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Review

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# Membrane assembly of the cholesterol-dependent cytolysin pore complex $\stackrel{ m >}{\sim}$

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#### ABSTRACT

The cholesterol-dependent cytolysins (CDCs) are a large family of pore-forming toxins that are produced, secreted and contribute to the pathogenesis of many species of Gram-positive bacteria. The assembly of the CDC pore-forming complex has been under intense study for the past 20 years. These studies have revealed a molecular mechanism of pore formation that exhibits many novel features. The CDCs form large  $\beta$ -barrel pore complexes that are assembled from 35 to 40 soluble CDC monomers. Pore formation is dependent on the presence of membrane cholesterol, which functions as the receptor for most CDCs. Cholesterol binding initiates significant secondary and tertiary structural changes in the monomers, which lead to the assembly of a large membrane embedded  $\beta$ -barrel pore complex. This review will focus on the molecular mechanism of pore forming proteins. This article is part of a Special Issue entitled: Protein Folding in Membranes.

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#### 1. Introduction

The cholesterol dependent cytolysins (CDCs) are produced by many species from the *Clostridium*, *Streptococcus*, *Listeria*, *Arcanobacterium*, *Gardnerella*, *Bacillus* and *Lactobacillus* genera. It is likely that additional

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species that express a CDC will be discovered. The CDC gene is probably one of the most widely disseminated toxin genes known, suggesting that many bacterial species find it useful to introduce large pores in cholesterol containing eukaryotic membranes. How these bacterial pathogens use these toxins to establish and/or contribute to the progression of an infection has been studied in only a handful of organisms. It is clear, however, that these pathogens use the CDCs in far more sophisticated ways than as simple cell lytic agents [1–4]. Many studies have shown that the CDCs can induce various cellular effects, usually at levels that do not lyse a cell [5–20], but the contribution of these effects to disease progression remains less well understood. Some disease causing pathogens express CDCs that lack the ability to form a pore [21,22], which suggests that there may be functions of the CDCs other than pore formation that can contribute to disease.

The pore forming mechanism of these toxins has been the object of intense scrutiny for the past 20 years, which has provided significant insights into their pore forming mechanism. The CDCs exhibit several hallmark features that include a complete dependence of their pore-forming mechanism on the presence of membrane cholesterol and the formation of extraordinarily large membrane pore complexes. The study of the CDC pore complex has focused on understanding how these soluble, monomeric proteins assemble into a large membrane spanning pore complex. The subject of this review will be confined to providing a description of the assembly of the CDC pore complex. As will be discussed below the study of the CDC pore forming mechanism has revealed new paradigms in the assembly of a membrane pore complex as well as providing insight into the mechanism of other pore forming proteins.

#### 2. The CDC structure

#### 2.1. The CDC primary structure

The first primary structure of a CDC was reported in 1987 by Walker et al. [23] for the Streptococcus pneumoniae CDC pneumolysin (PLY), which was followed shortly thereafter by the primary structure for the S. pyogenes streptolysin O (SLO) structure [24]. Since that time the primary structures of over 25 different CDCs have been reported in GenBank. These structures revealed that most CDCs are secreted as soluble monomers from the bacterial cell using a typical type II signal peptide. PLY is an exception to this rule since it lacks a signal peptide: how PLY is released from S. pneumonia remains unclear. An early study [25] suggested that it was released by cell lysis whereas more recent studies suggest that release is not associated with cell lysis but may be via a currently unidentified transport system [26,27]. Ultimately, the CDCs are released into the extracellular milieu as soluble monomers that eventually bind to and form a large pore complex in cholesterol containing membranes. The mass of the secreted CDCs range from about 50 to 72 kD. The core structure responsible for pore formation is about 50 kD: several CDCs, however, exhibit amino terminal peptide extensions that range from about a dozen amino acids to a 150 residue fucose binding lectin [28]. The function of these amino terminal extensions is generally poorly understood. A short amino terminal peptide extension in the Listeria monocytogenes CDC, listeriolysin O (LLO), appears to influence its stability [29,30]. Another short peptide extension found at the amino terminus of SLO appears to contribute to the SLO-mediated transport of another protein across the membrane of a eukaryotic cell [4,31]. The amino terminal fucose binding lectin of the Streptococcus mitis lectinolysin (LLY) appears to modulate its cytolytic activity, but how it does so is not known [28].

The primary structures of the CDC family show a high degree of conservation (40–70% identity). A highly conserved structural motif of 11 residues (ECTGLAWEWWR) near the carboxy terminus is a signature motif for the CDCs. This motif, which is termed the undecapeptide or tryptophan rich motif, contains the only cysteine

residue normally found in the secreted form of the CDCs and typically contains 3 tryptophans. As will be apparent from this review it's role in the pore forming mechanism has yet to be established unambiguously. The primary structure of the secreted form of the CDCs is remarkably hydrophilic with no obvious hydrophobic regions that could be potential transmembrane regions.

#### 2.2. The CDC crystal structure

The first crystal structure of the soluble secreted form of a CDC was solved in 1997 by Rossjohn et al. [32]. They crystallized and solved the structure of the secreted form of the CDC from Clostridium perfringens, perfringolysin O (PFO). In retrospect, the choice of PFO for crystallization trials was somewhat fortuitous, as the structures of the other three most well studied CDCs at that time, SLO, listeriolysin O (LLO) and PLY, have yet to be solved 14 years later. Other CDCs structures have been solved, however, and include Streptococcus intermedius intermedilysin (ILY) [33], Bacillus anthracis anthrolysin O (ALO) [34] and most recently Streptococcus suis suilysin (SLY) [35]. The structure of PFO (Fig. 1) revealed an elongated four-domain protein that is rich in  $\beta$ -sheet. The structures of the other CDCs mentioned above are highly similar to that of PFO with only small differences in their overall shape. Initially, we proposed that the domain 4 undecapeptide signature motif of PFO acted as a hydrophobic dagger to penetrate the membrane and form the pore [32]. This model of pore formation was based on the previous observation by Nakamura et al. [36] showing that the tryptophans of the undecapeptide entered the membrane. As described below, however, even though the undecapeptide enters the membrane it does not directly contribute to the formation of the membrane pore.

