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Glycan-Foraging Systems Reveal the Adaptation of *Capnocytophaga canimorsus* to the Dog Mouth

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**ABSTRACT** *Capnocytophaga canimorsus* is known to form two kinds of cells on blood agar plates (coccoid and bacillary), evoking phase variation. When grown in coculture with animal cells these bacteria appeared only as bacilli, but in the presence of vancomycin they were round, indicating that coccoid shapes likely result from weakening of the peptidoglycan layer. Polysaccharide utilization locus 5 (*PUL5*) and sialidase mutant bacteria, unable to retrieve glycans from glycoproteins, grew less than wild-type bacteria and also appeared polymorphic unless GlcNAc was added, suggesting that *C. canimorsus* is unable to synthesize GlcNAc, an essential component of peptidoglycan. Accordingly, a genome analysis was conducted and revealed that *C. canimorsus* strain 5 lacks the GlmM and GlmU enzymes, which convert glucosamine into GlcNAc. Expression of the *Escherichia coli* GlmM together with the acetyltransferase domain of GlmU allowed mutant bacteria to grow normally, indicating that *C. canimorsus* is a natural auxotroph that relies on GlcNAc harvested from the host N-glycoproteins for peptidoglycan synthesis. Mucin, a heavily O-glycosylated protein abundant in saliva, also rescued growth and the shape of *PUL5* mutant bacteria. Utilization of mucin was found to depend on Muc, a Sus-like system encoded by *PUL5*. Contrary to all known *PUL*-encoded systems, Muc cleaves peptide bonds of mucin rather than glycosidic linkages. Thus, *C. canimorsus* has adapted to build its peptidoglycan from the glycan-rich dog’s mouth glycoproteins.

**IMPORTANCE** *Capnocytophaga canimorsus* is a bacterium that lives as a commensal in the dog mouth and causes severe infections in humans. *In vitro*, it forms two kinds of cells (coccoid and bacillary), evoking phase variation. Here, we show that cell rounding likely results from weakening of the peptidoglycan layer due to a shortage of N-acetylglucosamine (GlcNAc). *C. canimorsus* cannot synthesize GlcNAc because of the lack of key enzymes. In its niche, the dog mouth, *C. canimorsus* retrieves GlcNAc by foraging glycans from salivary mucin and N-linked glycoproteins through two different apparatuses, Muc and Gpd, both of which are related to the *Bacteroides* starch utilization system. The Muc system is peculiar in the sense that the enzyme of the complex is a protease and not a glycosylhydrolase, as it cleaves peptide bonds in order to capture glycan chains. This study provides a molecular genetic demonstration for the complex adaptation of *C. canimorsus* to its ecological niche, the oral cavity of dogs.

*C. canimorsus* is a caphnophylic gliding Gram-negative bacterium from the *Bacteroidetes* phylum that is part of the normal flora of the dog mouth (1–5). Although it has not been reported to cause infections in dogs, it causes rare but severe infections in humans who are in contact with dogs (6, 7). The usual syndrome is septicemia, with mortality in the range of 50%. Patients are generally older than 40 years old, and roughly half of them have had a splenectomy or abused alcohol. However, the other half have had no medical history (for review, see references 8 to 10). *C. canimorsus* 5, a strain isolated from a patient with septic shock (11), has a lipopolysaccharide that triggers little inflammation (2, 11–13) and confers significant resistance to phagocytosis (2), antimicrobial peptides (12), and complement killing (2). Its genome, which consists of a single 2.5-Mb replicon (14), contains 13 so-called polysaccharide utilization loci (*PUL*) (15), a hallmark of *Bacteroidetes* bacteria (16). *PUL* typically encode polysaccharide-harvesting complexes made of several lipoproteins anchored in the outer membrane and that face the outside of the cell, as well as a TonB-dependent transporter. Some lipoproteins are devoted to glycan binding, while others are specific hydrolases. The archetype of such complexes is the *Bacteroides thetaiotaomicron* starch utilization system (*Sus*), initially described by Salzers and colleagues (17–20). *C. canimorsus* grows readily on blood agar plates and in coculture with animal cells but, surprisingly, it does not grow as well in the absence of animal cells or when it is separated from animal cells by a membrane, indicating that it relies on cells for nutrition (21). Unexpectedly, *C. canimorsus* does not damage the cells but instead deglycosylates the cellular...
surface-exposed N-glycoproteins via its Gpd complex, encoded by PUL5 (15, 21, 22). The Gpd complex consists of a SusC-like porin (GpdC), a SusD-like binding protein (GpdD), two ancillary lipoproteins (GpdEF), and an endo-β-N-acetylgalactosaminidase (GpdG) which cleaves the N-linked oligosaccharide after the first N-acetylglucosamine (GlcNAc) residue (22). The Gpd complex deglycosylates not only surface glycoproteins but also soluble glycoproteins, like human IgG (22) and transferrin (23). Deglycosylation is required for growth in coculture with animal cells but also for survival in the mouse (15, 21). In the initial description of the species, Brenner et al. (1) reported that C. canimorsus cultivated on blood agar plates appeared as bacilli but also in different forms, including longer rods, filaments that are often curved, spindle shaped, and coccoid (1), which suggests that C. canimorsus undergoes some form of phase variation.

In the present manuscript, we show that C. canimorsus 5 is a natural auxotroph for GlcNAc that is missing the glmM and glmU genes, which enable bacteria to convert glutamine into GlcNAc. The lack of GlcNAc leads to impaired peptidoglycan (PG) assembly and bacterial death, explaining the polymorphism seen on blood agar plates in the original description of the species (1). C. canimorsus harvests GlcNAc not only from host N-linked glycoproteins through the PUL5-encoded Gpd complex but also from mucin through the PUL9-encoded Muc complex. We propose that the GlcNAc auxotrophy, which contributes to virulence (21), results from the adaptation of C. canimorsus as a commensal of the guscan-rich dog mouth.

