Anionic lipids enriched at the ExPortal of Streptococcus pyogenes

Jason W. Rosch  
*Washington University School of Medicine in St. Louis*

Fong Fu Hsu  
*Washington University School of Medicine in St. Louis*

Michael G. Caparon  
*Washington University School of Medicine in St. Louis*

Follow this and additional works at: https://digitalcommons.wustl.edu/open_access_pubs

Please let us know how this document benefits you.

**Recommended Citation**

https://digitalcommons.wustl.edu/open_access_pubs/2489
Anionic Lipids Enriched at the ExPortal of \textit{Streptococcus pyogenes}

Jason W. Rosch, Fong Fu Hsu and Michael G. Caparon
Published Ahead of Print 1 December 2006.
Anionic Lipids Enriched at the ExPortal of Streptococcus pyogenes

Jason W. Rosch,¹ ¶ Fong Fu Hsu,² and Michael G. Caparon¹ *

Department of Molecular Microbiology, Washington University School of Medicine, Box 8230, St. Louis, Missouri 63110-1093,¹ and Department of Internal Medicine, Division of Endocrinology, Diabetes, Metabolism and Lipid Research, Washington University School of Medicine, Box 8127, St. Louis, Missouri 63110-1093²

Received 4 October 2006/Accepted 18 November 2006

The ExPortal of Streptococcus pyogenes is a membrane microdomain dedicated to the secretion and folding of proteins. We investigated the lipid composition of the ExPortal by examining the distribution of anionic membrane phospholipids. Staining with 10-N-nonyl-acridine orange revealed a single microdomain enriched with an anionic phospholipid whose staining characteristics and behavior in a cardiolipin-deficient mutant were characteristic of phosphatidylglycerol. Furthermore, the location of the microdomain corresponded to the site of active protein secretion at the ExPortal. These results indicate that the ExPortal is an asymmetric lipid microdomain, whose enriched content of anionic phospholipids may play an important role in ExPortal organization and protein trafficking.

Secretion of proteins across the single cellular membrane plays a key role in the pathogenesis of infections caused by gram-positive pathogens. These virulence-associated proteins have a number of distinct postsecretion trafficking fates. Some must be delivered into the bacterial membrane, while others are routed to the bacterial cell wall for display on the cell surface. Yet others are secreted into the extracellular milieu, and a subset of these are routed to receptors on the host cell membrane. In certain cases, secreted bacterial effector proteins are translocated across a host cell membrane into the host cell’s cytosolic compartment (26). How a gram-positive cell coordinates protein trafficking and the specific signals responsible for correct routing are only beginning to be understood.

A unique solution to the coordination problem has been observed with Streptococcus pyogenes, a pathogen responsible for a number of diseases ranging from superficial localized infection (e.g., impetigo) to life-threatening invasive diseases (e.g., necrotizing fasciitis). This bacterium possesses an ExPortal, a distinct membrane microdomain that contains a high concentration of the translocons of the general secretory (Sec) pathway, such that it is the primary cellular site for protein secretion (33). In addition, the microdomain also accumulates a high concentration of at least one protein (HtrA) that functions as an accessory factor in postsecretion folding (34), suggesting that the ExPortal functions to spatially couple secretion with protein maturation. Localized secretion has also been observed for two surface adhesins, M protein and protein F, though the mechanisms responsible for site-specific targeting remain unclear (7). Restricted distribution of the secretion machinery is not unique to S. pyogenes, since the translocons of Bacillus subtilis have been found to localize to specific clusters that follow a spiral-like pattern around the cell independently of any known helical structures (3, 6). Taken together, these data suggest that the ExPortal may play an important role in coordinating the trafficking of polypeptides to a number of different postsecretion fates.

An understanding of protein trafficking should also include an analysis of the signals responsible for retention of proteins at the ExPortal itself. The observation that the F1F0 ATPase of S. pyogenes is found in the peripheral membrane, rather than the ExPortal (34), shows that retention is not an intrinsic fate of all membrane proteins. The signals may be unique, since the localization patterns for M protein and protein F are distinct, indicating that different signals may be playing a role in the observed localization patterns (7). Interestingly, the ability of the translocons of B. subtilis to cluster to discrete sites was dependent on the presence of anionic phospholipids in the membrane (3). Thus, it was of interest to determine the distribution of anionic phospholipids in relation to the ExPortal in S. pyogenes.

