Molecular Characterization of Caveolin-Induced Membrane Curvature

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* Running Title: Dissecting caveolin-induced curvature

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Background: Caveolin-1 (Cav1) requires the caveolin scaffolding domain for caveola formation.

Results: The Cav1 scaffolding domain and oligomerization domain are tightly juxtaposed to the membrane in caveolae.

Conclusion: Concerted membrane association of the oligomerization, scaffolding and intramembrane domains are critical for caveola biogenesis and membrane deformation.

Significance: Understanding the membrane association of Cav1 is critical for dissecting how the protein regulates caveola formation and achieves regulation over cellular signaling.

ABSTRACT
The generation of caveolae involves insertion of the cholesterol-binding integral membrane protein caveolin-1 (Cav1) into the membrane, however the precise molecular mechanisms are as yet unknown. We have speculated that insertion of the caveolin scaffolding domain (CSD), a conserved amphipathic region implicated in interactions with signaling proteins, is crucial for caveola formation. We now define the core membrane-juxtaposed region of Cav1 and show that the oligomerization domain (OD) and CSD are protected by tight association with the membrane in both mature mammalian caveolae and a model prokaryotic system for caveola biogenesis. Cryo-electron tomography reveals the core membrane-juxtaposed domain to be sufficient to maintain oligomerization as defined by polyhedral distortion of the caveolar membrane. Through mutagenesis we demonstrate the importance of the membrane-association of the oligomerization domain/CSD for defined caveola biogenesis and furthermore, highlight the functional significance of the intramembrane domain (IMD) and the CSD for defined caveolin-induced membrane deformation. Finally, we define the core structural domain of Cav1, constituting only 66 amino acids and of great potential to nanoengineering applications, that is required for caveolin-induced vesicle formation in a bacterial system. These results have significant implications for understanding the role of Cav1 in caveola formation and in regulating cellular signaling events.

Caveolae are small (50-80 nm in diameter) bulb-shaped invaginations of the plasma membrane (PM) that function in a multitude of fundamental cellular processes, including the
mechanosensation response, cholesterol homeostasis and the regulation of cellular signaling(1-6). An essential structural protein of caveolae is Caveolin-1 (Cav1)(7), a 22kDa cholesterol-binding membrane protein that has been hypothesized to adopt a hair-pin like structure in the membrane, with both the N- and C-termini facing the cellular cytoplasm(8).

The C-terminus of Cav1 is composed of two separate domains, the intramembrane domain (IMD - amino acids 102-135) and the remaining residues (amino acids 136-178). It has been hypothesized that the C-terminus functions in lipid association as truncations consisting of this region in Cav1 and Caveolin-3 (Cav3) were constitutively associated with intracellular lipid droplets(9), and the C-terminus is triply palmitoylated during exportation from the Golgi. These post-translational modifications are not important for caveola formation but have been shown to exert local effects on the insertion of Cav3 within the membrane(10). Additionally, the C-terminus has been implicated in Cav1 homo-oligomerization as this region can co-immunoprecipitate both the N- and C-termini of the protein(11,12). However, expression of a protein lacking the final 31 amino acids of Cav1 (Cav11-147) resulted in normal caveolar formation when expressed in Cav1-/- mouse embryonic fibroblasts(13). The IMD is a highly hydrophobic domain that is critical for caveolar formation (13,14). NMR spectroscopy and chemical shift indexing demonstrated residues 97-136 adopt a “horseshoe” structure within a micelle membrane(15); residues 97-107 comprising α-helix 1, separated from α-helix 2 (residues 111-129) by an unstructured region (amino acids 108-110) and flanked by a second unstructured region (residues 130-136). This horseshoe conformation has been postulated to form a “wedge” between the inner and outer leaflet of the membrane allowing for membrane deformation, and is potentially responsible for the characteristic caveolar morphology(15-17).

Structural analysis of the N-terminus of Cav1 has remained elusive, with conflicting reports suggesting both α-helical(16,18-21) and β-sheet(22,23) organizations. The N-terminus encompasses amino acids 1-101 and is commonly represented with three separable domains comprising the membrane-associated caveolin scaffolding domain (CSD - residues 81-101), the oligomerization domain (OD - amino acids 61-101) that partially overlaps with the proposed N-terminal soluble domain (residues 1-80)(24). In cell systems, de novo expression of Cav1 truncations demonstrated the deletion of the first 48 amino acids of the N-terminus does not affect caveolar morphology but further deletion of the next 11 amino acids abrogates caveola formation(13). The CSD is a critical domain for caveolin oligomerization and caveola formation(13) but has also been proposed to regulate cellular signaling events. The interaction between the scaffolding domain and a consensus sequence termed the caveolin binding motif (CBM - proposed to have the sequence $\Omega X\Omega XXXX\Omega$, $\Omega XXXX\Omega XX\Omega$ or $\Omega X\Omega XXXX\Omega \Omega$ where $\Omega$ is an aromatic residue and X is any amino acid) has been hypothesized to regulate signaling via direct protein-protein mediated sequestration of interacting proteins in an inactive state within caveolae(2,25-27). However, a number of observations have questioned the likelihood of Cav1 regulating signaling in this manner. Most critically, many of the proposed CBMs appear unavailable for interaction with caveolin, often comprising the highly hydrophobic core structural backbone of the proposed binding protein(28,29). Putative CBMs within proteins for which structural information is available do not adopt specific structures and no particular enrichment of CBMs has been observed in organisms with caveolae compared to those without(28,29). Finally, the constitutive and irreversible association of signaling proteins with the caveolin CSD does not explain how reversible interactions with signaling proteins can occur. The availability of the CSD for interaction with CBMs has received less attention yet remains another crucial aspect of the caveolin signaling hypothesis.