The crystal structure of PFO was an important discovery that was critical to the subsequent elucidation of the pore forming mechanism



**Fig. 1.** Molecular structure of *Clostridium perfringens* perfringolysin O. Shown is a ribbon representation of the PFO crystal structure (PDB ID: 1PFO) [32]. Specific features of the structure are designated as D1–D4, domains 1–4; TMH1 and TMH2 (orange), transmembrane hairpins 1 and 2;  $\beta$ 1,  $\beta$ 4 and  $\beta$ 5,  $\beta$ -strands 1, 4 and 5;  $\alpha$ 1,  $\alpha$ -strand 1; L1–L3, loops L1–L3. The double glycine motif is shown as purple space filled atoms. All structures were generated using VMD [126].

of the CDCs. A combination of the PFO monomer structure with biophysical and biochemical studies revealed significant insights into the remarkable structural transitions in the CDC monomer that are required to form the pore.

#### 3. A brief introduction to the CDC pore-forming mechanism

Before proceeding into the details of the pore forming mechanism the reader will be provided a brief description of the general features of the CDC pore forming mechanism as it relates to the PFO crystal structure in Fig. 1.

Bacteria release CDCs as soluble monomers: typically by a type II secretion pathway. The soluble monomers then bind to the eukaryotic cell surface via cholesterol, although a few CDCs have evolved to use human CD59 as their receptor [37,38]. As described below the CD59 binding CDCs still require the presence of membrane cholesterol to function [39-41]. The initial interaction of the cholesterol binding CDC monomers with the cell surface is mediated via the tip of domain 4 (Fig. 1) where the conserved undecapeptide and loops L1-L3 have been shown to anchor the monomer to the membrane [42,43]. The monomers are bound via the tip of domain 4 in an upright position that is perpendicular to the membrane [44,45]. Membrane binding initiates changes within the monomer structure that leads to the formation of intermolecular contacts between membrane-bound monomers [46]. A major transition that is required for monomer-monomer contact is the rotation of the loop comprised of  $\beta$ 5 and  $\alpha$ 1 (Fig. 1) away from  $\beta$ 4 of the domain 3 core  $\beta$ -sheet. This action frees up the edge of  $\beta$ 4 to pair with  $\beta$ 1 of another monomer [46]. The monomers continue to extend the oligomeric structure until a ring shaped structure is achieved. This structure is termed as the prepore complex and is defined as the completed ring complex that has not yet inserted its  $\beta$ -barrel pore. The prepore state likely has several intermediate states that are currently defined by whether the ring complex is SDS-sensitive or resistant [47–49]. During assembly of the ring-shaped prepore complex two transmembrane  $\beta$ -hairpins (TMHs) from each monomer are derived from the two domain 3  $\alpha$ -helical bundles (Fig. 1) and contribute to the formation of the large  $\beta$ -barrel structure [50,51]. To assemble the PFO pore approximately 36 monomers form a completed ring complex, although this number can vary by a few monomers either way [45]. Therefore, PFO assembles and coordinates the insertion of a β-barrel structure that contains approximately 144 membranespanning  $\beta$ -strands or 72  $\beta$ -hairpins, which results in the formation of a pore with a diameter of 250–300 Å. The CDC pore is currently the largest known toxin pore.

#### 4. Membrane recognition

### 4.1. Characteristics of lipid, sterol and membrane structure that influence CDC binding

Most CDCs bind to membranes by using the cholesterol as their receptor. One of the first hallmark traits to be identified for the CDCs was the ability of added cholesterol to inhibit the hemolytic activity of these toxins, presumably by occupying a receptor binding site (reviewed in [52]). The interaction of these toxins with cholesterol and its role as the receptor have been difficult to elucidate due to the insoluble nature of cholesterol and the inability to generate cocrystal structures of the CDCs with cholesterol. Early studies by Prigent et al. [53] showed that an intact 3- $\beta$ -hydroxyl group, an aliphatic sidechain of appropriate length at carbon 17 of the D ring, a methyl group at carbon 20 and an intact B ring were important to its recognition by CDCs. Sterols with a 3- $\alpha$ -hydroxyl or modifications of the 3- $\beta$ -hydroxyl were not bound by these CDCs, thus demonstrating that recognition was restricted to the 3- $\beta$ -hydroxyl. Other minor changes in the ring structure, such as the saturation state of the B ring, did not affect

their ability to inhibit the hemolytic activity of the CDCs. These studies primarily determined the structures of purified sterols required to bind and inhibit the CDCs, but as described below the lipid environment of intact membranes also plays a major role in whether cholesterol will be recognized and bound by the CDCs. In a subsequent study Alouf and coworkers also showed that the stoichiometry of the interaction between the CDCs and cholesterol was 1:1 [54], suggesting a single cholesterol binding site was present on the toxins.

Ohno-Iwashita and colleagues have extensively studied the interaction of the CDC from Clostridium perfringens, PFO [36,55-67] with membranes and cholesterol. One of their first observations was that chemical modification of the single cysteine in the conserved undecapeptide of PFO caused a significant loss of hemolytic activity [55]. The loss in activity resulted from an altered membrane binding affinity that was associated with a reversible conformational change in PFO. This was the first evidence that suggested the undecapeptide was involved in the interaction of PFO with the membrane. They also found that PFO exhibited biphasic binding to cholesterol-rich membranes: high and low affinity binding was observed [68]. Since both high and low affinity binding sites were dependent on cholesterol, they suggested that cholesterol exhibited a heterogeneous distribution in membranes and that only a fraction of the cholesterol was easily accessible and formed the high affinity binding sites. This insight was prescient as Heuck et al. [69] subsequently studied binding to cholesterol containing liposomes. Although they did not observe a biphasic binding they did find that binding of PFO to cholesterol-rich liposomes exhibits a sharp transition from undetectable binding at 40 mol% total membrane cholesterol to maximal binding at 55 mol%. Hence, not all cholesterol at the surface of the membrane forms a suitable receptor for PFO (and probably most CDCs). Typically, efficient binding of CDCs requires >30 mol% of the total membrane lipid to be cholesterol [69–71], therefore, only a fraction of the total membrane cholesterol likely serves as a suitable receptor.

The studies of Flanagan et al. [70] and Nelson et al. [72] showed that the accessibility of cholesterol to PFO is dependent on the phospholipid structure: lipids that could pack tightly with cholesterol, such as those with unsaturated acyl chains, tended to decrease binding to cholesterol. Lipids that did not pack tightly with cholesterol, such as those with unsaturated acyl chains, shifted the half maximal binding to lower cholesterol levels. They also found that the addition of epicholesterol, which contains a  $3-\alpha$ -hydroxyl group and is not bound by PFO, increased binding of PFO to the membrane. This apparently resulted from the displacement of cholesterol by epicholesterol from its interaction with the lipids, thus freeing up cholesterol from its association with lipid and increasing its availability to PFO. These studies suggested that when the cholesterol concentration exceeded the associative capacity of the lipid it was free to be recognized and bound by PFO. The phospholipid headgroup size also influences the availability of cholesterol at the membrane surface. Zitzer et al. [73] have shown for SLO that lipids with smaller headgroups increase the binding of SLO to membranes. In retrospect these observations make sense since tightly packed lipid-sterol mixtures or mixtures containing lipids with larger headgroups are more likely to sterically occlude the relatively small 3- $\beta$ -hydroxyl of cholesterol thereby restricting access of the CDC cholesterol binding motif (described below) to the cholesterol 3- $\beta$ -hydroxy headgroup.

