Myxococcus CsgA, Drosophila Sniffer, and human HSD10 are cardiolipin phospholipases

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Myxococcus xanthus development requires CsgA, a member of the short-chain alcohol dehydrogenase (SCAD) family of proteins. We show that CsgA and SocA, a protein that can replace CsgA function in vivo, oxidize the 2′-OH glycerol moiety on cardiolipin and phosphatidylglycerol to produce diacylglycerol (DAG), dihydroxyacetone, and orthophosphate. A lipid extract enriched in DAGs from wild-type cells initiates development and lipid body production in a csgA mutant to bypass the mutational block. This novel phospholipase C-like reaction is widespread. SCADs that prevent neurodegenerative disorders, such as Drosophila Sniffer and human HSD10, oxidize cardiolipin with similar kinetic parameters. HSD10 exhibits a strong preference for cardiolipin with oxidized fatty acids. This activity is inhibited in the presence of the amyloid β peptide. Three HSD10 variants associated with neurodegenerative disorders are inactive with cardiolipin. We suggest that HSD10 protects humans from reactive oxygen species by removing damaged cardiolipin before it induces apoptosis.

[Keywords: Myxococcus xanthus, CsgA, Drosophila Sniffer, HSD10, phospholipase; cardiolipin]

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Myxococcus xanthus displays a unique multicellular lifestyle when faced with amino acid limitation. Cells move in travelling waves or ripples (Shimkets and Kaiser 1982), form evenly spaced multicellular aggregates known as fruiting bodies [Xie et al. 2011], and differentiate into dormant spores within these aggregates [Holkenbrink et al. 2014]. Several differentiating cell types are produced, leading to programmed cell death (PCD), sporulation, and production of peripheral rods (O’Connor and Zunserman 1991; Boynton et al. 2013). Some mutations that abolish multicellular development can be transiently restored in the presence of wild-type cells [Hagen et al. 1978]. Extracellular complementation is due to either intercellular signaling or metabolic cross-feeding.

Perhaps the most interesting extracellular complementation group is produced by loss of the csgA gene (MXAN_RS06255), which fully eliminates development [Hagen and Shimkets 1990]. Increasing levels of csgA expression induce successive developmental behaviors [Li et al. 1992]. Low expression levels (10%–20% of final developmental levels) induce rippling. Higher levels of expression then entrain cell aggregation into fruiting bodies. Sporulation is achieved when csgA is expressed at the peak concentration, completing the developmental program. Addition of exogenous, purified CsgA can restore aggregation at low concentrations and sporulation at higher concentrations [Kim and Kaiser 1991].

In the current model for extracellular complementation of csgA mutants, often referred to as C-signaling, full-length p25 (kilodalton) CsgA is processed to a p17 form that is exported to the cell surface where it binds a receptor on an adjacent cell [Lobedanz and Sogaard-Andersen 2003]. Proteolytic processing of CsgA is mediated by serine protease PopC, and a popC mutant has abrogated development [Rolbetzki et al. 2008]. However, the current model fails to account for many critical pieces of data that are inconsistent with it. For example, CsgA has features associated with enzymatic function, such as a Rossman fold, shown to bind NAD⁺ in vitro, and conserved catalytic residues typical of short-chain alcohol dehydrogenases [SCADs], shown to be essential for development in vivo [Lee et al. 1995]. Membrane fractionation indicated that CsgA is an inner membrane protein [Simunovic et al. 2003], not an outer membrane protein as previously reported [Lobedanz and Sogaard-Andersen 2003]. Extensive proteomic studies have likewise never placed CsgA in the outer membrane or on the cell surface [Kahnt et al. 2010; Bhat et al. 2011], and no receptor has ever been identified. The p17 form lacks the N-terminal NAD⁺-binding domain, and genomic replacement of a truncated csgA gene encoding an 18.1-kDa protein abolishes development [Lobedanz and Sogaard-Andersen 2003].

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Finally, up-regulation of SocA, a distantly related homolog of CsgA with only 28% amino acid identity, restores development [Lee and Shimkets 1994, 1996]. Since SocA must be substantially up-regulated to rescue development in vivo, it was suggested that rescue is due to SocA having substrate overlap. SocA was shown to oxidize the phospholipid lysophosphatidylethanolamine (lyso-PE) in vitro [Avadhani et al. 2006].