Several studies have sought to ascribe both structural and biochemical descriptions to homooligomerized Cav1, obtaining different sized oligomers, of 200, 400 and 600kDa(30) up to 3.3MDa(31) and different organizations, from filamentous and ring-shaped(19) to ovoid complexes(30). Recently, it has been demonstrated that the heterologous expression of Cav1 in *Escherichia coli* resulted in the generation of caveola-like structures(32). It was demonstrated that heterologous caveolae (h-caveolae) could be formed efficiently to high concentrations, easily
purified and did not suffer from many of the drawbacks of studying caveolin proteins in mammalian cell systems – mutational analysis regularly results in exportation problems out of the Golgi complex(13,33). H-caveolae closely mimic the size of mammalian caveolae, possess similar numbers of Cav1 proteins and mutations in Cav1 that do not give rise to caveolae in mammalian cells systems, similarly, did not form h-caveolae(32). The intrinsic propensity for Cav1 to generate vesicles in a heterologous system underlies the basic function of the protein in the general cellular context. Cav1, like many other proteins in the mammalian cell, is involved in sculpting cellular membranes although how this deformation is achieved remains poorly understood(34).

In the current study we utilize this heterologous expression system and purified mammalian caveolae to analyse the insertion of Cav1 in the membrane, map the exact amino acids tightly associated with the membrane and analyse the effects of deletions on h-caveola formation and structural regulation. We define the minimal domain of Cav1 required for h-caveola formation in bacteria and show that, remarkably, uniform vesicle formation can occur upon the expression of less than 100 amino acids.

**EXPERIMENTAL PROCEDURES**

*Expression, purification and cleavage of h-caveolae - E. coli* (Rosetta pLysS DE3) were grown at 37°C to an OD$_{600nm}$ of 1 in Terrific Broth. Caveolin fusion protein constructs, under the T7 RNA polymerase promoter, were induced with 1mM IPTG (Astral Scientific) to express for either 3hrs (for EM) or overnight at 30 oC (for purification). E. coli were lysed using a Cell Disruptor (Constant Systems) and h-caveolae were purified using affinity chromatography with amylose resin (NEB), eluted with maltose (100mM) in phosphate buffered saline (150mM NaCl, 15mM K$_2$HPO$_4$/K$_3$PO$_4$ at pH7.4) and concentrated with 100,000 MWCO Amicon Ultracel Centrifugal Filter (Millipore). The MBP-CAV1-His$_6$ fusion protein was engineered with two Tobacco Etch Virus (TEV) protease cleavage sites between MBP and CAV1. Purified h-caveolae were incubated with AcTEV (Invitrogen) at 4°C overnight to remove the MBP affinity tag.

*Proteinase K digestion* - Purified h-caveolae were incubated with ProK (~20µg of ProK per mg of h-caveolae, New England Biolabs) at 37°C for 30 minutes.

*Cloning* - Site-directed mutagenesis was performed to generate the ΔN, ΔC and ΔNC truncation mutants and the S80E, F81E and W85E point mutations. Primer sequences were as follows:

ΔN-forward: GCTGATATCGGATCCATCGACCTGGTCAAC, ΔN-reverse: GTTGACCAGGTCGATGGAAGGTCTGATCATC,
ΔC-forward: GTGCATCAGCCGTGTCATCGAAGGTCGTCATC, ΔC-reverse: GATGACGCACTTCGATGACACGGCTGATGCAC,
S80E-forward: GAACCAGAAAGGAACACAGTTTGCAGATGCATCATG, S80E-reverse: CTTCCAGATGCACATCAACGTCGTGTGCATCATG,
F81E-forward: CAGAAGGAACACACAGTGAAGATGGCATCTGGAAAG, F81E-reverse: CCTTCCAGATGCACATCAACGTCGTGTGCATCATG,
W85E-forward: CAGTTTTGATGGCATCGAAAAGGCGAGCTTCACCAC, W85E-reverse: GTGGTGAAGCTGGCTTTTCGATGCCATCAAAACTG.

His$_6$ removal-forward: GCAGAAAGAAACAATCTAAGGTCGTCATCATC, His$_6$ removal-reverse: GATGATGACGACCTTAGATTTGTTCATCTGTCATC.

Polymerase chain reaction was performed using HF Phusion (NEB) DNA polymerase. The library of Cav1 truncations was designed in MacVector and sequences were synthesized as
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gBlocks by Integrated DNA Technologies (IDT) and inserted into a pNMTMA vector [described previously (32)] by cohesive-end cloning as per the manufacturer’s instructions.

Purification of caveolae - MDCK cells stably expressing GFP or GFP-Cavin-1 were scraped at 4°C in TNE buffer (50 mM Tris-HCl – pH 7.4, 100 mM NaCl and 0.1 mM EDTA), syringe lysed and spun at high speed (100,000G) in an ultracentrifuge. The supernatant was discarded and the pellet was resuspended in TNE buffer and incubated with MBP-tagged GFP-trap. GFP/GFP-Cavin1:GFP-trap complex was bound to amylose resin (NEB) by the MBP-tag. MBP-bound protein complex was washed repeatedly with high salt PBS and GFP/GFP-Cavin-1 protein was eluted with maltose and concentrated in 100,000 MWCO Amicon Ultracel Centrifugal Filters (Millipore). ProK treatment was performed as described previously, however, 0.5% SDS for 30 min at 37°C was utilized for detergent treatment as caveolae are resistant to disruption by TX-100.

Preparation of liposomes - E. coli extracted lipids (Avanti) in chloroform (Sigma) were deposited on the sides of a round-bottom flask by evaporating the organic solvent under a nitrogen atmosphere. To insure the complete removal of chloroform, a formed lipid layer was stored over night under vacuum. Dried lipids were then hydrated with an aqueous buffer (20 mM HEPES-KOH, pH 7.5, 150 mM NaCl) at a concentration 1 mg/mL. To prepare 50 nm liposomes, a mini-extruder block (Avanti) was heated to 65 °C. Liposomes were extruded 21 times through 50 nm polycarbonate filters (Avanti) and plunge frozen immediately after preparation.

Mass Spectrometry - Protein bands were excised and destained in 50% acetonitrile, 25 mM NH₄HCO₃, reduced with 20 mM dithiothreitol (DTT), alkylated with 50 mM iodoacetamide (IAA), adjusted to pH 8 with 50 mM NH₄HCO₃, dehydrated with 100% acetonitrile and dried with a Speed Vac. The dried gel pieces were rehydrated and incubated overnight with 0.2 ug of trypsin per gel band (in 10% acetonitrile, 50 mM NH₄HCO₃). The digest was terminated with 5% formic acid and peptides were extracted with a solution of 60% acetonitrile, 1% formic acid. The extracted peptides were dried and resuspended in 5% formic acid for mass spectrometric analysis.

