Collagen-Like Proteins in Pathogenic E. coli Strains

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Abstract

The genome sequences of enterohaemorrhagic E. coli O157:H7 strains show multiple open-reading frames with collagen-like sequences that are absent from the common laboratory strain K-12. These putative collagens are included in prophages embedded in O157:H7 genomes. These prophages carry numerous genes related to strain virulence and have been shown to be inducible and capable of disseminating virulence factors by horizontal gene transfer. We have cloned two collagen-like proteins from E. coli O157:H7 into a laboratory strain and analysed the structure and conformation of the recombinant proteins and several of their constituting domains by a variety of spectroscopic, biophysical, and electron microscopy techniques. We show that these molecules exhibit many of the characteristics of vertebrate collagens, including trimer formation and the presence of a collagen triple helical domain. They also contain a C-terminal trimerization domain, and a trimeric α-helical coiled-coil domain with an unusual amino acid sequence almost completely lacking leucine, valine or isoleucine residues. Intriguingly, these molecules show high thermal stability, with the collagen domain being more stable than those of vertebrate fibrillar collagens, which are much longer and post-translationally modified. Under the electron microscope, collagen-like proteins from E. coli O157:H7 show a dumbbell shape, with two globular domains joined by a hinged stalk. This morphology is consistent with their likely role as trimeric phage side-tail proteins that participate in the attachment of phage particles to E. coli target cells, either directly or through assembly with other phage tail proteins. Thus, collagen-like proteins in enterohaemorrhagic E. coli genomes may have a direct role in the dissemination of virulence-related genes through infection of harmless strains by induced bacteriophages.

Introduction

Enterohaemorrhagic E. coli (EHEC) is responsible for gastrointestinal disorders in humans that range from abdominal pain and diarrhoea to haemorrhagic colitis and haemolytic uraemic syndrome [1,2,3]. The EHEC serotype most often linked with outbreaks of severe disease is E. coli O157:H7. The genomes of the E. coli O157:H7 strains EDL933 and Sakai are 9.9 Mb larger than that of the non-pathogenic laboratory E. coli strain K-12 [4,5]. That extra genetic material is the result of horizontal gene transfer (HGT) probably mediated by bacteriophages: the Sakai strain genome includes 18 prophages and 6 prophage-like elements integrated into different sites of the bacterial chromosome [5,6], while the EDL933 genome contains 18 prophages and prophage-like elements [4]. Up to 463 phage-associated genes are present in the O157:H7 strains for only 29 in the K-12 strain [3,7].

Several virulence genes of the O157:H7 strain are located into these prophages and prophage-like elements, notably the Shiga toxin (verocytotoxin) genes stx1 and stx2 [8], and various effector proteins that are injected into the host cells by a type III secretion system [9,10]. Collectively, EHEC strains are considered new pathogens that have emerged from less virulent strains by progressive acquisition of virulence factors via HGT. There is significant evidence that variation of the prophage sequences is a main factor for the genomic and virulence diversity of EHEC [6,7,11,12,13]. The acquired specific virulent attributes allow EHEC strains to adapt to new niches and to broaden the spectrum of disease.

Intriguingly, the genomes of E. coli O157:H7 also include several open reading frames containing stretches of collagen-like sequences. Collagen proteins are principal components of the extracellular matrix of metazoa and amongst Earth’s most abundant biopolymers. Vertebrates have at least 28 collagen types described [14], with type I collagen being the main fibrous protein component of skin, tendon, bone and other connective tissues. All collagen proteins have at least one domain with a specific three-dimensional structure known as the collagen triple helix, in which three polypeptide chains wrap around a common...
helical axis and are connected through a ladder of intermolecular hydrogen bonds roughly perpendicular to that axis [15,16,17,18]. The conformation of the collagen triple helix imposes a repetitive amino acid sequence pattern where glycine residues (Gly, G) occur at every third position. This (Gly-X-Y)n pattern is recognized as the signature of collagen proteins and domains.

A surprising number of collagen-like sequences have been detected outside the metazoan realm, notably in bacterial and viral genomes [19,20,21]. These “prokaryotic collagens” exhibit in their (Gly-X-Y)n regions significant differences in residue content and distribution with respect to vertebrate collagens, and yet they seem to show the basic molecular characteristics of true collagen proteins [22,23]. The functions and potential contribution to virulence of these prokaryotic collagens are currently under study, but they seem to participate in pathogenesis in unexpected ways. Thus, collagen-like glycoproteins from Bacillus anthracis are components of the exosporium that are able to interact with integrin receptors on professional phagocytes [24,25], while collagen-like surface proteins from Streptococcus pyogenes are able to promote bacterial adhesion and internalization to respiratory epithelial cells [26,27,28].

Open reading frames with collagen-like sequences in the genomes of E. coli O157:H7 and other EHEC strains are automatically annotated as “hypothetical tail fibre proteins”. These collagen-like sequences seem a distinctive feature of EHEC strains and several bacteriophages, and have not been detected in K-12 or other non-pathogenic strains. They are normally included in the prophage or prophage-like elements of the EHEC genomes and would be expected to participate in phage morphogenesis during phage induction. Indeed there is evidence of changes in levels of expression for some of these collagen-like protein transcripts under certain experimental conditions, normally in association with other prophage genes [29,30,31,32].

While most of the prophages in the EHEC genomes appear to be defective, often lacking genes apparently critical for phage induction and viability, phage induction from EHEC strains has been demonstrated and Shiga-toxin converting phages can be defective, often lacking genes apparently critical for phage induction and viability, phage induction from EHEC strains has been demonstrated and Shiga-toxin converting phages can be detected free in the extraintestinal environment [12,33]. Furthermore, potentially defective phages have been shown to be inducible, to release virus particles of different morphologies and, after release, to infect other E. coli strains, [34]. The same study also suggests that recombination and other inter-prophage interactions may make possible the biological activation of defective prophages [34].

Thus, prophages embedded in EHEC genomes have the potential of disseminating virulence factors through bacterial infection and HGT. Their morphogenetic proteins are largely uncharacterized and deserve investigation. Here, we present a first biochemical analysis of the collagen-like proteins of EHEC prophages, which we will refer collectively as EPClPs (EHEC Prophage collagen-like Proteins).

Results

Domain Architecture of Collagen-like Proteins in EHEC Genomes

Several open reading frames potentially encoding collagen-like proteins have been identified by automatic sequence annotation in the genomes of EHEC strains. Those from the Sakai and EDL933 genomes will be discussed here, but many related sequences have been identified in other strains. Their primary structures show one or more collagen-like domains (Col) with the repeating collagen signature sequence (Gly-X-Y)n, flanked at both ends by a series of non-collagenous, conserved domains (Figure 1 and Table 1). Domains PIN, PI2 and PIC have been described on the basis of sequence conservation and are associated to fibre tail proteins from phages. They appear in automatic annotation of EPClPs. Figure 1 shows the different protein architectures and the nomenclature used here to refer to them, plus two representative sequences. The most common architecture (EPclA) appears in multiple copies in each genome, with more than 90% amino acid sequence identity across copies. Table 2 gives the complete list of EPClP sequences from the Sakai and EDL933 genomes, whereas representative examples of other architectures and strains are given in Table 3.

The EPclA architecture shows a single collagen triple helical sequence capped by PIN and PIC domains at the N- and C-termini, respectively. Between the PIN domain and the Col domain there is a region of low-complexity. Analysis of its amino acid sequence suggests a coiled-coil conformation (see below), and thus will be referred here as Pcoil domain. The EPclB architecture shares the presence of PIN, Pcoil and PIC non-collagenous domains, and contains two Col domains separated by a PI2-type repeat. Protein sequences within each type of architecture show variable lengths of their Col and Pcoil domains (Tables 2 and 3). Differences in length are typically multiples of three for Col domains and multiples of seven for Pcoil domains, which is consistent with the lengths of the repetitive motifs in collagen and coiled-coil sequences, respectively. In sequences with two Col domains it is common that the first one contains a single interruption of the (Gly-X-Y)n repeating pattern, with a conserved Gly-X-Pro-Gly-Gly-Pro-X-Gly sequence.

Only a few sequences conform to the EPclC and EPclD architectures (Table 3), which are characterized by different C-terminal regions with no sequence homology to the PIC domains of EPclA or EPclB architectures. Also, the Pcoil domain is often missing in EPclC architectures. The EPclD sequence Stx2-86_gp21, from the Stx2-86 prophage in the Shiga toxin-producing E. coli strain O157:H– (accession codes Q08J34, YP_794068), has a 322-amino acid Col domain (Table 3), by far the longest collagen-like sequence of all EPClPs. This long domain also shows a single Gly-X-Pro-Gly-Gly-Pro-X-Gly interruption. No examples of EPclC or EPclD sequences are found in the EDL933 or Sakai genomes.