RESULTS

C. canimorsus polymorphism is due to N-acetylgalactosamine starvation. As reported early on (1), wild-type (wt) C. canimorsus 5 bacteria grown for 2 days on blood agar plates have different morphologies, ranging from filaments to coccoid forms (Fig. 1A). The coccoid forms strongly resemble the cells formed by Escherichia coli mutant strains that are unable to synthesize aminosugars (24). When glutamine or GlcNAc is not provided in the medium, these mutants cannot synthesize glucosamine-6-P, and they form spheroplasts due to a reduction in PG synthesis (24). We thus speculated that the polymorphism observed with C. canimorsus 5 could be due to a similar reason. In order to see whether this alteration of cell morphology was due to aminosugar starvation and not some kind of phase variation, we supplemented the blood agar plates with 0.005% GlcNAc. As shown in Fig. 1A, the addition of GlcNAc significantly reduced the formation of abnormal forms, indicating that aminosugar starvation, rather than phase variation, could be responsible for the observed polymorphism of wt C. canimorsus bacteria grown on blood agar plates.

PUL5 mutants grown on HEK293 cells are morphogenic due to a defect in peptidoglycan synthesis. In coculture with HEK293 cells, C. canimorsus 5 bacteria deprived of the PUL5-encoded Gpd complex reach a biomass that is 10-fold lower than that of wt C. canimorsus 5 bacteria (15, 22) (Fig. 1B). Since this growth defect can be rescued by supplementation of the medium with GlcNAc (15) (Fig. 1B), we hypothesized it could mimic the growth defect of wt bacteria on blood agar plates, and we examined the morphology of wt and PUL5 mutant bacteria after 24 h of coculture with HEK293 cells. Interestingly, while wt bacteria showed a bacillary form, PUL5 mutants appeared elongated, enlarged, or completely round (Fig. 1C). As shown in Fig. S1A and B in the supplemental material, PUL5 mutants progressively changed their morphology while they stopped dividing, suggesting that growth arrest and aberrant morphology are two linked phenomena. In contrast, in the presence of GlcNAc, PUL5 mutant bacteria had the same morphology as wt bacteria (Fig. 1C), suggesting that GlcNAc starvation affects PG biosynthesis. In order to see whether a defect in PG biosynthesis determines the same growth arrest and cell rounding, we added vancomycin (25) in the coculture of HEK293 cells and wt C. canimorsus 5. As shown in Fig. 1D (see also Movie S1 in the supplemental material), the addition of vancomycin transformed wt C. canimorsus 5 bacteria into round cells, similar to those observed with PUL5 mutant bacteria (see Movie S2 in the supplemental material). These results suggest that PUL5 mutant bacteria, which cannot harvest host N-linked glycans, stop growing and change their cell shape because starvation of aminosugars leads to defects in peptidoglycan synthesis.

Sialidase mutants also have a peptidoglycan synthesis defect. The PUL5-encoded Gpd complex works in concert with a periplasmic sialidase (SiaC) whose activity is crucial for the subsequent processing of the cleaved glycan chains (22). Since a siaC deletion mutant also shows reduced growth on HEK293 cells which can be rescued by addition of aminosugars (21), we asked whether the bacterial cell shape of a siaC mutant would also be affected. Indeed, a siaC mutant cocultivated with HEK293 cells for 24 h also showed aberrant morphologies: elongated, enlarged, and completely round (Fig. 1E). Addition of GlcNAc restored the normal cell shape (Fig. 1E). We concluded from this result that the inability to metabolize host N-linked glycans, due to either blockade of glycan chain cleavage and import (PUL5 deletion) or blockage of the first step of glycan processing in the periplasm (siaC deletion), results in aminosugar starvation. This starvation is likely to affect PG synthesis, leading to the appearance of aberrant cell shapes and growth arrest.

C. canimorsus 5 is missing the genes glmM and glmU. Since C. canimorsus 5 appears to be unable to synthesize GlcNAc, we performed a Delta BLAST search of its genome (26) for homologs of E. coli enzymes involved in PG biosynthetic pathways. As shown in Fig. 2A, we could identify genes encoding all the PG biosynthetic enzymes except for three: GlnS, GlnU, and GlnM, which convert fructose-6-P into UDP-N-acetylglucosamine (UDP-GlcNAc). GlnS is a glucosamine-6-P synthase responsible for the conversion of fructose-6-P into glucosamine-6-P; GlnM is a phosphoglucosamine mutase that converts glucosamine-6-P into glucosamine-1-P; GlmU is a bifunctional enzyme, with a C-terminal glucosamine-1-P–N-acetyltransferase domain that converts glucosamine-1-P into N-acetylglucosamine-1-P and an N-terminal N-acetylgalactosamine-1-P uridyltransferase domain that converts GlcNAc-1-P into UDP-GlcNAc (27). Since the lack of any of these enzymatic activities is lethal in E. coli (27), we assumed that it is the lack of these enzymes that makes C. canimorsus 5 dependent on exogenous GlcNAc for growth. If this hypothesis were correct, expression of these three enzymes in PUL5 mutant bacteria would enable them to synthesize GlcNAc and restore growth and bacterial shape in coculture with HEK293 cells. We cloned the three E. coli glm genes alone and in combination in a C. canimorsus expression vector (23) and monitored the growth and morphology of recombinant PUL5 mutants in coculture with HEK293 cells. PUL5-deficient bacteria expressing both GlmU and GlmM grew to the same extent as wt and showed a normal shape, indicating that these enzymatic activities allow C. canimorsus to synthesize GlcNAc (Fig. 2C and D).
The fact that the expression of GlmM and GlmU alone was sufficient to make PUL5 mutant bacteria independent of GlcNAc suggested that the C. canimorsus 5 genome encodes a glucosamine synthase (GlmS) that was not identified by our BLAST search. Furthermore, since the addition of GlcNAc to the coculture of HEK293 cells completely restored the growth of PUL5 mutant bacteria, we inferred that C. canimorsus is able to synthesize UDP-GlcNAc from GlcNAc, implying that a C. canimorsus 5 gene encodes a GlcNAc-1-P uridyltransferase. If this is correct, the phosphoglucosamine mutase GlmM and only the glucosamine-1-P-N-acetyltransferase domain of GlmU would be sufficient to compensate for the PUL5 deletion. To test this hypothesis, we generated two truncated GlmU protein variants: one with the first 77 residues deleted (GlmU del78), consisting only of the glucosamine-1-P-N-acetyltransferase activity, and one with the last 125 residues deleted (GlmU Tr331), consisting only of the N-acetylglucosamine-1-P-uridyltransferase activity (28) (Fig. 2B). We expressed in PUL5 mutants either GlmM and GlmU del78 or GlmM and GlmU Tr331.
Tr331, and we monitored the growth and bacterial shapes in coculture with HEK293 cells. As shown in Fig. 2C and D, the expression of GlmM and the GlmU glucosamine-1-P–N-acetyltransferase domain (GlmU del78) was sufficient to allow PUL5 mutant bacteria to behave similarly to wt. Expression of GlmM and the uridyltransferase domain of GlmU (GlmU Tr331) restored neither the growth nor the bacterial shape of PUL5 mutants (Fig. 2C and D). Quite expectedly, the expression of GlmM and GmU del78 had the same rescuing effect on siaC mutants (Fig. 2E and F). The slight but significant differences between the growth and the cell shape observed in PUL5 and siaC mutants expressing GlmM and GlmU del78 and the wt could be ascribed to the reduced enzymatic activity of GlmU del78 compared to the full-length GlmU, as already described by Pompeo and colleagues (28).