Tryptic peptides were subjected to liquid chromatography-tandem mass spectrometry using an Agilent 6520 QTOF coupled with a Chip CUBE and 1200 HPLC. The nano pump flow was set at 0.3 µL/min and the capillary pump flow at 4 µL/min. The HPLC-chip contained a 160 nl C18 trapping column, and a 150 mm C18 resolving column (Agilent G4240-62010). Buffer A was 0.1% formic acid and buffer B was 0.1% formic acid in 90% acetonitrile. The samples were analysed with a gradient of 5% to 50% Buffer B in 45 mins, and then 95% Buffer B at 46 mins for 3 mins, returning to 10% buffer B at 50 mins. The mass spectrometer was programmed to acquire 8 precursor MS1 spectra and 4 MS/MS spectra per second. Dynamic exclusion was applied after 2 spectra and released within 0.25 min. LC-MS/MS raw data generated by was processed with Agilent’s Spectrum Mill. Data were extracted with carbamidomethylated cysteine as fixed modification. The data was searched against species *Canis familiaris* of the SwissProt database, with carbamidomethylated cysteine as fixed modification, a maximum missed cleavage of 2, precursor mass tolerance of +/- 20 and product mass tolerance of +/- 50. Results were filtered by protein score of > 11.0, peptide score of >10, and % Scored Peak Intensity of >60.

Electron microscopy - Negative staining and immunolabeling of purified h-caveolae, caveolae and bacteria expressing different caveolin truncations were performed as previously described (32). Imaging was performed on a JEOL 1011 electron microscope at 80kV fitted with a Morada 4K X 4K soft imaging camera (Olympus) with 2-fold binning.

Cryo-ET was performed as described previously (32). Images were recorded using LC-1100 Direct Electron 4K x 4K camera (at a binning of 2) under the control of SerialEM(35). Tilt-series were reconstructed using 10nm fiducial markers and weighted back-projection in IMOD(36). Cryo-tomograms were subjected to a mild de-noising, non-linear anisotropic diffusion filter, to improve signal-to-noise ratio(37).

Electron tomography was performed on E. coli preserved by high-pressure freezing (EMPact2, Leica) and freeze-substitution (AFS2, Leica) in 1% osmium tetroxide OsO₄ and 0.1% UA then subjected to temperature ramping and cells were embedded in epon. Thick sections (300nm) were
cut on an ultramicrotome (UC6, Leica). Dual-axis tilt series were acquired on a TECNAI F30 (FEI) electron microscope encompassing a tilt range of -60° to +60° at 2° increments. Image acquisition was performed with a Direct Electron LC-1100 4K x 4K camera with 2-fold binning under the control of SerialEM(35). Reconstructions were performed using weighted back-projection in IMOD. Semi-automated segmentation was employed(38,39) to generate a three-dimensional representation of h-caveolae. Morphometric analysis of h-caveola diameters was performed as follows; individual h-caveola were scanned through the “z” dimension to determine the central region and measured, outer-membrane to outer-membrane, such that the bilayer was spanned twice (excluding the protein coat). Two-tailed Student t-tests were performed to determine significance.

RESULTS

Mapping the membrane-protected domains of Cav1 - The C-terminal domain (residues 102-178) and the CSD (residues 81-101) of Cav1 have been shown to possess membrane-binding affinity(11,15,40,41) and have been predicted by multiple different studies to adopt a predominantly α-helical structure(14,16,22,42,43). Despite these observations, the exact residues tightly associated with the membrane have remained unclear. A summary of predicted and experimental structural data including membrane associated domains is represented in Fig. 1. Using our model heterologous system and biochemical approaches coupled with enzymatic degradation of the soluble domains of Cav1, we investigated the exact residues tightly associated with the membrane. For all subsequent experiments, unless otherwise stated, MBP-Cav1-His6 h-caveolae (~72 kDa - the standard caveogenic fusion protein characterized previously(32)) were purified using the maltose binding protein tag (MBP) then treated with a Tobacco etch virus protease to specifically cleave the MBP-tag from the membrane associated Cav1-His6 (~22-24 kDa). Purified Cav1-His6 h-caveolae were treated with a non-specific protease, Proteinase K (ProK), to degrade accessible Cav1 epitopes.

After ProK treatment several small molecular weight bands, of approximately 16-17 kDa and 11-14 kDa, can be observed by Coomassie staining (Fig. 2A). To determine if these remaining small molecular weight bands were Cav1 domains, liquid-chromatography tandem mass spectrometry (LC MS/MS) was performed on the largest and most abundant fragment after ProK treatment. Control bands (undigested) showed a high abundance of tryptic peptides of Cav1 (Table 1) comprising amino acids 5 to 172, although the IMD (residues 102-136) was undetected. This highly hydrophobic domain does not possess tryptic cleavage sites and therefore is likely to result in a peptide too large for the detection by LC MS/MS analysis employed here. The undigested bands after ProK treatment were Cav1 peptides, but only constituted 41.6% of the protein (Table 1). Mapping of these domains revealed Cav1 lacked the immediate N-terminal domain but highly abundant peptides comprising a complete sequence from amino acids 48-102, as well as 135-148, were observed (Table 1). Interestingly, residues from the C-terminus of Cav1, amino acids 165-172, were also highly abundant (Table 1), suggesting that this peptide may be involved in lipid interactions.

To confirm our proteomic analysis, the small molecular weight bands were screened with a panel of antibodies directed against different domains of Cav1. An α-MBP antibody was utilized as a control to assess the effectiveness of ProK treatment, after which almost all reactivity was lost (Fig. 2B). The protected fragment after ProK digestion lacked reactivity to the VIP-N antibody (recognizes the first 20 amino acids of Cav1) after ProK digestion suggesting the first 20 residues of Cav1 are not tightly associated with the membrane and confirming our proteomic analysis. However, reactivity was observed with both the polyclonal α-caveolin antibody (directed against multiple recognition sites between amino acids 1-97) and the Concav α-caveolin antibody that recognizes a region within the OD immediately adjacent to the CSD (amino acids 67-81) (Fig. 2B). To determine if these domains are protected from ProK mediated digestion by their membrane association, we performed ProK treatment before and after membrane disruption with the non-ionic detergent Triton X-100 (TX100). A complete degradation of the protected small molecular weight fragments of Cav1 was observed when ProK treatment was performed after TX100 treatment (Fig. 2C), which corresponded with a complete disruption of h-caveola structure (Fig. 2B).
Furthermore, immunoelectron microscopy (immunoEM) of purified h-caveolae demonstrated that the Concav epitope was unavailable for binding in fully formed h-caveolae unless pretreated with TX-100 (data not shown). These data suggest that tight association with the membrane results in protection from ProK mediated digestion. The protected residues of Cav1 are schematically represented in Fig. 2E - this analysis closely mimics previous observations that demonstrated the minimal caveogenic domain of Cav1 to encompass amino acids 49-147(13).