The PIN and PI2 domains are not exclusive to collagen-like proteins from E. coli prophages and were first identified in the side tail fibre protein coded by the stf gene from λ bacteriophages [35,36] (accession code P03764). Virions with a functional stf gene show jointed tail fibres, expanded receptor specificity, and adsorb more rapidly to E. coli cells. Homologus proteins have been identified in embedded prophages of many E. coli strains, including the laboratory reference strain K12 (protein strR/ynaB, accession code P76072).

Other than their consistent presence in prophage tail fibre proteins, little is known about the structure and function of the PIN and PI2 domains. Some sequence similarity between PIN and a regulatory domain of eukaryotic carboxypeptidases may be indicative of a proteolysis-related function, but to date there is no experimental evidence for this. Due to the presence of these domains, EPClP sequences are automatically annotated as putative tail fibre proteins.

Amino Acid Composition and Positional Preference in the Collagen Domains of EPClPs

Collagen domains in EPClPs show amino acid preferences in the X and Y positions that differ from those seen in other collagen proteins. By far the most common residue in the X position is proline (Pro, P), which occurs there close to half of the time.
Pro is relatively infrequent in the Y position. Both X and Y positions also show a strong preference for charged amino acids, aspartate/glutamate (Asp/Glu, D/E) in the X position and lysine/arginine (Lys/Arg, K/R) in the Y position. Alanine (Ala, A) is also relatively frequent at both X and Y positions, and both glutamine (Gln, Q) and threonine (Thr, T) show a clear preference for the Y position. Interestingly, cysteine, phenylalanine, histidine, tryptophan and tyrosine are absent in collagen domains from EPclPs. At the triplet level, the most common are GPK (15%), GPQ (12%), GPA (12%), GER (8%) and GET (8%). The triplet pattern GP(Q/P)–GPK–G(D/E) is repeatedly observed in the collagen domains of EPclPs.

Figure 1. Collagen-like proteins from prophages embedded in the genomes of E.coli O157:H7 and other EHEC strains, referred here as EPclA to EPclD (EHEC Prophage collagen-like A to D). (A) Domain architectures. The collagen triple helical domains are labelled “Col”, and domains predicted to adopt an α-helical coiled-coil conformation (see text) are labelled “PCoil” (for phage coiled-coils). Key to other domain labels (Table 1): PfN, phage fibre N-terminal domain; PfC, phage fibre C-terminal domain; PfC2, phage fibre C-terminal domain, variant 2; Pf2, phage fibre repeat 2. (B) Sequence of a representative collagen-like protein with EPclA architecture (ECs2717), from the genome of E. coli O157:H7 Sakai. (C) Sequence of a representative collagen-like protein with EPclB architecture (Z1483), from the genome of E. coli O157:H7 EDL933. Amino acid sequences corresponding to the different predicted domains are colour-coded as in (A).

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residues in the Y position are often modified post-translationally to 4-hydroxyproline (Hyp, O), which contributes to the thermal stability of the collagen domains. Charged residues are also frequent in the X and Y positions of human collagens, with the same positional preferences as in collagen domains from EPclPs (Asp/Glu more often in X, Lys/Arg more often in Y). However, they are overall less frequent and their preferential position is less strict (Table 4). Other amino acids significantly contribute to the sequence variability at each position.

The expected average conformational parameters of the triple helical Col domains can be calculated from the distribution of imino acids along their sequences [38]. The expected values are –106° for the average twist and 2.88 Å for the average height, same as those predicted for human fibrillar collagens [38]. Thus, despite the differences in amino acid composition and positional preference, the overall conformation of the triple helical Col domains is expected to be very similar to that of human fibrillar collagens.

Collagen sequences found in other viral proteins are more similar to those from EPclPs, although the preference for Pro in the X position is not that strong. Viral collagens share with EPclPs the low proportion of Pro residues in the Y position, large number of charged amino acids, and relatively common occurrence of Gln and Thr in the Y position. Collagens from gram-positive bacteria, which include the well-studied examples of *Bacillus anthracis* or *Streptococcus pyogenes* [22,24,39], show a lower presence of Pro residues in the X and Y positions, much lower proportion of charged amino acids, and a higher proportion of Ala residues in the X position and Gln and Thr in the Y position.

The main difference between human collagens and the three groups of non-animal collagens in Table 4 is the lack of preference for Pro in the Y position (as already noted in an earlier analysis of viral and bacterial collagen structural motifs [21]). Bacteria and viruses do not have the prolyl-hydroxylase enzymes required for hydroxylation of Pro in the Y position of a collagen triple helix, and therefore there is no contribution to the stability of their

### Table 1. Domains observed in collagen-like proteins from the genomes of EHEC strains.

| Domain | Names | InterPro | Pfam |
|--------|-------|----------|------|
| Col    | Collagen triple helix | IPR008160 | PF01391 |
| PCol   | Putative coiled-coil region (see text) |          |      |
| Pfn    | Phage fibre N (Phage_tail_N; Prophage tail N-terminal) | IPR013609 | PF08400 |
| Pfc    | Phage fibre C (Phage_fiber_C; Putative prophase tail fiber C-terminus) | IPR009640 | PF08620 |
| Pfc2   | Phage fibre C, variant 2 (see text) |          |      |
| P2     | Phage fibre 2 (Phage_fiber_2; Phage tail fiber repeat 2) | IPR005068 | PF03406 |

Currently available accession codes on the InterPro and Pfam databases are given. Domain architectures are shown in Figure 1.

### Table 2. Collagen-like proteins from the genomes of *E. coli* O157:H7 EDL933 and Sakai strains, and their corresponding prophage locations.

| Strain | Gene name/locus | Type | Prophage | Length (aa) | PCol length (aa) | Collagen lengths (aa) | Reference |
|--------|-----------------|------|----------|-------------|------------------|----------------------|-----------|
| O157:H7 EDL933 | Z0982 | EPcIA | CP-933K | 437 | 116 | 111 | [4] |
|         | Z1382 | EPcIA* | CP-933M | 437 | 116 | 111 |
|         | Z1483 | EPcIB | BP-933W | 645 | 109 | 157, 114 |
|         | Z2147 | EPcIA | CP-933O | 437 | 116 | 111 |
|         | Z2340 | EPcIA | CP-933R | 437 | 116 | 111 |
|         | Z3074 | EPcIA | CP-933U | 437 | 116 | 111 |
|         | Z3309, Z3307 | EPcIA* | CP-933V | 437 | 116 | 111 |
|         | Z6027 | EPcIA | CP-933P | 437 | 116 | 111 |
| O157:H7 Sakai | ECS0844 | EPcIA | Sp3 | 437 | 116 | 111 | [5] |
|         | ECS1123 | EPcIA | Sp4 | 437 | 116 | 111 |
|         | ECS1228 | EPcIB | Sp5 | 645 | 109 | 157, 114 |
|         | ECS1808 | EPcIA | Sp9 | 437 | 116 | 111 |
|         | ECS1992 | EPcIA | Sp10 | 437 | 116 | 111 |
|         | ECS2159, ECS2158 | EPcIA* | Sp11 | 407 | 116 | 81 |
|         | ECS2231 | EPcIA | Sp12 | 407 | 116 | 81 |
|         | ECS2717 | EPcIA | Sp14 | 437 | 116 | 111 |
|         | ECS2941 | EPcIA | Sp15 | 437 | 116 | 111 |

*Defective protein sequence, with frame-loss between domains. The original gene sequence may correspond to two open reading frames.

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collagen domains by Hyp residues. Collagen domains from EPclPs appear to compensate this lack of prolyl hydroxylation with a larger proportion of Pro residues in the X position. The high ratio of charged amino acids and the relatively high occurrence of Ala, Gln and Thr in EPclPs and bacterial and viral collagens may be indicative of different mechanisms for stability of their collagen domains [40,41,42].

Interestingly, the metazoan collagen sequence closest to EPclPs comes from a sea anemone, *Nematostella vectensis* (NCBI accession code XP_001625905, incomplete sequence), with 56% identity to the collagen domain of the EPclA protein ECs2717 and containing a repetitive [GP(Q/E)-GPK-GDT-GIT]_{12} sequence, reminiscent of the commonly observed triplet pattern mentioned above.

