Functional characterization of exopolyphosphatase/guanosine pentaphosphate phosphohydrolase (PPX/GPPA) of Campylobacter jejuni

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Abbreviations: PPX, exopolyphosphatase; GPPA, guanosine pentaphosphate phosphohydrolase; poly-P, inorganic polyphosphate; (p)ppGpp, guanosine tetra/pentaphosphate; CPS, capsular polysaccharide; LOS, lipooligosaccharide; MEM, minimum essential media

The inorganic polyphosphate (poly-P) is a key regulator of stress responses and virulence in many bacterial pathogens including Campylobacter jejuni. The role of exopolyphosphatases/guanosine pentaphosphate phosphohydrolases (PPX/GPPA) in poly-P homeostasis and C. jejuni pathobiology remains unexplored. Here, we analyzed deletion mutants (Δppx1, Δppx2) and the double knockout mutant (dkppx), all Δppx mutants exhibited increased capacity to accumulate poly-P; however only Δppx1 and dkppx mutants showed decreased accumulation of ppGpp, an alarmone molecule that regulates stringent response in bacteria, suggesting potential dual role for PPX1/GPPA. Nutrient survival defect of Δppx mutants was rescued by the supplementation of specific amino acids implying that survival defect may be associated with decreased ppGpp and/or increased poly-P in Δppx mutants. The ppk1 and spoT were upregulated in both Δppx1 and Δppx2 suggesting a compensatory role for SpoT and Ppk1 in poly-P and ppGpp homeostasis. The lack of ppx genes resulted in defects in motility, biofilm formation, nutrient stress survival, invasion and intracellular survival indicating that maintaining a certain level of poly-P is critical for ppx genes in C. jejuni pathophysiology. Both ppx1 and ppx2 mutants were resistant to human complement-mediated killing; however, the dkppx mutant was sensitive. The serum susceptibility did not occur in the presence of MgCl₂ and EGTA suggesting an involvement of the classical or lectin pathway of complement mediated killing. Interestingly, the chicken serum did not have any effect on the Δppx mutants’ survival. The observed serum susceptibility was not related to C. jejuni surface capsule and lipooligosaccharide structures. Our study underscores the importance of PPX/GPPA proteins in poly-P and ppGpp homeostasis, two critical molecules that modulate environmental stress responses and virulence in C. jejuni.

Introduction

Campylobacter jejuni is a microaerophilic gram-negative curved bacterium that persistently colonizes the intestine of various livestock especially, poultry.¹ However, C. jejuni can infect humans and its infection is considered to be a prevalent cause of bacteria-mediated diarrheal disease worldwide.² Campylobacteriosis is often a self-limiting disease characterized by fever, abdominal pain, vomiting, and sometimes bloody diarrhea. In addition, fatal post-infectious complications like Guillain–Barre Syndrome, Miller Fisher Syndrome, and reactive arthritis can also occur.³ Efforts to reduce campylobacteriosis in humans are directly linked to a better understanding of the pathobiology of C. jejuni; however, the molecular basis of such virulence mechanisms are not fully elucidated yet.

Inorganic polyphosphate (poly-P), a polymer of ten to hundreds of phosphate residues, linked by high-energy phosphoanhydride bonds, is conserved in every cell in nature.⁴ It can serve as a source of energy for synthesis of sugars, nucleosides, and proteins as well as an activating precursor for fatty acids, phospholipids, polypeptides, and nucleic acids. Poly-P is also a key regulator of bacterial survival, stress responses, host colonization, and virulence in many pathogenic bacteria including C. jejuni.⁵,⁶ Poly-P metabolism is governed by several specialized enzymes; Polyphosphate kinase-1 (PPK1) is responsible for the synthesis of long chain poly-P from ATP. Although the reaction

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is reversible, the synthesis of poly-P is favored. Polyposphate kinase-2 (PPK2) utilizes poly-P to generate GTP at a rate 75-fold greater than that of the poly-P synthesis from GTP. Exopolypophosphatases (PPXs) degrade poly-P into a smaller branch of inorganic phosphate. PPXs have been classified into two kinds; the classical PPX and the dual function PPX/GPPA. Classical PPX enzymes are primarily monofunctional and their essential role is in poly-P homeostasis. Classical PPXs have been shown to be involved in motility, biofilm formation, sporulation and providing resistance to complement-mediated killing in different bacteria. On the other hand, PPX/GPPA enzymes have both exopolypophosphatase and guanosine pentaphosphate (pppGpp) phosphorhydrolase activities to generate guanosine tetraphosphate (ppGpp), an alarmone that regulates stringent response in bacteria. In particular, PPX/GPPA enzymes play a supplemental role in ppGpp metabolism related to the bacterial starvation induced stringent response.

C. jejuni 81–176 encodes two potential dual function exopolypophosphatases ppx1/gppa (CJ81176_0377) and ppx2/gppa (CJ81176_1251). To date, no studies describe the role of these enzymes in poly-P hydrolysis and/or ppGpp synthesis as well as in C. jejuni pathophysiology. Since poly-P homeostasis is critical, and PPX/GPPA enzymes are involved in poly-P metabolism and possibly in ppGpp synthesis, we hypothesize that PPX/GPPA enzymes may have an important role in C. jejuni pathobiology and persistence in different environmental conditions. To test our hypothesis, we generated deletion mutants of ppx1 and ppx2 (Δppx1 and Δppx2) as well as the double knockout for both genes (Δppx1 Δppx2) and the analyses of these mutants suggest that ppx1/gppa contributes to both poly-P and ppGpp pools in C. jejuni and ppx/gppa genes are important for the C. jejuni stress responses and virulence related traits.