The OD of Cav1 is protected from ProK digestion in mature mammalian caveolae - To examine whether this model of membrane insertion/association is consistent between h-caveolae and mature mammalian caveolae, caveolae were purified from Madin-Darby Canine Kidney (MDCK) cells using a novel pull-down method (schematic Fig. 3A) exploiting stable expression of GFP-tagged Cavin-1, a specific marker of PM caveolae(44). Previous observations have demonstrated Cav1 is present in multiple different subcellular domains and undergoes different conformational changes during its maturation/export to the PM(8,45). Cav1 is also present in non-caveolar pools at the PM suggesting Cav1 itself is an unsuitable candidate for the immunoprecipitation of mature PM caveolae(46). Cavin1, however, is exclusively localized to caveolae when present at the membrane and therefore is ideal for the isolation of mature PM caveolae(44). An 8-fold enrichment of purified caveolin-positive vesicles was observed comparing MDCK cells expressing Cavin1-GFP with cells expressing GFP alone (Fig. 3B – D). Purified mammalian caveolae were subjected to ProK treatment, after which, the remaining fragment of Cav1 demonstrated a shift in molecular weight from ~23kDa to ~16-17kDa, and demonstrated strong reactivity to both the Concav and polyclonal α-caveolin antibodies. Reactivity to both the VIP-N epitope and the GFP-tag from Cavin1 was completely lost after ProK treatment, indicating degradation was effective and confirming the first 20 amino acid of Cav1 are exposed for degradation (Fig. 3E). No reactivity to Cav1 was observed in purified caveolae pretreated with 0.5% SDS for 30min prior to ProK digestion (SDS was used as mammalian caveolae are resistant TX-100 disruption). Previous work has demonstrated the Concav α-caveolin antibody is only reactive against the intracellular pool of Cav1 in mammalian cells and not against Cav1 at the PM unless pre-treated with cholesterol perturbing agents(47). Together with our data this indicates that the OD adjacent to CSD is tightly associated with the membrane in both mature mammalian caveolae and h-caveolae.

Membrane-associated domains of Cav1 are sufficient for membrane-deformation - We have previously shown that formation of h-caveolae in bacteria depends on caveolin oligomerization and gives rise to the polyhedral tertiary structure of h-caveolae(32). High-resolution cryo-electron tomography (ET) revealed distinct membrane organization, defined as clear angles in the membrane through the medial plane of each h-caveola, and indicate that caveolin oligomers (directly or indirectly) cause distortion of the h-caveolar membrane (Fig. 4A). This is particularly evident in comparison to liposomes of similar bacterial lipid composition and size (Fig. 4C) that show a completely circular profile through the medial plane. To test whether membrane distortion was a feature of the membrane-associated domains of Cav1 or were imposed on the membrane by interactions on the exposed cytoplasmic face of h-caveola, h-caveolae treated with ProK were examined by cryo-ET. ProK-treated h-caveolae showed similar membrane distortion when compared to the control samples (Fig. 4B) Proteinase treatment resulted in a reduction in the density of the h-caveola coat, likely the MBP affinity tag, when compared to the control. These results indicate that the proteolytically protected fragments of Cav1 are sufficient to maintain three-dimensional structural integrity of h-caveolae.

The data obtained from our proteomic, biochemical and cryo-ET analyses indicated the importance of the central domain of Cav1 for maintaining the three-dimensional structure of h-caveolae. Therefore, we generated a series of truncation mutants of the caveolin fusion protein: an N-terminal truncation, ΔN, (Cav149-178), a C-terminal truncation, ΔC, (Cav11-147) and an N- and C-termini truncation, ΔNC, (Cav149-147) (Fig. 4D) to analyse the function of these domains in regulating the formation of h-caveolae. A timecourse of the expression of the Cav1 truncations demonstrated efficient and comparable

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protein expression after 3 h of induction (Fig. 4E). In order to further understand the role these domains have in the oligomeric arrangement of Cav1, the ability of these constructs to form h-caveolae was assessed using immunoEM of cryosections. The full-length MBP-Cav1-His\textsubscript{6} construct generated h-caveolae (Fig. 4F) as previously described\cite{32}. Both the ΔN and ΔNC constructs efficiently generated h-caveolae of consistent size and shape (Fig. 4G & I). The ΔC truncation, however, generated more pleomorphic structures with larger diameters and with reduced efficiency of formation (Fig. 4H; see also Fig. 5C & E). It was confirmed by ProK-mediated digestion that the ΔNC truncation represents the core membrane associated domain by western blot analysis, as ProK degradation did not result in a reduction in the size of the molecular weight of Cav1 (Fig. 4J).

ET was utilized on high-pressure frozen freeze-substituted E. coli expressing the various truncations after 3 h of induction to dissect changes in the three-dimensional architecture of h-caveolae in response to the N- & C-terminal truncations. Consistent with our previous observations, ΔC Cav1 h-caveolae were significantly larger [average diameter of full-length = 39.3 nm (n = 559) and ΔC = 43.3 nm (n = 300)] and also demonstrated greater variation in size (standard deviation of full-length = 11.4 nm compared to ΔC truncation = 13.0 nm) than full-length Cav1 h-caveolae (Fig. 5A, C & E). The N-termini truncation mutants, however, demonstrated a significant reduction in both the diameter, ΔN = 29.0 nm (n = 545) and ΔNC = 31.0 nm (n = 356) and the degree of variation in diameter (standard deviation of ΔN h-caveolae = 5.8 nm and ΔNC h-caveolae = 7.1 nm) (Fig. 5B, D & E). Through high-resolution electron tomography-based approaches we have determined that the deletion of the N-terminus, while not essential for h-caveola formation, results in a more monodisperse population of h-caveola diameters.