**A Low Complexity Region is Predicted as α-helical Coiled-coil Domain (PCoil)**

The region between the predicted PfN and Col domains in the most common architectures, EPclA and EPclB, shows an unusual low-complexity sequence with predominance of Ala (32%), Ser (19%) and Glu (13%) amino acids that often appear in tandems or in stretches of up to four consecutive identical residues (Figure 1). Different coiled-coil predicting algorithms (PCoils, Marcoil, Multi-Coil) give high scores for the region between residues 101 and 245 in both EPclA and EPclB (Figure 2). This region includes the low-complexity sequence between the PfN and Col domains plus the last 34 residues of PfN, and shows a loose seven-residue Ala-X-X-

### Table 3. Examples of collagen-like proteins from other *E. coli* and *Shigella* strains, and their prophage locations.

| Strain          | Gene name/locus | Type     | Phage/Prophage | Length (aa) | PCoil length (aa) | Collagen lengths (aa) | Reference |
|-----------------|-----------------|----------|----------------|-------------|-------------------|-----------------------|-----------|
| O157:H7 EC970520 | YYZ_gp70       | EPclA    | YYZ-2008       | 389         | 116               | 63                    | [80]      |
| O157:H7 EC508   | ECH7EC508_5575 | EPclA    | U              | 407         | 116               | 81                    | [81]      |
|                 | ECH7EC508_3568 | EPclA    | U              | 470         | 116               | 144                   |           |
| O26:H11         | ECO26_0887     | EPclA    | P02            | 404         | 116               | 78                    | [10]      |
|                 | ECO26_1634     | EPclA    | P06            | 440         | 116               | 114                   |           |
| O157:H7 Okayama | Stx2p020       | EPclB    | Stx2p-1        | 678         | 109               | 190, 114              | [82]      |
| ECOR-9          | ORF-401        | EPclC    | P-Eiba         | 479         |                   | 117, 75               | [83]      |
| O81 ED1a        | ECED1_1681     | EPclC    | U              | 493         |                   | 106, 102              | [84]      |
|                 | ECED1_2081     | EPclC    | U              | 520         |                   | 118, 117              |           |
|                 | ECED1_1138     | EPclC*   | U              | 566         | 95                | 48, 135               |           |
| S5989/EAEc      | EC55989_1078   | EPclC*   | U              | 467         | 116               | 140                   | [84]      |
| O86H–           | Stx2-86_gp21   | EPclD    | Stx2-86        | 673         | 88                | 322                   | [85]      |
| EC41008         | ECol_05072     | EPclD    | U              | 519         | 116               | 135                   | [86]      |
| *Shigella* dysenteriae | SDB_03138 | EPclD    | U              | 519         | 116               | 135                   | [87]      |
| *Shigella* boydii B5S12 | SbB5S12_E1096 | EPclD    | U              | 411         | 116               | 27                    | [88]      |

U: unassigned prophage.

*EPclC variant, containing a P Coil domain and either one or two collagen domains.

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### Table 4. Position-specific amino acid preferences in collagen triple-helical domains of EPclPs, human collagens, and collagen-like proteins from different groups of organisms.

| Collagen group | X                | Y                |
|----------------|------------------|------------------|
|                | Asp, Glu, Arg, Lys | Ala, Gln, Thr, Other |
| Pro            | 48.1             | 35.6             |
| Virus*         | 22.1             | 35.3             |
| Bacteria, gram positive | 31.0         | 14.1             |
| Human collagens |                  |                  |
| Fibrillar      | 31.1             | 18.5             |
| Non-fibrillar  | 24.6             | 19.7             |
| Human collagen-like proteins* | 27.5     | 21.6             |

The X and Y letters refer to the consensus sequence pattern (Gly-X-Y), characteristic of collagen triple-helical domains. The numbers indicate percentage occupation of the X or Y position by a given amino acid type.

*Excluding EPclPs from bacteriophages.

*Include molecules such as C1q, mannose binding proteins, collectins, macrophage scavenger receptors, or acetyl cholinesterase, which contain in their sequence a collagen domain but are not formally classified as collagen types.

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Collagens of Pathogenic *E. coli*
EPclA is a Trimeric Protein that Dissociates When Denatured

The quaternary structure of rEPclA was investigated by sedimentation equilibrium analytical ultracentrifugation (AUC) at increasing concentrations of guanidinium chloride (GuHCl) (Figure 3). The relative molar mass of rEPclA at 0 M GuHCl was 158±2.6 kDa, corresponding to the predicted molecular weight of a trimer of rEPclA molecules (3×47 kDa). As the concentration of GuHCl increased, a transition from trimer to monomer was observed and the relative molar mass of rEPclA at 5 M GuHCl was 43 kDa, which is consistent with the predicted molecular weight of the rEPclA monomer. Thus, rEPclA trimers dissociate into monomers as the GuHCl concentration increases; the trimer-to-monomer transition point was estimated at around 2.5 M GuHCl.

An independent measurement of the molecular weight of rEPclA was carried out by size exclusion chromatography followed by multilangle laser light scattering (SEC/MALLS) (Figure S1). The molecular weight obtained from MALLS is consistent with a trimer of rEPclA (Table 3). A proteolytic fragment from EPclA that included only the Col and PIC domains (Col–PIC fragment, Figures S2 and S4) could be produced in enough amounts for biophysical characterization. Analysis by SEC/MALLS of fractions containing the Col–PIC fragment (Figure S1) showed it to be trimeric as well (Table 5), with a molecular weight of 64 kDa consistent with three times the molecular weight of monomer Col–PIC (21–22 kDa, predicted from the peptide fingerprinting data obtained from mass spectrometry, Figure S2).

The molecular weights obtained from AUC and MALLS experiments were consistent with the predicted values for non-glycosylated rEPclA trimers and monomers. Lack of glycosylation of rEPclA was confirmed by periodic acid-Schiff staining analysis (data not shown).

Domains PCoil and PFC from EPclA are Trimerization Modules

Molecular weights of several recombinant fragments containing different combinations of domains were determined by SEC/MALLS (Figure S3 and Table 5). The data indicates that PIC is a trimerization domain, forming trimeric assemblies both when fused to a thioredoxin tag (Trx–PIC) or after removal of thioredoxin by thrombin digestion. The PIC–PCoil fragments were also trimeric, whereas the PFC domains were mainly in the monomer state (Figure S3). This data suggests that PCoil is also a trimerization domain.

EPclA Shows a CD Spectrum Consistent with Collagen and α-helical Conformations

The secondary structures of rEPclA and the Col–PIC fragment were analysed by circular dichroism (CD). Interpretation of the results is easier if the Col–PIC CD data is considered first. A sample of Col–PIC was purified from a preparation of full-length rEPclA by nickel-affinity and size exclusion chromatographies. Its CD spectrum was measured at different temperatures between 195 and 260 nm. The concentration of the Col–PIC sample was calculated as 0.2 mg/ml from its UV absorption at 280 nm and an estimated molar extinction coefficient ε = 11000 M⁻¹ cm⁻¹. The CD spectrum of Col–PIC at 4°C (Figure 4A) shows the characteristic features of triple-helical collagen: a small maximum of positive ellipticity at 220 nm and a deep minimum of negative ellipticity at around 199 nm [43]. Both these features are associated with the polyproline II conformation [44] characteristic of the collagen triple helix. The Col–PIC fragment includes mainly the collagen domain (Col) of EPclA and the C-terminal PIC domain, and thus its CD spectrum suggests that the Col domain adopts indeed a collagen-like, triple helical structure. The triple helical features disappeared from the CD spectrum upon increase of temperature, as shown by the CD curve at 55°C (Figure 4A). Interestingly, immediate cooling of the same sample back to 4°C recovered completely the triple helical structure, with a CD spectrum practically indistinguishable from the initial one (Figure 4A).

To examine the thermal denaturation of the collagen triple helix in the Col–PIC domain, the CD of another sample of purified Col–PIC fragment [also 0.2 mg/ml] was monitored at 220 nm as a function of continuously increasing temperature, from 4°C to 60°C. The thermal curve showed a single sharp transition at 42°C, which typically corresponds to the decrease of ellipticity at 220 nm and loss of collagen triple helical structure (Figure 4B).

The CD spectrum of rEPclA is different. A diluted sample of rEPclA was purified by nickel-affinity and size exclusion chromatographies and its CD spectrum was measured between 195 and 260 nm at 4°C (Figure 5A). The concentration of the rEPclA sample was calculated as 0.04 mg/ml from its UV absorption at 280 nm and a molar extinction coefficient ε = 17000 M⁻¹ cm⁻¹, deduced from the amino acid sequence of rEPclA. The CD spectrum showed two minima of negative ellipticity at 205 nm and 224 nm, the first one being deeper, and a small local maximum between the two minima, at 216 nm. To investigate this region in more detail a second sample with higher concentration, 0.3 mg/ml, was analyzed at different temperatures. When the sample was heated to 45°C, the height of the 216 nm maximum decreased significantly, the intensities of the two minima became more similar to each other, and their positions shifted to 210 nm and 222 nm, respectively (Figure 5A). This spectrum resembled more that of an α-helical coiled-coil conformation. Upon further increase of the temperature the overall ellipticity became less negative, and the two minima started to disappear and vanished completely when reaching 55°C (Figure 5A). The spectrum did not change upon further increase of temperature to 65°C. The slight decrease in ellipticity at 216 nm around 45°C and the more similar intensities of the two minima at that temperature suggest changes in the secondary structure that are consistent with the loss of the triple helical conformation in the Col domain while maintaining an α-helical conformation (Figures 4 and 5). Such α-helical structure appears to be more stable and does not disappear completely until 55°C. Immediate cooling of the same sample from 65°C back to 4°C recovered approximately half of the initial CD spectrum (Figure 5A).