It was also shown by Waheed et al. [66] that PFO preferentially partitioned into the detergent-resistant membrane (DRM) fraction of cells. Since microdomain structure is often rich in cholesterol it made sense that PFO partitioned with the detergent resistant membrane fraction. This observation, however, has been complicated by the recent studies of Flanagan et al. [70]. They showed that increasing the molar fraction of sphingomyelin, a constituent of rafts, in liposomes decreased PFO binding to cholesterol. Sphingomyelin contains unsaturated acyl chains and would tend to pack more tightly with cholesterol. Since sphingomyelin is concentrated in DRMs these data suggested that PFO would not preferentially bind to sphingomyelin/ cholesterol-rich microdomains in natural membranes. More problematic, however, is the observation by Flanagan et al. that the PFO oligomer alone exhibited a propensity to partition into the detergent resistant fraction, therefore detergent fractionation cannot be reliably used to determine if CDC complexes are associated with DRMs.

#### 4.2. The CDC cholesterol recognition/binding motif (CRM)

The identification of the CDC cholesterol recognition/binding motif (CRM) has been an elusive problem for nearly three decades. Many studies have shown that regions and residues in domain 4 of the CDC structure affect binding of the CDCs to the membrane [36,55,62,65,67,74-76]. More recent studies have narrowed the regions of domain 4 that interact with the membrane to the undecapeptide [43] and three loops (L1–L3) at the tip of domain 4 [42] (Fig. 1). The rest of domain 4 is surrounded by water throughout the assembly of the oligomeric pore complex [42]. Investigators have generally accepted that the conserved undecapeptide motif contributed to cholesterol recognition. This was not an unreasonable theory since the undecapeptide is highly conserved in the CDCs and studies showed that one or more of the tryptophans in the undecapeptide inserted into the bilayer [36], although not deeply [43]. Furthermore, Ohno-iwashita and colleagues [55,74] showed that chemical modification of the undecapeptide cysteine of PFO decreased binding, although the modification also induced a conformation change in the structure of PFO, which could have affected binding indirectly. Furthermore, alanine mutants within nearly every residue of the PFO undecapeptide resulted in significant perturbations in its hemolytic activity and membrane binding [33]. Jacobs, et al. showed that monoclonal antibodies to the undecapeptide inhibit binding [65], however a large antibody bound at the tip of domain 4 would likely disrupt any interaction mediated by this region with the membrane. In total, these studies provided compelling evidence that the undecapeptide was the cholesterol-binding motif.

A series of relatively recent studies, however, suggested the CRM resided within loops L1–L3 rather than the undecapeptide. Soltani et al. [77,78] showed that they could uncouple the cholesteroldependent membrane insertion of the undecapeptide from the cholesterol binding function of the CDCs, which suggested that the CRM resided somewhere in loops L1–L3. Furthermore, the solution of the crystal structures for other cholesterol binding CDCs showed that the undecapeptide 3-dimensional structure in the crystal structures of PFO, ILY, *Bacillus anthracis* anthrolysin O (ALO) and more recently *Streptococcus suis* suilysin (SLY) adopt different structures (Fig. 2A), which makes it difficult to envision how they could recognize and bind to a common receptor motif. These differences were not due to crystal packing forces, but appear to be the result of differences in specific sidechain interactions between conserved residues of the undecapeptide and residues outside of the undecapeptide.

Farrand et al. [40] performed an alanine scan of the residues in loops L1-L3 of PFO and determined that two mutations affected hemolytic activity and binding to a greater extent than any other residues in these loops. These residues are a threonine-leucine pair (residues 490 and 491 of PFO) in loop L1 (Fig. 2B). These residues are conserved in all known CDCs, even those that use human CD59 rather than cholesterol as a receptor. As described below (Section 4.3) the Thr-Leu CRM is also necessary to the cytolytic mechanism of the human CD59 binding CDCs, although it does not function as a receptor-binding motif in those proteins. Farrand et al. showed that alanine and/or glycine substitutions for both residues in PFO, SLO and PLY abolished binding to cholesterol rich liposomes, purified cholesterol and the cholesterol rich membranes of eukaryotic cells. They also showed that conservative substitutions for either residue significantly decreased binding and activity: substitution of serine for Thr-490 and either valine or isoleucine for the leucine-491 of PFO



**Fig. 2.** The structure of the undecapeptide motif and location of the CRM. Shown in panel A is an overlay of domain 4 structures from *Clostridium perfringens* PFO (red), *Streptococcus suis* SLY (PDB ID: 3HVN) [35] (yellow), *Bacillus anthracis* ALO (PDB ID: 3CQF) [34] (cyan) and *Streptococcus intermedius* ILY (PDB ID: 1S3R) [33] (blue). Shown in B is the location of the cholesterol recognition/binding motif of the same CDCs as in panel A (colors assignments are the same as in panel A).

nearly completely abolished binding. Therefore, even the most conservative substitutions are not well tolerated.

The cholesterol recognition/binding motif of the CDCs is surprisingly simple and inviolable. Neither trait should be unexpected since the headgroup of cholesterol is also comparatively simple when compared to other receptor molecules. A complex binding site on the CDCs for cholesterol would not be expected since cholesterol has a relatively limited headgroup exposed at the surface if the membrane. Only the 3- $\beta$ -hydroxyl and possibly portions of the A ring of cholesterol are exposed at the surface. One possible binding mechanism is that the leucine inserts into the bilayer and interacts with the A ring and the threonine forms a hydrogen bond with the 3- $\beta$ -hydroxyl group. The leucine may constantly sample the membrane surface and only upon the correct orientation with the cholesterol headgroup does the threonine lock in the interaction by forming a hydrogen bond with the 3-B-hydroxyl. This interaction is also stereo-specific, as the CDCs do not interact with epicholesterol, which has a  $3-\alpha$ hydroxyl.