Dissecting the membrane association of the OD and CSD - Our data suggest that the OD and the CSD are protected from ProK enzymatic digestion by their tight association with the membrane. To further examine the legitimacy of this claim we mutated a single amino acid, serine\textsuperscript{80}, to a negatively charged glutamate residue. Serine\textsuperscript{80} is located in the middle of the OD and adjacent to the CSD, which is critical for lipid binding, caveola and h-caveola structural integrity\cite{13,32,48}. As previously demonstrated, the introduction of this S80E point mutation disrupted h-caveola formation with the full-length MBP-Cav1-His\textsubscript{6} fusion protein (Fig. 6A)\cite{32}. We confirmed this result with the ΔN, ΔC and ΔNC truncations by immunoEM of cryosections; the S80E point mutant resulted in complete loss of defined h-caveola formation and dramatic tubulation of the cytoplasmic membrane (Fig. 6B - D, respectively). Additionally, electron tomography and segmentation analysis demonstrated an increased number of connections maintained with the cytoplasmic membrane (Fig. 6E & F) when compared to a non-mutated control (Fig. 6G) suggesting the introduction of the S80E point mutation results in a reduction in the potential to perform fission from the bulk membrane. Furthermore, ET revealed a complete loss of defined h-caveola formation (average diameter ΔN S80E = 89.9 nm, standard deviation = 30.6 nm: average diameter ΔC S80E = 110.1 nm and standard deviation = 41.2 nm) (Fig. 6H – J). These data are consistent with a model whereby normal association of the OD and the CSD with the membrane is critical for defined h-caveola formation and the addition of a single negatively charged residue at serine\textsuperscript{80} results in the forced separation between the CSD/OD and the membrane.

To confirm the importance of the membrane insertion of this domain in defined h-caveola structure formation we generated additional mutants; a ΔN construct lacking the polyhistidine tag and two additional point mutants in aromatic side-chains of the OD/CSD predicted to insert into the lipid bilayer, ΔN phenylalanine\textsuperscript{81} to glutamate (F81E) and ΔN tryptophan\textsuperscript{85} to glutamate (W85E). The ΔN construct that lacked the polyhistidine tag generated morphologically typical h-caveolae (Fig. 7A) confirming the potentially charged residues that comprise the His-tag do not affect formation. The two point mutants however resulted in a loss of defined vesicular h-caveola structure (Fig. 7B & C) resembling the tubular invaginations generated by the ΔN S80E mutant (Fig. 6B).
To further test this hypothesis, we analysed the effect and extent of ProK degradation on the ΔN construct in comparison to ΔN S80E, ΔN F81E and ΔN W85E point mutants (the ΔN and ΔN point mutants were used because these constructs expressed equivalently). In agreement with our hypothesis, the substitution of a glutamate residue at serine\textsuperscript{80} resulted in almost complete proteolytic degradation of Cav1 by western blot analysis of the OD/CSD with the Concav antibody (Fig. 7D) and a significant reduction in band intensity was observed comparing the fold changes of ΔN S80E ± ProK but not ΔN ± ProK (Fig. 7E). The extrusion of ΔN F81E and ΔN W85E OD/CSD domains were confirmed by western blot as these domains were largely digested by protease treatment (Fig. 7D & E). These results suggest that the disruption of the association/insertion of the OD/CSD with the membrane by the addition of the S80E, F81E or W85E point mutations is causal in abrogating defined h-caveola structure (a schematic is represented in Fig. 7F).

The IMD and CSD are critical for h-caveola formation - Next we investigated the minimal required structural domains for h-caveola formation. We generated an extensive library of Cav1 truncations and analysed the ability of these different truncations to generate h-caveolae by immunoEM of frozen sections. For this assay we have defined the formation of morphologically typical h-caveolae by the appearance of gold-labeled circular membranous profiles within the cytoplasm of expressing \textit{E. coli} - as in Fig. 4F to I. This is indicative of formation of the uniformly shaped spherical free vesicles typical of h-caveolae as shown by tomography [Fig. 5 and (32)]. As both the IMD and the CSD have previously been demonstrated to be essential domains for caveolar formation(13,14,33) we first interrogated the structural outcomes of deleting these domains. The removal of the IMD (Cav1\textsuperscript{48-97,134-178}) completely abrogated h-caveola formation and membrane association (Fig. 8A) whereas the removal of the CSD (Cav1\textsuperscript{49-81,97-178}) resulted in the accumulation of bacterial membranes in the cytoplasm of expressing cells but did not result in a defined structure (Fig. 8B).

Maintaining the core structural domains for h-caveola formation, we then performed sequential truncation of the C-terminus and defined two different phenotypes; the removal of 13 further residues from the core membrane-associated domain (Cav1\textsuperscript{89-134}) caused a loss of defined h-caveolae structure and extensive tubulation (Fig. 8C), however, the additional removal of the IMD resulted in a completely soluble protein (Cav1\textsuperscript{49-101}, Cav1\textsuperscript{60-101} and Cav1\textsuperscript{81-97}; Fig. 8D – F). Intriguingly, the further removal of the N-terminus did not significantly affect h-caveola formation as both Cav1\textsuperscript{60-147} and Cav1\textsuperscript{81-147} retained the ability to generate morphologically typical h-caveolae (Fig. 8G & H), suggesting, in our model prokaryotic system, the first 20 residues of the OD are not critical for h-caveola formation. Finally, the truncation of the remaining domains completely abrogated h-caveola structure; the CSD and IMD (Cav1\textsuperscript{81-134}) alone were unable to localize to membranes (Fig. 7I) whereas constructs consisting of the IMD alone (Cav1\textsuperscript{97-147} and Cav1\textsuperscript{101-134}) were localized to the cytoplasmic membrane but did not bend the membrane (Fig. 8J & K). These data suggest that both the IMD and the CSD are essential for h-caveola structure and for the first time we define a minimal h-caveogenic domain – a short sequence of only 66 amino acids (Cav1\textsuperscript{81-147}) that can generate vesicles when expressed in \textit{E. coli}.