CsgA homologs are found in eukaryotes, and genetic studies have provided compelling evidence that they prevent neurodegenerative disorders. A unifying feature of these neurodegenerative disorders is a marked increase in reactive oxygen species (ROS) [Gardner 1989]. Once a lipid radical is formed, it perpetuates production by creating a new radical on a neighboring fatty acid at the site of bis-allylic hydrogens. Formation of new radicals terminates only when two radicals react to produce a fatty acyl peroxide. An insertion in the csgA homolog sniffer from Drosophila produces oxidative stress-related neurodegeneration that includes enhanced apoptosis and reduced locomotor fitness [Botella et al. 2004]. Sniffer can reduce by a product of lipid peroxidation, (E)-4-oxonon-2-enal (4ONE), to (E)-4-hydroxynon-2-enal (4HNE) in vitro, which is less neurotoxic [Martin et al. 2011]. However, 4HNE is still a potent electrophile with deleterious effects on proteins and nucleic acids, suggesting that Sniffer has another role in neuroprotection.

Another interesting CsgA homolog is the human SCAD 17β-hydroxysteroid dehydrogenase type 10 (HSD10), the product of the HSD17B10 gene located on chromosome Xp11.2. HSD10 is imported into the mitochondria by a pathway involving Parkin, a protein whose function is abrogated in patients with Parkinson’s disease [Bertolin et al. 2015]. A wide variety of enzymatic activities have been associated with HSD10 [Yang et al. 2014], and their clinical significance has been examined extensively. Mutations in the HSD17B10 gene have been reported in 19 families [Zschocke 2012]. In the classical infantile form of the disease caused by mutation p.R130C, symptoms emerge ∼6–18 mo after birth, and males show progressive neurodegenerative disease, leading to death a few years later. Substantially up-regulated levels of HSD10 are also found in hippocampal synaptic mitochondria of βAPP transgenic mice [He et al. 2002] and human patients with Alzheimer’s disease [Kristofikova et al. 2009], suggesting a correlation with this neurodegenerative disorder. HSD10 (also known as amyloid β [Aβ] peptide-binding alcohol dehydrogenase) is a strong binding partner of the Aβ peptide, the principle component of the extracellular plaques indicative of Alzheimer’s disease [Yan et al. 1997]. Binding of Aβ to HSD10 can be detected at nanomolar concentrations, and micromolar concentrations inhibit HSD10 activity [Borger et al. 2013]. Altered HSD10 levels have also been observed in patients with multiple sclerosis [Kristofikova et al. 2009].

Here we show that CsgA, SocA, HSD10, and Sniffer have a novel phospholipase C-like activity. They oxidize the 2′-OH in the head group of cardiolipin (CL) and phosphatidyglycerol (PG). The enzymatic product is unstable and fragments into diacylglycerol [DAG], dihydroxyacetone [DHA], and orthophosphate [P]. We propose that developing Myxococcus cells use this reaction to convert phospholipids into triacylglycerol [TAG] as the cells shorten during sporulation. Drosophila and humans instead use this reaction to remove damaged CL before it causes apoptosis.

Results

Expression and purification of CsgA yields an active enzyme

To facilitate production of p25 CsgA, the csgA gene [MXAN_RS06225; MXAN_1294] was codon-optimized for Escherichia coli [Supplemental Fig. S1] and cloned with an N-terminal six-histidine tag. For the p17 form, only the 168 C-terminal codons were used. Purified recombinant protein [Supplemental Fig. S2A] rescues csgA mutant [LS2442] development to produce spore-filled fruiting bodies comparable in size and number with those produced by wild-type DK1622 cells [Supplemental Fig. S2B]. Our p17 failed to restore development in the same assay.

Enzymatic activity of the recombinant enzyme was examined with a modified version of the SocA assay [Avadhani et al. 2006]. Since a known substrate of SocA is lyso-PE, whole-cell lipids were extracted with chloroform and methanol. This extract was used as a CsgA substrate, and activity was measured as the conversion of NAD+ to NADH (Fig. 1A). CsgA displays no cofactor preference (Avadhani et al. 2006). Our p17 failed to restore development in the same assay.