MATERIALS AND METHODS

Strains, media, and culture conditions. Strains used included Escherichia coli DH5α, B. subtilis JH462 (32), and S. pyogenes HSC5 (14). S. pyogenes strain ΔcowR is derived from HSC5 (8), and the construction of several additional mutant derivatives of HSC5 is described below. Routine culture of S. pyogenes was at 37°C and employed Todd-Hewitt medium (BBL) supplemented with 0.2% yeast extract (Difco). Culture of E. coli and B. subtilis utilized LB broth at 37°C with shaking. Proteolytic assays employed strains grown in C medium that were cultured under the conditions previously described (25). When required, antibiotics were used at the following concentrations: erythromycin, 750 µg/ml for E. coli and 1 µg/ml for S. pyogenes; spectinomycin, 100 µg/ml for both E. coli and S. pyogenes.

DNA and computational techniques. Plasmid DNA was isolated via standard techniques and used to transform S. pyogenes as described previously (5). Restriction endonucleases, ligases, and polymerases were used according to the manufacturer’s recommendations. The fidelity of all constructs derived by PCR was confirmed by DNA sequencing analyses. All references to genomic loci are based on the genome of S. pyogenes strain SF370 (12). Gene assignments were based on the information available in the Kyoto Encyclopedia of Genes and Genomes (www.genome.jp) and were supported by subsequent interrogation of the SF370 genome using BLAST (1) and query sequences derived from Bacillus subtilis gene products with experimentally confirmed activities, as noted.
was performed as previously described (18). An internal standard was added (120,000 at 45 s, repeated four or five times. Membranes were collected by centrifugation with a reciprocating shaking device (FastPrep; Qbiogene) at a speed setting of 4.5 for appropriate sequences. An insertion mutation in confirmation of mutant genome structure by PCR using primers of the appropriate forward (clsIFDF-R (AAG AAT TCC TGG ATA TGA TTA TTT TAA AGG AGC TGC TGG CAA TGG AAT TGG)) and inverse primers pgsARXho (AAC TCG AGG GGA TTT CTT CCC ATT ACT AGG) gene encoding cardiolipin synthase (Spy1212), which was subsequently inserted into pJRS233 using the restriction sites embedded in the primers (underlined). An in-frame deletion was constructed by an inverse PCR technique (23) to delete an internal fragment encompassing approximately 1 kb of the 1.6-kb sequence using the primers clsIFDF (AAC TGC AGG CTA AGG TAA ACT TAG ACT ATC GTA GCC TTT ATC). Similarly, an attempt to construct a mutant lacking phosphatidyl glycerol synthase (Spy2196) was made using amplification primers pgsABam (AAG GAT CCG CTG ACT ACA AAC GTC TCT ATT CAG ATG AGG) and clsRXhoI (AAC TCG AG G CAA TGG TAA TAT ATC CAG ATG AAT CCG GCC ACG CCC TTC AAA AGG AGT CCC TGC TTG ATA AG) and pgsARXho (AAC TGC AGG GGA TTT CTT CCC ATT ACT TGC TGG CAA TGG AAT TGG) and inverse primers pgsAIFD-F (AAG AAT TCC TGG ATA TGA TTA TTT TAA AGG AGC AAG CCT T). Allelic replacement proceeded as described previously (23), with confirmation of mutant genome structure by PCR using primers of the appropriate sequences. An insertion mutation in covR was introduced into JWR1 as described previously (8).

Measurement of cell-free protease activity. The amount of SpeB cysteine protease activity in cell supernatants was determined using the substrate fluorescent protein filter set and are shown in grayscale. WT S. pyogenes was also stained with Nile red (panel D). The bar in each panel is equivalent to 0.5 μm.