Results

The PPX/GPPA proteins of C. jejuni

The phylogenetic analysis using MEGA-5 (Center for Evolutionary Medicine and Informatics) showed that the C. jejuni PPX1/GPPA clustered with Helicobacter and Acrobacter whereas PPX2/GPPA with E. coli and Mycobacterium tuberculosis species (Fig. S1). Further ClustalW2, structure based sequence alignment using reference sequence of Aquifex aeolicus also showed that the C. jejuni PPX1/GPPA (CJ81176_0377) has both the catalytic residue E116 required for PPX activity and arginine residues (R16 and R258) necessary for GPPA activity (Fig. S2A), suggesting that this enzyme might mediate both poly-P and ppGpp hydrolysis. Conversely, C. jejuni PPX2/GPPA (CJ81176_1251) possesses the catalytic residues E103 and R12 but not R254 (Fig. S2A), indicating this enzyme is less likely a mediator of ppGpp hydrolysis. Moreover, the predicted three-dimensional (3D) structure of C. jejuni PPX1/GPPA showed closer alignment with the reference sequence from A. aeolicus PPX/GPPA than the PPX2/GPPA further suggesting a role for (Fig. S2B and C) PPX1/GPPA in ppGpp hydrolysis.

The PPX1/GPPA and PPX2/GPPA proteins are conserved across the different strains of C. jejuni with amino acid sequence identity ranging from 97% to 100%. In addition, PPX/GPPA enzymes are also conserved among other sequenced Campylobacter spp. (PPX1/GPPA sequence identity 35% to 97%; PPX2/GPPA sequence identity 46% to 99%). However, the PPX1/GPPA and PPX2/GPPA share 30% identity and 49% similarity in amino acid content as also noted in Corynebacterium glutamicum (PPX1; PPX2 share only 25% identity; 45% similarity) and M. tuberculosis (Rv0496-PPX/GPPA; Rv1026-PPX/GPPA share only 27% identity; 42% similarity).

The ppx1/gppa genes contribute to exopolysphosphatase function

In order to verify that the deletion of ppx1/gppa genes did not affect the overall growth properties of the mutants, we assessed the Δppx1 mutants growth by determining CFUs at different time interval in MH broth. Our result confirmed that there was no significant difference between CFU forming ability of Δppx1 mutants (Δppx1, Δppx2, dkppx) and wild type (Fig. S3).

To investigate ppGpp role in exopolysphosphatase activity, we measured the intracellular poly-P levels in Δppx1 mutants (Δppx1, Δppx2, and dkppx). Our results indicated that the deletion of ppx genes alters the intracellular poly-P levels when compared with wild type. The Δppx1 and dkppx mutants showed significant (P ≤ 0.05, P ≤ 0.001) increase in the poly-P levels (45.5 nM and 48.5 nM/mg of total cellular protein [TCP], respectively) when compared with wild type (35 nM/mg TCP). The Δppx2 mutant also exhibited an increased poly-P levels (39.5 nM/mg TCP) but it was not significant (P = 0.079) (Fig. 1). The complemented strains (Cppx1 and Cppx2) displayed levels of poly-P (37 nM/mg TCP) similar to wild type. Our findings confirm that the ppx1/gppa and ppx2/gppa mutants have elevated poly-P and thus may encode exopolysphosphatase activity. Furthermore, these results indicate that ppx1/gppa gene has a major role in poly-P degradation and both genes may have additive effect in poly-P hydrolysis as observed with dkppx mutant.

The ppx1/gppa gene contributes to ppGpp pool

It has been previously reported that metabolism of ppGpp in C. jejuni is primarily governed by SpoT; however, PPX/GPPA proteins in C. jejuni are structurally similar and possess conserved amino acid residues required for the bifunctional activity as seen in A. aeolicus, suggesting that these proteins may have a subsidiary role in ppGpp metabolism. To address this, we determined the levels of ppGpp in the wild type and Δppx mutants starved in MOPS for up to 24 h. Our results showed that there was no difference in ppGpp accumulation at 3 h but differences were observed at 6 h and 24 h. At 24 h, the Δppx1 and dkppx mutants displayed significantly decreased levels of ppGpp 6% (P < 0.05) and 12% (P < 0.001) respectively, compared with wild type. Conversely, the level of ppGpp in the Δppx2 mutant was significantly higher (30%) than wild type (Fig. 2). However, complemented strains (Cppx1 and Cppx2) did not show significant change in the intracellular ppGpp pool (data not shown). The lack of complementation is likely due to chloramphenicol used to grow these strains, since chloramphenicol has been shown to inhibit ppGpp synthesis. Further, the ppGpp levels were increased in both ppx1 (4.5%) and dkppx (5.76%) mutants compared with the wild type suggesting a dual
role for ppx1. These results suggest that ppx1/gppa contribute to both poly-P and ppGpp pools in C. jejuni. This is consistent with the predicted 3D structure of C. jejuni PPX/GPPA proteins (Fig. S2B and C). Protein sequence and predicted 3D structure of C. jejuni PPX1/GPPA resembles the bi-functional PPX/GPPA protein of A. aeolicus. Only C. jejuni PPX1/GPPA possesses two conserved arginine residues (R16 and R258 corresponding to R22 and R267 of A. aeolicus) necessary for GPPA activity suggesting that the PPX1/GPPA of C. jejuni may be associated with both PPX and GPPA function as shown in our study.

C. jejuni ppx1/gppa genes are essential for survival in basal media

Previous studies have shown that poly-P is essential for C. jejuni to survive under low nutrient stress. Our results indicate that C. jejuni Δppx mutants have altered poly-P homeostasis. Thus, we further studied the mutants’ ability to withstand nutrient limitation stress by monitoring the Δppx mutants’ nutrient stress survival in MEM. Results indicate that Δppx1, Δppx2, and Δppx2 mutants were significantly (P ≤ 0.05) defective in survival after 36 h. Trans-complementation of Δppx1 and Δppx2 mutants partially restored the nutrient stress tolerance ability of mutants comparable to wild type (Fig. 3A–C).