Taken together, these results indicate that the inability of C. canimorsus 5 to synthesize GlnNAc is due to the lack of two enzymes, a phosphoglucosamine mutase and a glucosamine-1-P–N-acetyltransferase.

C. canimorsus 5 encodes a new N-acetylglucosamine-1-P–uridylyltransferase. In order to identify the unknown N-acetyl-
glucosamine-1-P-uridyltransferase of *C. canimorsus* 5, we searched the genome for gene products with an enzymatic domain similar to that of the *E. coli* GlmU. It turned out that Ccan_15070 encodes a protein with a conserved glycosyltransferase domain (Fig. 3A). Attempts to knock out Ccan_15070 were unsuccessful unless *C. canimorsus* 5 could synthesize the *E. coli* GlmU Tr331 protein, suggesting that Ccan_15070 is an essential gene and indeed encodes an N-acetylglucosamine-1-P-uridyltransferase.

We then generated a PUL5 Ccan_15070 double-knockout strain complemented with the *E. coli* glmM and glmU genes and monitored growth and cell morphology. As shown in Fig. 3B and C, the strain grew to the same extent as wt and had a normal cell shape, indicating that the deletion of Ccan_15070 could be fully complemented by the N-acetylglucosamine-1-P-uridyltransferase activity of the *E. coli* GlmU.

Finally, to test the Ccan_15070 enzymatic activity, we expressed the protein with a C-terminal Strep tag in *E. coli* and added purified Ccan_15070 to a solution of GlcNAc-1-P and UTP. The analysis of the reaction products via high-performance liquid chromatography (HPLC) confirmed the production of UDP-GlcNAc (Fig. 3D). Thus, Ccan_15070 encodes an N-acetylglucosamine-1-P-uridyltransferase, which we named GlmUCc.

GlcNAc is also harvested through a PUL-encoded system targeting mucin. The loss of the capacity to synthesize aminosugars during evolution implies that *C. canimorsus* relies on aminosugars from the dog mouth. In the oral cavity, one of the most abundant glycoproteins is mucin, which is heavily glycosylated and harbors mainly O-linked glycans. The PUL5-encoded Gpd system can thus not harvest the mucin sugars. In order to determine if the aminosugars from mucin O-glycans can be used to synthesize PG, we took advantage of the fact that PUL5 mutant bacteria grow less well than wt in coculture with HEK293 cells. Hence, we tested whether the addition of purified bovine submaxillary mucin restored the growth of PUL5 bacteria. As shown in Fig. 4G and H,
FIG 4  PUL9 gene products allow mucin degradation and deglycosylation. (A) Schematic representation of the PUL9 locus. (B) Schematic drawing of the PNA lectin target oligosaccharides. (C) PNA lectin staining of human saliva after incubation in the presence of wt C. canimorsus 5 and the 13 PUL knockout strains. The upper band represents mucin, while the lower band represents sIgA. (D) Saliva PNA staining after incubation with wt bacteria, PUL9, and Ccan_17430 knockout bacteria. (E) Saliva PNA staining after incubation with wt bacteria, PUL9, or PUL9 bacteria expressing the only Ccan_17430 protein. (F) Western blot analysis of human mucin 7 in saliva after incubation with wt bacteria, PUL9, or PUL9 bacteria expressing the only Ccan_17430 protein. See also Fig. S3 in the supplemental material. (G) Growth after 23 h of wt (C. canimorsus 5), PUL5 (ΔPUL5), PUL9 (ΔPUL9), and PUL5/PUL9 (ΔPUL5 ΔPUL9) mutant bacteria in the presence of HEK293 cells with or without bovine submaxillary gland mucin (1 mg/ml) or GlcNAc (0.005%) (MOI, 0.005). The averages from three independent experiments are shown. Error bars represent 1 standard deviation from the mean. (H) Bright-field microscopy picture of PUL5 (ΔPUL5) mutant bacteria grown for 23 h on HEK293 cells in the presence of bovine mucin (1 mg/ml) (MOI, 0.05). (I) Functional model of mucin degradation and deglycosylation. Mucin is bound at the bacterial surface by the MucCDEG surface protein complex. The MucG protease cleaves mucin into glycosylated peptides that are imported into the periplasm through the MucC pore. Terminal sialic acid is cleaved by sialidase (SiaC), and terminal sulfate groups are removed by the sulfatase. The glycans are further processed by the sequential activity of several periplasmic exoglycosidases that allow the liberation of monosaccharides.
addition of bovine mucin fully restored the growth and the morphology of PUL5 mutant bacteria. To ascertain that salivary mucins are deglycosylated by C. canimorsus 5, we incubated bacteria with filtered fresh human saliva and we analyzed the salivary glycoproteins before and after incubation, by using sodium dodecyl sulfate-polyacrylamide gel electrophoresis followed by staining with peanut agglutinin (PNA). PNA recognizes terminal galactose linked to GalNAc in O-linked oligosaccharides, a glycan structure named T-antigen (Fig. 4B) that is abundant in mucins (29). A saliva glycoprotein with a molecular mass around 100 to 140 kDa turned out to be deglycosylated after incubation with wt C. canimorsus 5 (Fig. 4C). In contrast, the glycosylation state of a protein with a molecular mass of around 60 kDa was unaffected (Fig. 4C). LC-mass spectrometry (LC/MS) analysis of the two proteins identified the higher-molecular-mass protein as salivary mucin (MUC7 or MG2; Uniprot accession number Q8tax7) (data not shown). This indicated that C. canimorsus 5 specifically targeted MG2 glycans rather than the slgA ones. We reasoned that one of the 13 PUL mutants could be responsible for the T-antigen cleavage from MG2, and we tested the 13 C. canimorsus 5 PUL knockout mutants (15). Of all the PUL mutants, only PUL9 knockout bacteria were impaired in MG2 T-antigen deglycosylation (Fig. 4C). To ensure that mucin utilization was PUL9 dependent, we generated a PUL5 PUL9 double-knockout strain and tested whether addition of mucin restored the growth of this mutant. As expected, addition of mucin did not restore the growth defect of the double mutant, indicating that mucin degradation and utilization are PUL9 dependent (Fig. 4G).