Once the Thr-Leu pair establishes an interaction with the membrane cholesterol the residues within loops 2 and 3 insert into the bilayer and firmly anchor the monomer to the membrane [40]. At some point, perhaps coincident with insertion of the L2–L3 loops, portions of the undecapeptide also enter the membrane. The undecapeptide is located near the Thr-Leu pair and so many of the previously described undecapeptide mutations [33,55,62] may have altered its structure in a way that interfered with the interaction of the CRM with cholesterol and/or membrane insertion of the loops L2 and L3.

#### 4.3. Human CD59 binding CDCs

*Streptococcus intermedius* produces ILY, which was shown by Nagamune et al. [79] to lyse only human erythrocytes: nonhuman primate erythrocytes were 100-fold less sensitive to ILY whereas all animal erythrocytes tested were resistant to lysis. They further showed that the hemolytic activity on human erythrocytes could be decreased by pretreatment of the erythrocytes with a protease, suggesting that a protein contributed to ILY binding. This observation was in stark contrast to what was generally understood about CDCs, which were thought to use only cholesterol as their receptor. Nearly a decade after its discovery the basis for the highly selective behavior of the ILY cytolytic activity was revealed by the studies of Giddings et al. [37] who showed that human CD59 was the receptor for ILY. Since then two more CDCs have been discovered that also use human CD59 as their receptor, Gardnerella vaginalis vaginolysin (VLY) [38] and Streptococcus mitis lectinolysin (LLY) [80]. Interestingly, CD59 is a GPI-anchored surface protein and is an inhibitor of the complement membrane attack complex (MAC) [81,82]. CD59 protects host cells from complement-mediated lysis during times of bacterial infection when complement is activated. It does so by binding to complement proteins  $C8\alpha$  and C9, which are two proteins of the complement system that are necessary for the formation of the complement membrane attack complex (MAC) pore (reviewed in [83]). CD59 is homologously restricted [82]: i.e., it only functions to inhibit the host's complement and not that of other species. The fact that ILY only bound and lysed human cells was shown to result from its ability to bind to the site on human CD59 that also endowed it with its speciesspecific inhibition of the complement MAC [37,82].

The ILY binding site for CD59 was recently mapped by Wickham et al. [80]. The CD59 binding site is located in domain 4 and contains a signature motif of **Y**X**Y**1<sub>4</sub>**RS** that is found in VLY and was used to identify LLY as a member of the CD59 binding family of CDCs. This motif is located on the outer surface of the ring on the side of domain 4 that is predicted to face away from the lumen of the pore. As indicated above, the normal cellular function of CD59 is to inhibit the assembly of the MAC pore complex by binding to complement proteins C8 $\alpha$  and C9. The ILY consensus sequence is not present in C8 $\alpha$  or C9. More details on the interaction of ILY and the complement proteins with CD59 are described below in Section 7.2, as the CDCs and C8 $\alpha$  and C9 may share mechanistic and evolutionary relationships.

For most CDCs binding to the cholesterol receptor initiates changes in the monomer structure that lead to the formation of the pore complex. In ILY this function has been transferred to the CD59 binding site [77]. Binding to human CD59 rather than cholesterol initiates these same structural transitions [77]. Binding to CD59 still triggers these changes in cholesterol-depleted membranes, but without cholesterol the pore itself does not form [41]. Therefore membrane cholesterol remains important to the mechanism of ILY, even though it has switched receptors from cholesterol to CD59. Why then does ILY still require cholesterol for its pore forming activity and why did it retain the CRM if its receptor function was replaced by CD59?

These questions were addressed in two studies: the first by Lachapelle et al. [39] and the second by Farrand et al. [40]. Lachapelle et al. [39] first showed that ILY disengaged from CD59 upon prepore to pore conversion. This study raised the question, however, that if ILY disengaged from its receptor how did it maintain contact with the membrane so that it could insert its  $\beta$ -barrel pore? Farrand et al. [40] subsequently showed that ILY containing a CRM knockout remained bound to the membrane via its interaction with CD59 prior to conversion of the prepore to pore, but upon conversion of the prepore to the pore the oligomeric complex formed by the CRM knockout lost contact with the membrane as it disengaged from CD59. Therefore, interaction between ILY and CD59 performs two important functions: (1) it initiates the structural changes that are necessary for pore formation and (2) it positions the CRM so that it can bind to cholesterol and initiate the insertion of loops L1-L3 into the membrane. The additional membrane anchor provided by the cholesteroldependent insertion of loops L1-L3 is necessary to keep the prepore complex attached to the membrane as it disengages from CD59 during the critical prepore to pore transition. If the CRM is knocked out then the pore complex detaches from the membrane as it disengages from CD59 and tries to insert the  $\beta$ -barrel pore [40].

Why does ILY disengage from CD59 during its transition to the pore complex? It appears that the adoption of CD59 by ILY as its receptor did not occur without consequences. The disengagement of ILY from CD59 is apparently necessary for the transition from the prepore to pore. Wickham et al. [80] showed that mutations in either CD59 or ILY that increased the binding affinity of the interaction decreased the rate of pore formation. Their studies strongly suggested that the increased affinity of the ILY-CD59 interaction slowed the rate at which ILY disengaged from CD59, therefore slowing the rate of the prepore to pore transition [80]. This predicted that if the binding affinity was sufficiently high it would prevent the prepore to pore transition. Therefore the binding affinity between ILY and CD59 must be balanced such that it is not too high, which would slow the rate of pore formation, but not so low as to affect the on-rate of binding, which would also slow pore formation.

It is also interesting to note that in the absence of the CRM the residues of loops 2 and 3 do not independently insert into the membrane and anchor ILY to the membrane surface. The CRM of ILY cannot bind it to the cell surface if CD59 is absent [37], which suggests that binding to CD59 must orient the CRM near the membrane so that it can interact with cholesterol. Presumably binding to CD59 also brings loops 2 and 3 near the membrane since they are juxtaposed to the CRM (Fig. 2B). Yet, in the absence of the CRM these hydrophobic loops alone cannot insert into the membrane and anchor ILY. Therefore the interaction of the CRM with cholesterol is a necessary prerequisite for the insertion of these loops.