Growth of C. jejuni depends upon availability of free amino acids. Further studies have shown that the preferential utilization of serine, aspartate, glutamate and proline in vitro by C. jejuni. It is also known that perturbation in ppGpp levels is associated with amino acid starvation in E. coli, we hypothesize that specific amino acids supplementation to Δppx mutants may rescue ppGpp-induced growth defect in basal media (MEM-α). Supplementation of amino acids serine, proline, glutamate, and aspartate at 20 mM in basal media significantly rescued the survival defect of all Δppx mutants at t36, t48, and t60 h except for aspartate in Δppx2 and Δppx2 at 60 h (Fig. 3D). Growth of the Δppx2 mutant, even though not defective in intracellular ppGpp accumulation, was also rescued by specific amino acids supplementation. However, although the amino acid supplementation in wild-type C. jejuni enhanced its survival, it was not significant. These results suggest that altered poly-P and/or ppGpp levels in ppx mutants may contribute to C. jejuni nutrient starvation stress survival.

The ppx1/gppa genes are essential for motility and biofilm formation in C. jejuni

Our results showed that C. jejuni Δppx1 and Δppx2 mutants showed no impairment in biofilm formation where the amount of biofilm formation for the Δppx1 (OD560 = 0.154) and for Δppx2 (OD560 = 0.119) was similar to the wild type (OD560 = 0.145). However, the deletion of both genes (Δppx1 and Δppx2) did not show a significant change in the intracellular ppGpp pool compared with wild type as chloramphenicol has been shown to inhibit ppGpp synthesis (data not shown). Each bar represents the average from 2 independent experiments performed using duplicate samples in each experiment.

We next investigated the role of ppx1/gppa genes in C. jejuni biofilm formation. Results in Figure 4C and D indicate that Δppx1 and Δppx2 mutants showed no impairment in biofilm formation where the amount of biofilm formation for the Δppx1 (OD560 = 0.154) and for Δppx2 (OD560 = 0.119) was similar to the wild type (OD560 = 0.145). However, the deletion of both genes (Δppx1 and Δppx2) did not show a significant change in the intracellular ppGpp pool compared with wild type as chloramphenicol has been shown to inhibit ppGpp synthesis (data not shown). Each bar represents the average from 2 independent experiments performed using duplicate samples in each experiment.
in transmission related phenotypes such as motility and biofilm formation in *C. jejuni*.

**Deletion of ppx/gppa genes affects *C. jejuni* survival during osmotic stress but not oxidative stress**

The ability of *C. jejuni* ppx mutants to resist osmotic stress was determined in the presence of NaCl as an osmotic stressor. Our results indicate that there was no significant difference in the osmotic stress tolerance response when tested on MH agar containing 0.17 M NaCl in both Δppx1 and Δppx2 mutants compared with wild type. Conversely, the dkppx mutant did exhibit a significant (*P* ≤ 0.05) sensitivity to osmotic stress in comparison to wild type strain (Fig. 5A and B). Furthermore, our CFU data indicated that the survival of the dkppx mutant under liquid osmotic stress (MH broth containing 0.25 M NaCl) was also significantly (*P* ≤ 0.05) affected at 48 and 60 h. Since, Δppx1 and Δppx2 mutants did not show defect in osmotic stress tolerance on solid media, we did not test these strains in liquid culture. These results indicate that the ppx/gppa genes may play a role in the *C. jejuni* osmotic stress tolerance.

The Δppx mutants showed no significant difference between zones of sensitivity to oxidative stressors (0.3% H₂O₂ or 20 mM Paraquat) compared with wild type (Fig. S4A and B).

**The Δppx mutants exhibit defect for invasion and intracellular survival**

To assay virulence associated phenotypes of invasion and intracellular survival, *C. jejuni* Δppx mutants and wild-type strains were allowed to infect INT 407 human intestinal epithelial cells at different MOIs. Our results indicated that the invasion profile of the Δppx mutants and wild type were similar at higher MOI; however, at low MOI the mutants were consistently less invasive. The intracellular survival was also assessed using a MOI of 100:1, although both Δppx2 and dkppx mutant were defective...
in intracellular survival, only the dkppx mutant was significantly ($P \leq 0.001$) defective in comparison to wild type (Fig. 6A and B). These data suggest that both PPX/GPPA enzymes are important for the invasion and intracellular survival.

The $ppx1$ and $ppx2$ mutants display resistance to complement-mediated killing

The bactericidal activity of complement factors present in serum is an innate defense mechanism against intruding
pathogenic bacteria. Since the lack of PPX in Neisseria meningitidis increased its resistance to complement-mediated killing, we next asked whether PPX/GPPA may have similar role in C. jejuni. To assess the complement mediated killing we compared the growth and survival of mutant and wild type strains in pooled normal (N) or heat inactivated (HI) human and chicken serum (NHS/NCS), respectively. The Δppx1 and Δppx2 mutants showed significant ($P \leq 0.05$) resistance to killing by human serum complement compared with wild type; surprisingly, the deletion of both the genes (dkppx) significantly ($P \leq 0.05$) increased the sensitivity to complement mediated killing. Trans-complementation of ppx1 and ppx2 genes restored the complement mediated killing to levels similar to wild type (Fig. 7A). However, surface structures such as LOS and CPS could not be attributed to complement mediated resistance/sensitivity observed for the mutants (Fig. S5A and B). This was further confirmed by qRT-PCR analysis for the expression of genes involved in LOS and CPS (kpsM, cstII, waaF, and lgtF) biosynthesis, which was similar in all mutants compared with wild type (Fig. S6).