PUL9 encodes a bona fide Sus-like complex. PUL9 encodes seven proteins (Ccan_17420 to Ccan_17490) (Fig. 4A). The neighboring Ccan_17470 and Ccan_17450 genes encode typical SusC-like (TonB-dependent receptor) and SusD-like proteins. Ccan_17480 is annotated as a β-galactosidase, while Ccan_17490 is a putative sulfatase. Both carry a signal peptide I and hence are predicted to be periplasmic. Ccan_17420 is a predicted lipoprotein annotated as a carboxypeptidase. Ccan_17430 is predicted to be a mucinase (30) with two BACON (Bacteroidetes-associated carbohydrate often N-terminal domain) modules (31) and one F5/8 type C (32) carbohydrate binding module. Ccan_17440 is the most abundant PUL-encoded protein of the C. canimorsus 5 sur- 
forme (15) and also harbors an F5/8 type C carbohydrate binding module. Ccan_17430 (mucinase) and Ccan_17450 (SusD-like) were also found in the surforme of C. canimorsus 5 (15). In order to determine whether all these proteins form a PUL5 complex, we tagged the C terminus of the porin Ccan_17470 and carried out a His-Strep tandem affinity purification (see Fig. S2 in the supplemental material). The LC/MS analysis of the eluate identified 35 proteins (see Table S1 in the supplemental material), among which were Ccan_17450, Ccan_17440, and Ccan_17430. Thirteen proteins were identified in the mock eluate, of which 11 were also present in the Ccan_17470–Strep-His eluate. No PUL9-encoded product was identified in the mock eluate (see Table S1). These data suggest that Ccan_17470, Ccan_17450, Ccan_17440, and Ccan_17430 form a complex at the bacterial surface. Hence, PUL9 encodes a bona fide Sus-like complex, and we propose to call MucC the SusC-like protein, MucD the SusD-like protein, MucG the putative mucinase, and MucE the four surface proteins (Fig. 4A). Interestingly, among the proteins that eluted with the Muc complex, we also identified a sialidase. This periplasmic enzyme was previously found to be associated with the Gpd complex and involved in N-glycan processing (22). Thus, we could also envision involvement of this protein in mucin glycan processing.

The hydrolytic enzyme from the complex is not a glycan hydrolase but a mucin peptidase. As already mentioned, Ccan_17430 is predicted to be a mucinase (30) with two BACON (31) and one F5/8 type C (32) carbohydrate binding modules.

Since PUL-encoded complexes are usually devoted to the hydrolysis of complex carbohydrates (33), the presence of a protein hydrolase was unexpected. In order to verify the function of Ccan_17430, we incubated human saliva with wt and Ccan_17430 mutant bacteria and monitored mucin glycosylation via PNA staining. Deletion of Ccan_17430 abolished mucin deglycosylation (Fig. 4D). Furthermore, expression of Ccan_17430 in a PUL9 deletion background led to a complete loss of mucin staining by PNA, suggesting that Ccan_17430 alone is responsible for the observed phenotype (Fig. 4E). We then monitored the presence of mucin MG2 by Western blotting and found that it was dramatically decreased, indicating that Ccan_17430 indeed encodes a protease that cleaves mucin (Fig. 4F). In agreement with this, purified recombinant Ccan_17430 fully degraded MG2 from human saliva (see Fig. S3 in the supplemental material). Addition of EDTA completely blocked the mucinase activity, indicating that Ccan_17430 is a metallopeptidase. Removal of the N-terminal conserved BACON and F5_F8 type C glycan binding domains led to reduced mucin degradation activity, suggesting that these domains could play a role in mucin binding, even though the truncated protein was still able to bind and degrade MG2.

Thus, the PUL9-encoded system appears to be quite different from the canonical ones, such as the Sus (33) and Gpd (22) systems, where a glycosylhydrolase at the bacterial surface is responsible for the cleavage of complex sugars like starch or N-linked glycans. In the case of PUL9, it is a protease that attacks mucin at the bacterial surface, generating glycosylated peptides.

**DISCUSSION**

Here, we addressed the question of the phenotypic polymorphism of C. canimorsus (1), and we found that it is linked to the adaptation of C. canimorsus to its host. We showed that C. canimorsus does not undergo a kind of phase variation but, instead, endures aminosugar starvation, which leads to bacterial growth arrest and cell shape change. This phenotype is a consequence of the blockade of peptidoglycan synthesis. These findings are explained by the fact that C. canimorsus relies on exogenous aminosugars because it is unable to synthesize its own GlcNAc. Bacteria generally synthesize GlcNAc starting from fructose–6-P through a series of metabolic steps (Fig. 2A). In *silo* analysis of the genome of C. canimorsus 5 did not identify orthologs of all the necessary genes. Furthermore, coexpression of the *E. coli* GlmM and the GlmU glucosamine-1-P–N-acetyltransferase domain completely restored the growth and cell morphology of PUL5 mutant bacteria. We conclude from these results that C. canimorsus is unable to synthesize GlcNAc because of the lack of these two enzymatic activities. Furthermore, we identified and characterized a novel N-acetylglucosamine-1-P–uridylyltransferase that is able to synthesize UDP-GlcNAc from GlcNAc-1-P. Differently from *E. coli* and other well-studied bacteria, where the glucosamine-
1-P- N-acetyltransferase and the N-acetylglucosamine-1-P-uridylytransferase activities are found in the same protein (GlmU) (27), the C. canimorsus enzyme (GlmU_Cc) is monofunctional, harboring only the N-acetylglucosamine-1-P-uridylytransferase activity and missing the glucosamine-1-P-N-acetyltransferase one.