To examine the thermal denaturation of rEPclA, the CD of another sample of purified rEPclA [concentration 0.3 mg/ml] was monitored at 216 nm as a function of increasing temperature from 20°C to 75°C. Two transitions were observed: a first transition at 42°C showed a sharp decrease in ellipticity, consistent with the loss of collagen triple-helical structure seen previously for the Col–PIC
Figure 2. Coiled-coiled predictions for the amino acid sequences of (A) EPcIA (ECs2717) and (B) EPcIB (ECs1228), using the PCoils [70] and Marcoil [71] algorithms. The graphs indicate regions of high probability for α-helical coiled-coil formation. Three different sequence window sizes were used with the PCoils algorithm: 14, 21 and 28 residues. Two different matrices were used in Marcoil: 9FAM, and MTK-based. doi:10.1371/journal.pone.0037872.g002
fragment at the same temperature; a second transition at 52°C showed a sharp and pronounced increase in ellipticity, corresponding to the loss of α-helical structure of the PCoil and PfN domains (Figure 5B). Thus, the α-helical structure of the PCoil and PfN domains is more stable than the collagen triple helix of the Col domain. The transition temperature of the Col domain is the same in rEPclA and its Col–PIC fragment, and seems unaffected by the presence of the PCoil and PfN domains. The melting transitions of the α-helical and collagen structures therefore appear to be largely independent.

The PfN–PCoil Region is Clearly α-helical and is Consistent with a Coiled-coil Structure

The secondary structures of the recombinant fragments PfN–PCoil and PfN were also studied by CD spectroscopy. Recombinant PfN–PCoil and PfN fragments were each purified with nickel-affinity and size-exclusion chromatographies. Concentrations of the PfN–PCoil and PfN samples were measured as 0.2 mg/ml and 0.3 mg/ml respectively, from their absorption at 280 nm and molar extinction coefficients ε = 7000 M⁻¹ cm⁻¹, initially calculated from the amino acid sequences of the PfN–PCoil and PfN recombinant fragments and adjusted using the observed UV absorption of samples in 8 M urea (see Materials and Methods). The CD spectrum of PfN–PCoil at 4°C in phosphate buffer (Figure 6A) shows the characteristic features of an α-helical protein, with two minima at 208 nm and 222 nm and a maximum at 195 nm. This spectrum is consistent with the prediction of an α-helical coiled-coil conformation for the PfN region. The α-helical features were still present in a spectrum measured at 45°C, although the signal intensities at the two minima started to decrease (data not shown). These features disappeared when reaching 60°C (Figure 6B), and were mostly recovered upon cooling the sample back to 20°C (data not shown). The spectrum of the PfN domain (Figure 6A) was also consistent with an α-helical structure but the intensity of the two minima at 208 nm and 222 nm was much lower than in the PfN–PCoil spectrum, indicating a lower α-helical content in this domain. This spectrum also vanished at 60°C (Figure 6C), but it was not recovered upon cooling back to 20°C (data not shown). Thus, the main contribution to the CD signal comes from the PCoil domain, most likely through the formation of a trimeric α-helical coiled-coil structure that disappears at a temperature of 60°C but is regained when the temperature is lowered again.

To examine the thermal transitions of the PfN–PCoil domain, the CD of two more samples of purified recombinant PfN–PCoil (0.35 mg/ml concentration) and PfN (0.1 mg/ml) were monitored at 222 nm as a function of continuously increasing temperature, from 5°C to 95°C and then cooling back to 5°C at the same speed (1°C per minute). The heating thermal curve showed a single sharp transition at around 49°C (Figure 7), corresponding to the loss of the strong α-helical CD spectrum. The cooling thermal curve also showed a single sharp transition at around 45°C, indicating partial re-gaining of the α-helical structure (the baseline in Figure 7 does not recover its initial value). This data suggests reversibility for the thermal transition of PfN–PCoil, mainly for the formation of the trimeric α-helical coiled-coil structure in the

Table 5. Average molecular weights of different recombinant fragments, calculated from the SEC/MALLS data.

| Molecule       | Average Mw (kDa) SEC/MALLS | Predicted Mw (kDa) from sequence | Oligomer state |
|----------------|---------------------------|-------------------------------|----------------|
| rEPclA         | 145                       | 47.3                          | Trimer         |
| Col–PIC        | 62                        | ~21                           | Trimer         |
| PfN–PCoil      | 88                        | 28.0                          | Trimer         |
| PIN            | 17                        | 17.2                          | Monomer        |
| Trx–PIC        | 64                        | 21.0                          | Trimer         |
| PfC            | 24                        | 7.2                           | Trimer         |

Domain compositions of each molecule, including protein fusion tags, are shown in Supplementary Figure S5.

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Figure 3. Analysis by analytical ultracentrifugation of the average molar mass of a sample of purified rEPclA as a function of increasing concentration of guanidinium chloride (GuHCl). Weight-averaged molar mass was determined using a single ideal species model (see Methods). Mean value masses for the upper and lower plateaux were 138±6 kDa and 43±1 kDa respectively (averages of three measures). The molecular mass of native rEPclA (0 M GuHCl) is consistent with three times that of denatured rEPclA (see text). The transition midpoint concentration is 2.38±0.09 M GuHCl.

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PCoil domain; the PfN domain does not recover completely after thermal denaturation and slow cooling. The actual value of the transition is sensitive to the heating or cooling rate. A repeat of the heating experiment with a slower speed (0.33 °C per minute) gave a transition temperature of 52 °C for the PfN–PCoil fragment (data not shown).

**Figure 4.** Far-UV CD analysis of the Col–PfC fragment after purification from rEPclA by SEC. (A) CD spectra at 4 °C, 55 °C and 4 °C after immediately cooling back the sample (see text). The vertical axis measures mean residue ellipticity θ in degrees cm² dmol⁻¹. The CD data was collected between 195 and 260 nm, with a protein concentration of 0.2 mg/ml in 10 mM Tris, 150 mM NaCl, pH 7.4. Measurements were taken in a 0.5 mm path length cell. (B) Thermal denaturation of the Col–PfC fragment, monitored by CD at 220 nm as a function of increasing temperature between 4 °C and 60 °C, with a protein concentration of 0.2 mg/ml in 10 mM Tris, 150 mM NaCl, pH 7.4, and a heating rate of 0.33 °C/min.

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**Figure 5.** Far-UV CD analysis of rEPclA after purification by SEC. (A) CD spectra at 4 °C, 45 °C, 55 °C and 4 °C after immediately cooling back the sample (see text). The vertical axis measures mean residue ellipticity θ in degrees cm² dmol⁻¹. The CD data was collected between 195 and 260 nm, with a protein concentration of 0.04 mg/ml (4 °C) or 0.3 mg/ml (the rest) in 10 mM Tris, 150 mM NaCl, pH 7.4. Measurements were taken in a 0.5 mm path length cell. (B) Thermal denaturation of rEPclA monitored by CD at 216 nm (the maximum between the two minima at 208 and 224 nm). The CD was measured as a function of increasing temperature between 20 °C and 75 °C, with a protein concentration of 0.3 mg/ml in 10 mM Tris, 150 mM NaCl, pH 7.4, and a heating rate of 0.33 °C/min.

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**Structural Organization of rEPclA and its Fragments Col–PfC and PfN–PCoil**

Full-length rEPclA was analyzed by rotary shadowing electron microscopy. Examination of the electron micrographs of rEPclA showed a “dumbbell-shaped” structure with two globular particles joined by a semi-flexible stalk, in which it is possible to distinguish two regions of different thickness (Figures 8 and 9). Sequence analysis of the P Coil region and the CD spectrum of PfN–PCoil
both suggest that the PCoil domain is a trimeric α-helical coiled-coil structure, and the observed thicker region of the stalk is consistent with such coiled-coil helical structure, which is known to have a larger cross-section than a collagen triple helix [45]. The remaining thinner region corresponds to the collagen triple-helical domain.

Inspection of rEPclA molecules from different micrographs suggests a hinge between the two regions of the stalk (collagen and coiled coil) that results in variable angles between the PCoil and Col domains and variable distances between the two globular domains (Figure 9). For most rEPclA molecules the distance between the centres of the PfN and PfC domains (N??C in Figure 9) has values around 50 nm, reaching 55 nm for the most extended, linear ones. The average N??C distance (48±5 nm) is probably less significant than the values shown by the extended molecules. The N-terminal globular structure, made of three PfN domains, has an approximate diameter of 8 nm and is slightly bigger than the C-terminal globular structure made of three PfC domains (approximate diameter 7 nm) (Figure 9). Taking into account the radii of the globular domains, rEPclA molecules can reach a length of 62 nm when totally extended. However, most rEPclA molecules appear slightly bent at the hinge between the PCoil and Col domains and are, overall, a bit shorter (55–60 nm, including the globular domains).