To further investigate the complement pathways involved, the activity of the classical and lectin pathways was inhibited in normal sera by adding 10 mM MgCl$_2$ and 10 mM EGTA. Addition of MgCl$_2$ and EGTA inhibited the observed relative serum resistance in both Δppx1 and Δppx2 and the sensitivity of dkppx (Fig. 7B) suggesting that serum resistance or sensitivity is likely mediated by the classical and/or the lectin pathway. Further, complement-mediated killing in human serum at 37 °C and 42 °C was similar; however, the killing efficiency was slightly pronounced at 37 °C (data not shown). In contrast to human serum, chicken serum had no effect on the killing of the Δppx mutants (Fig. 7C).

spoT and ppk1 are upregulated in Δppx mutants

Next, in order to better understand the mechanistic aspects of poly-P homeostasis in C. jejuni Δppx mutants, we performed qRT-PCR targeting known key genes involved in poly-P and ppGpp metabolism (spoT, ppk1, and ppk2). Our results show that spoT expression was upregulated 2-fold or more in all three mutants and was significant ($P \leq 0.05$) in Δppx1 mutant (>6-fold) (Fig. 8) compared with wild type. Similarly, ppk1 expression was also upregulated more than 2-fold in the ppx1 and dkppx mutants; whereas the expression of ppk2 was similar in all mutants compared with wild type. These data suggest that the upregulation of ppk1 and spoT might be important in regulating
poly-P and ppGpp levels thus avoiding excessive accumulation of poly-P in the absence of \(\text{ppx}\) genes.

Discussion

In contrast to other well studied enteric pathogens such as \textit{Salmonella}, \textit{Shigella}, and \textit{E. coli}, \textit{C. jejuni} lacks most of the classical virulence factors. Sequencing of several \textit{C. jejuni} strains, including one known to be highly virulent and invasive strain (\textit{C. jejuni} 81–176), did not uncover any of the classical virulence factors.\textsuperscript{33-35} Moreover, relatively small size genome (1.64 Mb) of \textit{C. jejuni} demands to exploit most of its genes efficiently to meet great challenges to survive and transmit between hosts. Hence, a better understanding of the host-microbe interactions that affect the \textit{C. jejuni} pathogenesis is of immense importance.

Previous studies have shown that the poly-P plays a critical role in virulence related phenotypic traits in pathogenic bacteria including \textit{C. jejuni}.\textsuperscript{6,36,37} In this work, we focused to expand our understanding of poly-P homeostasis in \textit{C. jejuni} by investigating the role of PPX enzymes. In \textit{M. tuberculosis}, deficiency of PPX leads to increased accumulation of poly-P and restricted the growth in auxenic cultures and in human macrophages.\textsuperscript{38} In \textit{A. aeolicus}, PPX/GPPA enzymes, in addition to poly-P degradation, also breaks down pppGpp to ppGpp,\textsuperscript{18,23} playing an important role in the starvation induced stringent response. As shown in the present study, \textit{C. jejuni} has two exopolyphosphatases which individually or additively act in maintaining the poly-P pool (Fig. 1) but are not redundant. In \textit{C. glutamicum}, deletion of \textit{ppx1} or \textit{ppx2} resulted in increased accumulation of poly-P and restricted the growth in auxenic cultures and in macrophages.\textsuperscript{30} In \textit{A. aeolicus}, PPX/GPPA enzymes, in addition to poly-P degradation, also breaks down pppGpp to ppGpp,\textsuperscript{18,23} playing an important role in the starvation induced stringent response. As shown in the present study, \textit{C. jejuni} has two exopolyphosphatases which individually or additively act in maintaining the poly-P pool (Fig. 1) but are not redundant. In \textit{C. glutamicum}, deletion of \textit{ppx1} or \textit{ppx2} resulted in increased accumulation of poly-P and restricted the growth in auxenic cultures and in macrophages.\textsuperscript{30} Determination of exopolyphosphatase activity and intracellular poly-P concentration in \textit{C. glutamicum} revealed that PPX2 is the major exopolyphosphatase effective on short chain poly-P. \textit{C. glutamicum} PPX1 and PPX2 share only 25% amino acids identity and it was suggested that non-overlapping substrate preference may be the reason for occurrence of two or more exopolyphosphatase within one species.\textsuperscript{15} Likewise, PPX1 and PPX2 of \textit{C. jejuni} share only 30% amino acid identity with the PPX1/GPPA likely serving as a major exopolyphosphatase; however, further analysis to detect poly-P molecules and their sizes are needed to determine

Figure 7. Complement dependent killing of \textit{C. jejuni} \(\Delta\text{ppx}\) mutants. (A) \(\Delta\text{ppx}\) mutants killing by human serum. (B) \(\Delta\text{ppx}\) mutants killing by human serum in the presence of MgCl\(_2\) and EGTA (C) \(\Delta\text{ppx}\) mutants killing by chicken serum. Log\(_{10}\) killing was determined by subtracting the difference between number of CFU determined in heat inactivated human or chicken serum (HINHS or HINCS) to normal serum (NHS or NCS). Bars represent the mean ± SE of 3 independent experiments performed in duplicates each time. *\(P\) ≤ 0.05.

Figure 8. qRT-PCR analysis of \textit{C. jejuni} \(\Delta\text{ppx}\) mutants for genes involved in poly-P and ppGpp homeostasis. Fold difference in transcript levels was assessed by \(\Delta\Delta\text{CT}\) method. The expression of target genes was normalized to the 16s-rRNA expression levels from the same strain and then the relative expression of target genes were determined by comparing to expression in wild type strain. Each bar represents the mean ± SE of the relative fold change in expression from 4 independent experiments performed in duplicates each time. *\(P\) ≤ 0.05.
C. jejuni PPX1 and PPX2 substrate specificity. The occurrence of two or more exopolyphosphatases within a species is not restricted to C. jejuni as is also seen in E. coli, Vibrio cholera, M. tuberculosis, and C. glutamicum. If we take into account that poly-P confers improved fitness under environmental stress, then the selective maintenance of ppGpp genes may be strongly associated with changing environments, while an immovable environment, such as a host cell, could cause gene/function loss, as suggested by the lack of ppGpp genes in Mycoplasma spp.