We reasoned that the loss of capacity to synthesize GlcNAc could be a common feature in the phylum Bacteroidetes. We performed an in silico search for orthologs of GlmU_Cc in bacteria and found that, indeed, in Bacteroidetes, 30 out of 45 genera have a monofunctional ortholog of GlmU_Cc (see Fig. S4 in the supplemental material). This suggests that, like C. canimorsus 5, most Bacteroidetes are unable to synthesize GlcNAc and so rely on exogenous aminosugars.

The loss of the capacity to synthesize N-acetylated aminosugars during evolution implies that C. canimorsus relies on aminosugars from its ecological niche, the dog mouth. In agreement with this hypothesis, we showed that C. canimorsus 5 degrades salivary mucin, a protein that is heavily O-glycosylated and very abundant in saliva. Mucin degradation was found to depend on a mucin protease (MucG), which is, unexpectedly, part of a Sus-like surface complex encoded by the PUL9 locus. Nevertheless, mucin degradation rescued the growth of a PUL5 mutant, indicating that mucin aminosugars can be taken up and metabolized. Given the structure of mucin, a protease is probably as efficient as a glycosyl hydrolase in liberating carbohydrates. This is nevertheless the first example of a PUL-encoded complex endowed with a protease activity. It is also the first example of an oral commensal that utilizes mucin via a PUL-encoded system. The C. canimorsus 5 PUL9 shows high similarity in gene composition and organization to the Bacteroides thetaiotaomicron BT4240-50 PUL, which is induced by the presence of mucin O-glycans (16). Like PUL9, besides the SusC and SusD-like proteins, the BT4240-50 PUL encodes a mucinase (BT4244) that has been shown to target bovine mucin in vitro (30). However, the C. canimorsus 5 and BT mucinases share only 27% identity. Both loci encode periplasmic β-galactosidases (Ccan_17480 and BT4241) that are 34% identical and likely to be involved in mucin glycans processing. In B. thetaiotaomicron, the β-galactosidase has been found to be constitutively expressed (16). PUL9 also encodes a putative periplasmic sulfatase (Ccan_17490), which is similar (47% identity) to the mucin-desulfating sulfatase BT3177 (34). We reason that these two enzymes could act on mucin glycans once the glycosylated peptides are transferred into the periplasm through the Ccan_17470 MucC pore. Mucin O-glycans are mainly composed of core 1- and core 2-type O-glycans that can harbor terminal sulfate groups (SO₃⁻) and sialic acid residues (29). One can thus envision a mucin degradation and deglycosylation model where mucin is first bound at the bacterial surface by the MucCDGE surface complex, then cleaved by the activity of the MucG mucinase, which liberates glycosylated peptides that are translocated into the periplasm through the MucC pore. Once in the periplasm, the glycopeptides are subsequently processed by periplasmic enzymes; the removal of terminal sialic acid residues by sialidase (Ccan_04780) and of terminal sulfate groups by the Ccan_17490 sulfatase make galactose accessible to β-galactosidases (Ccan_17480, Ccan_01530, and Ccan_15520) whose activities allow the β-N-acetylglucosaminidases (Ccan_16820, Ccan_20090, Ccan_03860, and Ccan_04040) to liberate the GlcNAc residues present in core 2-type O-glycans (Fig. 4I). The PUL9 locus also encodes, downstream from MucG, a putative lipoprotein with a carboxypeptidase domain (Ccan_17420) (Fig. 4A). The presence of this enzyme in the locus suggests an involvement in the processing of mucin glycopeptides once in the periplasm, but this was not investigated. Our data thus characterize a novel PUL-encoded system which allows harvesting of mucin glycan by the activity of a mucinase at the bacterial surface.

Overall, this work illustrates how a commensal that has evolved to adapt to a specific niche becomes dependent on that niche. C. canimorsus adapted to the dog mouth by acquiring or evolving two Sus-like machines that are tuned to harvest sugars from saliva and possibly from the oral epithelium. Since these machines provide abundant aminosugars, C. canimorsus could lose the capacity to synthesize GlcNAc. Interestingly, C. canimorsus has two Sus-like machines, allowing it to retrieve aminosugars. One of these, the Muc machine, can only be of use in the mouth, but the other one, the Gpd machine, is able to sustain growth in other compartments of a host and probably represents a virulence factor (15, 21).

Recently, it was shown that several members of the genus Bacteroides, which are able to utilize different polysaccharides, liberate polysaccharide breakdown products that can be consumed by bacteria from other species unable to grow on the polysaccharide alone (35). These findings indicate that, in the human gut microbial ecosystem, there are syntrophic interactions between different bacteria. Similar syntrophic interactions might well occur in the dog mouth, since the biology of C. canimorsus resembles that of B. thetaiotaomicron, but the dog oral ecosystem is quite different from the intestine in the sense that digestion does not start in the dog mouth and hence the microbial competition may be more fierce. In addition, PUL5-deficient C. canimorsus cannot be cross-fed by wt C. canimorsus (21), which can be explained by the mechanisms of uptake and processing of host glycans by the Gpd apparatus where N-glycans are cleaved and likely immediately imported into the periplasm (22). It is likely that the same applies to the Muc apparatus, but this was not investigated. For C. canimorsus, we can thus envision a scenario that is different from the one of the human gut. Capnocytophaga could have a more “selfish” behavior, as it harvests, takes up, and utilizes its own glycans, avoiding release of polysaccharide breakdown products, which could be utilized by competitor bacteria.

**MATERIALS AND METHODS**

**Bacterial strains and growth conditions.** (i) Conventional bacterial growth conditions and selective agents. The strains used in this study are listed in Table S2 in the supplemental material. Escherichia coli strains were routinely grown in LB broth at 37°C. C. canimorsus bacteria were routinely grown on heart infusion agar (HIA; Difco) supplemented with 5% sheep blood (Oxoid) for 2 days at 37°C in the presence of 5% CO₂. To select for plasmids, antibiotics were added at the following concentrations: 10 μg/ml erythromycin (Em), 10 μg/ml cefoxitin (Cf), 20 μg/ml gentamicin (Gm), 100 μg/ml ampicillin (Ap), 50 μg/ml kanamycin (Km), and 10 μg/ml tetracycline (Tc). GlcNAc (catalog number A8625; Sigma) was added at a final concentration of 0.005% when indicated.