#### 5. Formation of the prepore complex

The prepore complex is defined as a completed oligomeric complex on the verge of inserting its  $\beta$ -barrel pore. Whether a prepore state was an assembly intermediate of the large CDC oligomeric pore complex was initially an issue of debate. Palmer et al. [84] proposed a model of pore assembly in which a CDC dimer inserted in the membrane followed by the addition of monomers that enlarged the pore from a small pore to a large pore. They suggested that insertion of such a large  $\beta$ -barrel pore from a prepore complex was energetically unfavorable. In fact, the insertion of a naked amphipathic  $\beta$ -hairpin into a membrane is energetically unfavorable [85]. The membrane core lacks hydrogen bond donors or acceptors to which the polar atoms of the  $\beta$ -hairpin polypeptide backbone could hydrogen bond, therefore significantly raising the energetic cost of positioning the hairpins within the bilayer. Formation of a partial or wholly formed pre- $\beta$ barrel would be more energetically favorable for membrane insertion. It was subsequently shown in two studies by Shepard et al. [49] and Hotze et al. [48] that PFO formed a large prepore complex and that this complex then converted to the pore complex. The formation of the prepore complex may facilitate the organization of the large  $\beta$ -barrel pore prior to its insertion by allowing the formation of some or all of the interstrand hydrogen bonds between the transmembrane  $\beta$ hairpins (TMHs). The fact that insertion of the  $\beta$ -barrel after prepore formation was favorable was shown by the work of Hotze et al. [48]. They engineered a disulfide between TMH1 and domain 2 (Fig. 1) that prevented the extension of TMH1 and allowed PFO to form a SDSresistant prepore on membranes. If the disulfide was then reduced after oligomerization was completed the insertion of the B-barrel pore was rapid. These studies showed that (1) the rate limiting step in the formation of the pore complex is the assembly of the oligomeric prepore complex and (2) that once it is completed the insertion of the  $\beta$ -barrel pore is rapid.

More than one prepore state has been identified in PFO suggesting that monomer–monomer interactions proceed through various intermediate states during the assembly of an SDS-resistant prepore complex. Membrane binding primes the monomers for oligomerization: once a monomer has bound to the membrane its structure changes to allow monomer–monomer interactions. These interactions do not occur in solution, even at the high concentrations of protein required for crystallization [86]. The extent to which the structure of the CDC monomer changes upon binding, however, is incompletely understood. Ramachandran et al. [46] have shown that the disruption of the edge-on interaction of  $\beta$ 5 with  $\beta$ 4 in domain 3

of PFO is kinetically linked to membrane binding. This event frees up the edge of  $\beta$ 4 to then pair and form interstrand backbone hydrogen bonds with  $\beta$ 1 of another monomer. In addition to the formation of interstrand hydrogen bonds an intermolecular  $\pi$ -stacking interaction is also formed between a tyrosine in  $\beta 1$  and a phenylalanine in  $\beta 4$ (Tyr-181 and Phe-318 in PFO). This interaction is critical to the formation of a SDS-resistant prepore complex; loss of either residue, or moving one of these aromatics out of register on either strand results in the formation of an SDS-sensitive prepore complex that cannot form a pore [46,47]. It was suggested [46] that the  $\pi$ -stacking interaction locks  $\beta 1$  and  $\beta 4$  into the correct in-register pairing of their backbone hydrogen bonds. Although the great majority of CDCs maintain these two residues, the  $\pi$ -stacking interaction is not present in the CD59 binding CDCs and is also missing in the cholesterol binding CDCs pneumolysin, Lactobacillus iners inerolysin and Arcanobacterium pyogenes pyolysin. Regardless of the function of the  $\pi$ -stacking interaction in those CDCs that maintain this aromatic pair it is obvious that some CDCs have solved this problem in another wav.

Monomer–monomer contact may also help drive some of the domain 3 structural transitions necessary for pore formation. As indicated above, elimination of one of the aromatics that participate in the interstrand  $\pi$ -stacking interaction traps PFO in a SDS-sensitive prepore complex. If mixed with functional PFO this  $\pi$ -stacking mutant can be driven to form an SDS-resistant oligomeric complex and to insert its TMHs into the membrane [47]. This presumably occurs because the functional PFO monomers form cooperative interactions with the mutant PFO molecules that induce these monomers to undergo the necessary structural transitions to form a pore. Hence, membrane binding may induce some structural changes that prepare the CDC for oligomerization, but monomer–monomer interactions may propagate additional structural changes necessary for pore formation.

A second structurally important signature motif of the CDCs was also revealed by these studies: the presence of a conserved Gly– Gly pair (Gly-324 and 325 in PFO) that are located in the  $\beta$ -loop between  $\beta$ 4 and  $\beta$ 5 (Fig. 1). Residues with sidechains cannot be substituted for either glycine without blocking the disengagement of  $\beta$ 5 from  $\beta$ 4 [46], which suggests that sidechains at this location clash with other residues and prevent rotation of  $\beta$ 5 away from  $\beta$ 4. Interestingly, this motif is also positionally conserved within the recently solved crystal structures of several membrane attack/perforin (MACPF) proteins (Fig. 3). The MACPF and CDC proteins lack similarity at primary structure level, but the MACPF proteins exhibit structural similarity with domain 3 of the CDCs (described in Section 7 below) [87–90].

#### 6. Formation of the $\beta$ -barrel pore

#### 6.1. The structure of the $\beta$ -barrel pore

After solution of the PFO crystal structure one of the first features of the CDC mechanism to be revealed was the structure of the membrane-spanning domain. Originally, we proposed that the domain 4 undecapeptide acted as a hydrophobic dagger that plunged into the membrane, which was based on the available data that showed the undecapeptide tryptophans entered the bilayer [62]. Palmer et al. [91], however, identified three residues within domain 3 of streptolysin O that also appeared to also enter the membrane and proposed the presence of a membrane spanning amphipathic  $\alpha$ -helix. Shortly thereafter Shepard et al. [51] and Shatursky et al. [50] disproved both models by completely mapping the membrane spanning regions of PFO using a combination of cysteine scanning and fluorescence spectroscopy. Two sets of  $\alpha$ -helical bundles that flank the domain 3 core  $\beta$ -sheet, which extends from the domain 3 core  $\beta$ -strands 1–4 (Fig. 1), were identified that underwent a change in secondary structure to form two amphipathic β-hairpins. Shatursky et al. [50] further showed that these twin hairpins spanned the bilayer. Since the PFO oligomeric complex is composed of about 36 monomers (some CDCs may contain more monomers) up to 144 amphipathic  $\beta$ -strands are positioned in the membrane to form the CDC  $\beta$ -barrel pore.