CsgA and SocA share overlapping substrate specificities

To determine what lipids are potential substrates, we further assessed CsgA activity using a variety of synthetic C14:0 partial glycerides [Fig. 1B]. Recombinant p25 CsgA oxidizes sn-1-monoacylglycerol [MAG], sn-1,3-DAG, and lyso-PE, but not sn-1,2-DAG [structures depicted in Supplemental Fig. S4]. This data point to the glycerol 2′-OH being the CsgA target. CL and PG both contain this moiety and were examined as possible substrates, as CL is abundant in Myxococcus where it is found almost exclusively in the inner membrane [Orndorff and Dworkin 1980; Lorenzen et al. 2014]. Both CL and PG are excellent substrates for CsgA [Fig. 1B]. The initial rates with C14:0 CL and PG are about fivefold higher than those of the partial glycerides [Fig. 1D]. When substrate is in excess,
complete conversion of 100 μM NAD⁺ to NADH is observed, unlike reactions with the partial glycerides that reach ~50% conversion as expected for a reversible reaction. The CL and PG reactions thus appear to be irreversible and could not be driven in the opposite direction with NADH.

Since SocA can replace CsgA in vivo, their substrate specificity should overlap if enzymatic activity is responsible for bioactivity. While the previously reported SocA substrate lyso-PE gave the most activity, catalysis was also observed with sn-1-MAG, sn-1,3-DAG, PG, and CL substrates (Fig. 1C,D). As with CsgA, no activity was observed with sn-1,2-DAG. Kinetic analyses for both enzymes are shown for CL (Fig. 1E) and PG (Fig. 1F) substrates. Kinetic parameters are compared in Table 1.

Of note is the difference in affinity constants between CsgA and SocA for CL (62 μM and 603 μM, respectively) and PG (57 μM and 324 μM, respectively). CsgA is roughly an order of magnitude more efficient than SocA at using PG and CL. As controls, both p17 and K155R CsgA showed no activity in kinetic assays (Table 1).

**Oxidation of CL or PG constitutes a novel phospholipase C-like reaction**

To determine the consequence of the activity exhibited by these enzymes, the products of reactions using CsgA with C14:0 CL and PG were determined. Hydrophobic products were first extracted in chloroform and subjected to liquid chromatography/mass spectrometry and
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**Table 1. Kinetic parameters of enzymes**

| Enzyme | Lipid species | $K_m$ | $K_{cat}/K_m$ |
|--------|---------------|-------|--------------|
| CsgA (p25) | CL 14:0 | 62 μM | 2.0 min⁻¹ μM⁻¹ |
| CsgA (p25) | PG 14:0 | 57 μM | 1.8 min⁻¹ μM⁻¹ |
| CsgA (p17) | CL 14:0 | n.a. | 0 min⁻¹ μM⁻¹ |
| CsgA K155R | CL 14:0 | n.a. | 0 min⁻¹ μM⁻¹ |
| SocA | CL 14:0 | 603 μM | 0.15 min⁻¹ μM⁻¹ |
| SocA | PG 14:0 | 324 μM | 0.32 min⁻¹ μM⁻¹ |
| HSD10 | CL 14:0 | 140 μM | 0.74 min⁻¹ μM⁻¹ |
| HSD10 | CL 18:2 | 148 μM | 0.72 min⁻¹ μM⁻¹ |
| HSD10 | CL 18:2 (ox) | 40 μM | 4.0 min⁻¹ μM⁻¹ |
| HSD10 D86G | CL 14:0 | n.a. | 0 min⁻¹ μM⁻¹ |
| HSD10 R130C | CL 14:0 | n.a. | 0 min⁻¹ μM⁻¹ |
| HSD10 Q165H | CL 14:0 | n.a. | 0 min⁻¹ μM⁻¹ |
| Sniffer | CL 14:0 | 81 μM | 1.8 min⁻¹ μM⁻¹ |
| Sniffer | CL 18:2 | 87 μM | 2.0 min⁻¹ μM⁻¹ |
| Sniffer | CL 18:2 (ox) | 78 μM | 1.8 min⁻¹ μM⁻¹ |

[n.a.] No activity.

*Peroxidized CL 18:2.

Electron spectroscopic imaging [LC/MS-ESI]. C14:0 DAG was identified in both reactions. A representative peak was identified in reactions lacking enzyme [Supplemental Fig. S5B]. DAG was not identified in reactions lacking enzyme [Supplemental Fig. S5B]. The remaining soluble products were $P_i$ and DHA. $P_i$ is produced at a ratio of two per CL converted and one per PG converted relative to the amount of NADH produced [Fig. 2A,B, respectively]. For both substrates, one molecule of DHA was found per substrate oxidized [Fig. 2A, B]. Since it is possible that glycerol could be generated by hydrolysis, glycerol was assayed and not detected.