(ii) Growth of C. canimorsus 5 bacteria with HEK293 cultured cells. HEK293 cells were cultured in Dulbecco’s modified Eagle’s medium (DMEM; Invitrogen) with 10% (vol/vol) fetal calf serum (Invitrogen) and 1 mM sodium pyruvate (cDMEM). Cells were grown in medium without antibiotics in a humidified atmosphere enriched with 5% CO₂ at 37°C. Bacteria were harvested by gently scraping colonies off the agar surface and resuspended in phosphate-buffered saline (PBS). A total of 1 × 10⁴ bacteria were incubated with 2 × 10⁵ HEK293 cells (multiplicity of infection [MOI], 0.05) in a final volume of 1 ml medium devoid of antibiotics for 23 h.
(iii) Monitoring bacterial growth in HEK293 cell cultures. A total of $1 \times 10^6$ bacteria were added to $2 \times 10^5$ HEK293 cells (MOI, 0.05) in a final volume of 1 ml medium devoid of antibiotics. At different times, the medium containing bacteria and cells was collected and serially diluted. Dilutions were plated on HIA 5% sheep blood plates. After 48 h of incubation at 37°C with 5% CO₂, bacterial colonies were counted.

(iv) Growth in HEK293 cell cultures in the presence of bovine mucin. A total of $1 \times 10^6$ bacteria were incubated with $2 \times 10^5$ HEK293 cells (MOI, 0.05) in a final volume of 1 ml medium devoid of antibiotics and with 1 mg of mucin from bovine submaxillary glands (catalog number M3895; Sigma-Aldrich).

Mutagenesis and allelic exchange. Mutagenesis of C. canimorsus 5 wt was performed as described in reference 36 with slight modifications. The oligonucleotides used in this study are listed in Table S4 in the supplemental material. Briefly, replacement cassettes with flanking regions spanning approximately 500 bp homologous to the direct target gene-framing region were constructed with a three-fragment overlapping-PCR strategy. First, two PCRs were performed on 100 ng of C. canimorsus 5 genomic DNA with primers A and B for the upstream flanking regions and with primers C and D for the downstream regions. Primers B and C contained an additional 5’-20-nucleotide extension homologous to the resistance erm(F) insertion cassette or to the tet(Q) insertion cassette. The erm(F) resistance cassette was amplified from plasmid pmm106 DNA (36) with primers 5502 and 5503. The tet(Q) resistance cassette was amplified from plasmid pBl3 DNA (see Table S3 in the supplemental material) with primers 7180 and 7181. All three PCR products were cleaned and then mixed in equal amounts for PCR using Phusion polymerase (Finzymes).

The initial denaturation was at 98°C for 2 min, followed by 12 cycles without primers to allow annealing and then elongation of the overlapping fragments (98°C for 30 s, 50°C for 40 s, and 72°C for 2 min). After the addition of external primers (A and D), the program was continued with 20 cycles (98°C for 30 s, 50°C for 40 s, and 72°C for 2 min 30 s) and finally 10 min at 72°C. Final PCR products consisted of the target gene with the erm(F) or tet(Q) insertion cassettes and were then digested with PstI and SpeI for cloning into the appropriate sites of the C. canimorsus suicide vector pMM25 (36). Resulting plasmids were transferred by RP4-mediated conjugative DNA transfer from E. coli S17-1 to C. canimorsus 5 to allow integration of the insertion cassette. Transconjugants were then selected for the presence of the erm(F) or tet(Q) cassettes on erythromycin or tetracycline plates, respectively, and then checked for sensitivity to cefoxitin. Correct cassette insertions were confirmed by sequencing.

Construction of complementation and expression plasmids. Plasmids and oligonucleotides used in this study are listed in Tables S3 and S4 in the supplemental material.

(i) Construction of GlmM-, GlmU-, and GlmS-expressing vectors. Full-length glmS and glmU were amplified from E. coli MG1655 DNA with primers 7412/7413 and 7406/7407, respectively, and cloned into plasmid pPM5 (23) into NcoI and XhoI restriction sites, leading to plasmids pFR11 and pFR12. Full-length glmU and glmS were coamplified from E. coli MG1655 DNA by using primers 7406 and 7413 and cloned into pPM5 by using NcoI and XhoI restriction sites, leading to plasmid pFR13. Full-length E. coli glmM was amplified from E. coli MG1655 DNA with primers 7415 and 7416 and cloned downstream of the glmS gene into pFR11 by using XhoI and SpeI restriction sites, leading to plasmid pFR14. Full-length E. coli glmM was amplified from E. coli MG1655 DNA with primers 7415 and 7416 and cloned downstream of the glmS gene into pFR13 by using XhoI and SpeI restriction sites, leading to plasmid pFR15. Full-length E. coli glmM was amplified from E. coli MG1655 DNA with primers 7415 and 7416 and cloned downstream of the glmU gene into pFR12 by using XhoI and SpeI restriction sites, leading to plasmid pFR16. E. coli glmU with the first 77 codons deleted (GlM(U del78)) was amplified with primers 7422 and 7407 and cloned into pFR16 by using NcoI and XhoI restriction sites, leading to plasmid pFR17. E. coli glmU with codons 332 to 456 deleted (GlM(U Tr331)) was amplified from E. coli MG1655 DNA with primers 7406 and 7421 and cloned into pFR16 by using NcoI and XhoI restriction sites, leading to plasmid pFR18.

(ii) Construction of GlmUcC- and MucG-overexpressing vectors. To generate GlmUcC- with a C-terminal Strep tag, the glmUcC gene (Ccan_15070) was amplified with primers 7442 and 7450 and cloned into pET22B (+) by using NcoI and XhoI restriction sites, leading to plasmid pFR19.

mucG (Ccan_17430), devoid of the first 21 codons and encoding a C-terminal Strep tag, was amplified with primers 7020 and 7021 and cloned into pET22B (+) by using NcoI and XhoI restriction sites, leading to plasmid pFR22. mucG (Ccan_17430), devoid of the first 355 codons and encoding a C-terminal Strep tag, was amplified with primers 7019 and 7021 and cloned into pET22B (+) by using NcoI and XhoI restriction sites, leading to plasmid pFR23.