Rotary shadow electron micrographs of Col–PfC fragments showed molecules very reminiscent of those of rEPclA but with only one globular domain (PfC) connected to a stalk (Col) (Figure 10A,B). The stalk region appears slightly unravelled where the α-helical coiled coil and the PfN domain have been removed by endogenous proteolysis of full-length rEPclA. The observed morphology confirms the previous assignment of N- and C-terminal domains for rEPclA images and that the thin region of the stalk corresponds to the collagen triple-helical domain.

Rotary shadowing electron micrographs of PfN–PCoil fragments identified molecular shapes consistent with the N-terminal half of rEPclA molecules (Figure 10C,D). PfN domains (approximately the first 134 residues of EPclA) form a trimeric globular structure attached to an elongated stalk containing the PCoil domain. The micrographs show several instances of PfN–PCoil fragments apparently joined at the tails of their PCoil domains (Figure 10C). This association may result from some interaction between partially unravelled or unfolded chains at the terminal end of the PCoil domains.

From measures on the electron micrographs of the Col–PfC and PIN–PCoil fragments it is possible to estimate the lengths of the PCoil and Col regions as approximately 16 nm and 30 nm respectively (Figure 9). The length of the collagen domain is consistent with the predicted length of a collagen triple helix of 111 residues (111×2.9 Å = 32 nm, where 2.9 Å is an approximate measure of the height of an individual residue in a collagen triple helix [38]). Similar estimates can be obtained from measures on the thick and thin regions of the stalk in the rEPclA micrographs (14 nm for the PCcoil region and 28 nm for the Col domain). The slightly shorter values suggest some overlap between domains in

Figure 6. Far-UV CD spectra of the PIN–PCoil and PfN fragments after purification by SEC: (A) PIN and PIN–PCoil at 20°C; (B) PIN–PCoil at 20°C and 60°C; (C) PIN at 20°C and 60°C. In all panels the vertical axis measures mean residue ellipticity θ in degrees cm² dmol⁻¹. The CD data was collected between 195 and 260 nm, with protein concentrations of 0.2 mg/ml (PIN–PCoil) or 0.3 mg/ml (PIN), in 20 mM phosphate buffer (Na₂HPO₄/NaH₂PO₄), 100 mM NaCl, pH 7.4. Measurements were taken in a 0.5 mm path length cell. doi:10.1371/journal.pone.0037872.g006
the rEPclA stalk that cannot be resolved in the rotary shadowing micrographs. The apparent thickness of the P Coil and Col domains in the electron micrographs over-estimates the true cross-section dimensions of these domains, as it includes the thickness of the shadowing replica. Nevertheless, a slight difference is observed between these estimates for the P Coil (3.3±0.3 nm) and Col (2.7±0.3 nm) domains. Cross-section values seen in high-resolution micrographs of these domains are closer to 1.3–1.7 nm for a collagen triple helix and 2.2–2.7 nm for a trimeric α-helical coiled coil.

Structural Organization of rEPclB High-molecular Weight Aggregates

Expression of rEPclB both by IPTG induction or auto-induction produced a good yield of soluble protein. However, purification of rEPclB from the soluble fraction by nickel-affinity and size exclusion chromatography showed that all the protein went to form soluble, high-molecular weight aggregates that eluted in the void volume of the size exclusion columns (data not shown). The protein aggregated to such an extent that it was not possible to identify any additional peak or shoulder suitable for molecular mass determination by MALLS. Aggregation was worse in samples produced by auto-induction (presumably due to the increased protein production). Attempts to reduce the degree of aggregation by lowering the protein concentration, adding EDTA (to rule out His6-mediated metal chelation aggregation), adding glucose in high concentration, or changing the ionic strength of the buffers, were all unsuccessful: it was not possible to obtain enough non-aggregated rEPclB for molecular weight determination or for CD studies. Interestingly, the high molecular weight aggregates were soluble and the protein did not precipitate out of solution, even after high-speed centrifugation. The bands observed in SDS-PAGE experiments suggest that SDS treatment extracts monomeric rEPclB from the high molecular weight aggregates.

To investigate a possible structural organization of these aggregates, a sample of IPTG-induced rEPclB was used for rotary shadowing electron microscopy. The sample contained exclusively high-molecular weight aggregates that appeared in the electron micrographs as large masses of protein that, nevertheless, appeared to have a relatively narrow size distribution (300–500 nm in diameter, data not shown). Close inspection of the smallest aggregates (probably at an early stage of formation) revealed an internal structure that could be reconciled with entangled, multiple flexible linear beaded molecules (Figure 11). In the vicinity of these aggregates it was possible to discover individual features reminiscent of the rEPclA molecular morphology, but with three globular “domains” instead of two, connected by two flexible stalks (Figure 11). The terminal globular structures would correspond to the PNI and PIC domains, and the internal one would include the PIC domain. The flexible stalks would correspond to the two Col domains and the PCol domain predicted in the rEPclB sequence. All these structures are presumed to be trimeric due to the presence of the PIC, Col and P Coil domains, all shown to trimerize in rEPclA. The structural organization for rEPclB would therefore be similar to that seen for rEPclA. However, an effective protocol to increase the amount of non-aggregated protein will be necessary to demonstrate these assumptions and to properly characterize the molecular architecture of rEPclB (work in progress).

Discussion

Multiple open reading frames with collagen-like amino acid sequences have been identified automatically in the genomes of several EHEC strains. These open reading frames are incorporated in the sequence regions of prophage and prophage-like...
elements embedded in these EHEC genomes (Tables 2 and 3) and presumably code for proteins involved in phage morphogenesis. Two recombinant proteins rEPclA and rEPclB, representative of the most common domain architectures EPclA and EPclB (Figure 1), were amplified from a sample of genomic DNA from the O157:H7 Sakai strain, cloned into appropriate protein expression vectors, and the resulting recombinant proteins analyzed biochemically. The aims were to demonstrate their biochemical viability and structural integrity, to confirm the presence of molecular characteristics typical of collagen-like proteins, to investigate their quaternary structure, conformation, morphology and thermal stability, and to analyze some of their individual domains.

Both rEPclA and rEPclB are produced as soluble proteins in E. coli, although rEPclB has a strong tendency to form large, soluble aggregates. Thus, most of the biochemical analysis has been done on rEPclA. Our data demonstrate that EPclA shows the main characteristics of collagen-like proteins: it forms stable trimers in solution that dissociate upon denaturation (Figure 3), and its collagen-like sequence adopts a collagen triple helical conformation, as demonstrated by CD spectroscopy (Figure 4). These data confirm that the collagen-like sequence (Col) of EPclA is a true collagen domain, and strongly suggests that collagen-like sequences in other EPclPs will adopt the same conformation.

The molecular morphology of rEPclA has been visualized by rotary shadowing electron microscopy. rEPclA is a trimeric dumbbell-shaped molecule, with two globular domains joined by a semi-flexible “stalk”, or rod-shaped domain (Figures 8–9). This connecting stalk is made in fact of two triple-helical domains: the collagen triple-helical domain (Col) and a trimeric α-helical coiled coil (PCoil) encompassing the region between the Col domain and the N-terminal PfN domains. The CD spectrum of rEPclA (Figure 5) is largely dominated by the combination of CD spectra from an α-helical coiled coil and a collagen triple helix. Coiled-coil prediction algorithms give high scores to the last 30 amino acids of the PfN domain and to the region between the PfN and Col domains, both for EPclA and EPclB (Figure 2). Our data indicates that, at least in EPclA, the region between PfN and Col forms indeed a trimeric α-helical coiled coil, as demonstrated by SEC/MALLS (Figure S3) and CD spectroscopy (Figure 6). We predict that similar PCoil domains in EPclB and other EPclPs will also form trimeric α-helical coiled coils. The combination of collagen triple helices adjacent to trimeric coiled coils is not unusual and there are many proteins for which such structural arrangement is predicted [45,46], where the α-helical coiled coil functions mainly as an oligomerization domain.