These products suggest a novel phospholipase C-like reaction [Fig. 2C]. In the proposed reaction mechanism, CL is first oxidized by CsgA to form a ketone intermediate. The aqueous environment and neighboring phosphates leave the molecule susceptible to hydration at the 2′ position of the glycerol moiety. The resulting rearrangement removes the two DAG side chains. A proposed cyclic intermediate derived from the head group is rearranged in the presence of water to form DHA bisphosphate. $P_i$ appears to be released through another hydration step. The PG reaction is proposed to occur in a similar fashion.

Addition of a partial glyceride extract restores lipid body production

Myxococcus cells generate lipid bodies directly from their membrane lipids as cells shorten to become spores [Bhat et al. 2014]. csgA mutants fail to synthesize these structures, and cells remain long. Lipid bodies are principally composed of TAG species, and one in particular is known to induce sporulation of an esg mutant [Hoiczyk et al. 2009]. We sought to determine whether CsgA catalysis of CL and PG could be a limiting step in the conversion of phospholipids to TAGs. The rationale behind this idea is that feeding a csgA mutant the product of its reaction should bypass the mutational block. However, addition of commercial C14:0 DAG or DAG generated from enzymatic conversion of C14:0 CL failed to stimulate development of a csgA/socA double mutant [LS3931]. Since fatty acid composition may be important to bioactivity and since DAGs with the unusual fatty acids found in Myxococcus are not commercially available, we extracted a lipid fraction containing partial glycerides [MAGs and DAGs] from developing cells 24 h post-induction. Supplementation of LS3931 with this lipid fraction restored development [Fig. 3]. Fruiting bodies of a size similar to that of wild type appeared by 24 h, and sporulation occurred at levels greater than half of wild type compared with ~1% in the unsupplemented mutant [Fig. 3A]. Addition of the lipid extract also restored lipid body production. Representative cells can be seen in Figure 3B, stained with the lipophilic fluorescent dye Nile red. Lipid bodies are seen as red aggregates within cells. Quantification of this complementation was assessed by comparing the average lipid body composition of 30 cells at each lipid concentration [Fig. 3C]. Lipid body induction displays sigmoidal kinetics over a narrow concentration range. Complementation could not be achieved with lipid extracts from vegetative cells or LS3931, indicating that it is developmentally specific.

CL oxidation is a widespread SCAD activity

Representative homologs of CsgA and SocA localize into three distinct clades among SCAD enzymes [Fig. 4A]. The first group, which includes SocA, is annotated as 3-oxoacyl-acyl carrier protein reductases based on homology with E. coli FabG, an essential enzyme required for fatty acid biosynthesis. Other members of this group include HetN, a protein of unknown function required for heterocyst differentiation in Anabaena and Nostoc (Black and Wolk 1994; Liu and Chen 2009) and NodG, which is essential for nodulation of legumes by Rhizobium (Lopez-Lara and Geiger 2001). Interestingly, HetN contains a signaling motif separate from its catalytic site [Higa et al. 2012] that is not conserved in other homologs. The second clade contains enzymes annotated as hydroxysteroid dehydrogenases in humans, Comamonas, and Streptomyces, including HSD10. The final clade includes CsgA and Drosophila Sniffer.

Both Sniffer and HSD10 oxidize CL in vitro with NAD⁺ [Table 1]. Sniffer displays kinetics similar to CsgA, while HSD10 activity is intermediate between CsgA and SocA. These results point to a diverse class of SCADs in both prokaryotes and eukaryotes capable of oxidizing CL. Three HSD10 mutants known to cause mitochondrial disease in humans—D86G, R130C, and Q165H—are all unable to oxidize CL. Sniffer can replace NAD⁺ with NADP⁺ as a cofactor during oxidation of CL, while HSD10 cannot [Supplemental Fig. S3].

CL peroxidation is required for mitochondrial-induced apoptosis. Since HSD10 and Sniffer are thought to be neuroprotective and since mutations result in mitochondrial disrepair, we assayed these enzymes for catalysis using CL substrates containing oxidized fatty acids. In the presence...
of O₂, polyunsaturated fatty acids form lipid radicals. CL bearing four tetralinoleoyl (C18:2) fatty acids was oxidized by exposure to air, verified by colorimetric assay, and assessed as a potential substrate. HSD10 activity increases greatly in the presence of the oxidized C18:2 (CLox) species compared with unoxidized fatty acids (Fig. 4B). The catalytic efficiency under these conditions was twice that of CsgA due to greater affinity for the substrate (Table 1). HSD10 displayed activity with the unoxidized C18:2 species equivalent to that seen with the C14:0 species. Sniffer was also examined in the same manner (Fig. 4C), but no difference was seen, both yielding kinetics similar to that of the C14:0 form (Table 1). Sniffer does not appear to prefer one state over another, but the efficiency for all CL substrates is comparable with that of CsgA.

