XcpG Type II secretion system pseudopilus was based on the PulG filament structure from *Klebsiella oxytoca*¹⁵ (PDB code 5WDA) using Pymol. Modeling was divided in the following steps: (1) adding missing transmembrane helices to XcpH_pI_pJ_pK_p proteins, (2) positioning the XcpHIJK complex onto the PulG filament (3) fitting XcpHIJK helices to the position of PulG transmembrane helices within the filament (4) fitting XcpG⁶³ units to the PulG filament and (5) relaxing the whole system to remove atomic clashes. Missing transmembrane helices in the XcpH_pI_pJ_pK_p complex were initially modeled using the Phyre2 server using the respective amino acid sequence of the complete alpha helix. This modeled alpha helix was aligned to the soluble domain alpha helix structure in the HADDOCK model. Specific to XcpH, the helix was modeled on the PulG template from the EM reconstruction to include the unraveled section. Finally, the modeled helices were added to the XcpH_pI_pJ_pK_p complex PDB file using custom Python scripts. For XcpG a similar procedure was performed, in this case a model of the transmembrane helix was based on PulG helix (including the unraveled helix section), which was added to the XcpG_p solution NMR structure⁶³ (PDB code 2KEP). The XcpHIJK complex with the helices added was positioned at the tip of PulG filament by aligning the XcpH soluble domain with

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the first PulG unit. Then, XcpHIJK model transmembrane helices were modeled semi-manually to fit the interior cavity of the PulG filament. Custom Python scripts were used to modify ϕ and ψ for this purpose. To guide the modeling, the transmembrane helices of PulG units were used as a reference. Finally, PulG subunits were replaced with XcpG to complete the pseudopilus filament. To achieve this, XcpG with the added helix was aligned to a PulG unit and XcpG helix was modeled to closely match the PulG transmembrane helix. Then, the modeled XcpG unit was copied and aligned to the PulG units to complete the filament. Finally, the whole model was relaxed with pyRosetta⁵². During this procedure, the known hydrogen bond between E5 carboxyl group and N-terminal amino group of the next subunit in the filament was enforced as a distance restraint of 2.8 Å. Other restraints to maintain the quaternary tip complex and XcpG filament assembly are shown **Supplemental Tables 5 & 6**. Ten filament models were produced, from which the best was selected based on favourable XcpG subunit packing and the lowest RMSD of the XcpH_pI_pJ_pK_p tip structure with respect the HADDOCK model.

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AUTHOR CONTRIBUTIONS

RV, LQ, and KTF conceived the studies. CAE and BD subcloned, made site directed mutants, and purified soluble proteins. CAE carried out CSP and PRE experiments, and all quaternary structure

and filament modeling. BD performed cysteine cross linking experiments. GB constructed *P. aeruginosa* strains and carried out secretion assays. SA carried out NMR experiments in Supp. Fig S4. BB was responsible for coevolution calculations in Supp. Fig S5. LQ performed all acidic cross linking methods development and experiments. RV, KTF, and CAE drafted the manuscript. All authors discussed and edited manuscript and prepared figures.

COMPETING INTERESTS The authors declare no competing interests.

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FIGURE LEGENDS

Fig 1. Minor pseudopilins are required for Type II Secretion. a. Secretion of the principal Xcp T2SS-dependent exoprotein (LasB) assayed by Coomassie blue stained SDS-PAGE of equal OD equivalent loading of culture supernatants of the wild type PAO1, Δxcp (DZQ40), $\Delta xcpH$, $\Delta xcpI$, $\Delta xcpJ$ and $\Delta xcpK$ strains harboring empty vector (EV), or a complementing plasmid encoding XcpH (pxcpH), XcpI (pxcpI), XcpJ (pxcpJ) or XcpK (pxcpK). b. LasB protease activity on a skim milk plate is indicated by a halo around a colony, corresponding to casein degradation by secreted LasB. Colonies representing WT (PAO1), Δxcp , or complemented Δxcp strains are labeled accordingly. Note $\Delta xcpH$ is complemented by both pxcpH and by pJN105 expressing the XcpH deletion of the $\beta 3$ - $\beta 4$ loop, pxcpH $_{\Delta 34}$. c. Secretion assay, as in panel a. showing secretion of LasB by $\Delta xcpH$ /pxcpH $_{\Delta 34}$.