The thermal stability of the Col domain of EPclA is higher than expected when compared to eukaryotic collagens, especially after considering the lack of prolyl hydroxylation (discussed below). The thermal denaturation data of rEPclA (Figures 4, 5, 6, 7) shows two sharp transitions, a first one at 42°C and a second one at 52°C (Figure 5). These two transitions correspond respectively to the loss
of the collagen triple helix and the loss of the \( \alpha \)-helical coiled coil. The sharpness of both melting curves indicates highly cooperative transitions for each case (including local chain dissociation as a result of conformational change). Nevertheless, EPclA will remain trimeric at temperatures between the melting of its Col (collagen triple helical) and PCoil (\( \alpha \)-helical) domains. Identical transition temperatures are observed separately in two fragments: Col–PfC shows a single transition at 42 \( ^\circ \)C, and PfN–PCoil shows a single transition at around 50 \( ^\circ \)C, both consistent with the loss of conformation of their helical domains. The coincidence in

**Figure 9.** Molecular dimensions of \( \epsilon \)EPclA and its domains. (A) Magnified view of a representative \( \epsilon \)EPclA molecule from a rotary shadowing electron micrograph. (B) The same molecule with the background masked out showing the different molecular dimensions analyzed below. (C) Average dimensions obtained from multiple measures on electron micrographs: N–C and \(<\text{Col}–\text{Col}\) are averages of 224 measures on eight \( \epsilon \)EPclA micrographs; \( D_N \), \( L_{\text{Col}} \) and \( T_{\text{Col}} \) are averages of 76, 35 and 74 measures, respectively, on three Col–PfC micrographs (Figure 10A); \( D_N \), \( L_{\text{Col}} \) and \( T_{\text{Col}} \) are averages of 200 measures on one PfN–PCoil micrograph (Figure 10B). (D) Histograms showing the distribution of N–C and \(<\text{Col}–\text{Col}\) values on the sample of 224 \( \epsilon \)EPclA molecules.
transition temperatures with those of full-length rEPclA suggests that the two transitions are essentially independent of each other, and that the presence or absence of either the Col or PCoil domains does not change the thermal stability of the other one.

The thermal stability of the Col domain of EPclA appears remarkable. To remain functional at body temperatures, mammalian collagens have a high proportion of imino acids in their collagen domains (>20% in humans) and require a complicated mechanism of post-translational hydroxylation of proline residues [47,48,49]. The enzymes required for prolyl hydroxylation are not present in E. coli and yet, the melting temperature of EPclA Col domain (42°C) is higher than that of the much longer mammalian collagens (~37°C, around body temperature [50]). Glycosylation of threonine residues, another mechanism of collagen stabilisation [40], is not observed in rEPclA either. Thus, the higher than expected thermal stability of the Col domain of EPclA could be the consequence of an unusually high proportion of Pro residues in the X position (Table 4), a high proportion of charged residues Arg, Asp, Glu and Lys (Table 4), and possibly a stabilizing effect by the PfC domains (see below). A stabilization mechanism based on a high proportion of charged residues has been proposed for the collagen triple helical domain of S. pyogenes collagen-like protein Scl2 [42], but the overall proportion of charged amino acids in the collagen domain of Scl2 (30%) is higher than the overall proportion of charged amino acids in the collagen domains of EPclPs (22%).

The thermal stability of the PCoil domain in EPclA is even higher. It is currently unknown if these high transition temperatures for the Col and PCoil domains have any functional significance. As EPclPs are likely to participate in phage morphogenesis, such high thermal stability may be required to ensure appropriate assembly of the phage particles during prophage induction from EHEC, while inside the host intestine, or to facilitate survival of free phages in the extraintestinal environment.

Electron micrographs of rEPclA show a degree of variable bending in the connecting stalk (Figure 9) that suggests the presence of a “molecular hinge” between the PCoil and Col domains. This hinge could be simply an area of structural discontinuity and increased flexibility between the collagen triple helix and the three-stranded α-helical coiled coil. Some discontinuity is expected: the transition between the two helical domains needs to account for a change in axial chain staggering, from three chains in register in the α-helical coiled coil to the one-residue stagger characteristic of the collagen triple helix [45]. It is also likely that the PCoil to Col transition changes its superhelical handedness, from a right-handed collagen superhelix to the most common left-handed coiled-coil superhelix. Although right-handed coiled coils have been described, they show undecad (11) or pentadecad (15) residue periodicity instead of the canonical heptad (7) periodicity of left-handed coiled coils [51,52]. The sequence of the PCoil domain is unusual for an α-helical coiled coil, almost devoid of hydrophobic residues like leucine or isoleucine and with
Figure 11. Rotary shadowing electron microscopy of rEPclB. (A) Internal structure of a small aggregate of rEPclB molecules. The micrograph suggests multiple flexible molecules, reminiscent of those observed for rEPclA, with dark globular structures (presumably globular domains of PfN, PfC and Pf2), and poorly defined linear structures (presumably stalks containing the PCoil and Col domains). The rEPclB molecules seem to aggregate heavily through one of the globular domains. (B) Possible examples of individual rEPclB molecules observed, isolated from the large aggregates, in some electron micrographs. (C) Interpretation of the observed in terms of three globular domains (PfN, Pf2 and PfC) connected by two stalk regions. Approximate molecular dimensions are shown for comparison purposes with rEPclA.

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The rapid reversibility of Col–PfC thermal denaturation is consistent with a degree of recombination in both domains (Figure 9). Thus, both the PIC and PCoil domains can be considered trimerization modules in EPclPs, and in the case of PIC, it may contribute in part to the high melting temperature of the Col domain.

Recent studies on collagen-like proteins of some pathogenic bacteria have revealed a variety of functions for these proteins, including binding extracellular matrix proteins or adherence to mammalian cells [25,27,28]. It is not currently known if EPclPs have any functional role for E. coli itself, or they are strictly phage morphogenetic proteins. Currently available evidence of EPclP expression in EHEC seems to be linked to phage induction [29,30,31], and the presence of common domains in sequences of EPclPs and side tail fibre proteins of lambda phages is a strong indicator of EPclPs as structural proteins in tail fibres from phages. The high thermal stability of EPclPs is also consistent with such a role.

All prophages and phages containing EPclP sequences are described as Colavirales, or tailed bacteriophages. Bacteriophage 933W, isolated from the O157:H7 strain EDL933, was amongst the first EPclP-containing phages to be studied [53,54,55,56]. Particles of 933W observed under the electron microscope show regular hexagonal heads (probably icosahedral), about 70 nm wide, and short tails 22–28 nm long and 13–17 nm wide [55,56]. Often, 933W virions clump together through some form of tail-tail interaction [56]. Phage induction from EHEC strains results in virion particles with different morphologies [12,34,57,58], that are usually classified as members of the Podoviridae or Siphoviridae families, with short or long non-contractile tails, respectively. Phages of these families often show tail fibres extending laterally from the sides of the tails (Figure 12), although none of the published electron micrographs of phages from EHEC strains has sufficient detail for their visualization. Tail fibres are often used for the phage particles to attach to the target cells, and this attachment triggers further events leading to infection of the viral DNA in the host cell periplasm.

The presence of PIN and PI2 domains both in EPclP sequences and in side tail fibre proteins of λ bacteriophages [35,36] strongly suggests (although do not absolutely prove) that EPclPs are major components of prophage side tail fibres and probably participate in adhesion to the E. coli cell surface, either directly or through assembly with other prophage tail fibre proteins. At least one instance of direct interaction has been proposed, between EPclB proteins of short-tailed Shiga toxin-carrying phages and the conserved E. coli protein YacT. The study, which refers to EPclB as tail-spine protein, concludes that YacT is the surface molecule recognized by the majority of these phages [59].

The use of the collagen triple helix by bacteriophages to build trimeric fibrillar proteins is remarkable. Trimerization is a highly prevalent characteristic of bacteriophage tail proteins and adhesins [60,61] and novel trimeric folds have been discovered in bacteriophage fibre proteins [62,63,64,65]. The α-helical coiled coil is present in bacteriophage fibre proteins such as fibrin [66] and our data confirms that phages have added collagen helices to their armoury of trimeric folds, in combination with α-helical coiled coils and other capping domains like PIN and PIC, the latest being a trimerization module itself. Furthermore, similarity with other viral fibrous tail proteins would suggest that the PIN domain contains the viral attachment site and the PIC domain is involved in binding to E. coli cell surface proteins [63]. The modular nature of EPclPs and other side tail fibre proteins, with different combinations of the same domains present in both closely and distantly related genomes, is consistent with a degree of recombina-
nation between multiple prophages on the same bacterial genome that often results in novel phages with an expanded host range and in new bacterial strains [34]. Prophage recombination and the efficiency of bacteriophages as HGT vehicles are responsible in part of the heterogeneity amongst EHEC strains and their rapid evolution. Thus, understanding the mechanisms of phage morphogenesis and phage interaction with the E. coli or EHEC cell surfaces may be more important than previously thought and deserve further investigation.

Materials and Methods

Sequence Retrieval and Analysis

Sequences of prophage collagen-like proteins from EHEC strains (EPcPs) were retrieved from the UniProt database [67]. These sequences were classified into different domain architectures (EPcA, EPcB, EPcIC and EPcID, Figure 1) that were defined according to the occurrence and relative location of several conserved non-collagenous domains described in the InterPro [68] and Pfam databases [69] (Table 1). The probability of coiled-coil conformation and its oligomerization state in EPcIP sequences were calculated with different prediction algorithms: PColis [70], Marcoil [71], MultiCoil [72] and SCORER 2.0 [73]. Secondary structure predictions for EPcIP sequences were obtained from the Jpred3 prediction server [74]. Default settings were used for all prediction algorithms.