FIG. 1. S. pyogenes concentrates anionic phospholipids at discrete microdomains. Various bacterial species were stained with NAO and examined by fluorescence microscopy as described in Materials and Methods. Species included WT S. pyogenes (several representative images are shown in panels A1 to A6), E. coli (panel B), and B. subtilis (panels C1 and C2). Images presented were captured using the green fluorescent protein filter set and are shown in grayscale. WT S. pyogenes was also stained with Nile red (panel D). The bar in each panel is equivalent to 0.5 μm.

Construction of mutants. A mutant (JWR1) with an in-frame deletion of the gene encoding cardiolipin synthase (Spy1212) was constructed as follows: Primers clsBamHH (AAG GAT CCG CTG ACT ACA AAC GTC TCT ATT CAG ATG AGG) and clsRXhoI (AAC TCG AGG GAA CAA TGG TAA TAT ATC CAG CAT CCA TCA AAC G) were used to amplify a DNA fragment carrying Spy1212, which was subsequently inserted into pJRS233 using the restriction sites embedded in the primers (underlined). An in-frame deletion was constructed by an inverse PCR technique (23) to delete an internal fragment encompassing approximately 1 kb of the 1.6-kb sequence using the primers clsIFDF (AAC TGC AGG CTA AGG TAA ACT TAG ACT ATC GTA GCC TTT ATC). Similarly, an attempt to construct a mutant lacking phosphatidyl glycerol synthase (Spy2196) was made using amplification primers pgsABam (AAG GAT CCG CTG ACT ACA AAC GTC TCT ATT CAG ATG AGG) and clsRXhoI (AAC TCG AG G CAA TGG TAA TAT ATC CAG ATG AAT CCG GCC ACG CCC TTC AAA AGG AGT CCC TGC TTG ATA AG) and pgsARXho (AAC TGC AGG GGA TTT CTT CCC ATT ACT TGC TGG CAA TGG AAT TGG) and inverse primers pgsAIFD-F (AAG AAT TCC TGG ATA TGA TTA TTT TAA AGG AGC AAG CCT T). Allelic replacement proceeded as described previously (23), with confirmation of mutant genome structure by PCR using primers of the appropriate sequences. An insertion mutation in covR was introduced into JWR1 as described previously (8).

Measurement of cell-free protease activity. The amount of SpeB cysteine protease activity in cell supernatants was determined using the substrate fluorescent protein filter set and are shown in grayscale. WT S. pyogenes was also stained with Nile red (panel D). The bar in each panel is equivalent to 0.5 μm.

RESULTS AND DISCUSSION

Anionic lipid domains in S. pyogenes. Anionic lipids have long been known to play an important role in protein translocation across the bacterial membrane (11) and in determina-
tion of the topology of membrane proteins (40). Visualization of anionic lipids using the fluorescent dye NAO revealed that *Escherichia coli* and *B. subtilis* membranes contain anionic lipid-rich domains at their septal regions and at their poles (19, 28). For examination of *S. pyogenes*, cells from exponentially growing liquid cultures were stained directly in the growth medium with NAO and examined essentially as described previously (28). This revealed a nonuniform punctate pattern of membrane staining (Fig. 1A1 to A6). However, unlike the case with *E. coli* (Fig. 1B) or *B. subtilis* (Fig. 1C), where staining was observed primarily at the poles (Fig. 1C1) or occasionally in a spiral-like pattern in *B. subtilis* (Fig. 1C2), the single anionic lipid-rich domain of each *S. pyogenes* cell was located at a hemispherical position reminiscent of the location of the ExPortal (Fig. 1A). The NAO stain specifically recognized this hemispherical domain, since treatment with the nonpolar lipid stain Nile red did not stain a particular domain but rather the entire circumferential membrane (Fig. 1D).

Analysis of genomic information (12) revealed that *S. pyogenes* has the capacity to synthesize at least two anionic phospholipids, phosphatidylglycerol (PG) and cardiolipin (CL). When bound to PG, NAO emits green fluorescence. However, due to its greater density of negative charge, CL arranges the NAO molecules in a manner that results in a shift to red fluorescence (28, 29). The NAO-stained polar domains of *E. coli* and *B. subtilis* exhibit predominantly red fluorescence (19, 28; also data not shown). In contrast, the NAO-stained domain of *S. pyogenes* consistently fluoresced green (Fig. 1A), suggesting that it is enriched for PG.