(iii) Construction of mucG complementation vector. The 340-bp region upstream of the mucC (Ccan_17470) starting codon sequence, containing the putative mucC promoter, was cloned into pMM47A.1 vector (36) by using SalI and Ncol restriction sites, leading to plasmid pFR20. Full-length mucG (Ccan_17430) was amplified with primers 6923 and 6925 and cloned into pFR20 by using NcoI and XhoI restriction sites, leading to plasmid pFR21.

(iv) Engineering MucC-Strep-His-D-, E-, and -G and MucC-, D-, E-, and -G expression vectors. To engineer MucC (Ccan_17470) with a C-terminal Strep tag, the gene mucC (Ccan_17470) was amplified with primers 6842 and 6786 and cloned into pFR20 with NcoI and XhoI restriction sites in frame with the 6× His in the vector, leading to plasmid pFR24.

Full-length mucD, mucE, and mucG (Ccan_17450-30) genes were amplified with primers 6846 and 6845 and cloned into pFR24 downstream from mucC-Strep-His by using Spel restriction sites, leading to plasmid pFR26. Full-length mucC (Ccan_17470) was amplified with primers 6842 and 6805 and cloned into pFR20 with NcoI and XhoI restriction sites, leading to plasmid pFR25. Full-length mucD, mucE, and mucG (Ccan_17450-30) were amplified with primers 6846 and 6845 and cloned into pFR25 downstream from mucC with Spel restriction sites, leading to plasmid pFR27.

(v) Construction of a TetQ expression vector. The tet(Q) gene was amplified from pMM104 (36) DNA with primers 7156 and 7157 and cloned into pPM5 by using NcoI and XhoI restriction sites, leading to plasmid pFL38.

Microscopy pictures and movies. For each experiment, bacterial suspensions were deposited onto 1% agarose pads. All microscopy images were captured with an Axioscop (Zeiss) microscope with an Orca-Flash 4.0 camera (Hamamatsu) and Zen 2012 software (Zeiss). Images were processed using ImageJ software.

(i) Growth curves of bacteria in cocultures with HEK293 cells. For each time point, 10-μl aliquots of bacterium-cell suspensions were transferred onto a 1% agarose pad with PBS, and images were captured on the microscope.

(ii) Vancomycin treatment of wt bacteria in cocultures with HEK293 cells. wt bacteria were grown for 12 h in HEK293 cell cultures. Vancomycin hydrochloride was then added at a final concentration of 5 μg/ml. Bacteria were collected after 4 h, 10 μl of the bacterium-cell suspension was transferred onto a 1% agarose pad made with PBS, and images were collected on the microscope.

(iii) Microscopy movies of wt bacteria treated with vancomycin. Bacteria were grown to exponential phase on HEK293 cell cultures for 12 h in 1-ml final volumes and then resuspended. Ten microliters of the bacterium-cell suspension was transferred onto a 1% agarose pad made with DMEM with 10% (vol/vol) heat-inactivated human serum (S1-Liter; EMD Millipore) and vancomycin hydrochloride (Sigma-Aldrich) at a final concentration of 5 μg/ml. Time-lapse movies were generated by imaging the cells every 2 min for 12 h. During all microscopy experiments, the stage was kept at 37°C to allow bacterial growth. Movies were generated from time-lapse images using Zen 2012 software (Zeiss).
(iv) Microscopy movies of PUL5 mutant bacteria. PUL5 bacteria were grown to exponential phase in HEK293 cell cultures for 12 h in 1-ml final volumes and then resuspended. Ten microliters of the bacterium-cell suspension was transferred onto a 1% agarose pad made with DMEM with 0.5% (vol/vol) heat-inactivated human serum (S1-Liter; EMD Millipore), a condition we found allowed a few bacterial generations before leading to growth arrest and shape change. Time-lapse movies were generated by imaging the cells every 2 min for 12 h. During all microscopy experiments, the stage was kept at 37°C to allow bacterial growth. Movies were generated from time-lapse images by using Zen 2012 software (Zeiss).

Recombinant Ccan_15070 (GlmU_Cc) and Ccan_17430 (MucG) overexpression and purification. E. coli BL21 bacteria harboring plasmid pFR19, expressing a C-terminal Strep-tagged Ccan_15070, or pFR22, expressing a C-terminal Strep-tagged Ccan_17430 or harboring plasmid pFR23, expressing a C-terminal Strep-tagged Ccan_17430 devoid of the BACON and F5/8 type C domains, were inoculated at an optical density at 600 nm (OD600) of 0.05 in 300 ml LB medium with ampicillin and grown for 2 h at 37°C with agitation. To induce protein expression, isopropyl-β-D-thiogalactopyranoside was added to a final concentration of 1 mM, and cells were grown for 2 more hours. Bacteria were collected by centrifugation at 5,000 × g, resuspended in PBS, and lysed in a French press. Lysates were centrifuged at 15,000 × g for 15 min, and supernatants were collected.

Each supernatant was loaded into a column containing 1 ml of a 50% slurry (0.5 ml CV) Strept-Tactin Superflow resin (catalog number 2-1200-02; IBA). The flowthrough was reloaded into the resin 2 more times. The resin was then washed 4 times with 10 CV of buffer W (100 mM Tris, 150 mM NaCl, 1 mM EDTA; pH 8), and proteins were eluted in 3 steps with 0.5 ml elution buffer (100 mM Tris, 150 mM NaCl, 1 mM EDTA, 2.5 mM dethiobiotin; pH 8). The proteins present in the elution fractions were visualized by Coomassie staining and immunoblotting with mouse anti-Strep tag antibodies (MCA2489P; AbD Serotec).