DISCUSSION

This study of caveolae produced in a model prokaryotic system has facilitated novel insights into Cav1, how it associates with the membrane and the role different domains have in regulating h-caveola formation and structure. Using a variety of techniques, including negative staining electron microscopy, electron tomography, biochemical approaches, mass spectrometry, cryosectioning and immunolabeling and cryo-electron tomography, we have identified Cav1 peptides that are protected from protease degradation and determined the core structural domain required defined h-caveola formation. These results have fundamental implications for understanding the topology of Cav1 in the membrane and how Cav1 could regulate signaling events in the cell.

Cav1 has, to date, been represented as a protein with two separable domains, the largely soluble N-terminal domain\textsuperscript{(11,41)} and the hydrophobic C-terminus\textsuperscript{(12,41)}. The results presented here indicate that a reassessment of these boundaries is required. We have shown that the core membrane
associated domain of Cav1 constitutes residues 49-147; close to the minimal domain for caveola formation in mammalian cells(13).

To investigate the importance of the core membrane-associated domain in h-caveola structure formation we generated a library of truncations and assayed h-caveola formation concurrently with the membrane attachment by immunoEM. As Cav1 is an integral membrane protein, the conservation of membrane association is critical for elucidating domain function but is often overlooked in biochemical assays of caveolin oligomerization(11,12). Our results demonstrated that both the mutation of the CSD and the deletion of this domain abrogated defined vesicular structure, reiterating the importance of the CSD for caveola formation(13). Additionally, we demonstrated the IMD is critical for h-caveola formation. In our assays the IMD, in concert with amino acids 135-147, mediated the retention of the protein within the bacterial membrane; the removal of these domains resulted in the association of Cav1 to the cytoplasm of expressing cells. It is known that mutation of this region abrogates caveola formation in mammalian cells and inhibits the export of the protein from the Golgi complex to the PM(13,14,33). Intriguingly, large regions of the N-terminus could be removed without affecting h-caveola formation - in some cases resulting in a more regular structure. It is possible that the removal of the non-membrane associated domains (amino acids 1-48) may result in the tighter packing of caveolin molecules in the membrane, as fluorescence–based techniques do not show significant differences in the number of caveolins per vesicle comparing the full-length and the ΔN construct (data not shown).

Understanding how Cav1 associates with the membrane is critical for understanding the function of the protein in the general cellular context. The results presented here demonstrate a tight association between the OD/CSD and the lipid bilayer in both mammalian caveolae and our bacterial system. Our observation that the membrane association of this domain, and caveola formation, can be disrupted by the insertion of charged residues at serine S80, phenylalanine S81 or tryptophan S85 strongly links caveola formation to the stable insertion of this domain of caveolin within the membrane - consistent with a ‘wedge’ mechanism for cytoplasmic leaflet expansion and curvature generation(34). The disruption of OD/CSD membrane association by the phosphomimetic S80E construct is suggestive of a potential regulation mechanism for the disassembly of caveolae. While no in vivo data suggests serine S80 is directly phosphorylated [for example(49)], Cav1 can be phosphorylated at multiple different residues(50) and several studies have proposed serine S80 as a potential phosphorylation site(16,48). As these mutations increase the availability of the CSD for protease-mediated degradation and concurrently disrupt caveola formation, it is possible that the extrusion of this site from the membrane is a critical step in destabilizing the caveola unit in response to a mechanical stress(6). This hypothesis is strengthened by the observation that the S80E point mutant cannot form caveolae(13) and this region is masked by the membrane in mature PM-associated caveolae but exposed after the disruption of caveola structure by cholesterol depletion(47). This region is also exposed in Golgi-localized caveolin(45).

While the introduction of charged residues within the crucial serine S80 to tryptophan S85 region abrogates the formation of spherical vesicles of regular size and shape (characteristic of the wild-type caveolin), the mutated caveolin molecule still induces the formation of tubular elements that show limited fission from the bacterial cytoplasmic membrane. We have previously proposed a model in which a polyhedral caveolin cage is generated through a two-step process – linear extension and subsequent trimerization of preassembled linear oligomers(32). This can arise through distinct oligomeric signals within the membrane-associated region that we now pinpoint to residues 81-147. An analogy can be made between the combined activity of epsin (helix insertion) and clathrin (cage formation) to the activity of caveolin, which we propose acts as a hybrid protein with combined membrane insertion and cage assembly properties. We hypothesize in wild-type Cav1, the membrane association of the amphipathic helices of oligomerized CSDs function to promote vesiculation through destabilization of the membrane, in a manner similar to the coordinated function of epsin and clathrin in the internalization of clathrin coated pits(51). The mutation of the CSD in the critical serine S80 to tryptophan S85 region compromises the
insertion of these helices, which is reflected by the protease sensitivity of these mutants. As such, the extrusion of the CSD from the membrane alters the membrane remodelling properties of Cav1 resulting in a BAR-domain like phenotype - greater tubulation and reduced membrane fission (51).

These findings have wide-reaching implications for the control of cellular signaling events as the CSD is often presented as the key binding domain of Cav1. There is a large body of literature discussing the scaffolding domain and how it potentially participates in signal regulation via the “caveolin signaling hypothesis” (2-4,52-54). This proposes Cav1 mediates signal regulation through a series of direct protein-protein interactions between binding motifs on signaling proteins and the CSD, which, in turn facilitates the sequestration of these proteins in inactive forms within caveolae. Our findings argue against this model and suggest that the CSD is likely to be inaccessible for interaction with cytoplasmic proteins as this region is inaccessible to proteases and antibodies unless caveola membranes are disrupted with detergents. While detergent treatment could also potentially affect Cav1-Cav1 interactions that are dependent on membrane integrity (and thus make this domain accessible to proteases indirectly) the increased protease digestion observed upon replacement of serine80, phenylalanine81 and tryptophan85 with glutamate residues argues for the membrane insertion model of this region in the wild type protein and thus against direct interactions between the CSD and a caveolin binding motif on a cytoplasmic protein.