The Aβ peptide inhibits oxidation of CL by HSD10

Some previously characterized HSD10 activities are inhibited upon binding of the Aβ peptide. Aβ binds tightly to HSD10 and is thought to cause inhibition by altering the active site conformation (Yan et al. 2007). To determine whether inhibition of HSD10 occurs with CLox, we assayed HSD10 activity in the presence of the Aβ peptide (amino acids 1–42) (Fig. 5A). Addition of 10 μM Aβ greatly reduced enzymatic activity and displayed noncompetitive inhibition. Inhibitory kinetics were determined over a course of Aβ concentrations, and the inhibitor constant (Kᵢ) was determined to be 2.5 μM (Fig. 5B), in agreement with previous reports of HSD10 inhibition using a soluble substrate [Muirhead et al. 2010].

Figure 2. Enzymatic activity with CL and PG constitutes a novel lipase-like reaction yielding DAG, DHA, and P₃. (A) Quantification of soluble products from a CsgA enzymatic reaction using CL as a substrate. For each molecule of CL turned over, one molecule of NADH (●) is produced as well as one DHA molecule (▼) and two P₃ (○). Error bars indicate standard deviation of three replicates. (B) Quantification of soluble products from a reaction using PG as a substrate. For each molecule of PG turned over, one NADH (●) is produced along with one DHA (▼) and one P₃ (○) molecule. (C) Proposed enzymatic reaction mechanism for CL. Catalytic activity removes the hydrogen from the 2′-OH to form a ketone intermediate that undergoes nucleophilic attack by water. Rearrangement releases two DAG moieties and forms an unstable cyclic head group. The cyclic molecule is further hydrated to form DHA bisphosphate. Further hydration produces free P₃ and DHA. Oxidation of PG is proposed to occur similarly but with single DAG and P₃ products.
We describe a widespread and novel phospholipase C-like enzyme activity with some members of the SCAD family. A single proton abstraction from the 2′-OH of the CL or PG head group leads to the release of Pi, DAG, and DHA. As an added benefit, cells recover one NADH molecule in comparison with a conventional phospholipase C reaction. This activity appears to be essential for the development of M. xanthus. This enzymatic reaction seems to be repurposed in eukaryotes, where it is likely to be important in mitochondrial function.

CsgA is required for lipid body synthesis in Myxococcus

Myxococcus cells lacking CsgA fail to enter multicellular development. These mutants have an unusual property, however, by which mixture with wild-type cells efficiently restores mutant development. The nature of what is transferred from the wild type to stimulate complementation is a matter of debate. The current model suggests that extracellular complementation is achieved through cell surface recognition of a truncated CsgA protein (Lobedanz and Sogaard-Andersen 2003). We propose a new model built around the enzymatic activity of CsgA and show that SocA and CsgA have common substrates in vitro, albeit with different kinetic parameters. CsgA has the highest catalytic efficiency with CL and PG, components of the Myxococcus inner membrane (Orndorff and Dworkin 1980), where CsgA has been shown to reside (Simunovic et al. 2003). The SocA affinity constant (Kₘ) for CL is nearly 10-fold higher than that of CsgA, leading to a 13-fold reduction in catalytic efficiency (Table 1). This lower catalytic efficiency of SocA may explain the requirement for SocA up-regulation in order to replace CsgA.

During development, wild-type M. xanthus cells produce lipid bodies, while csgA mutants do not. We show that an extract of partial glycerides, including DAGs from developing wild-type cells, is capable of restoring lipid body production to csgA cells. From these results, we propose that TAGs are constructed in two steps. CL and PG are converted to DAGs by removal of the head group with CsgA. Next, an acyltransferase adds the final fatty acid to complete the TAG (Fig. 6A). Production of DAGs within a phospholipid membrane can induce negative curvature to initiate vesicle formation in eukaryotes (Szule et al. 2002). CsgA could initiate vesicle formation of lipid bodies in Myxococcus by a similar mechanism.