The primary structure of the TMHs is some of the least conserved primary structure of the CDCs, although they maintain the typical amphipathic structure of a membrane spanning  $\beta$ -hairpin. Other than the amphipathic nature of the TMHs, no other conserved features are apparent in either TMH. Each TMH is approximately 30 residues long or 15 residues per  $\beta$ -strand. A few residues are highly conserved within each TMH between the CDCs, although no reason for their conservation has been identified. Leu-207 and Phe-211 of PFO are the



**Fig. 3.** The crystal structures of MACPF family proteins showing a CDC domain 3 like structure. The crystal structures of the MACPF proteins mouse perforin (PDB ID: 3NSJ) [88], complement C8 (PDB ID: 30JY) [120] and the *Photorhabdus luminescens* Plu-MACPF (PDB ID: 2QP2) [89] are shown with that of PFO [32]. In each MACPF structure the PFO-like domain 3-like fold is highlighted. Shown in purple ribbon structure in each MACPF structure is the structure analogous to the PFO  $\alpha$ 1- $\beta$ 5 loop that rotates away from  $\beta$ 4. Shown in red space filled atoms is the conserved twin glycine motif of each protein that is present at the junction between  $\beta$ 4 and  $\beta$ 5, which is conserved in all known CDCs [46].

most well conserved in TMH1 among the CDCs (96% identity), but can be changed to cysteine with comparatively little loss in pore forming activity [51]. Ala-293 in TMH2 is also highly conserved (96% identify) among the CDCs. Substitution of Ala-293 with cysteine did not significantly affect its pore forming activity, however, the subsequent modification of the cysteine thiol with a fluorescent probe decreased the activity to less than 10% of native PFO suggesting that a bulky group at this position is not tolerated [50].

#### 6.2. The CDC oligomeric pore complex

CDCs are often seen to form a mixture of rings and incomplete, or arc-shaped complexes by electron microscopy (EM) [49,92-94]. Whether these incomplete rings can insert a partial  $\beta$ -barrel remains unknown: the available data suggests that they do not insert into the membrane and may be artifacts of the EM process, dead end complexes or simply a matter of partially inactive protein preparations. If incomplete rings inserted into the membrane with the same probability as a complete ring then the likelihood of forming complete rings would be comparatively low. It is possible that membrane inserted fragments could associate to form a complete ring but they would require, like a puzzle, pieces that would be just the right size to assemble the ring. This type of assembly mechanism would be a comparatively inefficient method for assembling a ring, especially if formation of a ring complex was necessary. A method of quantifying the formation of the CDC prepore and pore complex was developed by Shepard et al. [49]. They used SDS-agarose electrophoresis [49], which separated the very large CDC pore complex from the monomer and intermediates in the assembly process. They showed that the only SDS-resistant form of the PFO oligomeric complex was a fairly homogeneous high molecular weight species shown to be ring complexes: intermediates in the assembly of the ring complex were rare, even at early stages of assembly when they were stabilized by crosslinking. Furthermore, Heuck et al. [43] have shown that at limiting dilutions of PFO where, on average, a single pore is formed in a liposomal membrane the release of low and high molecular weight markers remains the same as observed at higher concentrations of pores. Hence, the available data suggests that a very large complex, which is presumably a complete ring, is the functional pore-forming species.

#### 6.3. CDC activation: disruption of the domain 2-3 interface

Disruption of the domain 2–3 interface is at the heart of the CDC pore forming mechanism. This interface primarily exists between the  $\alpha$ -helical bundle that ultimately forms TMH1 and domain 2 (Fig. 1), although regions of domain 1 also form some contacts with the domain 3  $\alpha$ -helical bundle. In order to unravel this  $\alpha$ -helical bundle to extend and form TMH1 it must disengage from its interface with domains 2 and 3. Before it was understood that these  $\alpha$ -helical bundles formed the TMHs Rossjohn et al. [32] had noted in the crystal structure of the PFO monomer that this interface exhibited poor complementarity, which was consistent with the fact that this interface must be disrupted to extend TMH1.

How this interface is disrupted is incompletely understood. Heuck et al. [69] showed that membrane insertion of the domain 4 undecapeptide was conformationally linked to the insertion of the TMHs, demonstrating a link between structural changes in domain 3 that are necessary for insertion of the  $\beta$ -barrel and membrane insertion of the domain 4 undecapeptide. Based on different crystal forms of PFO Rossjohn et al. [95] showed that rotational torsion between domains 1–3 and domain 4 broke several contacts at this interface: presumably some torsion may be induced by binding of PFO to the membrane. Whether binding alone can induce sufficient torsional stress to disrupt this interface and extend TMH1 remains unknown. How this conformation signal is communicated from the tip of domain 4, which interacts with the membrane, to this interface is also unknown. As indicated above in Section 5 monomer–monomer contact is also important in driving some domain 3 structural changes [47]. Hence, it is likely that a combination of membrane binding and monomer–monomer contact may be important in driving this transition.

#### 6.4. Membrane insertion of the $\beta$ -barrel pore

Shepard et al. [51] and Shatursky et al. [50] had clearly demonstrated that both TMH1 and TMH2 crossed the bilayer, but the length of the extended TMHs were calculated to be only long enough to reach the surface of the bilayer, but not cross it. This conundrum results from the fact that the bottom of domain 3 is suspended about 40 Å above the membrane, which is approximately the length of the extended hairpins. Therefore, how did the  $\beta$ -barrel pore cross the membrane? Czajkowsky et al. [45] solved this problem by using atomic force microscopy to measure the height of the prepore and pore complexes. They first measured the height of a PFO mutant that was trapped in the prepore state by an engineered disulfide between domain 2 and the domain 3  $\alpha$ -helical bundle that inhibited the complete disruption of the domain 2–3 interface [48]. The prepore complex was determined to extend about 113 Å above the membrane, which closely corresponded to the length of the monomer from the tip of domain 4 to the top of domain 1. Upon reduction of the disulfide, which allowed the prepore complex to convert to the pore complex, they showed that the pore complex extended only 73 Å above the membrane. These data showed that insertion of the  $\beta$ -barrel was associated with a 40 Å vertical collapse of the prepore complex, the distance required to bring domain 3 sufficiently near the membrane surface to allow the hairpins to extend across the bilayer. This was confirmed shortly thereafter by Ramachandran et al. [44] who used Förster resonance energy transfer (FRET) based distance measurements between fluorescent probes within domains 1 and 3 in PFO and the membrane surface. They showed that domains 1 and 3 moved about 35 Å closer to the membrane when the prepore was unlocked and allowed to convert to the pore complex.

Czajkowsky et al. [45] also proposed that once the domain 2–3 interface was disrupted that the domain 2 structure collapsed or folded thereby allowing domains 1 and 3 to move towards the membrane. Tilley et al. [96] later supported this scenario using cryoEM data of the PLY pore, which they used to generate a 3D reconstruction of the pore complex that was fitted with the crystal structure of the PFO monomer. Their data showed that the vertical collapse of the prepore complex was associated with the appearance of a bulge on the outer ring surface that was consistent with a folded domain 2 when fitted with the PFO crystal structure.