Previous studies of PPX/GPPA proteins in A. aeolicus and E. coli have shown that these enzymes possess bifunctional activity. However, in M. tuberculosis two PPX-GPPA homologs were shown to have distinct biochemical activities. To our knowledge, this is the first study in C. jejuni depicting its PPX/GPPA possible dual role and its significance in numerous transmission and virulence related phenotypes. Our densitometry data for ppGpp production indicated that both Δppx1 and Δppx2 mutants have decreased ppGpp (Fig. 2), while ppx2 mutant has increased ppGpp levels when compared with wild type. This suggests that the ppGpp metabolism in C. jejuni is also regulated by factors other than PPXs. In this context Gaynor et al. showed that C. jejuni stringent response is mediated by SpoT which mediates both ppGpp synthesis and pppGpp hydrolysis and lack of this protein impairs C. jejuni resistance to rifampicin. Our qRT-PCR data support the concept of SpoT upregulation as compensation mechanism in poly-P and ppGpp homeostasis (Fig. 8). However, sensitivity to rifampicin was not affected in the ppx mutants (data not shown) which suggest that ppGpp levels are not below the threshold levels to have any effect on rifampicin sensitivity. Further studies are needed to understand the complex mechanisms behind regulation of (p)ppGpp by the PPX-GPPA proteins in C. jejuni.

C. jejuni’s ability to withstand the nutrient limited conditions in environment is critical for waterborne transmission. The biofilm formation and viable but not culturable (VBNC) state are known defense mechanism that a bacterium can exploit. The poly-P homeostasis in C. jejuni is shown to be important for biofilm formation, nutrient stress survival, osmotic stress tolerance, and for the maintenance of the VBNC state. Furthermore, C. jejuni flagellar motility has also been implicated in biofilm formation. As shown in B. cereus, our data indicate that PPX/GPPA enzymes are required for the motility and biofilm formation. The motility was affected in both mutants, a pronounced defect was seen in the Δppx2 mutant, suggesting that C. jejuni ppx1/gppa and ppx2/gppa genes additively contribute to motility. Conversely biofilm formation was only affected in the Δppx2 mutant (Fig. 4A and B) indicating a complex interconnected network of genes regulating this process in C. jejuni.

Complement-mediated bacterial killing is a key component of the humoral arm of the innate immunity necessary to control systemic infections from mucosal surfaces. Thus encapsulated bacteria counteract the antibacterial effect of complement by preventing deposition of the membrane attack complex on bacterial membranes. In N. meningitidis, lack of PPX increases its resistance to complement-mediated killing independently of changes in surface structures such as CPS, LPS, and the factor-H-binding protein. Instead N. meningitidis lack of PPX function modifies the interaction with the components of alternative pathway of complement activation. Our findings are in agreement with this previous report, as the deletion of individual ppx genes conferred resistance to complement-mediated killing. Consistent with this finding, in a previous study it was reported that poly-P by itself can inhibit complement-mediated killing. Strikingly, our result also showed that in C. jejuni deletion of both genes (Δppx2) results in sensitivity to complement-mediated killing (Fig. 7A) suggesting a pleotropic effect of the double knockout contributing to its sensitivity even in the presence of higher levels of poly-P.

Analyses of the CPS and LOS profiles of C. jejuni ppx mutants as well as qRT-PCR analysis of selected LOS/CPS biosynthetic genes suggested that these surface structures may not play a role in complement-mediated killing. Additionally, our results revealed that either classical or lectin mediated complement activation pathway are likely to play a role in C. jejuni ppx-mediated resistance or sensitivity to human serum (Fig. 7B). Perhaps the increased resistance of ppx1 and ppx2 mutants may suggest that C. jejuni uses it genes optimally to suite its lifestyle in humans and thus facilitating further spread to new host. However, additional studies are required to resolve the mechanism(s) behind observed susceptibility to the complement mediated killing. Interestingly, there was no effect of chicken complement factors on ppx mutants’ killing (Fig. 7C). This could be explained if the functionality of PPX/GPPA proteins depends on several factors such as temperature, host restricted expression of ppx genes, and/or the host immune system tolerance, among others. To our knowledge, this is first study showing differential contribution of ppx genes to complement mediated killing.

In our study in general the significant phenotypic changes (motility, biofilm formation, osmotic stress survival, and serum sensitivity) were observed in the double mutant (Δppx2) compared with the single mutants. However at present, we do not know what molecules are involved in these phenotypic differences. We believe that it is either due to additive effect of ppx1 and ppx2 deletion leading to higher poly-P levels or decreased ppGpp in the double knockout compared with single mutant; for example, motility was affected more in the double mutant compared with the single mutants. However, we observed osmotic stress defect, biofilm formation defect, and serum sensitivity only in the double mutant, we believe that these results observed in the double mutant is most likely a pleotropic effect of increased poly-P or decreased ppGpp to a threshold levels that is not seen in single mutants. It is also likely that poly-P and/or ppGpp molecules act separately or in combination toward specific phenotypes. In this regard, a recent study has demonstrated that ppGpp and poly-P signaling molecules uniquely and/or commonly regulated developmental regulators with cell type specific activities in Caulobacter crescentus.