HSD10 is a multifunctional enzyme involved in mitochondrial disease

Reduction or loss of HSD10 in Xenopus, mice, or humans diminishes mitochondrial integrity and enhances apoptotic death (Rauschenberger et al. 2010). HSD10 was initially described in isoleucine catabolism (Zschocke 2012),
and the original hypothesis was given that patients with 
HSD17B10 mutations suffered from buildup of pathway 
intermediates. This particular activity is not directly cor-
related with human disease, however, since an isoleucine-
restricted diet did not relieve symptoms. Furthermore, the 
R130C and D86G mutations associated with severe symp-
toms still produced 64% and 30%, respectively, of wild-
type activity, while Q165H, which causes milder clinical 
symptoms, has <3% activity. HSD10 is also capable of 
steroid metabolism, where allopregnanolone (ALLOP), a 
potent ligand of the GABAA receptor, is oxidized to the 
neuro-inactive molecule 5α-dihydroprogesterone (Mel-
cangi and Panzica 2014). HSD10 could play a role in brain 
development and cognitive function, but ALLOP metabo-

Figure 4. CsgA represents a class of enzymes capable of CL ox-
idation, some of which may be important during oxidative stress 
by removing CL peroxides. [A] Cladogram showing CsgA homo-
logs clustering into three groups, with either SocA, HSD10, or 
CsgA. [B] Kinetic analyses of HSD10 using CL 18:2 (*) or CL 
18:2 peroxide (○) as a substrate. [C] Kinetic analyses of Droso-
phila sniffer using CL 18:2 (*) or CL 18:2 peroxide (○) as a sub-
strate. Error bars indicate standard deviation of three replicates.

Figure 5. The enzymatic activity of HSD10 is inhibited by Aβ. 
[A] Kinetic activity of HSD10 with the substrate CLox in the pres-
ence (*) or absence (○) of 10 mM Aβ [1–42] peptide. [B] Aβ inhibi-
tion kinetics as measured by apparent Vmax changes relative to 
wild type. Aβ displayed noncompetitive inhibition on HSD10 ac-
tivity with a KI = 2.5 µM. Error bars indicate standard deviation of 
three replicates.
HSD10 may mediate CL homeostasis

Mitochondria are massive oxygen consumers that provide vast amounts of energy but leave toxic waste products in their wake. Between 0.15% (St-Pierre et al. 2002) and 2% (Chance et al. 1979) of mitochondrial oxygen is converted into the potent ROS hydrogen peroxide. We propose that HSD10 is part of a previously unrecognized pathway to reduce the ROS burden of stressed cells and ameliorate the damage.

In eukaryotic cells, CL is found predominantly in the mitochondria, where it induces negative curvature to help form the highly folded cristae. Proper formation of cristae requires extensive deacylation and acylation of CL [Mejia and Hatch 2015]. Within the cristae, CL stabilizes the respiratory complexes involved in electron transfer [Zhang et al. 2002]. Perturbation of these complexes increases production of ROS [Dudek et al. 2013] leading to peroxidation of CL and apoptosis. The major contributor to CL peroxidation is cytochrome c (CytC). During respiration, CytC functions to transfer an electron from respiratory complex III to respiratory complex IV. The CytC structure in this role is well-studied and involves a heme moiety covalently attached by two thioether linkages. In a separate function, CytC can also bind CL and induces a disordered structure that has presented a challenge to study but has also led to an appreciation that these changes provoke the transition from electron carrier to peroxidase [Muenzner and Pletneva 2014]. Once bound to CytC, the polyunsaturated fatty acids of CL are oxidized in the presence of hydrogen peroxide [Kagan et al. 2005, 2009].

The formation of CLox is essential for release of proapoptotic factors such as CytC from the mitochondria into the cytosol [Petrosillo et al. 2009]. For apoptosis to occur, CLox must first be translocated from the inner membrane to the outer membrane [Garcia Fernandez et al. 2002]. While controlled apoptosis is necessary to produce form and function in developing organisms (for example, the digitation of fingers and toes during embryogenesis), rampant apoptosis causes necrosis. In the latter case, the CytC–CL complex performs a seemingly subversive function. We suggest to the contrary that CytC peroxidase performs a valuable, regulated response to ROS by using HSD10 to remove CLox before oxidative stress progresses to apoptosis (Fig. 6B). This is a dangerous proposition given the autocatalytic nature of the lipid oxidation pathway and can be easily derailed by diminished HSD10 activity or abundance. In addition, HSD10 would create DAGox, a product that can activate cell signaling pathways outside of the mitochondria, such as protein kinase C, a well-known enzyme that blocks further effects of oxidative stress [Gopalakrishna and Jaken 2000; Cosentino-Gomes et al. 2012]. DAGox is known to activate these pathways similarly and in some cases in greater magnitude than DAG [Kambayashi et al. 2007; Takekoshi et al. 2014].