Cloning of EPcIA and EPcIB Sequences from E. coli O157:H7

Recombinant EPcIA and EPcIB proteins were produced using laboratory E. coli strains. Two DNA fragments coding for EPcIA and EPcIB were amplified from a sample of genomic DNA of Escherichia coli O157:H7 strain RIMD 0509952 (Sakai), which was a gift from Charles W. Penn (School of Biosciences, University of Birmingham, UK). Forward and reverse primers were designed from the nucleotide sequences of the open reading frames ECs2717 (EPcIA) and ECs1228 (EPcIB) of the E. coli Sakai strain genome [5], with accession numbers Q7ACX5 and Q8XAX7 (Uniprot), or NP_310744 and NP_309225 (NCBI), respectively. Appropriate NdeI and XhoI restriction sites were incorporated into the primer designs, with the resulting oligonucleotide sequences 5’-CAT ATG ATG GCA GTA AAG ATT TCA GGT GTA CTG-3’ (EpclA forward), 5’-CTC GTC GAG TCC TGT TCT GCC TGT ACT GCC-3’ (EpclA reverse), 5’-CAT ATG ATG ACG AAT GAT CAT GGC GGG GAT TAT GCG-3’ (EpclB forward), and 5’-CTC GAG TCA TCC TGT TCT GCC TGT ACT AC-3’ (EpclB reverse). The ECs2717 sequence was chosen amongst the six EPcIA open reading frames from the Sakai genome as it had a putative promoter sequence (TGTTATGAC) 38 nucleotides upstream of the predicted start of the coding sequence. The products of PCR amplification were ligated into the pET-28a+ (expression vector (Novagen)) and the correct frame and ligation of the clones were confirmed by sequencing. The PIC construct failed to produce any detectable expression and the PIC sequence was subsequently cloned into the fusion protein expression vector pHisTrx, an in house derivative of pET-32a (Novagen) encoding E. coli thioredoxin (txr) with an N-terminal hexahistidine tag and a thrombin cleavage site, followed by a unique multiple cloning site [75] (Figure S4).

Expression and Purification of Recombinant Proteins

Recombinant proteins and fragments were produced in E. coli BL21(DE3) cells (rEPcIA and rEPcB) or JM109(DE3) cells (rPN–Pcoil, rPN and Trx–rPfC), using both IPTG induction (small-scale test expression and large-scale expression) and auto-induction (only large-scale expression) [76]. Best expression conditions for IPTG induction were achieved by inoculating 5 ml cultures with single bacterial colonies expressing the recombinant proteins, followed by incubation at 37°C overnight, and then inoculating 500 ml cultures in 2-litre flasks with 1% overnight pre-culture, followed by incubation at 37°C until log phase, and induction with 1 mM IPTG followed by further incubation for 4 hours at 30°C. Best expression using auto-induction was achieved by inoculation with 1% overnight pre-culture of 500 ml cultures supplemented with auto-induction solutions (1 M each of Na2HPO4, KH2PO4, NH4Cl, Na2SO4, and MgSO4, 20 mM CaCl2, 50% glycerol, 1 M glucose and 20% lactose) followed by incubation overnight at 37°C. All recombinant products mainly localised to the soluble fraction, auto-induction expression being more effective in producing larger amounts of soluble protein. Recombinant proteins were purified by nickel-affinity chromatography (QIAGEN) using a 5 ml column and following the manufacturer protocols, followed by size-exclusion chromatography using a 120-ml HiLoad 16/60 Superdex 200 column (GE Healthcare), with 10 mM Tris, 150 mM NaCl, pH 7.4 as elution buffer. SDS-PAGE analysis showed that the bands corresponding to rEPcIA and rEPcB migrated with apparent molecular weights of ~66 kDa and ~100 kDa, respectively (Figure S2 and data not shown), which are higher than their predicted molecular weights of 47 kDa and 66 kDa. Glycosylation was ruled out by a negative periodic acid-Schiff stain analysis (not shown), and thus the anomalous gel migration of purified rEPcIA or rEPcB must relate to their collagen-like sequences, a behaviour commonly observed in collagens and collagen-like proteins. Overproduction of rEPcIA
by auto-induction produced significant amounts of two endo-
genous proteolytic fragments whose partial sequences were
identified by peptide mass spectrometry (see below and Figure
S2). A fragment containing only the Col and PCiC domains from
\( \varepsilon \)EPclA (Col–PCiC fragment, Figures S2 and S4) could be isolated
in enough quantities from full-length \( \varepsilon \)EPclA for subsequent
biophysical characterisation. Conditions for consistent production
of the two proteolytic fragments could not be established, and
most samples of \( \varepsilon \)EPclA produced by auto-induction and IPTG
induction did show one major band in SDS-PAGE analyses of
purified samples, corresponding to full length \( \varepsilon \)EPclA (Figure
S2A). In these cases, levels of the proteolytic fragments were too
low for characterization and isolation.

**Gel electrophoresis, Immunoblot Analysis and Protein
Identification**

Protein samples were denatured by heating at 90°C for 10
minutes and electrophoretically separated in 0.1% SDS, 4–12% gradient NuPAGE Bis-Tris polyacrylamide gels (Invitrogen),
Proteins were visualised via Coomassie Brilliant Blue staining,
and their electrophoretic migrations compared to those of
prestained molecular weight markers (Precision Plus All Blue
standards, BioRad). All proteins and fragments containing the
Col or PCoil domains showed a slower than normal electropho-
retic migration and higher than expected apparent molecular
weights in the gels. Thus, identities of individual protein bands
had to be confirmed by in-gel trypsin digestion followed by
reverse phase chromatography of the tryptic peptides and
sequence identification with mass spectrometry in the Biomolec-
ular Analysis Facility of the Faculty of Life Sciences, University
of Manchester (BAF-FLS). Bands of interest were excised from the
gels, reduced with 10 mM dithiothreitol and alkylated with
55 mM iodoacetamide. Samples were digested overnight with
trypsin at 37°C and then analysed using a CapLC (Waters)
nanoLC system coupled to a Q-TOF Micro Mass Spectrometer
(Waters). Peptides were separated by reverse phase chromatog-
raphy using a 0.075×150 mm PepMap column (Dionex) and an
acetonitrile gradient in 0.1% formic acid. Peptides eluting from the
column were selected automatically for fragmentation. Data
were searched against the UniProt database using the Mascot
engine (Matrix Science). Further confirmation by immunoblot-
ing was performed via transfer of SDS-PAGE separated proteins
onto nitrocellulose membranes (Whatman) followed by incuba-
tion with the appropriate primary antibody and horseradish
peroxidase-coupled Fab-specific anti-mouse IgG (Sigma) was
used as secondary antibody, and detection was achieved with
SuperSignal West Pico Chemiluminescent Substrate (Thermo
Scientific).

**Protein Concentration**

Protein concentration was determined by measuring UV
absorption at 280 nm with a NanoDrop ND-1000 spectropho-
tometer (Labtech International), using molar extinction coeffi-
cients calculated from the amino acid sequences of the different
recombinant proteins and fragments [77]. For recombinant
fragments without tryptophan residues in their sequences, a
corrected value of the molar extinction coefficient was derived
from the UV absorption at 280 nm of equal concentration samples
in aqueous buffers and in 8 M urea. In these cases the extinction
coefficient calculated from sequence was considered accurate
enough for determination of the protein concentration in 8 M
urea [77], and an adjusted extinction coefficient was derived for
the protein in water-based buffers.

**Analytical Ultracentrifugation**

Sedimentation equilibrium analysis of \( \varepsilon \)EPclA was performed
using an Optima XL-A ultracentrifuge (Beckman Instruments)
from the BAF-FLS. The protein was in a 10 mM Tris, 150 mM
NaCl, pH 7.4 buffer that was supplemented with increasing
concentrations of guanidinium chloride (0–6 M). Sample volumes
of 110 μl were used in six-sector Epon-filled centrepiece cells
equipped with quartz windows. All experiments were conducted at
20°C and sedimentation equilibrium data were collected at
10,000, 18,000 and 28,000 rpm. Weight-averaged molar mass
\( (M_w) \) was determined using *Hetero* analysis (developed by J. Cole
and J. Lary at the University of Connecticut), using a single ideal
species model. A value of \( r = 0.7139 \) was used for the partial
specific volume of \( \varepsilon \)EPclA, calculated from its amino acid sequence
using Sednterp, version 1.09.