### Analysis of *S. pyogenes* anionic phospholipids

Lipid analysis of streptococci has indicated the presence of two acidic phospholipids, CL and PG, with a majority of the membrane being composed of glycolipids (37). The genome of *S. pyogenes* contains a single copy of *pgsA* (Spy2196), encoding the enzyme responsible for converting phosphatidic acid to PG, and a single copy of *cls* (Spy1212), which encodes the enzyme that converts PG to CL (38). Of the total phospholipid content of bacterial cells, cardiolipin typically comprises approximately 5% and PG about 20% (22). An in-frame deletion mutant lacking *cls* (herein referred to as the Cls−/H11002 mutant) was constructed to lack both the region encoding the enzyme’s predicted active site (38) and approximately 1 kb of additional
Membranes were purified from both wild-type (WT) and Cls/H11002 strains and subjected to Bligh-Dryer extraction with addition of a (12:0) 4-CL internal standard. The subsequent negative-ion electrospray ionization/mass spectrometry analysis of the Cls/H11002 mutant extracts showed only the [M-H] ion of the internal standard (12:0) 4-CL at m/z 1239.9 and no detectable CL ions in the mass range from m/z 1300 to 1500 (Fig. 2A). This contrasted with the abundant ions corresponding to CL that were detected in the same mass range for the WT control (Fig. 2B). These data confirm that S. pyogenes contains a single gene, cls, responsible for the biosynthesis of cardiolipin.

The lipid extracts from both strains were then analyzed for the presence of PG species. In the mass range from m/z 600 to 800, where the [M-2H]+ ions of CL and the [M-H]- ions of PG lie, major ions at m/z 691, 717, 745, 747, and 773, arising from PG species, were observed for the Cls/H11002 cell extract with the CL species seen only as an [M-2H]+ ion at m/z 619.8 [(1240.9-2)/2], arising from the internal standard (Fig. 2C). In contrast, the WT sample had similar PG species in addition to the corresponding [M-H]- ions of CL, as seen in Fig. 1B. The levels of PG appeared elevated in the Cls/H11002 mutant compared to the internal standard as a relative measure (Fig. 2D), although it is possible that this increase may result from an enhanced efficiency of PG extraction from membranes of the Cls/H11002 mutant. The absence of CL and the increase of PG in the Cls/H11002 mutant sample compared to data observed for the WT sample are consistent with the notion that in the prokaryotic biosynthesis pathway, two molecules of PG are involved in the synthesis of each molecule of CL (16). The blockage of the synthetic pathways of CL from PG in the Cls/H11002 mutant is consistent with the apparent increase in the amount of PG and the absence of CL (22).

Cls/H11002 mutant phenotypes. Overall, the Cls/H11002 mutant showed a rather modest phenotype under all conditions examined. For example, the mutant grew at near-WT rates with all media examined and secreted normal amounts of several toxins, including streptolysin O and the SpeB cysteine protease, although the mutant did display a defect in processing the proform of the latter to the active enzyme (approximately 25% of WT rate; data not shown). Transcription regulator CovR (CsrR) mutants overexpress the hyaluronic acid capsule and produce colonies with a characteristic mucoid appearance (8). Interestingly, a construction of a double CovR/Cls/H11002 strain showed no apparent defect in the ability to produce mucoid colonies compared to results for the CovR/H11002 mutant alone (data not shown), suggesting Cls is dispensable for production of capsule, despite the observation that the activity of the hyaluronate synthase enzyme is enhanced by CL in vitro (36). These data suggest that increased levels of PG may compensate for the absence of CL in membranes of the mutant. Consistent with the spectral characteristics of NAO staining in the WT, depletion of CL did not alter the staining pattern of the anionic lipid-rich microdomain (data not shown). These data are consistent with a lipid microdomain enriched in PG. However, analysis of a PG-deficient mutant was not possible, since deletion of pgsA was never recovered. The method for mutagenesis proceeds via the generation of a tandem duplication of mutant and wild-type alleles in the genome that can resolve to either allele by recombination. Typically, chromosomes with