In conclusion, we report that C. jejuni ppx/gppa genes are important for maintaining critical levels of poly-P and ppGpp in the cell. Besides their role in stress related phenotypic traits, they have a role in complement mediated killing. Our study expands the multi-factorial regulation of poly-P metabolism in C. jejuni.
Table 1. Bacterial strains and plasmids used in this study

| Strain/plasmid | Relevant description | Source |
|---------------|----------------------|--------|
| **Strains**   |                      |        |
| *C. jejuni* 81–176 | Used as wild type strain of *C. jejuni* | Dr Qijing Zhang |
| *E. coli* DH5α | *E. coli* strain used for cloning | Invitrogen |
| Δppx1         | *C. jejuni* 81–176 derivative with deletion in ppx1/gppa gene; ppx1/gppa::Kan | This study |
| Δppx2         | *C. jejuni* 81–176 derivative with deletion in ppx2/gppa gene; ppx2/gppa::Kan | This study |
| dkppx         | *C. jejuni* 81–176 derivative with deletion of ppx1/gppa and ppx2/gppa; ppx1/gppa::Kan, ppx2/gppa::Cm | This study |
| Cppx1         | Δppx1 with pRY111- containing ppx1 coding region and the upstream promoter sequence for complementation; Cm | This study |
| Cppx2         | Δppx2 with pRY111- containing ppx2 coding region and the upstream promoter sequence for complementation; Cm | This study |
| **Plasmids**  |                      |        |
| pRY111        | *E. coli*-Campylobacter shuttle vector for complementation | Yao et al.25 |
| pRK2013       | Helper plasmid for conjugation, Kan | Akiba et al.38 |
| pZErO-1       | Cloning vector for making suicide vector; Zeo | Invitrogen |
| pUC4K         | Source for kanamycin gene, Kan | Amersham |
| pZErO-1-ppx1  | pZErO-1 containing ppx1/gppa region plus 1 kb upstream and downstream sequences from *C. jejuni* 81–176; Zeo | This study |
| pppx1-suicide vector | pZErO-1-ppx1 with ppx1/gppa replaced by kanamycin resistance gene from pUC4K; Zeo, Kan | This study |
| pZErO-1-ppx2  | pZErO-1 containing ppx2/gppa region plus 1 kb upstream and downstream sequences from *C. jejuni* 81–176; Zeo | This study |
| pppx2-suicide vector | pZErO-1-ppx2 with ppx2/gppa replaced by kanamycin resistance gene from pUC4K; Zeo, Kan | This study |
| pppx2-suicide vector-Cm | ppx2-suicide vector with kanamycin resistance marker replaced by chloramphenicol resistance gene from pRY111; Zeo, Cm | This study |
| pRY111-ppx1   | pRY111 containing ppx1 coding and upstream sequence for complementation, Cm | This study |
| pRY111-ppx2   | pRY111 containing ppx2 coding and upstream sequence for complementation, Cm | This study |

Kan, kanamycin; Zeo, zeocin; Cm, chloramphenicol

Materials and Methods

Bacterial strains and growth conditions

Bacterial strains and plasmids used in this study are listed in Table 1. All studies were performed with the highly invasive *C. jejuni* strain 81–176 originally isolated from a diarrheic patient.37 C. jejuni cultures were routinely grown on Mueller–Hinton (MH; Becton Dickinson and Company) agar under microaerobic condition (85% N₂, 10% CO₂, and 5% O₂) in a DG250 Microaerophilic Workstation (Microbiology International). For the growth curve and stress survival assays, *C. jejuni* strains were cultured in MH broth with or without specific antibiotics at 42 °C. Growth media were purchased from Promega. pZEro-1, the zero background cloning shuttle vector for *Campylobacter* species, was purchased from Epicenter. Restriction enzymes were purchased from New England Biolabs. pZErO-1, the zero background cloning shuttle vector for *Campylobacter* species, was purchased from Epicenter. Restriction enzymes were purchased from New England Biolabs. pZErO-1, the zero background cloning shuttle vector for *Campylobacter* species, was purchased from Epicenter. Restriction enzymes were purchased from New England Biolabs.

Cloning techniques

The primers for PCR and qRT-PCR were designed using Vector NTI® software (Invitrogen) or Integrated DNA Technologies scientific tool (IDT) and were commercially synthesized by IDT. All the primers used in the present study are listed in the Table S1. QIAquick® PCR purification and a spin miniprep kits for plasmid isolation were purchased from Qiagen. Masterpure® DNA purification and Fast-Link DNA ligation kits were purchased from Epicenter. Restriction enzymes were purchased from Promega. pZErO-1, the zero background cloning vector and *E. coli* DH5α competent cells were purchased from Invitrogen. Cloning and other molecular biology techniques were performed as previously described.38

Construction of *C. jejuni* 81–176 ppx deletion mutants and complemented strains

*C. jejuni* 81–176 genome possesses two exopolyphosphatase genes, *ppx1/gppa* (CJJ81176_0377) and *ppx2/gppa* (CJJ81176_1251) encoding 486 and 324 amino acids, respectively (Fig. S7). Further, BLAST search using *ppx/gppa* gene from *Aquifex aeolicus* as a reference sequence confirmed that these two genes likely represent exopolyphosphatases. The deletion mutants (Δppx1, Δppx2, and dkppx) were generated by double crossover homologous recombination as we described previously.41 Briefly, to construct the mutant, the target gene, along with its upstream and downstream sequences (approximately 1

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kb), was amplified from _C. jejuni_ 81–176 genomic DNA with specific primers. The PCR product was cloned into pZErO-1 plasmid, the resulting construct was amplified by inverse PCR ensuring that most of the _ppx1_ and _ppx2_ coding sequences were deleted. The inverse PCR product was ligated to a kanamycin gene from pUC4K and the final suicide vector was introduced into _C. jejuni_ 81–176 by electroporation as described. To generate a double knockout (dkppx) mutant, _ppx2_ suicide vector, in which the kanamycin resistance gene was replaced with chloramphenicol gene, was electroporated into ∆ppx1 mutant. Colonies grown on MH agar containing kanamycin and chloramphenicol were selected as the double knockout mutant and deletion of target genes were confirmed by PCR using primers specific to upstream and downstream sequences of the target gene.