**Size Exclusion Chromatography and Multiangle Laser-
Light Scattering**

Molecular weights of the recombinant proteins were determined
by size exclusion chromatography coupled to multiangle laser-light
scattering (SEC/MALLS) in the BAF-FLS. Depending on their
size, proteins were chromatographed in Superdex 200 10/300 GL
(10 to 600 kDa), Superose 6 10/300 GL (5 to 3000 kDa), or
Superdex 75 5/150 GL (5 to 70 kDa) columns, all from GE
Healthcare, using 10 mM Tris pH 7.4, 150 mM NaCl, 1 mM
EDTA as elution buffer. Elution from the column was continu-
ously analysed in-line with a light scattering (LS) detector Dawn
Helios II (Wyatt Technology) and an Optilab rEX refractometer
(Wyatt Technology). The LS intensity and eluant RI were
analyzed using ASTRA software (version 5.21, Wyatt) to give a
weight-averaged molar mass \( (M_w) \). Fractions of 0.5 ml were
collected for further analysis.

**Circular Dichroism (CD) Spectroscopy**

Secondary structures and thermal denaturation of the recom-
inant proteins were analyzed by CD spectroscopy. Samples were
eluted in 10 mM Tris, 150 mM NaCl, pH 7.4 and purified
via size-exclusion chromatography before analysis. For the PNP-
PCoil and PIN recombinant proteins a phosphate buffer with
20 mM Na2HPO4/NaH2PO4, 100 mM NaCl pH 7.4, was also
used for CD analysis. CD spectra were recorded with a Jasco J-810
spectrometer equipped with a peltier temperature controller.
Wavelength scans were performed using a 0.5 mm-path-length
Starna quartz cell of acceptable birefringence for CD, and data
were collected every 0.5 nm with a 1 nm bandwidth. Each
spectrum was obtained from the accumulation of data from 10
scans; baseline was corrected using the spectrum of a Tris/NaCl
buffer blank. Spectra were obtained at different temperatures, and
thermal transition profiles were recorded between 4°C and 90°C
at 220, 216 or 222 nm (depending on the protein being examined)
with a data pitch of 0.5 nm, bandwidth of 1 nm, detector response
time of 32 sec and temperature slope of 20°C/hr. Samples were
cooled back to 4°C after the different transitions and final spectra
were recorded at that temperature. Ellipticities in millidegrees
were converted to mean residue molar ellipticities (degree cm²
dmol⁻¹) by normalizing for the number of residues on each protein
or fragment.

**Rotary Shadowing Electron Microscopy**

The structural organization of the different recombinant
proteins was investigated under an electron microscope after
rotary shadowing using the mica sandwich technique [78]. Five μl
of sample (5 μg/ml) were adsorbed onto freshly cleaved mica.
Another freshly cleaved mica disc was then gently placed on top, causing the drop to spread evenly between the two mica discs. The sample was allowed to absorb for 1 min, after which it was washed with water and then freeze-dried at ~80°C in a Cressington CVE-50 freeze-fracture apparatus (Cressington Scientific Instruments). The dried sample was maintained under vacuum and its temperature was dropped to ~190°C. Rotary shadowing was then performed at an angle of 5° relative to the mica surface by electron-beam evaporation of platinum-carbon. The platinum film thickness in the plane of the substrate was measured by a quartz-crystal film-thickness monitor as 1.8 Å; this film was subsequently coated with a 7 nm carbon backing layer. Replicas were floated from mica on distilled water and placed on 400 mesh copper grids. Photomicrographs were taken with an FEI Tecnai Twin transmission electron microscope (FEI, Eindhoven, Netherlands) operated at 120 kV. Digital images (1024×1024 pixels) were recorded on a TVIPS F214 cooled CCD camera. Magnification was calibrated using a diffraction grating replica (2160 lines/min; Agar Scientific, Stansted, UK).

Supporting Information

Figure S1 Analysis of rEPclA and its Col-PIC fragment by SEC/MALLS. (A) Chromatogram showing the elution of nickel-affinity purified rEPclA from a Superose 6 10/300 GL size exclusion column; the red trace corresponds to the light scattering detector and the green trace to the UV absorption detector, both in arbitrary units. Peak 1 corresponds to the void volume and contains high molecular aggregates; peak 2 corresponds to native rEPclA. (B) Molar mass distribution or native rEPclA (peak 2 in A) measured by light scattering. The blue trace corresponds to the refractive index detector (in arbitrary units) and the dashed black line shows the weight-average molecular mass for each slice, as measured by the light scattering detector. The molar mass distribution is consistent with trimeric rEPclA (Table 5). (C) Chromatogram showing the elution of a nickel-affinity purified auto-induction sample of rEPclA from a Superdex 200 10/300 GL size exclusion column (traces as in panel A). Peak 1 corresponds to the void volume and contains high molecular aggregates; peaks 2 and 3 show molar mass distributions consistent with trimeric rEPclA and trimeric Col-PIC fragment, respectively (Table 5); peak 4 is consistent with monomeric rEPclA. (D) Molar mass distribution of peak 3 from C (Col-PIC) re-chromatographed in the same Superdex 200 column. The blue trace corresponds to the refractive index detector (arbitrary units) and the dashed black line shows the weight-average molecular mass for each slice, as measured by the light scattering detector. The molar mass distribution is consistent with trimeric Col-PIC (Table 5).

Figure S2 Large-scale expression of rEPclA: SDS-PAGE analysis of the different fractions after purification by nickel affinity chromatography. Individual peptides identified by mass spectrometry on each protein band are shown in red against the original EPclA sequence. (A) Overexpression of rEPclA by IPTG induction. Lane 1: molecular weight markers; lane 2: flow-through; lanes 3–4: fractions eluted with 5 mM and 100 mM imidazole (washed); lanes 5–10: fractions eluted with 1 M imidazole. The overexpressed band of rEPclA, confirmed by mass spectrometry, shows an apparent molecular weight of ~66 kDa (higher than the true molecular weight of 47 kDa). (B) Overexpression of rEPclA by auto-induction. Lane 1: molecular weight markers; lanes 2–10: fractions eluted with 500 mM imidazole. The rEPclA band runs at ~66 kDa, also confirmed by mass spectrometry. Two additional protein bands were identified by mass spectrometry as endogenous proteolytic fragments of rEPclA fragments. The mapped peptides reveal the extent and domain composition of each fragment. The band corresponding to the Col-PIC fragment shows an apparent molecular weight of ~80 kDa (higher than the predicted molecular weight of ~21 kDa). Another band at ~60 kDa seems to correspond to a fragment with a partial digestion of the PIN domain and including the Pcoil–Col–PIC domains.

Figure S3 Analysis of PIN, PIN–Pcoil and Trx–PIC fragments by SEC/MALLS. (A) Chromatogram showing the elution of nickel-affinity purified PIN–Pcoil fragment (green trace) or PIN fragment (red trace), from a Superdex 200 10/300 GL size exclusion column. Both traces correspond to the UV absorption detector, in arbitrary units. The dashed green and red lines show weight-average molecular masses for each slice of peaks 1 to 4, as measured by the light scattering detector. Peaks 1 and 2 correspond to trimeric and monomeric PIN–Pcoil fragment, respectively, whereas peaks 3 and 4 correspond to trimeric and monomeric PIN. Molar mass distributions on each peak are consistent with these oligomerization states (Table 5). The predominant species in the PIN–Pcoil sample is the trimer (peak 1), but a small amount of monomer (peak 2) can be detected. For PIN the predominant species is the monomer (peak 4), but a small amount of trimer (peak 3) can be detected. Elution volumes appear to be non-linear between the two purifications as the PIN–Pcoil monomer elutes at a lower volume than the PIN trimer. (B) Molar mass distribution or the Trx–PIC fragment, measured by light scattering. The blue trace corresponds to the refractive index detector (in arbitrary units) and the dashed black line shows the weight-average molecular mass for each slice, as measured by the light scattering detector. The molar mass distribution is consistent with trimeric Trx–PIC (Table 5).

Figure S4 Domain architecture of the different recombinant proteins and constructs used in this study. Key to domain labels: PIN, phage fibre N-terminal domain; Pcoil, phage coil domain; Col, collagen domain; PIC, phage fibre C-terminal domain; H, hexahistidine tag; Trx, thioredoxin tag.

Figure S5 Nucleotide and amino acid sequences of rEPclA from DNA sequencing of the product amplified from a sample of genomic DNA from E. coli O157:H7 Sakai and cloned into a pET-28a(+) expression vector (see Methods). Sequence colour code: red, PIN domain; orange, Pcoil domain; green, Col domain; blue, PIC domain; black, additional amino acids introduced by cloning to the protein expression vector, including N-terminal and C-terminal hexahistidine tags and a thrombin cleavage site preceding the PIN domain. Twelve nucleotide changes with respect to the most similar deposited EPclA sequence (ECs2717) are highlighted in yellow. Of those, eight are silent and four lead to changes in the amino acid sequence, also highlighted in yellow. All these amino acid changes correspond to normal sequence variability amongst EPclA sequences from different O157:H7 strains.

Author Contributions

Conceived and designed the experiments: NG IR JB. Performed the experiments: NG TJM TAJ MH HD DFH. Analyzed the data: NG TJM TAJ MH HD IR JB. Wrote the paper: NG JB.
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