UDP-GlcNAc synthesis by recombinant Ccan_15070 (GlmU_Cc). The activity and substrate specificity of the enzyme Ccan_15070 was assessed by incubating 30 µg of purified recombinant Ccan_15070 protein with 1,245 µg of UTP (catalog number U6625; Sigma) and 500 µg of GlcNAc-1-P (A2142; Sigma) in 1 ml 100 mM Tris-HCl (pH 7.5) and 10 mM MgCl2 solution for 16 h at 37°C. As a control, the same amount of substrate was incubated in the absence of recombinant purified Ccan_15070. UDP-GlcNAc production was detected by HPLC using a Waters 600 E device with a C18 Atlantis T3, 5 µm, 4.60- by 250-mm column. Elution was done with 50 mM triethylamine acetic acid, pH 6.8, 0.5% CH3CN; detection was at 262 nm and at a flow rate of 1 ml/min.

Human salivary mucin deglycosylation and degradation analyses. Fresh human saliva was collected from healthy volunteers and sterilized by filtration using 0.22-µm filters (Millipore). Bacteria were collected from blood agar plates and resuspended in PBS at an optical density at OD600 of 1. Aliquots of 100 µl of bacterial suspensions (corresponding to 5 × 107 bacteria) were incubated with 100 µl of human saliva for 240 min at 37°C. As a negative control, 200 µl of 1.2-diluted human saliva solution alone was incubated for 240 min at 37°C. Samples were then centrifuged for 5 min at 13,000 × g, and supernatant and collected and loaded in a 10% SDS gel. Samples were analyzed by lactic staining with PNA according to the manufacturer’s recommendations (DIG glycan differentiation kit; Roche). Mucin deglycosis was monitored by Western blotting with mouse anti-human Muc2 antibodies (ab55542; Abcam).

Mucinase assays with purified recombinant MucG. One hundred microliters of human saliva was incubated with 1 µg of purified MucG protein for 1.5 or 3.5 h at 37°C in the presence or absence of 50 mM EDTA. Samples were loaded onto 10% SDS-PAGE gels and analyzed by lactic staining with PNA according to the manufacturer’s recommendations (DIG glycan differentiation kit; Roche).

Muc proteins copurification. C. canimorsus 5 ΔmucC bacteria harboring plasmid pFR26 expressing a C-terminal Strep-ΔHis double-tagged MucC, or harboring plasmid pFR27, expressing MucC without any tag (mock), were grown on sheep blood plates for 2 days at 37°C in the presence of 5% CO2. Bacteria from plates were scraped and lysed in 35 ml of 25 mM Tris-HCl, 150 mM NaCl, 0.2% Triton, 1% NP-40, 1% sodium deoxycholate, pH 7.6.

For His affinity purification, the lysates were clarified by centrifugation (10 min at 18,500 × g at room temperature), and the supernatant was diluted 1:2 in PBS, 10 mM imidazole in the presence of protease inhibitor (cOmplete Mini, EDTA-free protease Inhibitor cocktail tablets; Roche). Aliquots of 3.5 ml of a 50% slurry of chelating Sepharose fast flow beads (GE Healthcare) were first coupled to Ni2+ according to the manufacturer’s instructions, and then 1.75 ml of resin was added to the solution and the mixture was incubated overnight at 4°C on a rotating wheel. The solution was then loaded into a column, and the resin was washed first with 25 column volumes (CV) of high-salt buffer (50 mM Tris, 500 mM NaCl; pH 8) and then with 5 CV of low-salt buffer (50 mM Tris, 100 mM NaCl; pH 8). Proteins were then eluted from the resin with 2 CV of elution buffer (50 mM Tris, 100 mM NaCl, 350 mM imidazole; pH 8). The material eluted from the Ni2+ column was then diluted 1:2 in PBS, and 1 ml of a 50% slurry (0.5 ml CV) of Strept-Tactin superflow resin (catalog number 2-1200-02; IBA) was added. The solution was then incubated overnight at 4°C on a rotating wheel. Next, the solution was loaded into a column, and the flowthrough was reloaded into the resin 2 more times. The resin was then washed 4 times with 10 CV of buffer W (100 mM Tris, 150 mM NaCl, 1 mM EDTA; pH 8), and proteins were eluted in 3 steps with 0.5 ml of elution buffer (100 mM Tris, 150 mM NaCl, 1 mM EDTA, 2.5 mM dethiobiotin; pH 8). Tagged MucC was detected by immunoblotting using mouse anti-His antibody (catalog number 27-4710-01; Amersham Bioscience), and the proteins present in the elution fractions were identified by LC/MS.

LC/MS identification of proteins in the elution fractions. Proteins eluted from the MucC-Strep-His and MucC (mock) pullouts were separated on a 12% SDS gel. The proteins were silver stained and excised with a razor blade. Proteins were trypsin digested and analyzed by LC/MS as described in reference 15.

Phylogenetic analyses of GlmU_Cc. The orthologous cluster of KEGG: ml:CanC_15070 (GlmU_Cc) was initially searched in the SSDB (KEGG) database within the bacterial kingdom with a best-best rule. Only hits above a threshold of 150 were further considered (137 hits). Because of low similarity levels between GlmU_Cc and most of its bifunctional homologs, the cluster of orthologs was extended by searching the SSDB with the bifunctional GlmU ortholog, with the highest similarity score (i.e., the closest bifunctional relative of GlmU_Cc: dте: Dester_1461, 27% similarity). The second search clustered 2,403 hits (including ml:CanC_15070). The two merged sets resulted in a cluster of 2,532 protein sequences that were submitted for Pfam domain identification (http://pfam.xfam.org/search#tabview=tab1). A total of 534 consensual 16S rRNA sequences (derived from 2 to 20,843 [Bacillus] individual sequences from the Ribosomal Database Project website [37]) from the genera identified here were used to determine a 16S rRNA-based phylogeny. The evolutionary history was inferred by using the neighbor-joining method (38). The bootstrap consensus tree inferred from 1,000 replicates (39) was taken to represent the evolutionary history of the taxa analyzed (40). Branches corresponding to partitions reproduced in less than 70% bootstrap replicates were collapsed. The analysis involved 554 nucleotide sequences. All positions with less than 50% site coverage were eliminated. That is, fewer than 50% alignment gaps, missing data, and ambiguous bases were allowed at any position. There were a total of 1,453 positions in the final data set. Evolutionary analyses were conducted with MEGA6 (41).

SUPPLEMENTAL MATERIAL

Supplemental material for this article may be found at http://mbio.asm.orglookup/suppl/doi:10.1128/mBio.02507-14/-/DCSupplemental.

Figure S1, TIF file, 3 MB.