Alzheimer’s disease, Parkinson’s disease, and Drosophila sniffer have increased ROS

A unifying theme in HSD10 and Sniffer disease states is increased levels of oxidative stress. In Drosophila, the P-element insertion into the X-chromosomal sniffer gene causes reduced life span, neurodegeneration, and sluggish walking that deteriorates with age [Botella et al. 2004]. sniffer is distinct from the Drosophila version of HSD17B10, called scully, which appears to be an essential gene [Torroja et al. 1998]. The sniffer phenotype worsens in a hyperoxic environment and is relieved upon overexpression of sniffer, suggesting oxidative stress as the major
determinant. The cellular location of Sniffer is unknown, but if it is located in the mitochondria and in proximity to CLox, it could provide protective value through removal of this radical. It might also convert CytC–CL into the soluble and energetically favorable form of CytC.

In humans, HSD10 is normally found in the brain and cerebral spinal fluid, where it becomes elevated in Alzheimer’s disease patients (Yang et al. 2014). HSD10 binds Aβ and is suspected of direct involvement in Alzheimer’s disease neurodegeneration (Yan et al. 1997). Interestingly, Aβ binding does not exhibit a deleterious effect on HSD10 involvement in the RNase P complex, suggesting that the leading theory of HSD10 function in mitochondria is not involved in Alzheimer’s disease (Vilardo and Rossmanith 2013). Our work shows that Aβ binding inhibits the CL phospholipase activity of HSD10 despite the substantially larger and more complex hydrophobic substrate in comparison with other previous known substrates. These results suggest that the Alzheimer’s disease state could be at least partially due to inactivation of HSD10, which would be expected to block removal of CLox and stimulate apoptosis (Fig. 6B).

HSD10 may play a somewhat analogous role in some forms of Parkinson’s disease. Nearly 40% of the Parkinson’s disease cases beginning before age 45 are due to PARK2 mutations. These mutations prevent ubiquitylation of HSD10, leading to reduced import into the mitochondria (Bertolin et al. 2015). Disruption of HSD10 import would effectively remove its ability to deplete CLox levels and could lead to rampant apoptosis (Fig. 6B). Thus, one could consider Parkinson’s and Alzheimer’s as diseases where HSD10 is depleted or inactivated, respectively. Mitochondrial dysfunction due to ROS has been proposed to play a major role in these two diseases (Nunomura et al. 2006; Schapira 2006). In a broad sense, oxidative stress is correlated with all age-related degenerative disorders. Unfortunately, it has been consistently ignored in mechanistic studies that attempt to discern a path to the disease state.

Conclusions
CsgA, Sniffer, and HSD10 have similar enzyme activities on CL but different purposes in their respective organisms. The Myxococcus saturated and monounsaturated fatty acids are not a ready target for peroxidation, so this organism has little need to repair/replace lipids whose fatty acids have become oxidized. However, CsgA phospholipase C activity appears to provide a direct path to shape change that accompanies sporulation. Our work predicts that CsgA does so by initiating the conversion of excess CL and PG into DAGs that can be converted to TAGs. Future experiments can be devoted to identifying the molecular complexes at work during sporulation, such as the conversion and relocation of membrane phospholipids into lipid body TAGs. Specific probes can be developed to quantify the carbon flux through these lipid metabolic pathways.

Humans have robust pathways for making lipid bodies and would have no need to derive lipid bodies from existing phospholipids except under dire nutritional conditions. Fatty acids are fed into mitochondria and oxidized as the principle energy source. Here, HSD10 appears to be a beneficial gatekeeper that removes damaged CL and in so doing blocks the apoptotic pathway. Our results suggest that quantification of CLox and perhaps DAGox in normal and disease states, particularly those molecules in proximity to CytC, should figure prominently into metrics and mechanisms regarding the maintenance of healthy cells and the development of neurodegenerative disorders.