#### 6.5. Regulation of the listeriolysin O pore forming activity by pH

The disruption of the domain 2–3 interface is also used to control the pH dependent activity of one CDC, listeriolysin O (LLO), from Listeria monocytogenes. For nearly 25 years it's been known that the pore-forming activity of LLO was sensitive to pH: LLO exhibits maximal activity at lower pH ranges (pH 5-6) and undetectable activity at pH values of 7 or higher [97]. LLO was also shown to be critical to the escape of L. monocytogenes from the vacuole into the cytosol, where it replicated and could undergo cell-to-cell spread without an extracellular phase, thereby protecting it from the immune system (reviewed in [98-100]). It is important that LLO cytolytic activity be tightly controlled: it must function at the low pH of the endosome to effect escape of the bacterial cell into the cytoplasm but must be switched off once escape has occurred to prevent lysis of the host cell membrane. If the host cell membrane is lysed it exposes the bacterial cells to the immune system, which results in an avirulent phenotype [1,101-103]. The structural basis for

the pH sensitivity of LLO was unknown, although it was generally assumed to be similar to other pH-sensitive toxins, such as diphtheria toxin and anthrax toxin. These toxins are inactive and stable at neutral pH, but upon being taken up into an endosome, which is acidified by the vacuolar ATPase, the low pH triggers the structural changes that allow them to insert into the membrane and translocate their enzymatic fragment [104–110].

This mechanism, however, is not utilized by LLO to control formation and insertion of its β-barrel pore. In contrast, acidic residues in domain 3 act as a pH sensor and prematurely trigger the disruption of the domain 2-3 interface in the soluble monomer at neutral pH above temperatures of 30 °C [111]. The premature unfurling of the domain 3  $\alpha$ -helical bundles in the soluble monomers inactivates LLO and causes it to aggregate in solution. The aggregation results from the exposure of hydrophobic residues that are normally protected in the soluble monomer, but which are exposed by the disruption of the domain 2-3 interface [111]. Disulfide locking TMH1 to domain 2 significantly decreased the rate of this inactivation, showing that it was the disruption of the domain 2-3 interface that results in this inactivation. These structural changes do not occur at acidic pH, which is found in the vacuole where the LLO monomers remain stable and capable of membrane binding and insertion of the  $\beta$ -barrel pore. Hence, LLO is not pH activated, but is pH inactivated at neutral pH at temperatures above 30 °C. This mechanism allows LLO to function in the acidic environment of the vacuole where it is required to effect release of the phagocytized bacterial cell, but then is inactivated to prevent lysis of the host cell as the bacteria cell escapes into the neutral environment of the cytoplasm.

The LLO pH sensor is composed of a triad of domain 3 acidic residues, Asp-208, Asp-320 and Glu-247. They are located in the domain 3 core  $\beta$ -sheet near the domain 2–3 interface and their carboxylates are predicted to point in at each other [111]. The close proximity of carboxylates is known to increase the pKa of carboxylates to pH 6–8 [112–115], therefore at acidic pH these residues are mostly protonated and domain 3 remains stable. However, as the pH increases the carboxylates deprotonate and charge repulsion between the carboxylate groups is predicted to destabilize the domain 2–3 interface and cause its disruption, which inactivates the soluble monomer. Replacement of these LLO residues by those present in PFO, a CDC that is stable at acidic and neutral pH, eliminated the pH sensitive nature of LLO [111].

Therefore, at the acidic pH of the vacuole the LLO monomers remain stable and can bind to the membrane and follow the normal mechanism of activation to form a pore, thus facilitating the escape of the bacterial cell in to the cytoplasm. Remaining or newly synthesized soluble LLO monomers that are present after escape of *L. monocytogenes* from the vacuole would be rapidly inactivated in the neutral pH environment of the cytoplasm at 37 °C. Hence, in LLO domain 3 plays a dual role, at acidic pH it functions normally to extend the TMHs and form the  $\beta$ -barrel pore after membrane binding, but at neutral pH it causes the premature unfurling of the  $\alpha$ -helical bundles in soluble monomers, which results in the inactivation of LLO thereby preventing the unwanted lysis of the host cell.

#### 7. The membrane attack complex/perforin family proteins

## 7.1. Structural similarities between the CDC and MACPF protein families suggest a pore forming mechanism for the MACPF proteins

The pore forming mechanism(s) of the complement membrane attack complex/perforin (MACPF) family of proteins has remained ambiguous due to the technical challenges of working with these proteins. The pore forming complement membrane attack complex (MAC) and perforin define the MACPF protein family (reviewed in [83,116]), but there are many members that are widespread among both eukaryotes and prokaryotes [117]. The terminal pathway of

complement, the MAC, is assembled from C5b, C6, C7, C8 and C9 proteins. C8 and C9 are the major proteins that form the MAC pore. This pathway is involved in the defense against invading Gramnegative bacteria by the formation of pores in their membrane [118]. Perforin is a major mediator of host cell destruction by natural killer cells and CD8<sup>+</sup> cytotoxic T-cells. Tumor cells, cells infected with viruses or intracellular bacterial pathogens can be targeted for destruction. Perforin forms pores in the host membrane, which can be directly cytotoxic. It appears, however, that its major function may be effect the delivery of granzymes to the cytosol of the targeted cells to induce apoptosis [119]. Regardless of the mechanism by which perforin effects cell killing the perforin pore is required.

The mechanism by which the MAC and perforin form pores in the membrane has languished due to the difficulty of working with these proteins and the difficulty in expressing functional recombinant forms of these proteins. Recently, however, the crystal structures have been solved for several members of the MACPF family of proteins. These include complement protein C8 [120] and C8 $\alpha$  [87,90], mouse perforin [88] and a complement C9-like protein, pleurotolysin, from the bacterium Photorhabdus luminescens [89]. The MACPF crystal structures revealed that they contained a protein fold strikingly similar to domain 3 of PFO (Fig. 3). The similarities of the MACPFs with the CDCs and a description of the MACPF family of proteins have been the subject of an excellent recent review by Rosado et al. [117]. This common protein fold, which is responsible for the formation of the CDC pore [50,51], was found only in the CDCs prior to the solution of the MACPF crystal structures. This revelation was a major breakthrough in elucidating a potential pore forming mechanism for the MACPF proteins. These investigators [87,89,90] suggested that like the CDCs the MACPF proteins also unravel two  $\alpha$ -helical bundles to form two amphipathic β-hairpins that combine to form an oligomeric  $\beta$ -barrel pore complex.