Fig 2. XcpH_p NMR assignments and secondary structure determination. a. XcpH_p assignments by solution NMR. 2D 1H - ^{15}N HSQC spectrum showing amino acid assignments for XcpH_p. NMR experiments were performed at 37°C using a sample of 500 μM ^{13}C - ^{15}N labeled XcpH_p in NMR buffers. b. Secondary structure analysis of XcpH_p. Secondary chemical shifts were calculated for XcpH_p using α and β carbon chemical shifts obtained after backbone sequential assignments. Stretches of amino acids with positive values indicate the presence of α helix while negative values correspond to β strands. Amino acids not assigned or overlapped are represented by a value of zero. Secondary structure elements in the XcpH_p model structures are shown above.

Fig 3. XcpH_p chemical shift perturbations due to interaction with XcpJ_p. a. Plot of CSPs calculated for XcpH_p amino acids. Each value was calculated using NH chemical shift differences obtained from the XcpH_p spectrum collected without XcpJ_p and or with 80 μM XcpJ_p. CSP values above 0.06 were considered significant. Amino acids presented with value zero correspond to residues unassigned or overlapped. b. CSP values were mapped onto an XcpH_p model based on *E. coli* GspH (PDB: 2KNQ⁶⁴). Non-significant changes are shown in white, while more significant values are depicted in a gradient of red. Amino acids with no data are shown in grey.

XcpH:I:J:K Interaction

Fig 4. *Cysteine cross linking suggests XcpH_p:XcpJ_p α1C packing.* Cysteine substitutions were made in indicated positions within XcpH_p (by column) and in XcpJ_p (by row). Purified proteins were incubated under oxidizing conditions and visualized by Coomassie blue staining of non-reducing SDS PAGE. In each of the six pairwise combinations, Lane 1 is XcpH_p alone, Lane 2 is XcpH_p+XcpJ_p, and Lane 3 is XcpJ_p alone. Monomer, dimer, and heterodimer sizes are indicated along the right-hand side for all three gels in each row.

Fig 5. *PRE experiments of XcpH_p in the presence of MTSL labeled XcpJ_p.* a. PRE effect of the two α1C MTSL-labeled XcpJ_p cysteine mutants on the XcpH_p spectra. Spectra were collected using 80 μM ¹⁵N XcpH_p and 24 μM XcpJ_pR46C-MTSL or XcpJ_pR53C-MTSL. Significant signal intensity changes are designated with a gradient of red. All amino acids with displayed data have an intensity ratio above zero, while residues with no assignments or overlapped have values of zero. b. PRE effects observed with XcpJ_pR46C-MTSL or XcpJ_pR53C-MTSL were mapped onto the structural model of XcpH_p.

Fig 6. *Acidic crosslinking of XcpH_pI_pJ_pK_p soluble complex.* a. SDS-PAGE of the cross-linked proteins with a box around the band identified by the mean of proteomics as the cross-linked XcpH_pI_pJ_pK_p tetramer. b. MS/MS spectrum of a particularly relevant cross-link pair between XcpH_p and XcpJ_p. The spectrum shows intense ions identified as y- and b-type ions, allowing an unambiguous identification of the two partners (mass accuracy for the parent < 2ppm and for the fragment < 10ppm). c. Circular representation of all significant cross-linked pairs detected in the tetramer.

Fig 7. *3D model of XcpH_pI_pJ_pK_p soluble complex.* A model of the XcpH_pI_pJ_pK_p quaternary complex (XcpH_p gray, XcpI_p blue, XcpJ_p green and XcpK_p wheat) was obtained via HADDOCK docking. Acidic cross-linking derived restraints are shown in yellow dotted lines. In the case of PRE restraints, XcpJ_p spin labeled residues R46 and R53 are shown as sticks while most affected XcpH_p residues N55 and D49 are represented as blue spheres at the amide nitrogen. XcpH_pJ_p residues forming salt bridges or hydrogen bonds are shown in stick model connected by yellow dotted lines. Residues from XcpH_p and XcpJ_p are labeled in dark grey and dark green respectively.

XcpH:I:J:K Interaction

932 **Fig 8.** *Model of Type II secretion system pseudopilus.* a. Model of XcpHIJK tip complex and
 933 XcpG filament shown as a solvent-accessible surface. (above) Top and bottom views of the
 934 models (below) Side views of the pseudopilus model. Color coding of each XcpG protofilament
 935 is a lighter shade of the respective tip subunit color from whence it originates. b. Transmembrane
 936 α 1N contacts between neighboring subunits include canonical F1-E5 salt bridge (included as
 937 restraint during minimization), and hydrophobic residues at position 16 and 10 previously
 938 described³⁷ but importantly not used as a modeling restraint. Images were generated using Pymol
 939 software.