Finally, we have characterized the core structural domain for h-caveola formation and, strikingly, this comprises only 66 amino acids (residues 81-147). The need for tissue-specific delivery of drugs, such as anti-cancer therapies, is critical to improve the efficacy of treatment and reduce non-specific deleterious side effects of these cytotoxic compounds(55). As a tool for biotechnology our prokaryotic h-caveolae system is a significant advance on currently available methods. It is cost-effective, easily manufactured and purified in a single-step but has the added advantages of being a genetically encoded nanovesicle that is easily genetically modified for directed targeting(32). This study has demonstrated significant reduction in the structural heterogeneity to h-caveolae by the removal of the soluble N-terminal domain of Cav1 - this system now generates more monodisperse nanovesicles. Further understanding at the atomic level of how caveolin-lipid interactions allow a 66 amino acid membrane-inserted polypeptide to generate a uniform nanovesicle should provide insights into both the fundamental properties of caveolin in forming caveolae and elucidate how further modifications can be made to optimize the properties of genetically encoded nanovesicle.

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CONFLICT OF INTEREST
The authors declare that they have no conflicts of interest with the contents of this article.

AUTHOR CONTRIBUTIONS
NA and RGP designed experiments, interpreted data and edited the manuscript. NA performed all experiments and wrote the manuscript. JR provided technical assistance for Fig. 7 & 8, SO for Fig. 3, NL for Fig. 4 and CF with EM. DL and MMH provided proteomic expertise. PW and BMC edited the manuscript. BM performed the alignments/predictions in Fig. 1. RGP supervised the work. All authors reviewed the results and approved the final manuscript.
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FIGURE LEGENDS

FIGURE 1. Alignments of observed and predicted structural organizations for full-length caveolin 1 and peptides of Cav1. Red = α-helix, light blue = β-sheet and grey = predicted transmembrane domain.

FIGURE 2. Cav1 is protected from ProK digestion by tight association with the bilayer. A) Coomassie stain of purified MBP-Cav1-His6 h-caveolae ± ProK. Small molecular weight bands of approximately, 16-17kDa and 11-14kDa remained. B) Western blot analysis of Cav1-His6 (after TEV cleavage) using the α-MBP, VIP-N, polyclonal and Concav α-caveolin antibodies. VIP-N recognizes amino acids 1-20, the polyclonal α-caveolin has multiple recognition domains between amino acids 1-97 and the Concav binds residues 67-81. C) Coomassie stain of purified Cav1-His6 h-caveolae (TEV cleaved) ± 0.5% TX-100 for 1 h prior to ProK digestion. The small molecular weight fragments of Cav1 are lost when h-caveolae are disrupted prior to ProK treatment. D) Negative staining (1% uranyl acetate) of purified h-caveolae show ProK-digestion does not disrupt h-caveola morphology, however a complete loss of morphology is evident in the TX-100 treated conditions. Scale Bar 100nm. E) Refined model of Cav1 association with the membrane.

FIGURE 3. The OD is protected by the PM in mature caveolae. A) Schematic of the method used for mammalian PM caveolae purification - described in detail in Experimental Procedures. B) Immunolabeling of GFP expressing MDCK cells shows very few caveolin positive vesicles (polyclonal α-caveolin primary, 10nm Protein A Gold secondary). C) Immunolabeling of immunoprecipitated vesicles from MDCK cells expressing Cavin1-GFP demonstrated abundant and specific labeling of vesicles within the size range of caveolae. D) Quantification of Cav1-positive vesicles. E) Purified caveolae from MDCK cells expressing GFP-Cavin1 digested with ProK alone or treated with 0.5% SDS for 30min at 37°C prior to ProK digestion. Scale bars 100nm.

FIGURE 4. Membrane-associated domains of Cav1 are sufficient for membrane-deformation. A) Optical slices through the medial plane of MBP-Cav1-His6 h-caveolae – abundant protein coat is observed around each h-caveola. Scale bar = 5nm. B) Optical slices through the medial plane of MBP-Cav1-His6 h-caveolae + ProK – no protein coat was observed to each h-caveola. Scale bar = 5nm. The medial planes of MBP-Cav1-His6 h-caveolae ± ProK possess definitive polyhedral tertiary structure as defined by the clear angled profile to the membrane. C) Comparative medial plane slices through liposomes demonstrating circular profiles. Scale bars = 5nm. D) Diagrammatic representation of Cav1 truncations generated for biochemical and structural analysis. E) Western blot time course of induction (α-MBP) of the Cav1 truncations. F) MBP-Cav1-His6 construct generated h-caveola consistent with previous observations(32). G) ∆N generated h-caveola of regular size efficiently. H) ∆C was less efficient in generating regular size and structurally uniform h-caveolae. I) ∆NC generated h-caveola of regular size efficiently. Labeling for F) to I), α-MBP antibody primary (1:100 dilution) and 10nm PAG secondary (1:70 dilution). Scale bars 100nm. J) Western blot analysis with the Concav antibody demonstrates ProK treatment of ∆NC does not reduce the molecular weight.

FIGURE 5. Three-dimensional characterization of h-caveolae generated by Cav1 truncations. A) An optical slice from a reconstructed tomogram of an E.coli expressing full-length MBP-Cav1-His6 h-caveolae. B – D) Optical slices of E. coli expressing ∆N, ∆C and ∆NC. Red asterisks highlight small h-caveolae. E) Distribution of the diameters of h-caveolae revealed significant changes in h-caveolae structure and size compared to the full-length protein. Significant differences between the means of populations are indicated by asterisks on right comparing two populations and illustrating differences with P<0.01 ** and P<0.001 ****, respectively. Diameters were measured from the outer leaflet to the corresponding outer leaflet through the medial plane of each h-caveola. IMODinfo was used to export diameter measurements from IMOD for two-tailed Student t-Test statistical analyses. Scale 50nm.
FIGURE 6. Mutating serine\textsuperscript{80} abrogates defined h-caveola formation. A – D) Immunolabeled frozen sections of *E. coli* expressing Cav1S80E, ΔN S80E, ΔC S80E and ΔNC S80E, respectively. Sections were labeled with an α-MBP antibody (1:100) and 10nm Protein A gold (1:70). A loss of defined h-caveola formation and a marked increase in the number of tubules were observed. Scale 100nm. E) Optical slice of reconstructed electron tomogram employed to analyse the structure of h-caveola generated by S80E point mutation in three dimensions. An increase in the number of cytoplasmic membrane connected h-caveola was observed. Black arrowheads indicate membrane-connected h-caveola. Scale 100nm. F) Segmentation of ΔC S80E, abundant surface connected h-caveola (solid white) were observed when compared to h-caveolae where no connections were resolved (red). Cytoplasmic membrane - translucent white. Scale 100nm. G) Segmentation of ΔC shows reduced number of surface connected membrane invaginations (surface connected h-caveola - solid white, disconnected h-caveola – red, cytoplasmic membrane - translucent white). Scale 100nm. H & I) The diameter of ΔN S80E (n = 337) and ΔC S80E (n = 178) h-caveola were analysed, respectively, at their mid-section. Scale 100nm. J) Quantification revealed a significant loss of defined h-caveola formation. Statistical significance was determined by two-tailed *t*-Tests by comparison to the distribution of diameters of ΔN (p<0.01) and ΔC (p<0.01) respectively.