For complementation, target genes along with the potential promoter regions were amplified from genomic DNA with specific set of primers; products were cloned into pRY111, an _E. coli–Campylobacter_ shuttle vector. Resulting constructs were introduced into the respective mutants by triparental conjugation. The complementation strain for the _dkppx_ mutant was not generated due to a limited availability of replicating plasmids as well as antibiotic markers of choice for use in _C. jejuni_. Complemented strains ( _Cppx1_ and _Cppx2_ ) were also confirmed by PCR.

**Growth curve assay**

The growth curve assay was performed as described previously. Cultures were grown to mid-log phase on MH agar, washed, suspended in MH broth, adjusted to OD₆₀₀ 0.05, and incubated microaerobically for 42 °C for 60 h with shaking at 200 rpm. For assessing the growth, CFUs were determined at different time points by plating 10-fold serially diluted cultures on MH agar.

**Extraction and quantification of poly-P**

The extraction of poly-P from _C. jejuni_ strains were performed using the glassmilk method as previously described. Briefly, _C. jejuni_ cultures were grown in MH broth to mid-stationary phase to reach OD₆₀₀ of 0.3, cells were harvested followed by lysis with buffer (guanidine isothiocyanate-GITC 4 mM, 500 mM TRIS-HCl pH 7.0). Total protein concentration in the lysate was estimated by Bradford analysis (BCA kit, Pierce Scientific) and poly-P content was analyzed using toluidine blue O dye. Briefly, for poly-P quantification, lysates were washed with buffer (5 mM TRIS-HCl [pH-7.5], 50 mM NaCl, 5 mM EDTA, 50% ethanol) and pellets were resuspended in a buffer (50 mM TRIS-HCl [pH 7.4], 10 mM MgCl₂, and 20 μg each of DNase and RNase per mL) and the poly-P was eluted by brief centrifugation at 9500 × g with elution buffer (50 mM TRIS-HCl pH 8.0).

The poly-P extracted from _C. jejuni_ lysates were incubated with toluidine blue O dye (TBO 6 mg/liter in 40 mM acetic acid) at room temperature followed by assessing absorbance at 630 nm and 530 nm, where the ratio of 530 nm/630 nm provides the amount of poly-P in a given sample. Levels of poly-P in _C. jejuni_ lysates were determined by direct comparison with a phosphorus standard curve and were expressed as mmol/mg of total cellular protein.

**ppGpp isolation and detection**

Isolation and detection of ppGpp was assayed as described previously. Briefly, bacterial strains were grown on MH agar for 16–18 h. Cultures were diluted to adjust OD₆₀₀ of 0.25 in MH broth; cells were pelleted, washed twice in MOPS-MGS at 9500 × g (50 mM MOPS, 55 mM mannitol, 1 mM MgSO₄·6H₂O, 0.25 mM CaCl₂, 19 mM glutamic acid, and 0.004 mM biotin) and suspended in 250 μL MOPS-MGS. 32P at 100 μCi per mL (3.7 × 10¹² Bq) was added to cells and incubated microaerobically for 3, 6, and 24 h at 37 °C. Previously we have observed that the prolonged exposure (starvation stress) in MOPS results in increased accumulation of ppGpp that can be readily detected. Labeled bacteria were harvested at the above specified times, washed and treated with lysozyme in 10 mM Tris (pH 8.0) for 20 min. The ppGpp was extracted with equal volume of 2 M formic acid and placed on ice for 15 min. Samples were spun for 5 min at 9500 × g, and 3 μL of supernatant was spotted directly onto cellulose TLC plates, dried, and developed in 1.5 M KH₂PO₄ and visualized by autoradiography. For TLC separation, loading concentrations were normalized across all strains and approximately equal number of cells were loaded on the TLC plate.

**Motility assay**

The motility assay was performed as described previously. Briefly, _C. jejuni_ grown overnight on MH agar microaerobically at 42 °C was harvested, washed with MH broth and OD₆₀₀ was adjusted to 0.05. Two microliter culture (~5 × 10⁵ cells) was stabbed onto 0.4% semi-solid agar and incubated microaerobically at 42 °C. After 24 h of incubation, the swarming zone (diameter) around the stabbed area was measured in centimeters.

**Biofilm assay**

The static biofilm formation was assayed as previously described. Briefly, cells were grown microaerobically overnight on MH plate at 42 °C, harvested in MH broth and OD₆₀₀ was adjusted to 0.05. Then the culture (100 μL) was inoculated to 2 mL of MH broth in borosilicate tubes and incubated microaerobically at 42 °C for 48 h without shaking. Static biofilm formation was visualized by staining with 1% (w/v) crystal violet for 15 min. To quantify amount of biofilms, 3 mL of dimethyl sulfoxide (DMSO, Sigma) was added and incubated at room temperature for 24 h. The absorbance was measured at 570 nm.

**Growth in chemically defined medium**

Briefly, _C. jejuni_ cultures were grown microaerobically on MH agar for 16–18 h at 42 °C. The cultures were washed twice by centrifuging at 9500 × g for 2 min with minimum essential media (MEM without glutamine) or stringent basal media (MEM-α) with or without supplementation of amino acids l-serine, l-aspartate, l-glutamate, and l-proline at 20 mM concentration. Bacteria were suspended in the defined media to adjust OD₆₀₀ of 0.05 (approximately 1 × 10⁸ CFU/mL) and incubated microaerobically at 42 °C with shaking at 200 rpm. CFUs were determined by plating on MH agar at different time intervals.