**FIG. 3.** The ExPortal is enriched in anionic lipids. WT S. pyogenes was stained with NAO and then examined in an assay which visualized the ExPortal as the site of secretion of active SpeB protease (33), as described in Materials and Methods. Cells were analyzed by fluorescence microscopy and examined for NAO staining and protease localization, as indicated. An overlay of NAO and protease images is shown on the right (Merge). Three representative groups of streptococcal cells, which are presented at various magnifications, are shown. Bars in leftmost panels are equivalent to 0.5 μm.
either allele are isolated at similar frequencies among the progeny (4). However, while it was possible to produce the merodiploid intermediate strain at the pgsA locus, all progeny recovered following resolution of the duplication contained a copy of the wild-type allele (30/30 tested from several independent pools). This bias towards recovery of the wild-type allele suggests that pgsA may be an essential gene in S. pyogenes, which would not be surprising, since depletion of PgsA in B. subtilis results in filamentous cells that eventually lysis (3, 22).

The ExPortal is a microdomain enriched in anionic phospholipids. The ExPortal can be observed in live cells on the basis of visualizing the site of secretion of active SpeB protease in cells embedded in agarose along with a protease substrate that is intramolecularly quenched but becomes fluorescent when cleaved (33, 35). Examination of the Cts− mutant showed that CL is not required for ExPortal integrity, since the WT and the Cts− mutant showed identical patterns of Sec secretion (data not shown). Examination of NAO-stained WT cells in the protease localization assay produced the expected single membrane domains revealed by both NAO staining (Fig. 3, NAO) and cleavage of SpeB protease substrate (Fig. 3, Protease). Furthermore, merging the NAO and protease localization images revealed a concordance between the positions of the anionic lipid-rich domain and the ExPortal (Fig. 3, Merge), indicating that the ExPortal is a microdomain enriched in anionic phospholipids.

The presence of a high concentration of anionic lipids at the ExPortal is consistent with the importance of anionic lipids in promoting protein secretion. A negatively charged microdomain could provide an efficient scaffold for targeting proteins to the membrane for secretion via interaction with conserved regions of positive charge in signal sequences (10, 20). In addition, the SecA component of the translocon has a high affinity for anionic lipids (39), and its ATPase activity is enhanced by the presence of acidic phospholipids (21). A localized region of negative charge could also provide a mechanism of retention of membrane proteins at the ExPortal via interaction with the positively charged residues.

It is unclear whether the ExPortal is restricted to streptococci or is shared by gram-positive bacteria with different morphologies. The lipid biosynthetic machinery appears to be concentrated in the septal regions of B. subtilis cells (30), and the anionic lipids themselves are predominantly localized to the poles and nascent division sites (19). Depletion of these anionic lipids results in the rapid mislocalization of Sec components from sites of concentration along a helical axis, though localization may also rely upon the expression levels of Sec translocons (3). This may suggest that in bacilli, anionic lipids play a vital role in the organization of the helical subcellular apparatus around which the Sec translocons may be organized. Gram-positive cocci, which generally lack any known subcellular architecture proteins, may have developed a simpler system of organization of the Sec translocons at a single anionic lipid-rich microdomain. How such a domain is maintained remains an open question; however, it is interesting to note that acidic phospholipids may also play a role in the initiation of DNA replication via interaction with the initiator protein DnaA (27). Thus, one possibility for organization of the ExPortal is that a high concentration of a positively charged scaffolding protein may assist in the segregation of anionic lipids into distinct domains.

The fact that many naturally occurring antimicrobial agents are also highly positively charged raises some interesting possibilities as to their mechanisms of action. For example, defensins are cationic peptides produced by numerous host tissues to protect against bacterial infection (13). Polymyxin B also has a high affinity for PG (2) and can greatly decrease the efficiency of protein translocation via the Sec pathway in inverted membrane vesicles (41). This raises the possibility that these compounds may directly interfere with ExPortal-mediated protein secretion via their abilities to recognize anionic lipids, a property that could be exploited for further analysis of protein secretion in the gram-positive cocci.

ACKNOWLEDGMENTS

This work was supported by Public Health Service grant 46433 from the National Institutes of Health. We thank Petra Levin for the gift of B. subtilis strains.

REFERENCES