FIGURE 7. The Serine\textsuperscript{80} to Tryptophan\textsuperscript{85} region is critical for efficient association of Cav1 with the bilayer. A) ImmunoEM of MBP-ΔN-Cav1 without the polyhistidine tag to confirm the potentially charged residues that comprise the C-terminal tag do not effect h-caveolae formation - h-caveolae were morphological identical to h-caveolae from the MBP-ΔN-Cav1-His6 construct. Scale bar = 100 nm. B) ΔN F81E results in heterogeneous h-caveola formation. Scale bar = 100 nm. C) ΔN W85E point mutation dramatically effects h-caveolae morphology with extensive tubulation of the membrane. Scale bar = 100 nm. D) Western blot of Concav α-caveolin antibody demonstrated reduced abundance of recognition domain in ΔN S80E, ΔN F81E and ΔN W85E h-caveolae treated with ProK compared to ΔN and confirms the accessibility of this domain for digestion after the introduction of a glutamate residue. E) Quantification of band intensities comparing ± ProK. Band intensities were not significantly reduced with ΔN ± ProK, however ΔN S80E ± ProK resulted in a significant reduction in the abundance of the epitope (p = 0.0003; n = 3). The recognition of this epitope was also significantly reduced in the ΔN F81E and ΔN W85E point mutants (p = 0.0007, p = 0.018 respectively; n = 2). Statistical significance was determined by two-tailed *t*-Tests. F) Schematic of Cav1 membrane association after introduction of the negatively charged residue.

FIGURE 8. A library of Cav1 mutants reveal the critical domains for h-caveola formation. ImmunoEM of frozen sections of *E. coli* expressing various Cav1 truncations. A schematic of each truncation is depicted above a representative image of each mutant. Dotted lines represent deleted domains. Red lines are representative traces of h-caveola morphologies generated by each truncation. A) Cav1\textsuperscript{49-97,134-178} B) Cav1\textsuperscript{59-81,97-178} C) Cav1\textsuperscript{49-134} D) Cav1\textsuperscript{49-101} E) Cav1\textsuperscript{60-101} F) Cav1\textsuperscript{51-97} G) Cav1\textsuperscript{80-147} H) Cav1\textsuperscript{81-147} I) Cav1\textsuperscript{81-134} J) Cav1\textsuperscript{97-147} K) Cav1\textsuperscript{97-134}. Scale bars = 100 nm.
**TABLE 1.** Summary of detected peptides after ProK treatment.

|                                      | Untreated | + ProK |
|--------------------------------------|-----------|--------|
| Number of detected peptides          | 11        | 7      |
| Percentage coverage of Cav1          | 74.6      | 41.6   |
| **Peptides**                         |           |        |
| MSGG(K)                              | -         | -      |
| (K)YVDSEGHLYTVPIR(E)                 | +         | -      |
| (R)EQGNIYKPNNK(A)                    | +         | -      |
| (K)AMAEEMSE(K)                       | -         | -      |
| (K)QVYDAHTK(E)                       | +         | -      |
| (K)EIDLVRDPEK(H)                     | +         | +      |
| (K)HLNDDVVK(I)                       | +         | +      |
| (K)IDFEDVIAEPEGTHSFDGIWK(A)          | +         | +      |
| (K)ASFTFTTVTK(Y)                     | +         | +      |
| (K)YWFYR(L)                          |            |        |
| (L)LSALFGIPMALSFLHIWAVVPCIK(K)       | -         | -      |
| (K)SFLIEIQCISR(V)                    | +         | +      |
| (R)VYSIYVHTFCDPFEAVGK(I)             |            |        |
| (K)IFSNIQ(I)                         | +         | +      |
| (I)INMQKET                           | -         |        |

MSGGK:YVDSEGHLYTVPIR:EQGNIYK:PNNK:AMAEEMSEK:QVYDAHT:K:EIDLVRD:PK:HLNDD:VVK:IDFEDVIAEPEGTHSFDGIWK:ASFTFTTVTK:YWFYR:LLSALFGIPMALIWGIYFAILSFLHIW:AVVPCI:K:SFLIEIQCISR:VYSIYVHTFCDPFEAVGK:IFSNIQ:INMQKET
FIGURE 1.

Dissecting caveolin-induced curvature

[Diagram showing the process of dissecting caveolin-induced curvature]
FIGURE 2.

A

MW (kDa)

72
55
40
33
24
17
11

ProK - +

Full-length fusion protein
MBP (cleaved)
Cav1-His6

B

MW (kDa)

72
55
40
33
24
17
11

ProK - +

α-MBP

Polyclonal α-caveolin

Concav α-caveolin

C

MW (kDa)

72
55
40
33
24
17
11

ProK - +

0.5% TX100 - +

D

0.5% TX100 - +

ProK - +

E

h-caveola membrane

Cav1

Key:
- Oligomerization Domain
- CSD
- IMD
- Membrane-protected
- ProK-digested
FIGURE 3.
FIGURE 4.

A

B

C

D

E

F

G

H

I

J

Dissecting caveolin-induced curvature
FIGURE 5.
FIGURE 6.

Dissecting caveolin-induced curvature
FIGURE 7.

A. ΔN (His*6)
B. ΔN F81E
C. ΔN W85E

D. MW (kDa) 17

E. Fold change in band intensity

F. h-caveola membrane

Key:
- Oligomerization Domain
- CSID
- IMD
- Negative charge/ phosphorylation
Molecular Characterization of Caveolin-Induced Membrane Curvature
Nicholas Ariotti, James Rae, Natalya Leneva, Charles Ferguson, Dorothy Loo, Satomi Okano, Michelle M. Hill, Piers Walser, Brett M. Collins and Robert G. Parton

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