Materials and methods

Bacterial strains, plasmids, and recombinant proteins
All strain and plasmids used in this study are listed in Supplemental Table S1. M. xanthus and E. coli strains were grown in CYE [1.0% Bacto Casitone, 0.5% Difco yeast extract, 10 mM 3-[N-morpholino]propanesulfonic acid at pH 7.6, 0.1% MgSO4] and LB, respectively, and supplemented with 50 µg mL−1 kanamycin or 50 µg mL−1 ampicillin where appropriate. M. xanthus and E. coli were grown at 32°C and 37°C, respectively.

pTOB10 contains the full-length csgA gene (MXAN_RS06225, MXAN_1294) codon optimized for E. coli and synthesized by Genscript bearing 5’ Nhel and 3’ HindIII restriction sequences [Supplemental Fig. S1]. The gene was ligated into the respective sites of the commercial vector pTrcHisB (Invitrogen) to create the p25 expression plasmid pTOB12. For the p17 expression vector pTOB24, the appropriate sequence was obtained from pTOB10 through PCR with 5’ Nhel and 3’ HindIII flanking restriction sequences and ligated into pTrcHisB. The beginning of the p17 sequence is denoted in Supplemental Figure S1 and is taken from the previously proposed start site for the processed protein (Lobedanz and Sogaard-Andersen 2003). For the CsgA K155R mutant expression plasmid, pTOB57 was constructed via QuickChange PCR [Agilent Technologies] using primers designed by PrimerX [http://www.bioinformatics.org/primers/] to change codon AAA to AGA.

Expression and purification of recombinant proteins
All recombinant CsgA proteins were expressed as 6xHis-tagged in E. coli grown overnight in 1 L of Circlegrow medium (MP Biomedicals) and then purified from inclusion bodies (IBs) resulting in a yield of ~5 mg of protein per liter of culture. Cells were lysed by sonication in solubilization buffer [50 mM Tris-buffered MOPS at pH 8.0, 100 mM KCl, 1% sodium cholate] and centrifuged at 100,000g for 30 min. The resulting pellet was resuspended in solubilization buffer containing 1% Triton X-100 and again harvested at 100,000g for 30 min to enrich IB purity. This pellet was resuspended in 40 mL of TE buffer [100 mM Tris-HCl at pH 8.0, 1 mM EDTA] and centrifuged at 1100g for 5 min. Supernatant was collected and centrifuged at 12,000g for 3 min to pellet IBs. IBs were washed three times with 40 mL of TE buffer containing 2% Triton X-114. The final pellet was resuspended in 3 mL of TE containing 1.5% sodium lauryl sulfate (SLs) and dialyzed overnight in solubilization buffer. Protein was then diluted in 50 mL of solubilization buffer and purified as previously describe using HisPur cobalt resin [Thermo Scientific] [Boynton et al. 2013]. SocA, HSD10 variant, and Sniffer expression was induced by 1 mM IPTG, and proteins were purified directly as soluble proteins. Cells were lysed in solubilization buffer and directly applied to cobalt affinity columns as above. The purity and integrity of
of all proteins were analyzed by SDS-PAGE and Coomassie blue staining. Protein analysis and quantification was carried out using UV-Vis spectrometry and the predicted extinction coefficients for each protein.

Development of M. xanthus

For the development of Myxococcus cells, cultures were first grown in CYE to a density of 5 x 10^9 cells per milliliter. Cells were then harvested by centrifugation at 10,000 rpm for 10 min and resuspended to a final concentration of 5 x 10^9 cells per milliliter. The resulting suspension was plated onto TPM agar (10 mM Tris-HCl, 8 mM MgSO_4, 1 mM K-PO_4 at pH 7.6, 1.5% agar) to initiate development and incubated for appropriate periods at 32°C.

Enzyme assays

CsgA activity was assayed in a manner similar to that previously described for the CsgA homolog SocA (Avadhani et al. 2006). One-milliliter reaction mixtures consisted of TE buffer (10 mM Tris-HCl at pH 8.0, 1 mM EDTA) containing 0.1% (w/v) sodium desferrioxamine B, 1 mM MnCl_2, 1 mM MgCl_2, and 1% DMSO. A band with an Ri value of 0.55 was extracted and dried under nitrogen. For complementation, various amounts (by weight) of lipids dissolved in DMSO were added 1:10 to cells prior to development. After 24 h, complementation was indicated by the presence of lipid bodies as previously described (Bhat et al. 2014). Lipid bodies were quantified by averaging the fluorescent intensity of individual cells normalized to cell length.

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