Escobar & Douzi, *et al.*, Figure 1

Fig 1. Minor pseudopilins are required for Type II Secretion. a. Secretion of the principal Xcp T2SS-dependent exoprotein (LasB) assayed by Coomassie blue stained SDS-PAGE of equal OD equivalent loading of culture supernatants of the wild type PAO1, Δxcp (DZQ40), $\Delta xcpH$, $\Delta xcpI$, $\Delta xcpJ$ and $\Delta xcpK$ strains harboring empty vector (EV), or a complementing plasmid encoding XcpH (*pxcpH*), XcpI (*pxcpI*), XcpJ (*pxcpJ*) or XcpK (*pxcpK*). b. LasB protease activity on a skim milk plate is indicated by a halo around a colony, corresponding to casein degradation by secreted LasB. Colonies representing WT (PAO1), Δxcp , or complemented Δxcp strains are labeled accordingly. Note $\Delta xcpH$ is complemented by both *pxcpH* and by pJN105 expressing the XcpH deletion of the beta3- beta4 loop, *pxcpH* $\Delta 34$. c. Secretion assay, as in panel a. showing secretion of LasB by $\Delta xcpH$ /*pxcpH* $\Delta 34$.

Escobar & Douzi, *et al.*, Figure 3

Fig 3. XcpHp chemical shift perturbations due to interaction with XcpJp. **a.** Plot of CSPs calculated for XcpHp amino acids. Each value was calculated using NH chemical shift differences obtained from the XcpHp spectrum collected without XcpJp and or with 80 μM XcpJp. CSP values above 0.06 were considered significant. Amino acids presented with value zero correspond to residues unassigned or overlapped. **b.** CSP values were mapped onto an XcpHp model based on *E. coli* GspH (PDB: 2KNQ64). Non-significant changes are shown in white, while more significant values are depicted in a gradient of red. Amino acids with no data are shown in grey.

Escobar & Douzi, *et al.*, Figure 4

Fig 4. Cysteine cross linking suggests XcpHp:XcpJp 1:1 packing. Cysteine substitutions were made in indicated positions within XcpHp (by column) and in XcpJp (by row). Purified proteins were incubated under oxidizing conditions and visualized by Coomassie blue staining of non-reducing SDS PAGE. In each of the six pairwise combinations, Lane 1 is XcpHp alone, Lane 2 is XcpHp+XcpJp, and Lane 3 is XcpJp alone. Monomer, dimer, and heterodimer sizes are indicated along the right-hand side for all three gels in each row.

Fig 5. PRE experiments of XcpHp in the presence of MTSL labeled XcpJp. **a.** PRE effect of the two α 1C MTSL-labeled XcpJp cysteine mutants on the XcpHp spectra. Spectra were collected using 80 μ M 15N XcpHp and 24 μ M XcpJpR46C-MTSL or XcpJpR53C-MTSL. Significant signal intensity changes are designated with a gradient of red. All amino acids with displayed data have an intensity ratio above zero, while residues with no assignments or overlapped have values of zero. **b.** PRE effects observed with XcpJpR46C-MTSL or XcpJpR53C-MTSL were mapped onto the structural model of XcpHp.

Escobar & Douzi, et al., Figure 6

Fig 6. Acidic crosslinking of XcpHlpJpKp soluble complex. a. SDS-PAGE of the cross-linked proteins with a box around the band identified by the mean of proteomics as the cross-linked XcpHlpJpKp tetramer. **b.** MS/MS spectrum of a particularly relevant cross-link pair between XcpHp and XcpJp. The spectrum shows intense ions identified as y- and b-type ions, allowing an unambiguous identification of the two partners (mass accuracy for the parent < 2ppm and for the fragment < 10ppm). **c.** Circular representation of all significant cross-linked pairs detected in the tetramer.

Escobar & Douzi, et al., Figure 7

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Escobar & Douzi, et al., Figure 8

Fig 8. Model of Type II secretion system pseudopilus. a. Model of XcpHIJK tip complex and XcpG filament shown as a solvent-accessible surface. (above) Top and bottom views of the models (below) Side views of the pseudopilus model. Color coding of each XcpG protofilament is a lighter shade of the respective tip subunit color from whence it originates. **b.** Transmembrane $\alpha 1N$ contacts between neighboring subunits include canonical F1-E5 salt bridge (included as restraint during minimization), and hydrophobic residues at position 16 and 10 previously described (37) but importantly not used as a modeling restraint. Images were generated using Pymol software.