If the MACPF  $\alpha$ -helical bundles do form extended transmembrane  $\beta$ -hairpins like the CDCS then there is at least one obvious difference in their structures when compared to the CDCs: the estimated length of these hairpins is significantly longer, in most cases, than those found in the CDCs. This suggests that these hairpins may be sufficiently long to span the bilayer without a significant collapse of a prepore structure, as is found in the CDCs. Law et al. [88] recently proposed that the putative hairpins of mouse perforin were sufficiently long to reach and cross the membrane. In fact, they proposed that instead of extending out on the same side of the perforin molecule they should extend to the opposite side of the molecule. If true this would represent a remarkable deviation from the CDC mechanism.

Are there any other features of the CDCs that suggest a functional relationship with the MACPF proteins? No sequence homology exists between the MACPF proteins and the CDCs, however, the double glycine motif that exists in the loop that connects  $\beta$ 4 and  $\beta$ 5 in the CDC domain 3 core  $\beta$ -sheet is also found in the MACPFs (Fig. 3). As indicated above in Section 5, these two glycines are conserved in all known CDC primary structures. They are also conspicuously present in a similar location in the MACPF monomer crystal structures (Fig. 3). We have shown that this glycine pair cannot be changed in PFO to residues with a sidechain without loss of pore forming activity [46]. If amino acids with sidechains are introduced at this location the rotation of  $\beta$ 5 away from  $\beta$ 4 is prevented, presumably due to steric clashes of the sidechains with nearby residues. As mentioned above in Section 5, this rotation frees up the edge of  $\beta$ 4 to pair with  $\beta$ 1 of another monomer. The presence of a glycine pair in a conserved location in the MACPF crystal structures suggests that these residues may also be necessary for the rotation of the MACPF structure that is analogous to the CDC  $\beta$ 5- $\alpha$ 1 loop away from  $\beta$ 4 [46] to allow the intermolecular pairing of  $\beta 1$  and  $\beta 4$ . Although it is compelling to suggest a mechanism of pore formation for the MACPF protein family based on the structural similarities with the CDCs, they must be viewed with caution until they are rigorously demonstrated.

#### 7.2. Human CD59: the yin and yang receptor

As described in Section 4.3 the CDCs ILY, VLY and LLY initiate the formation of the pore by binding to human CD59. Interestingly, the primary function of CD59 on human cells is to inhibit the formation of the complement MAC pore on the host cell membranes to prevent their lysis during times of complement activation. Complement activation during a Gram-negative bacterial infection results in the deposition of the complement MAC onto the bacterial outer membrane, thereby forming a pore in the outer membrane of the bacterial cell. Activated complement MAC, however, can also cause collateral damage to nearby host cells. In addition to CD59 host cells can express a number of other proteins that antagonize complement assembly at different stages (reviewed in [121]). CD59 is a GPI-anchored terminal complement inhibitor that binds to MACPF proteins C8 $\alpha$  and C9 and inhibits the formation of the terminal MAC pore [122,123]. In contrast, the CDCs ILY, VLY and LLY initiate formation of their pores on human cells by binding to human CD59. Therefore, CD59 serves to initiate the formation of the ILY, VLY and LLY pores whereas it inhibits the formation of the complement MAC pore.

Prior to formation of the pore ILY remains bound to CD59 [39]. While bound to CD59 it blocks the site on CD59 that interacts with complement proteins C8 $\alpha$  and C9. Therefore, CD59 cannot inhibit assembly of the MAC while ILY remains engaged with CD59 [39]. This results in the host cells being more prone to attack by their hosts own complement system. This observation and prior studies [37] suggested that both ILY and C8 $\alpha$  and C9 bound to similar sites on CD59. As discussed in Section 4.3 above, Wickham et al. recently mapped the binding site for ILY on CD59. Interestingly, the binding site for ILY exhibited a deep correspondence to the site on CD59 that binds to  $C8\alpha$  and C9 [80]. Most of the residues involved in the binding site for the CDCs were also involved in its interaction with C8 $\alpha$  and C9. As indicated above the ILY binding site for CD59 exhibits a consensus motif of YXYX<sub>14</sub>RS that is not found in C8 $\alpha$  or C9. In fact, neither  $C8\alpha$  nor C9 exhibits significant similarity in their binding sites for CD59 [122-125]. Therefore the MAC proteins and the CDCs bind to a highly conserved site on CD59, yet their binding sites for human CD59 exhibit a remarkable lack of similarity [122,123].

#### 8. Summary and future perspectives

The study of the CDC pore-forming mechanism has revealed new paradigms of how the large CDC oligomeric pore complex is assembled. The study of the CDCs has shown that they undergo significant secondary and tertiary structural changes in order to form the CDC pore. Furthermore, the CDCs employ a novel cholesterol-binding motif to recognize and bind cholesterol at the membrane surface. The study of the CDC mechanism has also led to new insights into the possible mechanism of pore formation for the large family of eukaryotic and prokaryotic MACPF proteins and suggests that they may be ancient ancestors of the CDCs.

Some major features of the CDC pore forming mechanism still remain elusive. For instance, we do not yet understand how the CDCs time the insertion of the large  $\beta$ -barrel pore. Evidence suggests that this happens upon completion of the ring shaped complex, but the nature of the mechanism that triggers/regulates the insertion remains unknown. After insertion of the  $\beta$ -barrel the fate of the lipid and any associated proteins from the lumen of the pore also remains unknown. Do the CDCs specifically carve out sections of membrane that reduces the level of specific membrane proteins that may have important roles in cellular physiology or signaling? Over a decade ago Caparon and colleagues revealed that the *Streptococcus pyogenes* CDC, SLO, specifically translocated another secreted protein from *S. pyogenes*, a NAD glycohydrolase enzyme (SPN), into human keratinocytes in an actin-independent process [4]. Remarkably, they recently showed that this translocation system functions in the absence of pore formation by SLO by showing that the prepore locked form of SLO translocates SPN as well or better than native SLO. Revealing the basis of this transport system will likely reveal an entirely new non-pore forming function for some CDCs. The study of the MACPF proteins also promises to reveal new insights into how they form a pore. Like the CDCs they may form a  $\beta$ -barrel pore from the  $\alpha$ -helical bundle-derived  $\beta$ -hairpins, but are other features of the assembly of the CDC pore complex conserved in the MACPF proteins? Future studies into these various problems will reveal new paradigms that will likely change our understanding of this fascinating group of toxins and provide new insights into the mechanism of other pore forming proteins.

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