**Osmotic stress assay**

Osmotic stress was determined as described previously. Briefly, mid log-phase grown bacterial cultures were suspended in MH broth and OD₆₀₀ adjusted to 0.05. Cultures were then serially diluted (10-fold) and a 10 μL samples were spotted on
MH agar containing 0.17 M NaCl and the growth was assessed after 48 h of incubation microaerobically at 42 °C. To test survivability of ppx/ppa mutants to osmotic stress in liquid media, cultures were adjusted to OD₆₀₀ 0.05 in MH broth containing 0.25 M NaCl and incubated microaerobically for 60 h at 42 °C with shaking at 200 rpm. CFUs were assessed at different time points by plating on MH agar.

**Oxidative stress assay**

The ability of *C. jejuni* ppx mutants to survive under oxidative stress was assessed as previously described. Bacterial strains (5 × 10⁷ cells) were spread for confluent growth on MH agar. At the center of MH agar, 6 mm diameter hole was made and well bottom was sealed with 50 μL of molten MH agar. Wells were filled with 30 μL 0.3% H₂O₂ or 20 mM paraquat and plates were incubated microaerobically for 24 h at 42 °C, clear zones of growth inhibition surrounding the wells were measured in centimeter.

**Rifampicin sensitivity assay**

Fresh overnight cultures of *C. jejuni* ppx strains were streaked onto MH plate containing different rifampicin concentration (50, 100, 150, 200, 250, and 300 μg/mL) as previously described. The plates were incubated at 37 °C, microaerobically for 24–48 h to determine the sensitivity.

**Invasion and intracellular survival in INT407 cells**

The invasion and intracellular survival in INT 407 cells were performed as previously described. INT 407 cells (1.4 × 10⁵) were washed with MEM twice, and 100:1 MOI was used for intracellular survival assay. To determine the sensitivity.

**Serum bactericidal assay**

Serum bactericidal assay was performed as described previously. Commercially available pooled normal human serum (NHS) and pooled normal chicken serum (NCS) (Innovative Research) were used. Brieﬂy, *C. jejuni* strains were harvested after 16–18 h of growth on MH plates, washed and suspended to 10⁶ cells/mL in Medium 199 with Hank’s balanced salt solution containing 0.01% glutamine. From each culture, 150 μL aliquots were transferred in duplicate to a 96-well microtiter plate. To each well, 50 μL of either 100% NHS/NCS in Medium 199, 100% heat-inactivated (56 °C for 30 min) HI-NHS/NCS or Medium 199 alone was added to obtain 25% final concentration of NHS/NCS or HI-NHS/NCS and the plate was incubated for 60 min microaerobically at 37 °C for human serum, and at 42 °C for chicken serum. Aliquots (100 μL) from each well were 10-fold serially diluted and plated to assess CFUs. Log₁₀ killing was determined by subtracting the number of CFUs between HI-NHS/NCS to NHS/NCS-treated samples.

**Detection of capsular polysaccharide (CPS) and lipooligosaccharide (LOS)**

CPS and LOS were analyzed as described previously. For CPS analysis, *C. jejuni* wild type and mutant cells were harvested after 48 h of growth, washed in PBS, standardized to an OD₆₀₀ of 2.0, then solubilized in 200 μL of sample buffer (2% SDS, 4% 2-mercaptoethanol, 10% glycerol, 1 M TRIS-HCl, pH 6.8, and 0.02% bromophenol blue), for 10 min at 100 °C, and then subsequently incubated with 60 μg of proteinase K for 1 h at 60 °C. Whole cell lysates were analyzed by 15% SDS-PAGE at 80 mA for 4 h. Pre-stained protein marker (HyperPAGE, Bioline) was used to determine the molecular mass and bands were resolved with periodic acid silver nitrate (PAS) stain. The presence of LOS (normalized by protein content) was analyzed by 1 dimensional silica gel thin layer chromatography (TLC) using the solvent system n-propanol-water-25% NH₄OH (60:30:10 v/v/v), developed with 10% (v/v) sulfuric acid in ethanol, and visualized by heating the TLC at 120 °C for few min as previously described.

**Quantitative RT-PCR**

qRT-PCR was performed targeting key genes involved in poly-P homeostasis, CPS, and LOS synthesis. Mid stationary phase grown bacterial cultures were used for total RNA extraction using RNeasy Mini Kit (Qiagen). RNA concentration and purity were determined using NanoDrop ND-2000c spectrophotometer (Thermo Scientific) and agarose gel electrophoresis. DNase treated RNA (200 ng) was used for cDNA synthesis using SuperScript® III First-Strand Synthesis SuperMix kit (Invitrogen). Following cDNA synthesis, cDNA concentration was normalized to 200 ng and gene specific primers were used to amplify the genes involved in poly-P homeostasis (*spoT, ppk1, cstII, waaF, and lgtF*), CPS and LOS synthesis (*kpsM, cstII, waaF*, and *lgtF*) along with 16S-rRNA as internal control. qRT-PCR was performed using SensiMixPlus® SYBR RT-PCR Kit (Quantace) in a replex® mastercycler (Eppendorf). The expression of genes were normalized using 16S-rRNA expression of the same strain and the relative difference in expression of genes was calculated using the comparative threshold cycle (ΔΔCt) method to yield fold-difference in transcript levels compared with wild type.

**Statistical analyses**

Statistical significance of data was determined using one-way analysis of variance (ANOVA) followed by the Tukey multiple comparison post-test or Student t test (paired 2-tailed). A P value of P ≤ 0.05 (α level) was considered statistically significant.

Disclosure of Potential Conflicts of Interest

No potential conflicts of interest were disclosed.
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